

# Bibliometric Analysis of Ongoing Projects

13th Report  
September 2022

Document reference: IMI.2018.OP.01.

Copyright©2022 Innovative Health Initiative

# TABLE OF CONTENTS

<b>Bibliometric Analysis of Ongoing Projects .....</b>	<b>1</b>
13th Report September 2022.....	1
<b>1 EXECUTIVE SUMMARY .....</b>	<b>4</b>
Summary of key findings.....	5
<b>2 OVERVIEW.....</b>	<b>8</b>
2.1 Innovative Medicines Initiative (IMI) .....	8
2.2 Clarivate.....	8
2.3 Scope of this report .....	9
<b>3 DATA SOURCES, INDICATORS AND INTERPRETATION.....</b>	<b>10</b>
3.1 Bibliometrics and citation analysis.....	10
3.2 Data source.....	11
3.3 Methodology.....	11
3.4 Data collation .....	13
<b>4 CITATION ANALYSIS – IMI SUPPORTED PUBLICATIONS OVERALL.....</b>	<b>14</b>
4.1 Publications from IMI-supported projects.....	14
4.2 Publications from IMI projects by document type.....	16
4.3 Trends in publication output.....	17
4.4 Publication output by country.....	19
4.5 Publication output by IMI project.....	22
4.6 Is IMI project research well cited?.....	24
4.7 In which journals do IMI project publications appear most frequently?.....	26
4.8 Which research fields account for the highest volume of IMI project publications?.....	30
4.9 IMI research fields with the highest volume of publications benchmarked against EU-28 publications of the same field .....	35
<b>5 CITATION ANALYSIS – AT IMI PROJECT LEVEL.....</b>	<b>37</b>
5.1 Trends in publication output by IMI funding call.....	37
5.2 Summary bibliometric analyses for imi 1 projects – call 1.....	42
5.3 Summary bibliometric analyses for IMI 1 projects – call 2.....	44
5.4 Summary bibliometric analyses for IMI 1 projects – call 3.....	47
5.5 Summary bibliometric analyses for IMI 1 projects – call 4.....	49
5.6 Summary bibliometric analyses for IMI 1 projects – calls 5-10.....	51
5.7 Summary bibliometric analyses for IMI 1 projects – call 11.....	54
5.8 Summary bibliometric analyses for IMI 2 calls 1-4 projects.....	56
5.9 Summary bibliometric analyses for IMI 2 calls 5-10 projects.....	58
5.10 Summary bibliometric analyses for IMI 2 calls 11-23 projects.....	62
<b>6 GEOGRAPHIC CLUSTERING ANALYSIS .....</b>	<b>65</b>
6.1 Locations where IMI-funded research takes place.....	65

<b>7 COLLABORATION ANALYSIS FOR IMI RESEARCH .....</b>	<b>81</b>
7.1 Collaboration analysis for IMI research.....	81
7.2 Collaboration analysis by IMI project.....	84
7.3 Collaboration metrics for IMI research .....	103
7.3.1 Metric 1 (cross-sector score): fraction of cross sector collaborative papers.....	104
7.3.2 Metric 2 (international score): fraction of internationally collaborative papers.....	106
7.3.3 Metric 3 (stability score): stability of institutional collaboration.....	108
7.4 Collaboration index.....	113
<b>8 BENCHMARKING ANALYSIS – IMI PROJECT RESEARCH AGAINST RESEARCH FROM SELECTED COMPARATORS .....</b>	<b>117</b>
8.1 Identifying comparators.....	117
8.2 Trends in output: IMI project research compared with selected comparators .....	118
8.2.1 Trends in output: IMI project research compared with selected comparators .....	118
8.2.2 Trends in field-normalised citation impact: IMI project research compared with selected comparators 121	
8.2.3 Trends in journal-normalised citation impact: IMI project research compared with selected comparators.....	123
8.2.4 Trends in raw citation impact: IMI project research compared with selected comparators.....	126
8.2.5 Trends in uncited research: IMI project research compared with selected comparators.....	129
8.2.6 Trends in highly cited research: IMI project research compared with selected comparators .....	131
8.2.7 Trends in open access research: IMI project research compared with selected comparators.....	134
8.3 Summary of bibliometric indicators: IMI project research compared with selected comparators.....	136
<b>ANNEX 1: Bibliometrics and citation analysis.....</b>	<b>137</b>
<b>ANNEX 2: Biomedically related journal categories.....</b>	<b>147</b>
<b>ANNEX 3: Total number of Web of Science Publications from IMI projects between 2010 and 2021 by country.....</b>	<b>148</b>
<b>ANNEX 4: Total number of Web of Science Publications, papers and open access papers between 2010 and 2021 by Project.....</b>	<b>152</b>
<b>ANNEX 5: Collaboration index for all IMI supported research projects.....</b>	<b>157</b>
<b>ANNEX 6: Bibliography of hot papers and highly cited papers.....</b>	<b>162</b>

# 1 EXECUTIVE SUMMARY

This report presents a bibliometric analysis of the Innovative Medicine Initiative Joint Undertaking's (IMI JU) research published between 2010 and 2021, using citations as an index of academic impact and co-authorship as an index of collaboration. This is the thirteenth report commissioned by IMI from Clarivate.

The data show that IMI continues to perform well. To date, IMI projects have produced 8,609 publications which have been matched to the Clarivate Web of Science™. This represents a 20% increase from the 7,177 publications matched to the Web of Science in the twelfth report, which covered IMI project research published between 2010 and 2021.

The number of IMI research publications has generally increased year on year, except for the year 2019 where output fell by almost 8% (likely due to some of the longest running projects e.g. BTCure coming to an end) as noted in the eleventh report. Publication growth has since recovered and in 2021, the most recent publication year, publication output has increased by nearly 10% from 2020. IMI's publication growth is showing signs of stabilisation, which is expected as the IMI programme matures. As a programme matures, longer running projects that have now closed observe lower publication outputs while the newer projects slowly see an increase in publication outputs. These changes in publication output cause the overall output to begin stabilising.

The majority of IMI research (65%) continues to be published in high impact journals, i.e., those journals in the highest quartile (Q1) when ranked by Journal Impact Factor, and the average Journal Impact Factor of all IMI project publications was 7.53. IMI research was wide-ranging from basic biological research to clinical practice. IMI project research has been published most frequently in the fields of Neurosciences, Pharmacology & Pharmacy, and Biochemistry & Molecular Biology.

The impact of IMI project research (as indicated by citation impact) remains twice (2.03) that of the world average (1.00), which indicates that the research was internationally influential. Between 2010 and 2021, the field-normalised citation impact of IMI papers was considerably higher (76%) than the European Union's (EU) average citation impact (1.15) in similar biomedical fields (journal subject categories). Around a quarter (25.3%) of IMI project papers were highly cited; that is, the papers were in the world's top 10% of papers (taking journal category and year of publication into account), when ranked by number of citations.

The output of individual IMI projects has also increased between 2010 and 2021. BTCure (Call 2) has remained the most prolific IMI project, with 719 publications as of this report. This is a 2% increase on the 703 publications attributed to BTCure in the previous report. It is also worth noting that AIMS-2-TRIALS is new to the Top 10 projects, ranking 6<sup>th</sup>, as its publication output has nearly doubled since the previous report (2020).

Projects funded by IMI are highly collaborative. Since the last report, an increasing percentage of IMI publications involve collaboration between researchers in different sectors. Two-thirds (67.0%) of all IMI project papers were co-authored by researchers working in different sectors, more than three-quarters (85.9%) involved collaboration between institutions and more than half (64.5%) were internationally collaborative. Internationally collaborative IMI project research had an average citation impact (2.73) well over twice the world average (1.00) and higher than domestically collaborative IMI project research (1.82).

Research in both Europe and North America tends to be clustered in major cities with an existing strong academic research base. The citation impact of IMI papers within these clusters is higher than

national averages and rates of international co-authorship are very high (70-100%) compared to the averages for EU-28 biomedical research (40%). The European and North American clusters with the highest proportion of open access papers are Lyon, France (98.8%) and Seattle (94.4%) respectively.

IMI's field-normalised citation impact (2.03) is two times the world average and is comparable to other well-established funding bodies such as the Medical Research Council (MRC) and the Wellcome Trust (WT) and is higher than all other comparators.

IMI's field-normalised citation impact increased more than the Critical Path Institute, Foundation for the National Institutes of Health (FNIH), and Grand Challenges in Global Health (GCGH) which saw no change or a decrease in their citation impact.

IMI's journal-normalised citation impact (1.23) is the second highest among the comparators only surpassed by CSIRO (1.25). IMI's percentage of highly cited papers (25.3%) outperforms all the comparators, except GCGH (26.5%). IMI publishes more open access papers than three out of the seven comparators (CSIRO, C-Path, and ICMR).

A more detailed summary of the key findings of this report (with cross-references to the relevant sections) is provided below.

## Summary of key findings

Since its first call for proposals in 2008, IMI has funded 182 projects from a total of 34 funding calls. Of the calls, 11 were from IMI's first phase (IMI 1), which ran from 2008 to 2013, and the rest from its second phase (IMI 2), which was launched in 2014 and ended in 2020. While the IMI 1 and 2 programmes have ended, many of the projects funded by these programmes are still ongoing.

It may take several months for a project to progress from inception to the point where it has generated sufficient data for a publication. It may take further months or years until it has produced its most valuable results. As some of the IMI projects analysed in this report are relatively young, the bibliometric indicators may not fully reflect their eventual impact.

- IMI projects have published a total of 8,609 unique Web of Science publications (Figure 4.1.1).
- IMI's publication growth is showing signs of stabilising as the programme matures (Figure 4.3.1).
- A quarter (25.3%) of IMI papers were in the world's top 10% of most highly cited papers in the relevant field and year of publication, suggesting very strong performance (Table 4.6.1).
- The field-normalised citation impact of IMI project papers was twice the world average (2.03) between 2010 and 2021, higher than last years (1.99). (Figure 4.6.1)
- More IMI project publications appeared in *Scientific Reports* (195 publications) and *PLOS One* (193 publications) than in other journals. Of the 20 journals in which IMI-funded projects published most frequently, nearly two-thirds (13 journals) rank in the top quartile by Journal Impact Factor (Table 4.7.1).
- The highest Impact Factor journal in which IMI research was published is the *Lancet* (6 publications), which has a Journal Impact Factor of 202.73. Of the Top 20 journals by Impact

Factor, IMI published most frequently in *Nature* (69.50) with 26 publications, followed by *Nature Medicine* (87.24) with 17 publications (Table 4.7.2).

- IMI project research was most frequently published in Neuroscience journals (Figure 4.8.1) a change from previous years where Pharmacology & Pharmacy was the most frequently published in category. Of the 875 papers published in Neuroscience, 28.1% were highly cited, 69.5% were open access, and the average citation impact of these papers was 2.12 times higher than the world average for the year and field of publication (Table 4.8.2 and Table 4.8.3).
- IMI research in the Clinical Neurology remains the category with the highest percentage of highly cited papers (37.5%). (Table 4.8.3)
- IMI project research had a citation impact well above the European (EU-28) average in all of the 20 journal subject categories to which most IMI publications are assigned (Table 4.9.1 and Figure 4.9.1).
- Early IMI 1 calls (1-4) follow a similar pattern of initial growth in publication output for 3 to 6 years followed by a decline as the projects end (Figure 5.1.1). Later IMI 1 calls published very few papers over the time period, normally less than 50 each year. The exception being IMI 1 call 11.
- The publication output of most of IMI 2 calls is currently growing. Especially Call 21 which had a 400% increase in publication output from last year's report. This is likely driven by the fact that many of the projects within this call are coronavirus related. IMI 2 Call 10 is also growing steadily. (Figure 5.1.3)
- Papers assigned to IMI 2 call 21 had the highest average field-normalised citation impact (3.71), more than three times the world average. Again, this is likely due to the projects within this call being coronavirus related. (Table 5.1.1)
- The largest geographic clusters of research supported by IMI in Europe are London (1,800 publications), Amsterdam (1,515 publications) and Stockholm (843 publications). The largest clusters in North America are Boston (392 publications), Toronto (368 publications) and New York (257 publications). (Table 6.1.1 and Table 6.1.3)
- IMI research in all the European and North American geographic clusters performs well above the national averages in terms of citation impact. The highest citation impact clusters in Europe are Maastricht (3.83) and Zurich (3.65), both more than twice their respective national averages which are both 1.71. (Table 6.1.2 and Table 6.1.4)
- Around 40% of all EU-28 biomedical research involves international co-authorship while in comparison rates of international collaboration for IMI project research are very high for most clusters, especially in North America where most clusters have around 90% international collaboration which is expected as IMI is a European funding organisation that primarily funds researchers working in EU-28. The European cluster with the highest rate of internationally collaborative papers was Basel, with 94.5% of its research involving international co-authorship. While the European cluster, Rome, was the lowest at 75.3% international collaboration. (Table 6.1.1 and Table 6.1.3).
- IMI project research is collaborative across sectors, institutions, and countries. Two-thirds (67%) of IMI project papers were co-authored by researchers from different sectors. More than three-

quarters (86%) of IMI project papers involved collaboration between different institutions. Nearly two-thirds (65%) of all IMI project papers were internationally collaborative (Table 7.1.1).

- IMI's collaborative research for sectors, institutions, and countries has an average field-normalised citation impact that is almost 50% higher than IMI's non-collaborative research. (sectors: 2.71 vs 1.82, institutions: 2.61 vs 1.60, and countries: 2.74 vs 1.82) (Figure 7.1.1)
- BTCURE, followed by EU-AIMS, had the highest number of papers with co-authors from more than one country, institution and sector. This may be due to these projects having the highest and second highest overall number of papers. (Table 7.1.1-Table 7.2.3)
- For those projects with at least 100 papers, BigData@Heart had the highest percentage of its papers with co-authors from more than one country (76.3%), sector (90.4%), and institution (98.5%), indicating the highly collaborative nature of this project. (Table 7.2.1-Table 7.2.3).
- King's College London is part of six out of the ten most productive pairs of collaborating institutions, including the second most productive pair where King's College London collaborated with Heidelberg University on 134 publications. (Figure 7.3.3.1)
- Karolinska University Hospital and Karolinska Institute were the top collaborating pair, collaborating on 148 publications.
- PROACTIVE has the highest collaboration index score of 2.66. (Table 7.4.1)
- IMI's research output grew faster between 2010 and 2021 than any of the seven selected comparators (Table 8.2.1.1).
- IMI's field-normalised citation impact (2.03) was lower than the Wellcome Trust's (2.09) and the MRC's (2.13) and higher than all the other comparators (Figure 8.2.2.2).
- IMI's percentage of uncited research has been the lowest of all the comparators since 2017, including the most recent year of 2021 (44.6%). (Table 8.2.5.1 ) However, it has the third highest percentage of uncited papers between 2010-2021.
- IMI has a higher percentage of highly cited papers (25.3%) than all the comparators except GCGH (26.5%). (Figure 8.2.6.2)

## 2 OVERVIEW

The Innovative Medicines Initiative (IMI) Joint Undertaking has commissioned Clarivate to undertake a yearly evaluation of its research portfolio using bibliometric indicators.

The commissioned evaluation comprises a series of reports focusing on research publications produced by IMI funded researchers. This report is the twelfth evaluation in the series.

### 2.1 Innovative Medicines Initiative (IMI)

IMI's purpose is to improve health by speeding up the development of, and patient access to, innovative medicines, particularly in areas where there is an unmet medical or social need. It does this by facilitating collaboration between the key players in healthcare research, including universities, pharmaceutical companies and other industries, small and medium-sized enterprises (SMEs), patient organisations, and medicines regulators.

IMI is a partnership between the EU and the European pharmaceutical industry, represented by the European Federation of Pharmaceutical Industries and Associations (EFPIA). IMI, as part of its second phase (IMI 2), has a budget of €3.3 billion for the period of 2014 to 2024. Half of this comes from the EU's research and innovation programme, Horizon 2020. The other half comes from large companies, mostly in the pharmaceutical sector; these organisations do not receive any EU funding, but contribute to the projects 'in kind', for example by donating their researchers' time or providing access to research facilities or resources. The first phase of IMI had a budget of €2 billion equally shared between EU and EFPIA.

To date, IMI has announced 11 calls for proposals under its first phase and a further 23 calls for proposals under its second phase. The first funding call was announced in 2008 and the final calls were launched in June 2020. In February 2021, the Innovative Health Initiative (IHI), a new public-private partnership in health was announced that will run under Horizon Europe, the new European framework programme for research and innovation. This new partnership will build upon the Innovative Medicines Initiative (IMI) but will have a greater focus on cross sectoral collaborations involving biopharmaceutical, medical technology, and biotechnology sectors. This report covers the research output (publications and papers) of a total of 61 projects from IMI phase one and 126 projects from IMI phase two.

### 2.2 Clarivate

Clarivate, provides reporting and consultancy services to enable customers to understand and interpret their research performance and to inform strategic decision-making. We have extensive experience with databases of research inputs, activity and outputs and have developed innovative analytical approaches for benchmarking, interpreting and visualising research impact.

Clarivate's Research Analytics is a suite of products, services and tools that provide comprehensive research analysis, evaluation and management. For over half a century we have pioneered the world of citation indexing and analysis, helping to connect scientific and scholarly thought around the world. Today, academic and research institutions, governments, not-for-profits, funding agencies, and all others with a stake in research, need reliable, objective methods for managing and measuring performance.



Our consultants have up to 20 years of experience in research performance analysis and interpretation. In addition, the Clarivate regional Sales team provide effective on-site support to maximise the value of our work.

Visit [Clarivate](#) or our [Professional Research Data Services](#) team online for more information.

## 2.3 Scope of this report

The analyses and indicators presented in this report have been selected to provide an analysis of IMI research published output for research management purposes:

- To identify excellence in IMI-supported research overall and at individual call or project level.
- To benchmark IMI project research performance against other funders research, the EU-28 biomedical research and world averages.
- To show that collaboration, at all levels (researcher, institutional and country), is being encouraged through the projects funded by IMI.

Outline of this report:

- Section 3 describes the data sources and methodology used in this report along with definitions of the indicators and guidelines to interpretation.

### **Bibliometrics**

- Section 4 presents analyses of IMI project publications overall, including trends in publications, frequently used journals, and top research fields. Where possible IMI research is benchmarked to EU-28 biomedical research.<sup>1</sup>
- Section 5 presents citation analyses of IMI publications at the call level, examining the citation impact and outputs of individual project. Where possible the IMI projects are benchmarked to world output and overall IMI output.
- Section 6 presents geographic clusters where IMI research activity occurs, including bibliometric data, the constituent institutions and top five journal subject categories within the clusters.

### **Collaboration**

- Section 7 presents collaboration analyses for IMI publications overall and at the project level, examining collaboration between different sectors, institutions, and countries.

### **Benchmarking**

- Section 8 presents analysis of IMI publications, benchmarked to similar funding organisations. The organisations are: Commonwealth Scientific and Industrial Research Organisation (CSIRO), Critical Path Institute (C-Path), Foundation for the National Institutes of Health (FNIH), Grand

---

<sup>1</sup> At time of publication, September 2022, the United Kingdom has left the European Union, however to date there has not been any large changes to the United Kingdom's participation in Horizon 2020 funded research therefore the United Kingdom is still included in the EU-28.

## 3 DATA SOURCES, INDICATORS AND INTERPRETATION

### 3.1 Bibliometrics and citation analysis

Research evaluation increasingly uses bibliometric data and analyses to assess performance. Bibliometrics is the analysis of data derived from publications and their citations. Publication of research outcomes is an integral part of the research process and is a universal activity. Consequently, bibliometric data have a currency across subjects, time and location that is found in few other sources of research-relevant data. The use of bibliometric analysis, allied to informed review by experts, increases the objectivity of, and confidence in, evaluation.

Research publications accumulate citation counts when they are referred to by more recent publications. Citations to prior work are a normal part of publication and reflect the value placed on a work by later researchers. Some papers get cited frequently and many remain uncited. Highly cited work is recognised as having a greater impact and Clarivate has shown that high citation rates are correlated with other qualitative evaluations of research performance, such as peer review.<sup>2</sup> This relationship holds across most science and technology areas and, to a limited extent, in social sciences and even in some humanities subjects.

Indicators derived from publication and citation data should always be used with caution. Some fields publish at faster rates than others and citation rates also vary. Citation counts must be carefully normalised to account for such variations by field. Because citation counts naturally grow over time, it is essential to account for growth by year. Normalisation is usually done by reference to the relevant global average for the field and for the year of publication.

Bibliometric indicators have been found to be more informative for core natural sciences, especially for basic science, than they are for applied and professional areas and for social sciences. In professional areas the range of publication modes used by leading researchers is likely to be diverse as they target a diverse, non-academic audience. In social sciences there is also a diversity of publication modes and citation rates are typically much lower than in natural sciences.

Bibliometrics work best with large data samples. As the data are disaggregated, so the relationship weakens. Average indicator values (e.g., of citation impact) for small numbers of publications can be skewed by single outlier values. At a finer scale, when analysing the specific outcome for individual departments, the statistical relationship is rarely a sufficient guide by itself. For this reason, bibliometrics are best used in support of, but not as a substitute for, expert decision processes. Well-founded analyses can enable conclusions to be reached more rapidly and with greater certainty and are therefore an aid to management and to increased confidence among stakeholders, but they cannot substitute for review by well-informed and experienced peers.

---

<sup>2</sup> Evidence Ltd. (2002) *Maintaining Research Excellence and Volume: A report by Evidence Ltd to the Higher Education Funding Councils for England, Scotland and Wales and to Universities United Kingdom (UK)*. (Adams J, et al.) 48pp.

## 3.2 Data source

For the bibliometric analysis, data will be sourced from the databases underlying the Clarivate **Web of Science**, which gives access to conference proceedings, patents, websites, and chemical structures, compounds and reactions in addition to journals. It has a unified structure that integrates all data and search terms together and therefore provides a level of comparability not found in other databases. It is widely acknowledged to be the world's leading source of citation and bibliometric data.

The **Web of Science Core Collection** is part of the Web of Science and focuses on research published in journals and conferences in science, medicine, arts, humanities, and social sciences. The authoritative, multidisciplinary content covers over 34,000 of the highest impact journals worldwide, including open access and over 205,000 conference proceedings. Coverage is both current and retrospective in the sciences, social sciences, arts, and humanities, in some cases back to 1900. Within the research community, these data are often still referred to by the acronym 'ISI'.<sup>3</sup> Clarivate has extensive experience with databases on research inputs, activity and outputs and has developed innovative analytical approaches for benchmarking and interpreting international, national, and institutional research impact.

## 3.3 Methodology

**Publications:** Many different document types are indexed in the Web of Science, including editorials, meeting abstracts, book reviews as well as research journal articles and reviews. In this report all documents regardless of type are referred to as 'publications'.

**Article:** Reports of research on original works. Includes research papers, features, brief communications, case reports, technical notes, chronology, and full papers that were published in a journal and/or presented at a symposium or conference.

**Review:** A renewed study of material previously studied. Includes review articles and surveys of previously published literature. Usually will not present any new information on a subject.

**Papers:** The terms 'paper' and 'publication' are often used interchangeably to refer to printed and electronic outputs of many types. However, in this report the term 'paper' is used exclusively to refer to articles and reviews - a subset of 'publications' that excludes all other document types.

Articles and reviews are the main way researchers communicate their results to the wider community and standards in methodology and interpretation are ensured by pre-publication peer-review by experts in the same field. Therefore, citation data for papers is the most informative for bibliometric evaluations and only citations to papers are used in calculations of the citation impact indicators presented in this report.

**Citations:** Papers mention earlier papers to acknowledge their intellectual contribution to a field of research. A paper receives a citation when it is mentioned or cited by another, usually more recent paper.

---

<sup>3</sup> The origins of citation analysis as a tool that could be applied to research performance can be traced to the mid-1950s, when Eugene Garfield proposed the concept of citation indexing and introduced the Science Citation Index, the Social Sciences Citation Index and the Arts & Humanities Citation Index, produced by the Institute of Scientific Information – ISI (now Clarivate).

**Citation count:** The number of citations received by a paper since it was published reflects the impact it has had on later research. Not all citations are necessarily recorded as not all the citing papers are indexed in the Web of Science. The material indexed by Clarivate, however, is estimated to attract about 95% of global citations.

**Citation impact:** Citations per paper is an index of academic or research impact (as compared with economic or social impact). For a single paper, raw citation impact is the same as its citation count. For a set of papers, it is calculated by dividing the sum of citations by the total number of papers in any given dataset. Impact can be calculated for papers within a specific research field such as Clinical Neurology, or for a specific institution or group of institutions, or a specific country.

Citation count declines in the most recent years of any time-period as papers have had less time to accumulate citations (papers published in 2007 will typically have more citations than papers published in 2010).

**Field-normalised citation impact:** Broadly the field-normalised citation impact compares the citation impact of a paper or set of papers to the average citation impact of all similar papers published worldwide in the same field and year.

As citation rates vary between research fields and with time, analyses must take both field and year into account. In addition, the type of publication will influence the citation count. For this reason, only citation counts of papers (as defined above) are used in calculations of citation impact. The standard normalisation factor is the world average citations per paper for the year and journal category in which the paper was published.

As field-normalised citation impact is normalised to global averages the performance of papers in different fields can be directly compared as the world average always equals 1.00. Therefore, a field-normalised citation impact exceeding 1.00 indicates papers have received more citations than the world average, conversely a value below 1.00 suggests papers are underperforming. See page 113 for a worked example of how field-normalised citation impact is calculated.

**Highly Cited Papers:** Highly cited papers are papers that are recognized as having a greater impact than other papers published in a similar year and field. For a paper to be considered highly cited they must be in the Top 10% in terms of citation frequency, considering the field and year of publication. High citation rates have shown to be correlated with other qualitative research performance evaluations, such as peer reviews.

**Web of Science journal categories or Clarivate InCites: Essential Science Indicators<sup>SM</sup> fields:** Standard bibliometric methodology uses journal category or ESI fields as a proxy for research fields. ESI fields aggregate data at a higher level than the journal categories – there are only 22 ESI research fields compared to 254 journal categories.<sup>4</sup> Journals are assigned to one or more categories, and every article within that journal is subsequently assigned to that category. Papers from prestigious, ‘multidisciplinary’ and general medical journals such as *Nature*, *Science*, *The Lancet*, *The BMJ*, *The New England Journal of Medicine* and the *Proceedings of the National Academy of Sciences* (PNAS) are assigned to specific categories based on the journal categories of

---

<sup>4</sup> Essential Science Indicators are defined by a unique grouping of journals with no journal being assigned to more than one field. These fields are focussed on the science, technology, engineering and medicine subjects and arts & humanities subjects are excluded. Customised analyses, however, can be designed to include these as an additional category.

the references cited in the article. The selection procedures for the journals included in the citation databases are documented here <http://mjl.clarivate.com/>.

**Journal-normalised citation impact:** Broadly the journal-normalised citation impact compares a paper or set of papers citation impact to all the other papers published in the same journal in the same year.

It is another bibliometric indicator which can be very useful in small datasets. This indicator is calculated from the citation impact relative to the specific journal in which the paper is published. For example, a paper published in the journal *Acta Biomaterialia* in 2005 that has been cited 189 times, would have an expected citation rate of 49.57 (the average number of citations per paper for this journal and publication year) and hence a journal-normalised citation impact of 6.3. This paper, therefore, has been cited more than expected for the journal.

Like the field-normalised citation impact a value exceeding 1.00 indicates that a paper or set of papers is receiving more citations than other papers in the same journal, and a value less than 1.00 indicates that a paper or set of papers is underperforming, receiving fewer citations than papers in the same journal.

**Open access publication:** Open access publications are publications that are made available online, at no cost to the reader. The Web of Science open access data come from the Directory of Open Access Journals (DOAJ) and collaborations with Impact Story and Our Research's Unpaywall services. The Web of Science therefore provides unrivalled coverage of open access publications that are published through DOAJ Gold, Other Gold, Green Published, Green Accepted or Bronze routes.

It is also possible that some publishers make publications available without following a recognised open access route. In these cases publications will not be indexed as open access in the Web of Science. Additionally, the analysis presented in this report covers all document types and not just papers, and some of these are not indexed as open access in the Web of Science databases.

The Web of Science open access data coverage is summarised at:  
[clarivate.com/webofsciencegroup/solutions/open-access/](http://clarivate.com/webofsciencegroup/solutions/open-access/)

### 3.4 Data collation

This analysis used a dataset comprising publications arising from IMI-supported projects. These publications were identified using grant acknowledgments, title, and abstract text searches, as well as other parameters developed in conjunction with IMI staff. There are currently 187 IMI projects. IMI staff validated the publications identified by this process and the list of projects to be analysed was provided by IMI staff.

## 4 CITATION ANALYSIS – IMI SUPPORTED PUBLICATIONS OVERALL

This section analyses the volume and citation impact of publications arising from IMI-supported projects, and where possible, benchmarks this against similar European research funders.

The datasets analysed in this, the thirteenth report, include IMI-supported publications identified in Clarivate Web of Science up to 31<sup>st</sup> December 2021. The census point for inclusion of publications into the twelfth report was 31<sup>st</sup> December 2020. Therefore, this report reflects changes in IMI activity between these points. Citations to these publications were counts up to 31<sup>st</sup> December 2021. Unless otherwise specified metrics are for all IMI-supported documents from all calls in IMI 1 and IMI 2, in aggregate.

When considering the analyses in this section, earlier caveats regarding paper numbers should be borne in mind ([Section 3](#)).

### 4.1 Publications from IMI-supported projects

Publications from IMI-supported projects were identified using bibliographic data supplied by IMI, and through specific keyword searches using funding acknowledgment data in the Web of Science. The process of identifying publications from IMI-supported projects that have Clarivate citation data is outlined in Figure 4.1.1.

The IMI project dataset started with 7,177 publications which were previously identified as IMI publications and used as the IMI publication dataset in the previous report. Separately, 1,630 new publications were identified as IMI-associated through keyword searches of funding acknowledgement text in databases which underlie Clarivate Web of Science. The combination of these two datasets led to a total of 8,807 unique publication records associated with IMI-supported projects. Of these 8,807 publications, 198 were eliminated as they were either published in 2022 or could not be distinguished as IMI from a manual review of the dataset. Therefore, 8,609 Web of Science publications remained.

The citation counts for this report were sourced from the citation databases which underlie Clarivate Web of Science and were extracted in June 2022. Normalised bibliometric indicators were calculated using standard methodology and the Clarivate National Science Indicators (NSI) database for 2021.

Figure 4.1.1 Process for IDENTIFYING PUBLICATIONS FROM IMI-SUPPORTED PROJECTS, 2010-2021

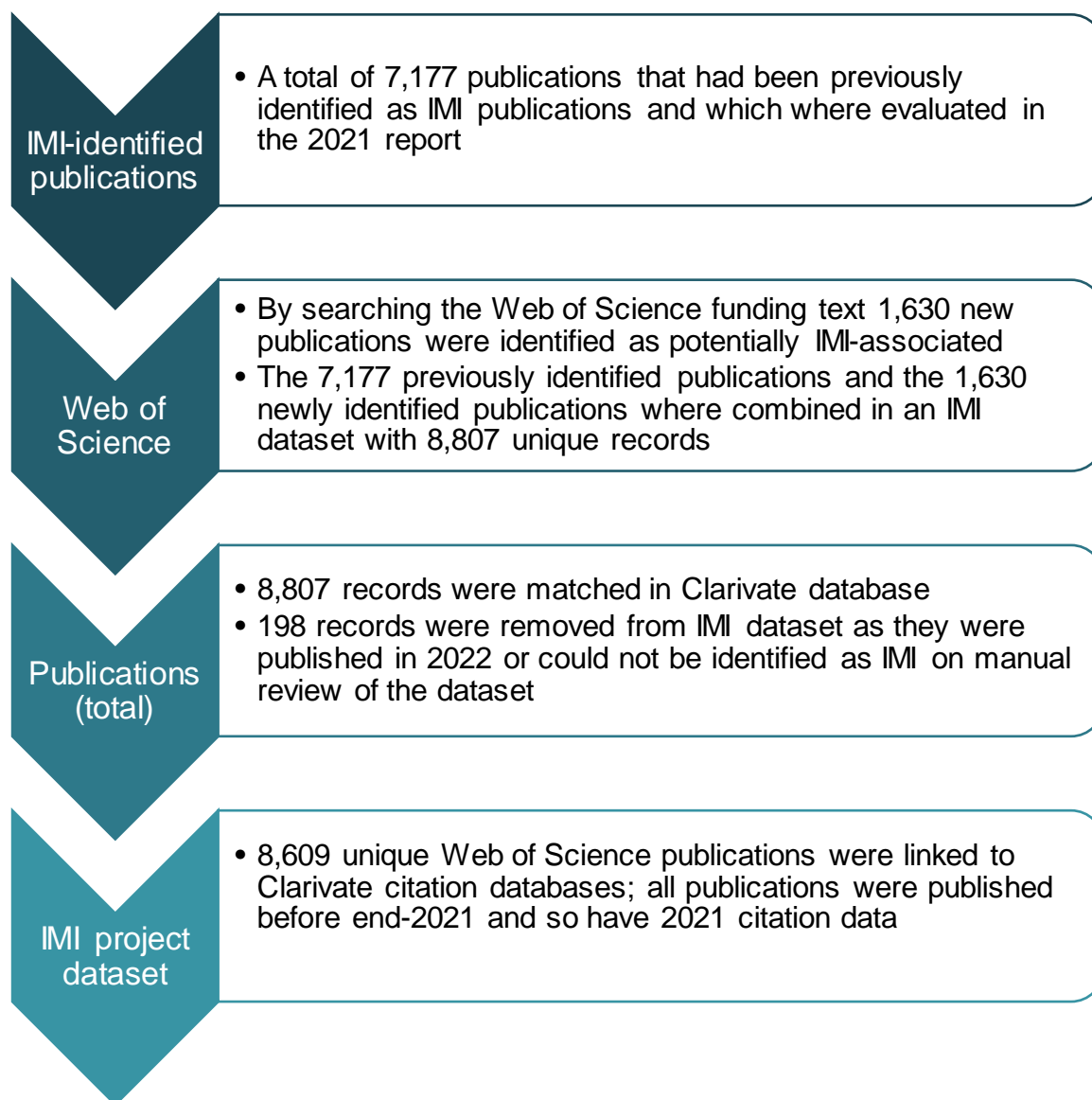


Table 4.1.1 NUMBER OF PUBLICATIONS FROM IMI PROJECTS, 2010-2021

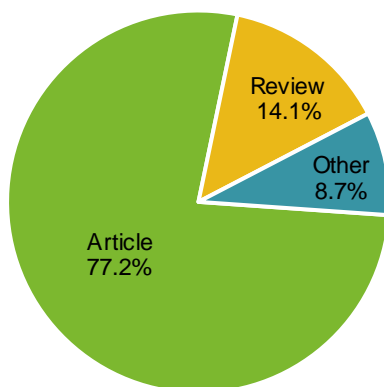
	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS
All IMI	8,609	7,856
IMI 1	6,323	5,896
IMI 2	2,382	2,070

Note that some publications belong to IMI 1 and IMI 2, therefore the total number of publications shown for All IMI is smaller than the sum of publications shown for IMI 1 and IMI 2.

## 4.2 Publications from IMI projects by document type

Figure 4.2.1 shows the percentage of Web of Science publications by document type and the same data is shown in Table 4.2.1.

Figure 4.2.2 PERCENTAGE OF IMI PROJECT PUBLICATIONS BY DOCUMENT TYPE, 2010-2021



Articles + Reviews = Papers, 91.3%

- IMI project research resulted in 8,609 unique Web of Science publications.
- Of these publications, 91.3% were articles (77.2%) and reviews (14.1%) which are collectively referred to as 'papers' in this report.
- A further 753 publications (8.7%) were not papers. These 'other' publications are composed of 174 editorials, 378 meeting abstracts, 87 proceeding papers, 92 letters, 15 corrections and three news items and four data papers.

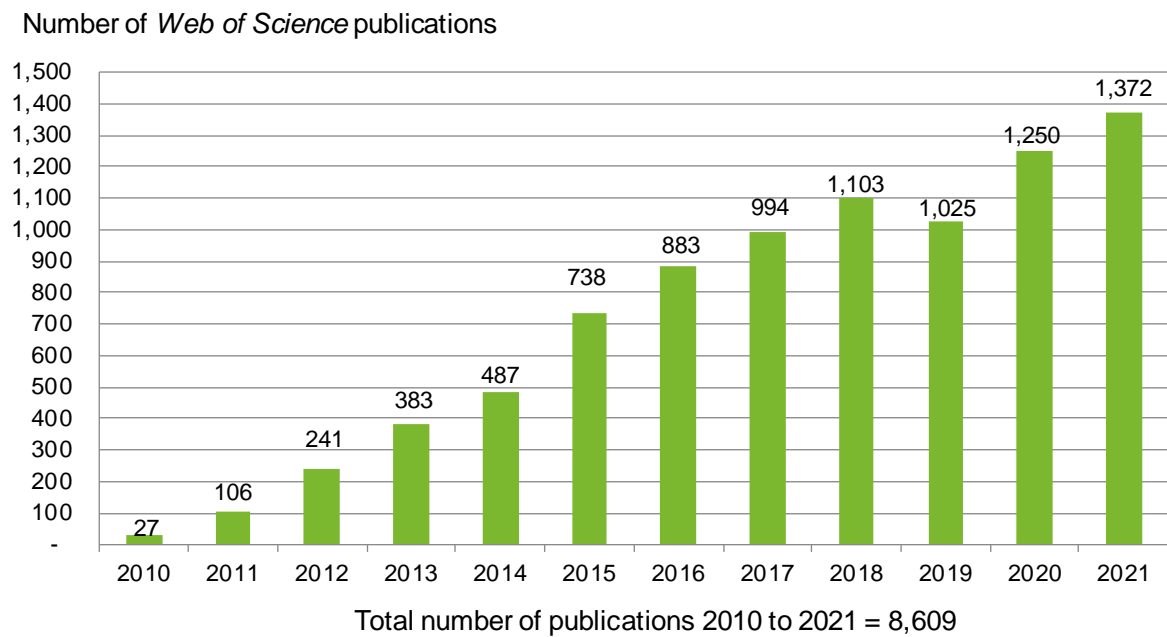
Table 4.2.1 NUMBER AND PERCENTAGE OF IMI PROJECT PUBLICATIONS BY DOCUMENT TYPE, 2010-2021

	DOCUMENT TYPE	NUMBER OF PUBLICATIONS	% OF IMI PUBLICATIONS
Papers	Article	6,643	77.16%
	Review	1,213	14.09%
Other document types	Meeting Abstract	378	4.39%
	Editorial Material	174	2.02%
	Letter	92	1.07%
	Proceedings Paper	87	1.01%
	Correction	15	0.17%
	Data Paper	4	0.05%
	News Item	3	0.03%



### 4.3 Trends in publication output

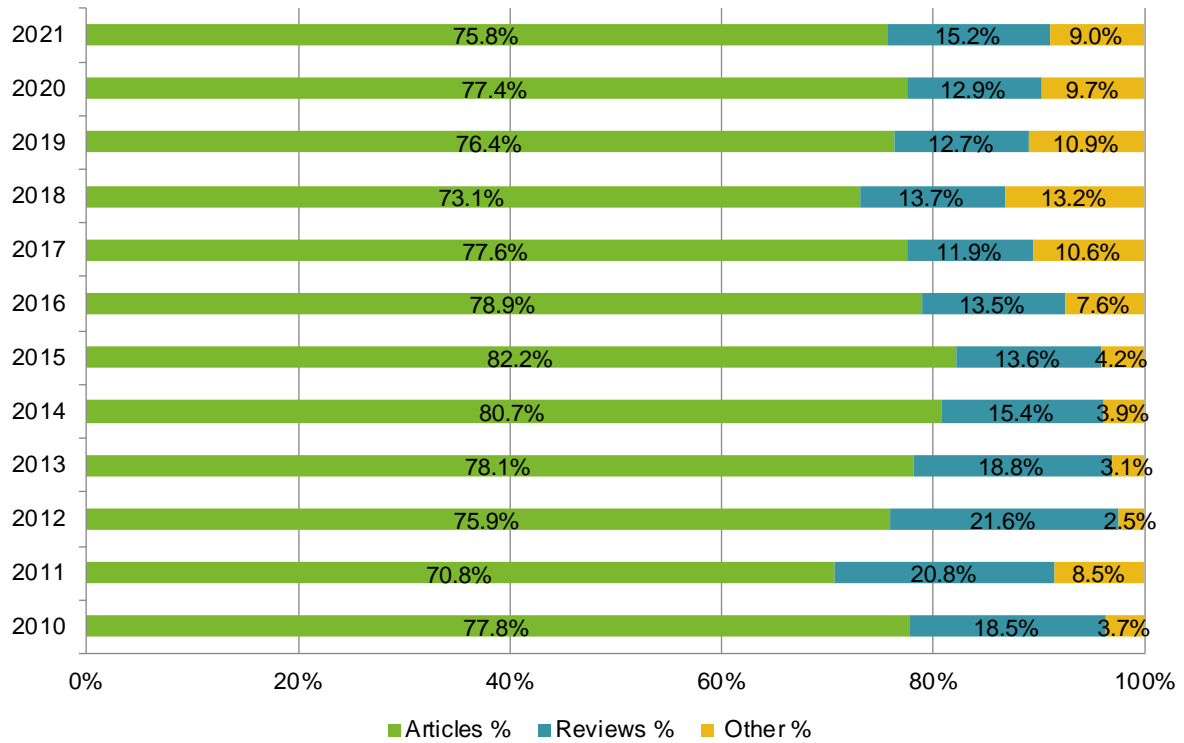
Figure 4.3.3 NUMBER OF WEB OF SCIENCE PUBLICATIONS FOR IMI PROJECTS BY YEAR, 2010-2021



- IMI project research output continued to increase in 2021, however at a slower pace with a change of nearly 10%. The stabilization of the rate of growth is expected as IMI programme matures.

Figure 4.3.2 shows the proportion of papers (articles and reviews) relative to other document types for IMI project research between 2010 and 2021.

Figure 4.3.4 PERCENTAGE OF IMI PROJECT PUBLICATIONS EACH YEAR BY DOCUMENT TYPE, 2010-2021

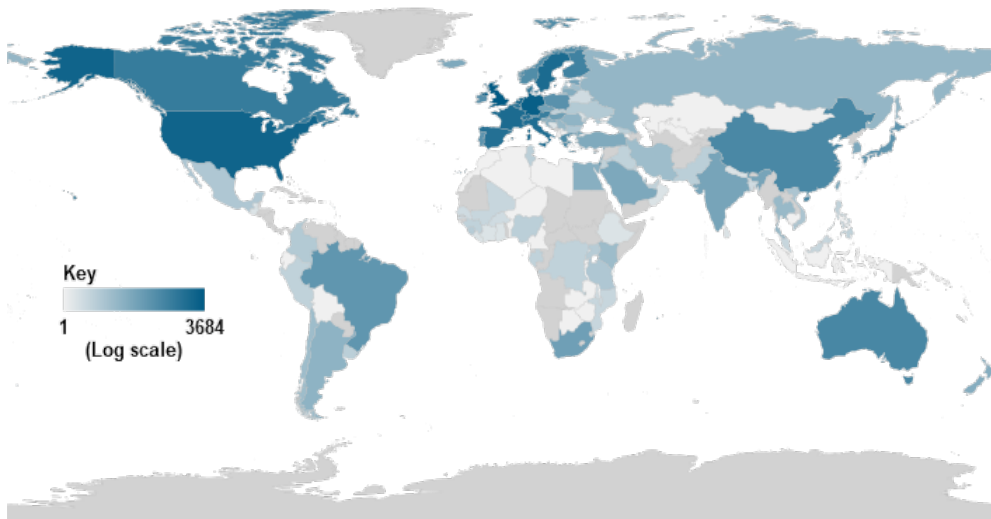


- IMI project research continued to generate a high proportion of papers relative to other document types. Articles accounted for around 75.8% of all publications in 2021, consistent with recent years.

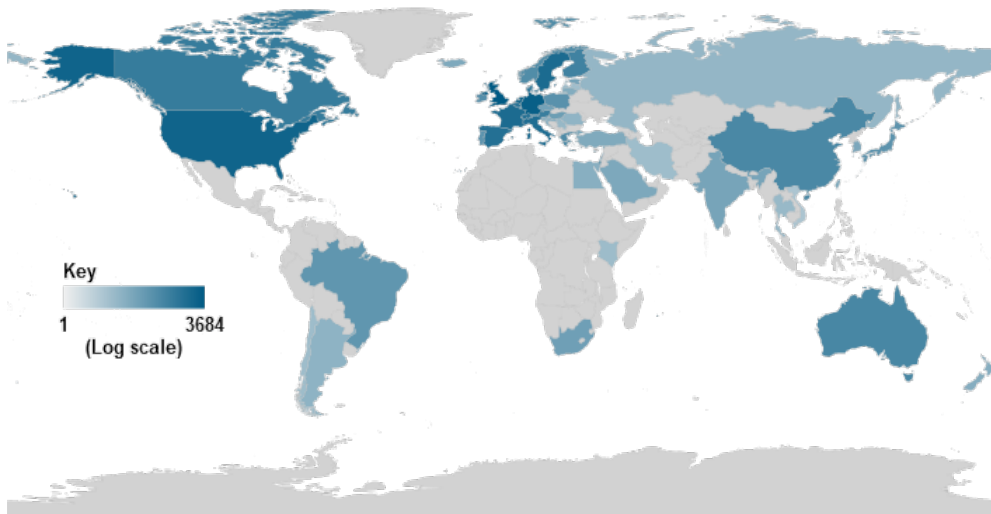
#### 4.4 Publication output by country

Figure 4.4.1 shows a map highlighting all countries with one or more publication from IMI projects between 2010 and 2021. Figure 4.4.2 shows a map highlighting all countries with at least ten Web of Science publications from IMI projects between 2010 and 2021. Table 4.4.1 and Figure 4.4.3 shows the corresponding data; the total number of publications for the 20 and 10 countries respectively with the highest number publications from IMI projects between 2010 and 2021. A full list of all countries output of publications is included in [Annex 3](#).

*Figure 4.4.5 MAP OF COUNTRIES WHICH HAVE AT LEAST ONE WEB OF SCIENCE PUBLICATION FOR IMI PROJECTS, 2010-2021*



*Figure 4.4.6 MAP OF COUNTRIES WHICH HAVE AT LEAST TEN WEB OF SCIENCE PUBLICATION FOR IMI PROJECTS, 2010-2021*



- In total 122 countries have at least one IMI publication and 55 countries have at least ten IMI publications.

Figure 4.4.7 TEN COUNTRIES WITH THE MOST IMI PROJECT PUBLICATIONS. [ANNEX 3](#) LISTS ALL COUNTRIES WITH AT LEAST ONE IMI PROJECT PUBLICATION, 2010-2021

Number of *Web of Science* publications

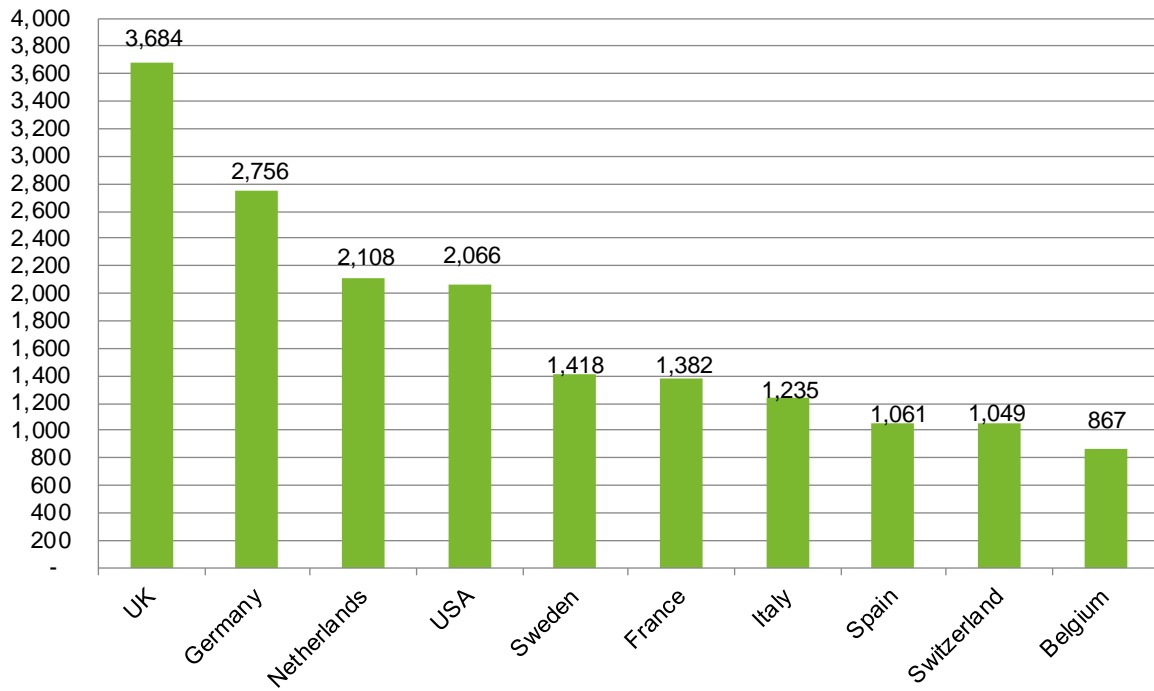


Table 4.4.2 TWENTY COUNTRIES WITH THE MOST IMI PROJECT PUBLICATIONS. [ANNEX 3](#) LISTS ALL COUNTRIES WITH AT LEAST ONE IMI PROJECT PUBLICATION, 2010-2021

COUNTRY	NUMBER OF PUBLICATIONS
UK	3,684
Germany	2,756
Netherlands	2,108
USA	2,066
Sweden	1,418
France	1,382
Italy	1,235
Spain	1,061
Switzerland	1,049
Belgium	867
Denmark	633
Canada	568
Austria	508
Finland	402
Australia	313
China	286
Greece	249
Norway	225
Ireland	203
Poland	172

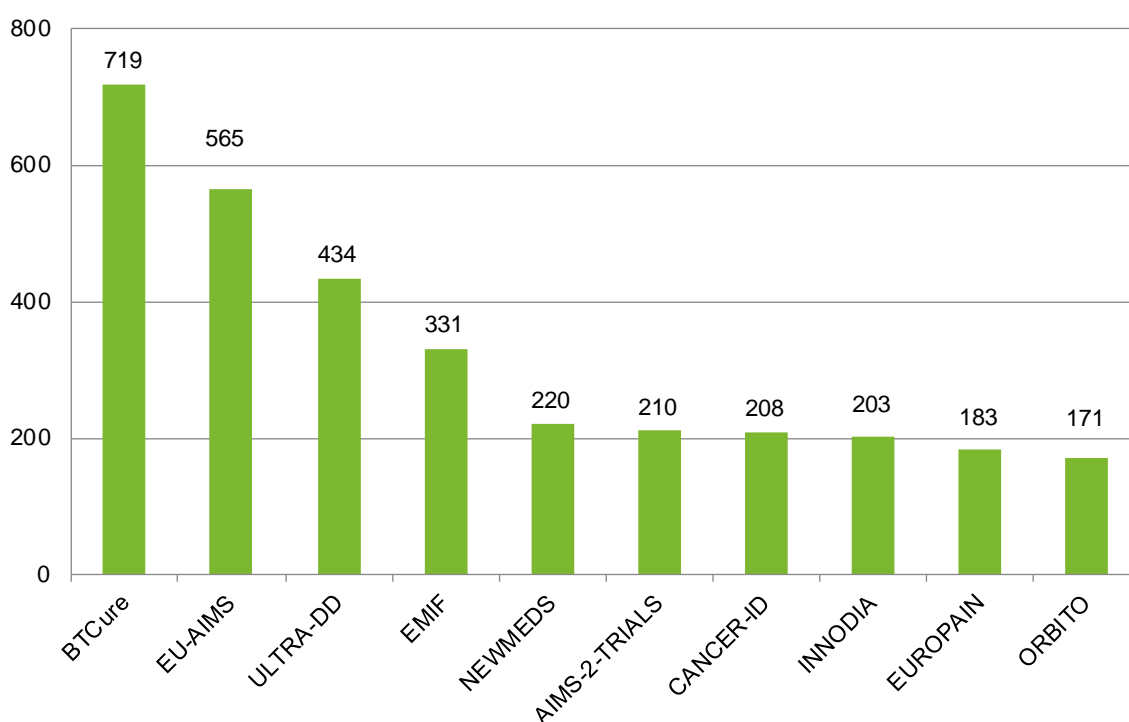
- Researchers affiliated to the United Kingdom authored the most IMI project publications (3,684 publications).
- Other EU-28 countries were among the countries with the highest output. The most productive exceptions are the USA (2,066 publications) and Switzerland (1,049 publications).

## 4.5 Publication output by IMI project

Figure 4.5.1 shows the ten IMI projects with the highest output of publications between 2010 and 2021. Table 4.5.1 expands upon Figure 4.5.1, listing the 20 IMI projects with the most publications, including the number and percentage of open access papers and the number of papers between 2010 and 2021. A full list of projects and the number of associated publications is presented in [Annex 4](#).

Figure 4.5.8 NUMBER OF WEB OF SCIENCE PUBLICATIONS FOR TEN IMI PROJECTS WITH THE HIGHEST OUTPUT OF PUBLICATIONS, 2010-2021

Number of *Web of Science* publications



- BTCure remains the most productive IMI project in terms of number of publications (719 publications) and the second most productive project is still EU-AIMS (565 publications).
- AIMS-2-TRIALS project continued its rapid growth nearly doubling the number of publications from last year and is now 6th in the top 10 projects with highest output of publications (210 publications), displacing Translocation (164 publications)
- AIMS-2-TRIALS and INNODIA are the only two projects from IMI phase 2 in the Top 10 projects by highest publication output.

Table 4.5.3 TWENTY IMI PROJECTS WITH THE MOST PUBLICATIONS, THE NUMBER OF PAPERS, NUMBER AND PERCENTAGE OF OPEN ACCESS PAPERS, 2010-2021.

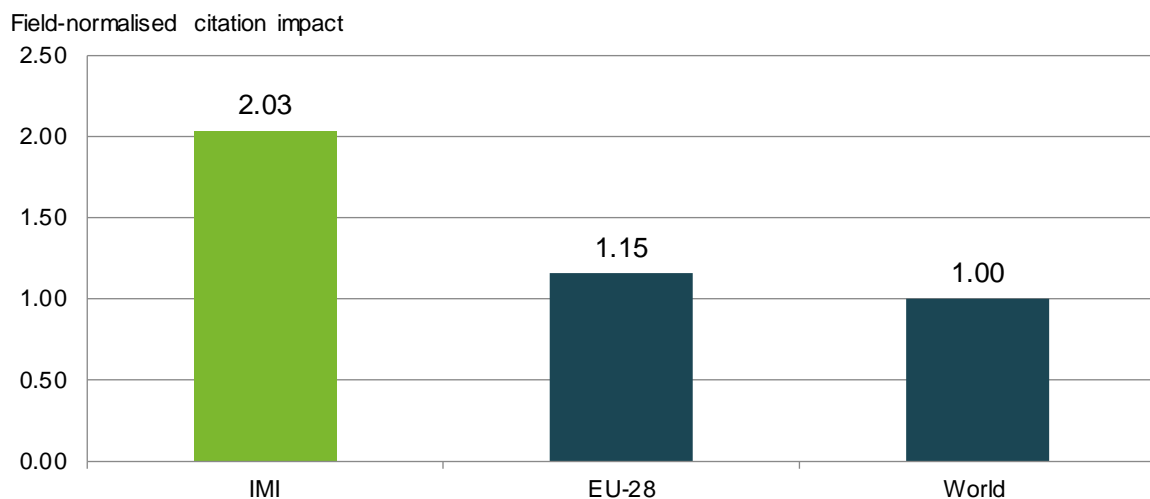
PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
<b>BTCure</b>	719	672	487	72.5%
<b>EU-AIMS</b>	565	546	453	83.0%
<b>ULTRA-DD</b>	434	425	362	85.2%
<b>EMIF</b>	331	310	261	84.2%
<b>NEWMEDS</b>	220	214	124	57.9%
<b>AIMS-2-TRIALS</b>	210	196	182	92.9%
<b>CANCER-ID</b>	208	180	132	73.3%
<b>INNODIA</b>	203	166	146	88.0%
<b>EUROPAIN</b>	183	181	73	40.3%
<b>ORBITO</b>	171	168	61	36.3%
<b>TRANSLOCATION</b>	164	164	110	67.1%
<b>BigData@Heart</b>	157	135	125	92.6%
<b>STEMBANCC</b>	153	147	120	81.6%
<b>IMIDIA</b>	151	141	121	85.8%
<b>U-BIOPRED</b>	148	93	68	73.1%
<b>RTCure</b>	146	131	112	85.5%
<b>SUMMIT</b>	141	136	106	77.9%
<b>ELF</b>	135	134	108	80.6%
<b>CHEM21</b>	131	128	64	50.0%
<b>PreDiCT-TB</b>	124	118	109	92.4%

## 4.6 Is IMI project research well cited?

The number of citations a paper receives (also known as its raw citation impact) is at least partly determined by the field to which it relates and the year in which it was published. Typically, papers published in disciplines such as biomedical research receive more citations than papers published in subjects such as engineering, and older papers tend to have higher citations counts on average than newer ones because they have had longer to accrue them. Therefore, citation impact is usually normalised to the relevant world average to allow comparison between years and fields; the resulting indicator is called the field-normalised citation impact.

Figure 4.6.1 shows the average field-normalised citation impact for all IMI papers compared to the average for EU-28 papers in relevant biomedical journal categories (see [Annex 2](#)) and all global papers published between 2010 and 2021. Table 4.6.1 and Table 4.6.2 present average citation impact indicators for all IMI papers.

*Figure 4.6.9 FIELD-NORMALISED CITATION IMPACT FOR IMI SUPPORTED RESEARCH PAPERS COMPARED TO THE AVERAGE FOR EU-28 AND WORLD PAPERS, 2010-2021*



- IMI's field-normalised citation impact remains twice that of the world average and is 2% higher than last year's report and 77% higher than the EU-28.



Table 4.6.4 SUMMARY CITATION ANALYSIS FOR IMI SUPPORTED RESEARCH PAPERS, 2010-2021

	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
<b>IMI projects</b>	7,856	2.03	1.23	35.0	25.3%
<b>IMI 1</b>	5,896	1.95	1.18	32.9	25.1%
<b>IMI 2</b>	2,070	2.19	1.34	41.6	25.6%

Table 4.6.5 SUMMARY OF IMI SUPPORTED RESEARCH PUBLICATIONS, 2010-2021

	NUMBER OF PUBLICATIONS	% OF OPEN ACCESS PAPERS*	NUMBER OF PAPERS	CITATIONS	RAW CITATION IMPACT
<b>IMI Projects</b>	8,609	77.2%	7,856	246,617	31.39
<b>IMI 1</b>	6,323	73.3%	5,896	218,147	37.00
<b>IMI 2</b>	2,382	89.2%	2,070	26,099	12.61

## Summary of key findings

- The field-normalised citation impact of IMI project papers was 2.03 for the twelve-year period, 2010-2021, 2% higher than last year's report and double the World average.
- The field-normalised citation impact of IMI project papers was 76% higher than the EU's average citation impact (1.15)<sup>5</sup> between 2010 and 2021, in similar biomedical journal categories.
- More than a quarter (25.3%) of IMI papers were highly cited, that is they were in the world's top 10% of most highly cited papers in the relevant journal category and year of publication.
- IMI 2 has a higher percentage of open access papers compared with IMI 1. This is likely due to the stipulation that IMI 2 funded researcher should publish open access articles.<sup>6</sup>

<sup>5</sup> EU-28 grouping of countries: Clarivate National Science Indicators 2021 database; similar research has been defined as biomedical journal categories listed in [Annex 2](#).

<sup>6</sup> Note that IMI 2 funded researchers are contractually obliged to make their scientific articles open access through Green or Gold routes. However, for some of other document types, such as editorials, reviews or conference proceedings open access publication is strongly encouraged but not mandatory.

## 4.7 In which journals do IMI project publications appear most frequently?

The 20 journals in which IMI project publications appeared most frequently (ranked by number of IMI publications) between 2010 and 2021, are listed in Table 4.7.1. Together, the 20 most frequently used journals account for 1,745 publications, 20.3% of IMI's publications

IMI project publications appeared most frequently in *Scientific Reports* which IMI published 195 publications. This was followed by *Plos One* where they published 193 publications. For most journals, papers (articles and reviews) were the most frequent publication type, however large collections of meeting abstracts were published in *European Respiratory Journal* (28 meeting abstracts).

IMI had a strong focus within Multidisciplinary Sciences and Pharmacology where four of the top 20 journals were assigned to each subject category. Followed by Rheumatology and Neurosciences which each had 3 journal titles in the top 20 titles.

Of the 20 most frequently used journals, nearly two thirds were in the top quartile (Q1) by Journal Impact Factor (JIF) while the rest were in the second quartile (Q2) ranked against other journals in the same category.

Overall, IMI project publications were published in a total of 1,555 journals. The average Journal Impact Factor for all IMI project publications is 7.53, a slight increase of 0.60 compared to the previous year.

The 20 highest Journal Impact Factor journals in which IMI project research was published are listed in Table 4.7.2. The journal with the highest Impact Factor is *Lancet*, with a Journal Impact Factor of 202.73 where IMI published 6 publications, four of which are papers. This is followed by *New England Journal of Medicine* with an Impact Factor of 176.08 where IMI published 1 publication. Of the top 20 journals by Impact Factor, IMI published the most publications (26) in *Nature* which has an Impact Factor of 69.50. IMI published a total of 132 publications in these top ranked journals by journal impact factor.

The 20 open access journals in which IMI projects publish most frequently (ranked by number of publications), are listed in Table 4.7.3. Of the top 20 open access journals, IMI published most frequently in *Scientific Reports* (195 publications) and the Journal with the highest Impact Factor was the *Annals of the Rheumatic Diseases*. 13 of these journals are ranked in the top quartile in their relevant journal categories, lower than last year's report of 16.

---

Nevertheless, it is obvious that fewer than all of IMI's publications are classified as open access in this analysis, and this is likely to be due to ancillary factors (such as challenges relating to definitions and coverage) as well as non-compliance. The Web of Science open access data come from the Directory of Open Access Journals (DOAJ) and collaborations with Impact Story and Our Research's Unpaywall services. The Web of Science therefore provides unrivalled coverage of open access publications that are published through DOAJ Gold, Other Gold, Green Published, Green Accepted or Bronze routes.

It is also possible that some publishers make publications available without following a recognised open access route. In these cases, publications will not be indexed as open access in the Web of Science or in this report

The Web of Science open access data coverage is summarised at: <https://clarivate.com/webofsciencgroup/solutions/open-access/>

Table 4.7.6 JOURNALS IN WHICH IMI PROJECT PUBLICATIONS WERE PUBLISHED MOST FREQUENTLY, TOP 20 RANKED BY NUMBER OF IMI PUBLICATIONS, 2010-2021

JOURNAL	NUMBER OF IMI PUBLICATIONS	NUMBER OF IMI PAPERS	JOURNAL IMPACT FACTOR (2021)	WEB OF SCIENCE JOURNAL CATEGORIES	QUARTILE
Scientific Reports	195	195	5.00	Multidisciplinary Sciences	Q2
Plos One	193	193	3.75	Multidisciplinary Sciences	Q2
Annals of the Rheumatic Diseases	191	126	27.97	Rheumatology	Q1
Diabetologia	150	79	10.46	Endocrinology & Metabolism	Q1
Nature Communications	117	116	17.69	Multidisciplinary Sciences	Q1
Frontiers In Immunology	96	95	8.79	Immunology	Q1
Journal Of Medicinal Chemistry	76	76	8.04	Chemistry, Medicinal	Q1
Diabetes	73	51	9.34	Endocrinology & Metabolism	Q1
Arthritis Research & Therapy	69	69	5.61	Rheumatology	Q1
Journal of Alzheimers Disease	67	66	4.16	Neurosciences	Q2
Arthritis & Rheumatology	64	55	15.48	Rheumatology	Q1
European Respiratory Journal	58	21	33.80	Respiratory System	Q1
Pain	57	55	7.93	Anesthesiology; Clinical Neurology; Neurosciences	Q1
International Journal of Molecular Sciences	54	54	6.21	Biochemistry & Molecular Biology; Chemistry, Multidisciplinary	Q2
European Journal of Pharmaceutics and Biopharmaceutics	50	50	5.59	Pharmacology & Pharmacy	Q1
European Journal of Pharmaceutical Sciences	48	46	5.11	Pharmacology & Pharmacy	Q2
Journal Of Antimicrobial Chemotherapy	48	47	5.76	Infectious Diseases; Microbiology; Pharmacology & Pharmacy	Q2
Translational Psychiatry	48	48	7.99	Psychiatry	Q1
Proceedings of the National Academy of Sciences of The United States of America	47	47	12.78	Multidisciplinary Sciences	Q1
Psychopharmacology	44	44	4.42	Neurosciences; Pharmacology & Pharmacy; Psychiatry	Q2

Table 4.7.7 JOURNALS IN WHICH IMI PROJECT PUBLICATIONS WERE PUBLISHED MOST FREQUENTLY, TOP 20 RANKED BY JOURNAL IMPACT FACTOR, 2010-2021

JOURNAL	NUMBER OF IMI PUBLICATIONS	NUMBER OF IMI PAPERS	JOURNAL IMPACT FACTOR (2021)	WEB OF SCIENCE JOURNAL CATEGORIES	QUARTILE
Lancet	6	4	202.73	Medicine, General & Internal	Q1
New England Journal Of Medicine	1	0	176.08	Medicine, General & Internal	Q1
Jama-Journal Of The American Medical Association	9	7	157.34	Medicine, General & Internal	Q1
Nature Reviews Molecular Cell Biology	1	1	113.92	Cell Biology	Q1
Nature Reviews Drug Discovery	15	8	112.29	Biotechnology & Applied Microbiology; Pharmacology & Pharmacy	Q1
Nature Reviews Immunology	4	2	108.56	Immunology	Q1
Lancet Respiratory Medicine	4	3	102.64	Critical Care Medicine; Respiratory System	Q1
BMJ-British Medical Journal	8	7	93.33	Medicine, General & Internal	Q1
Nature Medicine	17	16	87.24	Biochemistry & Molecular Biology; Cell Biology; Medicine, Research & Experimental	Q1
World Psychiatry	1	1	79.68	Psychiatry	Q1
Nature Reviews Microbiology	2	2	78.30	Microbiology	Q1
Lancet Psychiatry	6	4	77.06	Psychiatry	Q1
Nature Reviews Gastroenterology & Hepatology	4	3	73.08	Gastroenterology & Hepatology	Q1
Chemical Reviews	3	3	72.09	Chemistry, Multidisciplinary	Q1
Lancet Infectious Diseases	11	10	71.42	Infectious Diseases	Q1
Nature Reviews Cancer	2	2	69.80	Oncology	Q1
Nature	26	26	69.50	Multidisciplinary Sciences	Q1
Nature Biotechnology	3	1	68.16	Biotechnology & Applied Microbiology	Q1
Cell	6	6	66.85	Biochemistry & Molecular Biology; Cell Biology	Q1
Nature Reviews Disease Primers	3	3	65.04	Medicine, General & Internal	Q1

Table 4.7.8 OPEN ACCESS JOURNALS IN WHICH IMI PROJECT PUBLICATIONS WERE PUBLISHED MOST FREQUENTLY, TOP 20 RANKED BY NUMBER OF OPEN ACCESS WEB OF SCIENCE PUBLICATIONS, 2010-2021

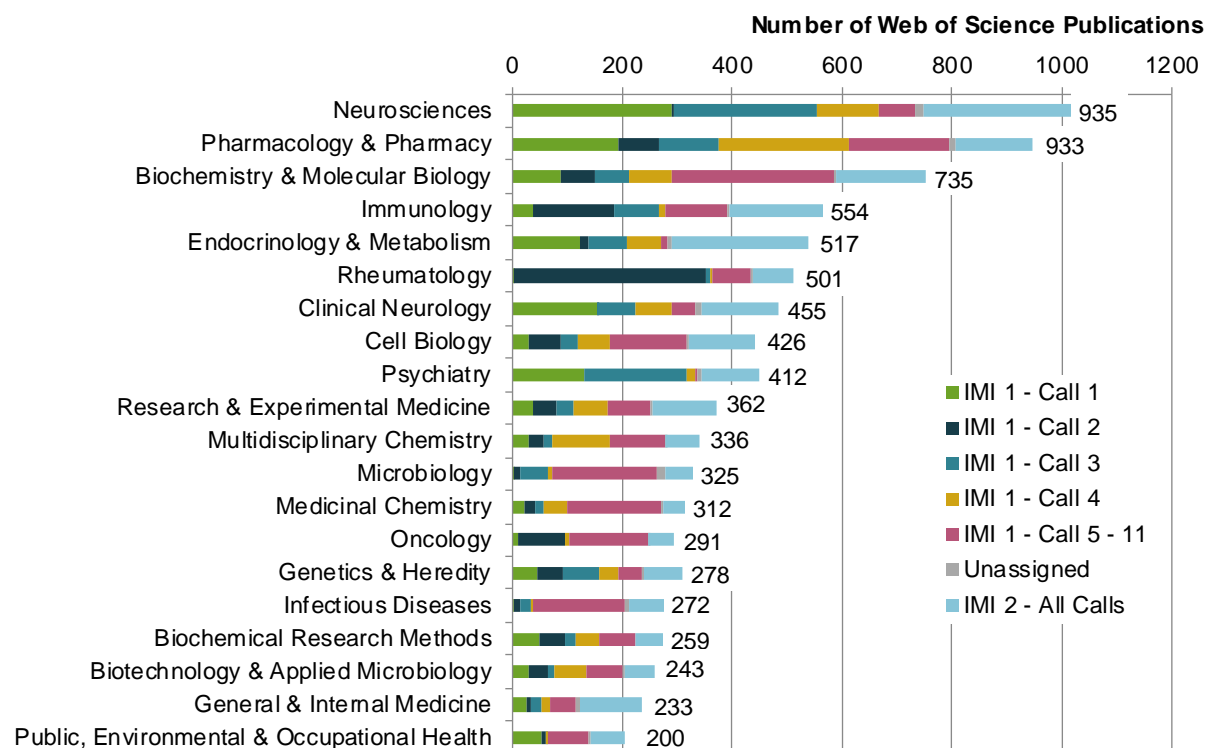
JOURNAL	NUMBER OF IMI PUBLICATIONS	NUMBER OF IMI PAPERS	JOURNAL IMPACT FACTOR (2021)	WEB OF SCIENCE JOURNAL CATEGORIES	QUARTILE
Scientific Reports	195	195	5.00	Multidisciplinary Sciences	Q2
Plos One	193	193	3.75	Multidisciplinary Sciences	Q2
Nature Communications	117	116	17.69	Multidisciplinary Sciences	Q1
Annals Of The Rheumatic Diseases	116	74	27.97	Rheumatology	Q1
Frontiers In Immunology	96	95	8.79	Immunology	Q1
Diabetologia	73	70	10.46	Endocrinology & Metabolism	Q1
Arthritis Research & Therapy	69	69	5.61	Rheumatology	Q1
International Journal Of Molecular Sciences	54	54	6.21	Biochemistry & Molecular Biology; Chemistry, Multidisciplinary	Q2
Journal Of Medicinal Chemistry	51	51	8.04	Chemistry, Medicinal	Q1
Journal Of Alzheimers Disease	51	50	4.16	Neurosciences	Q2
Translational Psychiatry	48	48	7.99	Psychiatry	Q1
Journal Of Antimicrobial Chemotherapy	46	45	5.76	Infectious Diseases; Microbiology; Pharmacology & Pharmacy	Q2
Arthritis & Rheumatology	46	45	15.48	Rheumatology	Q1
Proceedings Of The National Academy Of Sciences Of The United States Of America	45	45	12.78	Multidisciplinary Sciences	Q1
Antimicrobial Agents And Chemotherapy	43	42	5.94	Microbiology; Pharmacology & Pharmacy	Q1
BMJ Open	43	43	3.01	Medicine, General & Internal	Q2
Molecular Autism	42	41	6.48	Genetics & Heredity; Neurosciences	Q1
Diabetes	40	40	9.34	Endocrinology & Metabolism	Q1
Alzheimers Research & Therapy	40	40	8.82	Clinical Neurology; Neurosciences	Q1
Cell Reports	40	40	9.99	Cell Biology	Q1

## 4.8 Which research fields account for the highest volume of IMI project publications?

Figure 4.8.1 shows the twenty Web of Science journal categories<sup>7</sup> most frequently associated with IMI funded research between 2010 and 2021. IMI 1 calls 5-11 have a lower number of publications relative to calls 1-4 and for clarity of presentation these publications are shown as one group in Figure 4.8.1. Likewise, IMI 2 has far fewer publication compared to IMI 1 and so all IMI 2 publications are shown as one group in Figure 4.8.1. Publications that acknowledge IMI funding but do not specify a project, phase or call are classed as Unassigned. Note that some bars are longer than the total number of IMI publications in a journal category (indicated by the data labels) due to some papers being associated with multiple calls. Figure 4.8.2 shows the ten Web of Science journal categories most frequently associated with IMI 2 funded research.

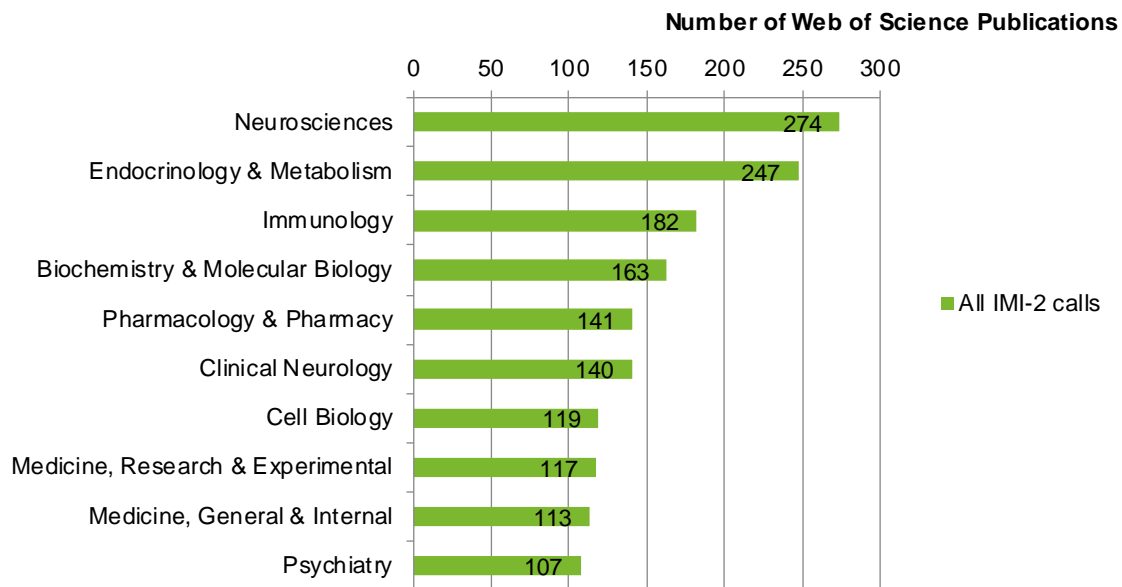
Table 4.8.1 shows the same data as Figure 4.8.1 and Figure 4.8.2 for the top twenty journal categories. It provides the number of publications assigned to each of the top twenty Web of Science journal categories in which IMI project research is published by IMI 1 calls and IMI 2 in total.

*Figure 4.8.10 TOP TWENTY WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI PROJECT RESEARCH WAS PUBLISHED MOST FREQUENTLY, 2010-2021. DATA LABELS SHOWS THE TOTAL NUMBER OF PUBLICATIONS PER JOURNAL CATEGORY*



<sup>7</sup> Journals can be associated with more than one Web of Science category.

Figure 4.8.11 TOP TEN WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI 2 PROJECT RESEARCH (ALL CALLS) WAS PUBLISHED MOST FREQUENTLY, 2010-2021. DATA LABELS SHOWS THE TOTAL NUMBER OF PUBLICATIONS PER JOURNAL CATEGORY



- IMI projects produced more publications in Neurosciences (935) than other journal categories which is a change from previous years where Pharmacology & Pharmacy (933 publications), now second, was the most productive journal category. They are followed by Biochemistry & Molecular Biology (735 publications) and Immunology (554 publications).
- Most publications in IMI 1 calls 5 to 11 belong to call 11.
- IMI 2 publications most frequently appeared in Neurosciences (274 publications) again a change from last year where Endocrinology & Metabolism journals (247 publications) was the most productive. They are followed by Immunology (182 publications) and Biochemistry & Molecular Biology.

Table 4.8.9 NUMBER OF PUBLICATIONS BY IMI 1 CALL AND IMI 2 FOR TWENTY WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI PROJECT RESEARCH WAS PUBLISHED MOST FREQUENTLY, 2010-2021. ORDERED BY TOTAL NUMBER OF PUBLICATIONS.

JOURNAL CATEGORY	NUMBER OF PUBLICATIONS BY IMI 1 CALL												IMI 2	Not assigned
	1	2	3	4	5	6	7	8	9	10	11			
Neurosciences	291	3	261	110	0	0	0	33	3	0	34	274	13	
Pharmacology & Pharmacy	194	71	110	239	14	44	9	17	50	0	71	141	12	
Biochemistry & Molecular Biology	89	61	63	77	30	44	0	36	16	1	172	163	5	
Immunology	37	146	83	12	1	9	17	16	10	35	31	182	4	
Endocrinology & Metabolism	121	19	68	63	0	0	0	1	2	0	8	247	8	
Rheumatology	2	348	11	2	0	0	1	36	0	0	34	75	2	
Clinical Neurology	154	1	68	68	0	0	0	13	0	0	30	140	11	
Cell Biology	29	59	30	61	2	6	0	23	11	1	95	119	5	
Psychiatry	132	0	185	14	0	0	1	1	1	0	4	107	5	
Research & Experimental Medicine	36	45	29	63	0	3	19	4	2	15	36	117	4	
Chemistry, Multidisciplinary	30	25	16	107	36	15	0	8	5	0	38	59	1	
Microbiology	2	12	52	5	2	90	1	11	60	8	77	56	16	
Chemistry, Medicinal	23	18	15	42	50	9	0	14	1	0	102	44	5	
Oncology	10	87	0	8	1	0	2	1	0	0	137	44	2	
Genetics & Heredity	45	47	65	36	0	2	0	10	1	0	33	69	3	
Infectious Diseases	4	8	23	2	2	60	2	7	65	10	82	71	7	
Biochemical Research Methods	48	46	20	43	2	9	0	16	1	1	40	49	0	
Biotechnology & Applied Microbiology	31	34	13	55	2	3	0	25	2	6	28	56	5	
General & Internal Medicine	25	7	20	16	0	7	2	4	24	0	20	113	8	
Public, Environmental & Occupational Health	53	7	2	4	0	12	20	0	28	1	24	65	3	



Table 4.8.2 and Table 4.8.3 show the citation impact, percentage of highly cited papers and percentage of open access papers for IMI project research in the top twenty journal categories.

*Table 4.8.10 FIELD-NORMALISED, JOURNAL-NORMALISED AND RAW CITATION IMPACT OF PAPERS FOR THE TWENTY WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI PROJECT RESEARCH WAS PUBLISHED MOST FREQUENTLY, 2010-2021. ORDERED BY TOTAL NUMBER OF PAPERS.*

JOURNAL CATEGORY	NUMBER OF PAPERS	CITATION IMPACT		
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL	RAW CITATION IMPACT
Neurosciences	875	2.12	1.40	31.94
Pharmacology & Pharmacy	875	1.36	1.09	21.10
Biochemistry & Molecular Biology	718	2.20	1.49	36.67
Immunology	526	1.65	1.21	25.89
Endocrinology & Metabolism	409	1.82	1.18	20.86
Rheumatology	408	1.90	0.90	28.89
Clinical Neurology	403	2.51	1.43	39.66
Cell Biology	408	2.08	1.31	36.49
Psychiatry	379	2.34	1.26	31.47
Medicine, Research & Experimental	352	1.97	1.03	27.32
Chemistry, Multidisciplinary	331	1.41	1.14	33.90
Microbiology	315	1.66	1.10	22.84
Chemistry, Medicinal	309	1.44	1.19	16.20
Oncology	262	2.54	1.41	50.11
Genetics & Heredity	259	2.14	1.26	40.28
Infectious Diseases	250	2.17	1.34	22.62
Biochemical Research Methods	254	1.36	1.16	22.87
Biotechnology & Applied Microbiology	223	1.66	1.26	26.68
Medicine, General & Internal	219	3.21	1.60	39.49
Public, Environmental & Occupational Health	178	1.54	1.28	14.47

Table 4.8.11 NUMBER OF PUBLICATIONS, NUMBER OF PAPERS, PERCENTAGE OPEN ACCESS AND PERCENTAGE HIGHLY CITED PAPERS FOR THE TOP TWENTY WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI PROJECT RESEARCH WAS PUBLISHED MOST FREQUENTLY, 2010-2021. ORDERED BY TOTAL NUMBER OF PUBLICATIONS.

JOURNAL CATEGORY	NUMBER OF PUBLICATIONS	% OF OPEN ACCESS PAPERS	NUMBER OF PAPERS	% OF HIGHLY CITED PAPERS
Neurosciences	935	72.9%	875	28.1%
Pharmacology & Pharmacy	933	59.1%	875	19.1%
Biochemistry & Molecular Biology	735	76.5%	718	25.9%
Immunology	554	79.7%	526	19.6%
Endocrinology & Metabolism	517	80.2%	409	20.5%
Rheumatology	501	68.9%	408	27.7%
Clinical Neurology	455	64.3%	403	37.5%
Cell Biology	426	84.6%	408	36.3%
Psychiatry	412	74.4%	379	25.6%
Research & Experimental Medicine	362	77.0%	352	26.7%
Multidisciplinary Chemistry	336	72.2%	331	19.6%
Microbiology	325	88.9%	315	24.1%
Medicinal Chemistry	312	66.0%	309	14.6%
Oncology	291	76.3%	262	35.1%
Genetics & Heredity	278	83.4%	259	27.0%
Infectious Diseases	272	86.8%	250	26.8%
Biochemical Research Methods	259	65.7%	254	23.6%
Biotechnology & Applied Microbiology	243	83.9%	223	26.9%
General & Internal Medicine	233	94.1%	219	30.6%
Public, Environmental & Occupational Health	161	57.8%	142	19.0%

- IMI project research was most frequently published in Neurosciences journals. Of the 875 papers published in this category, more than a quarter (28%) were highly cited.
- Clinical Neurology (403 papers) remains the category with the highest percentage of highly cited papers (37.5%), followed by Cell Biology with 408 papers of which 36.3% are highly cited.
- The percentage of open access papers is highest in General & Internal Medicine (94.1%), followed by Microbiology (88.9%) and Infectious Diseases (86.8%).

## 4.9 IMI research fields with the highest volume of publications benchmarked against EU-28 publications of the same field

Figure 4.9.1 shows the field-normalised citation impact of IMI funded research in the twenty Web of Science journal categories in which IMI project research was published most frequently between 2010 and 2021. These data are benchmarked against the average citation impact of all EU-28 research papers in the same journal categories. Table 4.9.2, expands on the data presented in Figure 4.9.1, showing the percentage of IMI and EU-28 papers in each journal category.

Figure 4.9.12 THE FIELD-NORMALISED CITATION IMPACT OF IMI PROJECT RESEARCH IN THE TOP 20 WEB OF SCIENCE JOURNAL CATEGORIES WHICH IMI PROJECT RESEARCH WAS MOST FREQUENTLY PUBLISHED, BENCHMARKED AGAINST EU-28 PAPERS IN THE SAME JOURNAL CATEGORIES, 2010-2021. ORDERED BY THE FIELD-NORMALISED CITATION IMPACT OF IMI RESEARCH.

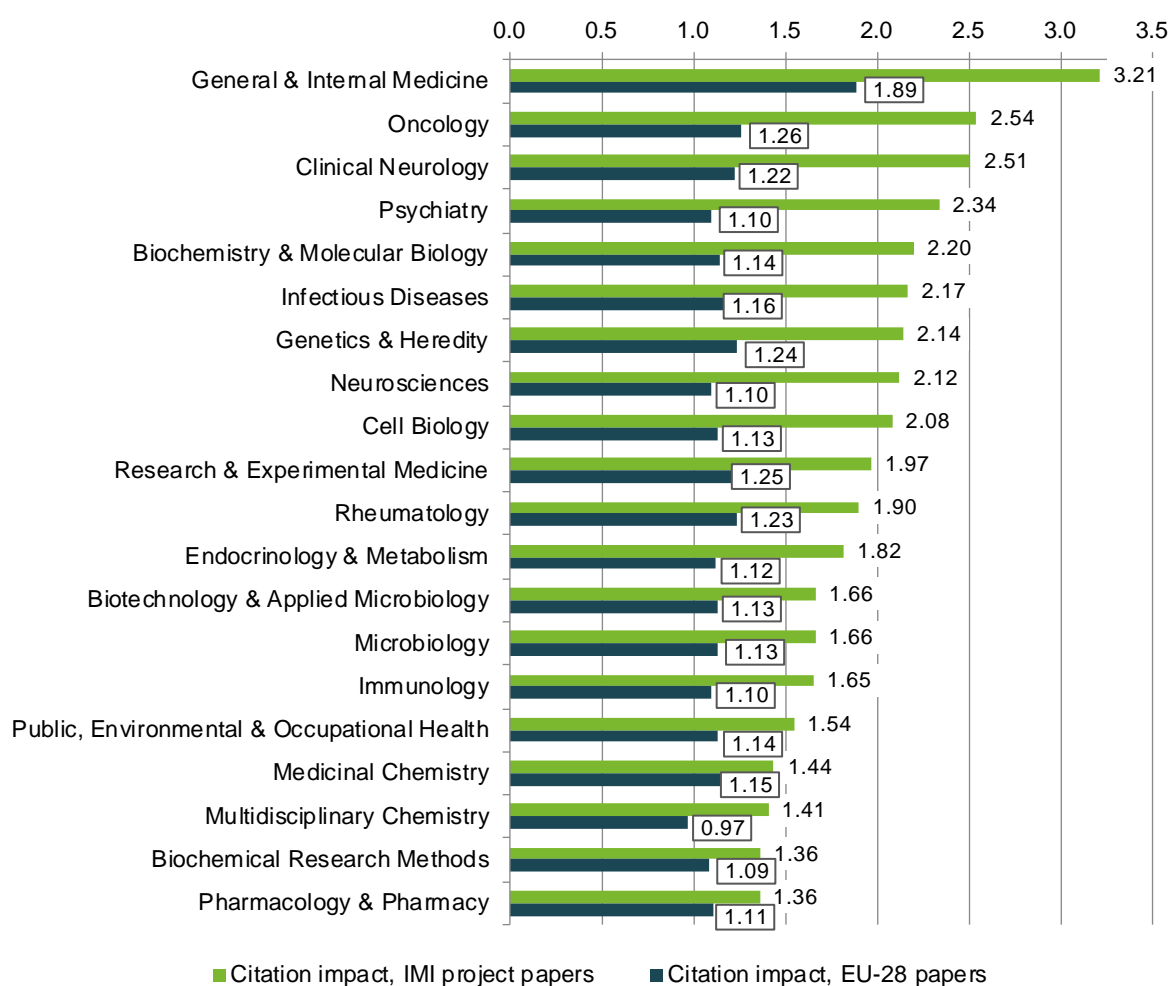


Table 4.9.2 CITATION IMPACT AND PERCENTAGE OF PAPERS IN TOP TWENTY WEB OF SCIENCE JOURNAL CATEGORIES IN WHICH IMI PROJECT RESEARCH WAS MOST FREQUENTLY PUBLISHED, BENCHMARKED AGAINST EU-28 IN THE SAME JOURNAL CATEGORIES, 2010-2021

JOURNAL CATEGORY	% OF IMI PAPERS	% OF EU-28 PAPERS	CITATION IMPACT NORMALISED AT FIELD LEVEL	
			IMI papers	EU-28
General & Internal Medicine	2.97%	0.24%	3.21	1.89
Oncology	3.70%	2.37%	2.54	1.26
Clinical Neurology	5.79%	1.94%	2.51	1.22
Psychiatry	5.24%	1.48%	2.34	1.10
Biochemistry & Molecular Biology	9.36%	3.60%	2.20	1.14
Infectious Diseases	3.46%	1.02%	2.17	1.16
Genetics & Heredity	3.54%	1.31%	2.14	1.24
Neurosciences	11.90%	2.71%	2.12	1.10
Cell Biology	5.42%	1.70%	2.08	1.13
Research & Experimental Medicine	4.61%	1.14%	1.97	1.25
Rheumatology	6.38%	0.45%	1.90	1.23
Endocrinology & Metabolism	6.58%	1.34%	1.82	1.12
Biotechnology & Applied Microbiology	3.09%	1.31%	1.66	1.13
Microbiology	4.14%	1.54%	1.66	1.13
Immunology	7.05%	1.51%	1.65	1.10
Public, Environmental & Occupational Health	2.55%	1.96%	1.54	1.14
Medicinal Chemistry	3.97%	0.67%	1.44	1.15
Multidisciplinary Chemistry	4.28%	3.13%	1.41	0.97
Biochemical Research Methods	3.30%	1.04%	1.36	1.09
Pharmacology & Pharmacy	11.88%	2.19%	1.36	1.11

- In all twenty journal categories listed, IMI project research had a higher field-normalised citation impact than EU-28 papers in the same field.
- General & Internal Medicine (3.21) and Oncology (2.54) were the top two journal categories in which IMI-supported research had the highest field-normalised citation impact. This is a change from last year where Clinical Neurology was the top journal category but is now third with a field-normalised citation impact of 2.51.
- The average field-normalised citation impact of EU-28 papers was also the highest in the same two categories of General & Internal Medicine (1.89) and Oncology (1.26).

## 5 CITATION ANALYSIS – AT IMI PROJECT LEVEL

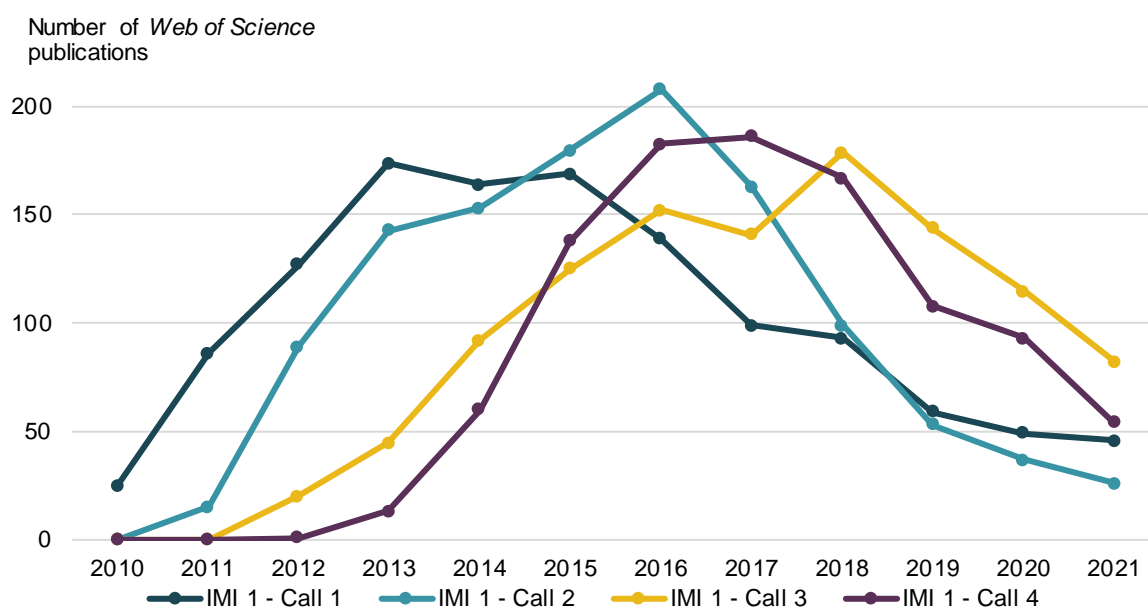
This section analyses the volume and citation impact of publications arising from different IMI-phases and calls.

### 5.1 Trends in publication output by IMI funding call

Figure 5.1.1 and Figure 5.1.2 show the number of Web of Science publications between 2010 and 2021 for IMI project research disaggregated by call. IMI 1 calls 1-4 (Figure 5.1.1) are shown separately from the more recent IMI 1 calls 5-11 (Figure 5.1.2) which tend to have fewer publications. Likewise, IMI 2 calls are shown separately in Figure 5.1.3 as individual IMI 2 calls has far fewer publication compared to most IMI 1 calls as the longest running IMI 2 projects only started publishing in 2015.

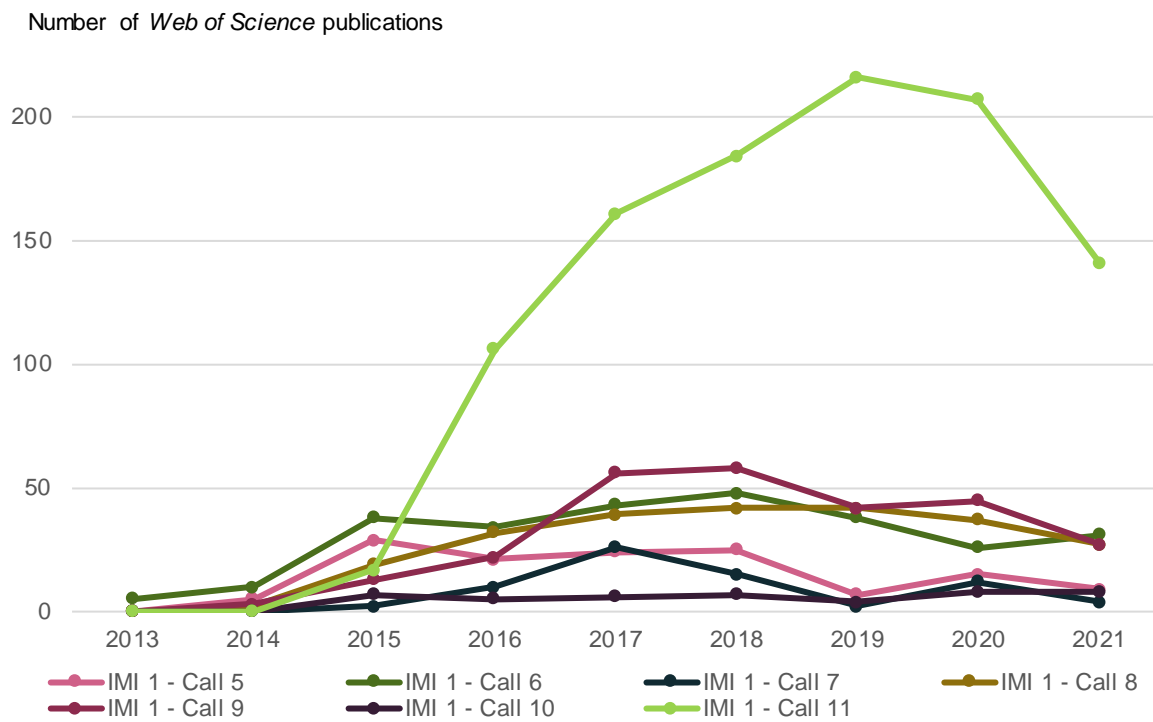
Table 5.1.1 presents summary bibliometric data for all IMI 1 and IMI 2 calls that have at least one publication, including the number of publications, numbers of papers, and citation impact indicators.

Figure 5.1.1 NUMBER OF WEB OF SCIENCE PUBLICATIONS BY YEAR AND FUNDING CALL, 2010-2021



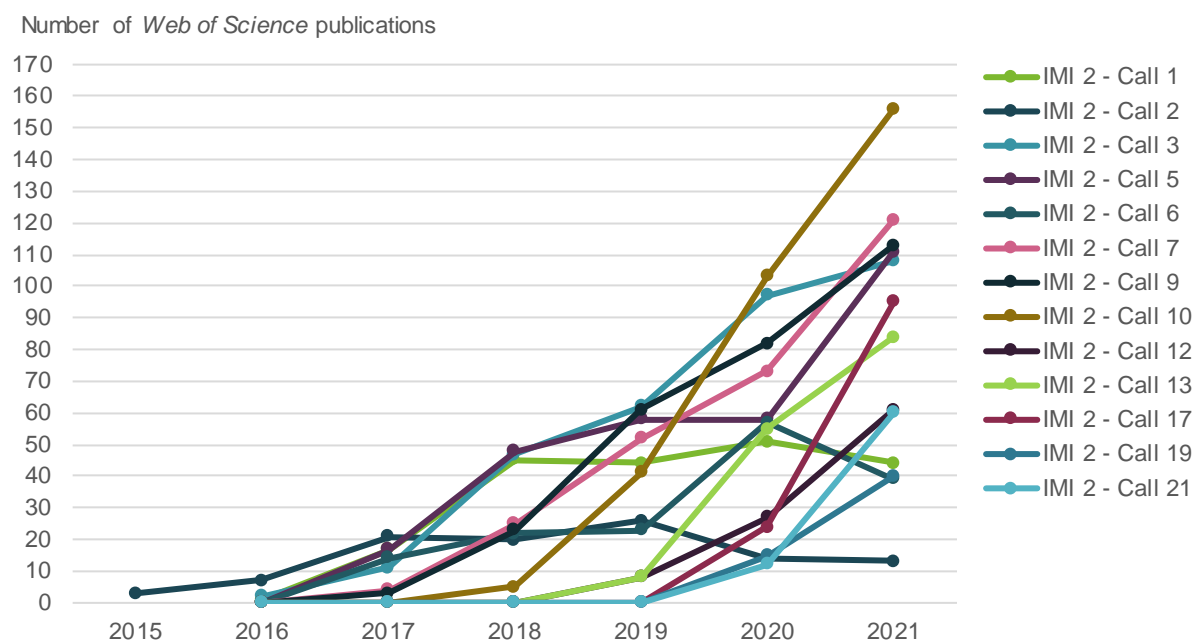
- Over the five years 2010 to 2014, IMI 1 call 1 had the highest output of publications, reaching a peak output of 174 publications in 2013.
- In 2015 and 2016, IMI 1 call 2 had the highest number of publications (180 and 208, respectively). In 2017 call 2's output fell (163 publications) and call 4 had the highest output of publications (186 publications).
- In 2021 all IMI 1 calls 1-4 continue to trend downward. Which is likely to continue since all the calls are now closed.
- Call 3 remains the call with the highest number of publications and appears to have a steeper decline than the other calls.

Figure 5.1.2 NUMBER OF WEB OF SCIENCE PUBLICATIONS BY YEAR AND FUNDING CALL, 2010-2021



- Overall, IMI 1 calls 5-10 have not grown as rapidly as IMI 1 calls 1-4, most calls produce fewer than 50 publications a year. Call 11 is the exception, with growth in output akin to IMI 1 calls 1-4. This growth has declined substantially in 2021 (-32%).
- Many of the call 11 projects closed in 2021 so it is likely that we will continue to see this type of decline for call 11.

Figure 5.1.3 NUMBER OF WEB OF SCIENCE PUBLICATIONS BY YEAR AND FUNDING CALL, 2010-2021. ONLY SHOWING IMI 2 CALLS WITH AT LEAST 50 PUBLICATIONS IN TOTAL.



- The output of publications from IMI 2 calls is growing. IMI 2 call 2 was the first to start publishing in 2015 and after plateauing between 2017-2019 it has now begun declining.
- IMI 2 Calls 12, 17, 19 and 21 are newly included to this analysis as the number of publications within these calls have increased rapidly, mostly driven by the project EHDEN, EubOPEN, INNODIA HARVEST, and DRAGON, respectively.
- Call 10's rapid increase continues and is still largely driven by the AIMS-2-Trials which published 101 publications in 2021 which is 64% of Call 10s publications in 2021.
- In contrast to last year, call 5 had a rapid increase in publications. While the rest of the calls continue on their trajectory from last year with the exception of Call 6 which has decreased from last year.

Table 5.1.1 SUMMARY BIBLIOMETRIC ANALYSES OF IMI PROJECTS AGGREGATED BY FUNDING CALL, 2010-2021

PHASE	CALL	NUMBER OF PUBLICATIONS <sup>8</sup>	% OF OPEN ACCESS PAPERS	NUMBER OF PAPERS	CITATION IMPACT		
					RAW CITATION IMPACT	NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL
1	1	1,230	61.6%	1,134	44.92	1.08	1.79
1	2	1,166	75.1%	1,098	50.05	1.19	2.16
1	3	1,095	80.9%	1,010	32.60	1.06	1.94
1	4	1,003	67.8%	957	36.57	1.27	2.02
1	5	135	80.6%	134	18.61	1.01	1.11
1	6	273	73.9%	264	20.68	0.95	1.28
1	7	71	85.9%	64	18.81	1.28	1.53
1	8	240	79.8%	213	31.21	1.36	2.18
1	9	266	72.5%	247	26.44	1.41	1.63
1	10	45	84.1%	44	15.34	0.95	1.58
1	11	1,032	81.9%	945	28.40	1.25	2.18
2	1	203	88.0%	166	15.39	1.15	1.62
2	2	104	89.9%	99	19.67	1.26	1.93
2	3	327	88.9%	270	14.66	1.21	2.04
2	4	4	50.0%	4	8.25	0.32	0.58
2	5	292	93.5%	261	15.27	1.23	2.41
2	6	155	92.2%	129	14.05	1.26	2.01
2	7	275	91.4%	244	19.62	1.37	2.72
2	8	26	91.7%	24	8.83	0.77	0.83
2	9	282	85.5%	241	16.72	1.41	2.84
2	10	305	90.0%	270	8.95	1.19	2.21

<sup>8</sup> Publications can be associated with more than one call.



PHASE	CALL	NUMBER OF PUBLICATIONS <sup>8</sup>	% OF OPEN ACCESS PAPERS	NUMBER OF PAPERS	CITATION IMPACT		
					RAW CITATION IMPACT	NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL
2	12	96	92.5%	80	4.64	1.28	1.74
2	13	147	90.3%	134	8.16	1.38	1.96
2	14	39	82.4%	34	6.00	1.12	2.30
2	15	47	85%	33	4.61	2.51	1.81
2	16	2	100%	2	10.00	5.21	4.39
2	17	119	79%	117	4.50	1.45	2.07
2	18	4	75%	4	4.00	4.20	3.97
2	19	55	98%	46	4.70	1.48	1.60
2	20	10	88%	8	2.75	0.83	2.70
2	21	72	97%	59	9.42	3.71	4.08
2	23	3	100%	3	0.67	0.34	0.50

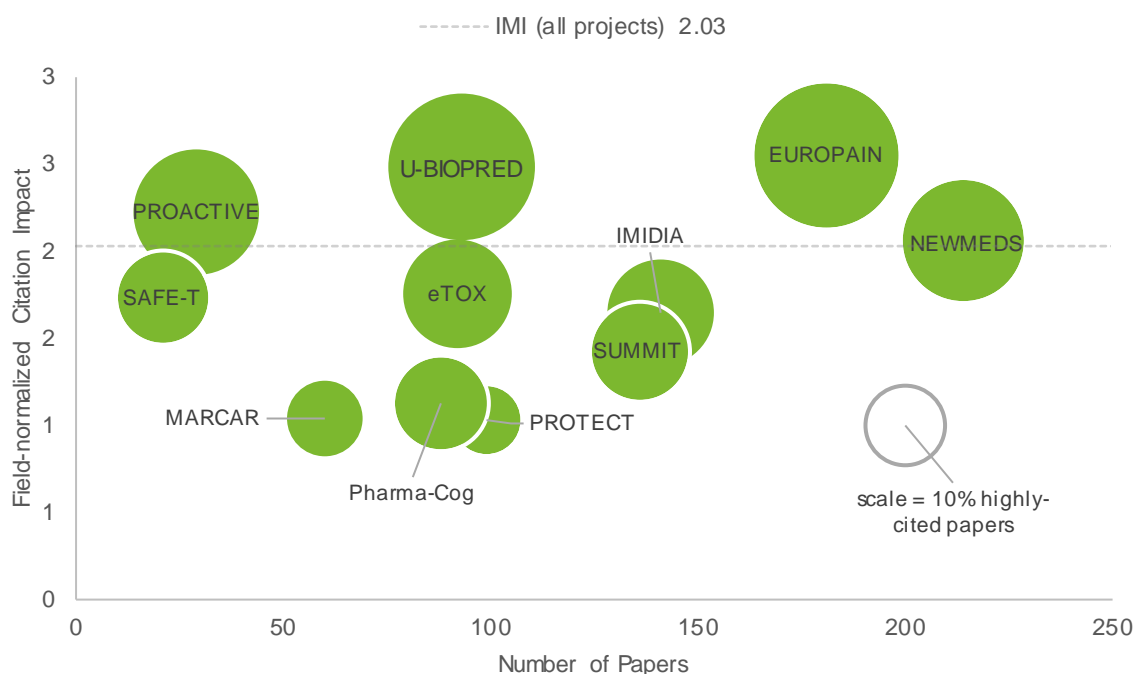
- IMI 1 call 1 remains the funding call that produced the highest number of publications (1,230), and papers (1,134). Although papers from IMI 1 call 2 had the highest raw citation impact (50.05).
- Papers assigned to IMI 2 call 21 had the highest average field-normalised citation impact (3.71)<sup>9</sup>, which is three times the world average and is likely driven by the fact that many of these projects in this call are Coronavirus related. This is driven by a few highly cited papers, mainly within the CARE and DRAGON projects, which were cited between 10 and 30 times the world average.
- The highest percentage of open access papers belongs to IMI 2 call 19 where 98% of the publications are open access<sup>9</sup>.
- Generally, IMI 2 calls have a higher proportion of open access papers compared to IMI 1 calls likely due to the mandate that papers in IMI 2 be published as open access.
- IMI 2 call 3 with 327 publications is IMI 2's highest output call.

<sup>9</sup> Only calls with at least 10 papers were considered

## 5.2 Summary bibliometric analyses for imi 1 projects – call 1

Figure 5.2.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 call 1 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.2.4 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS - CALL 1, 2010-2021



The data in Figure 5.2.1 shows that:

- The average field-normalised citation impact of all IMI 1 call 1 projects with at least 10 papers was above the world average (1.00). Furthermore, the percentage of highly cited research was also above or in line with the world average (10%) for all projects except for the PROTECT Project which 8.1% of its papers were highly cited. This indicates excellent research performance.
- Research associated with NEWMEDS, EUROPAIN, PROACTIVE and U-BIOPRED was cited more than twice the world average. These four projects also have an average citation impact greater than the average citation impact of all IMI project papers (2.03).

Table 5.2.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 1 publications. Table 5.2.2 shows the normalised citation impact (normalised against world average values) of IMI 1 call 1 projects and is an expansion of the data shown in Figure 5.2.1.

Table 5.2.2 BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALL 1, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
NEWMEDS	220	214	57.9%	12,350	57.71
EUROPAIN	183	181	40.3%	11,592	64.04
IMIDIA	151	141	85.8%	6,834	48.47
SUMMIT	141	136	77.9%	3,888	28.59
PROTECT	101	99	46.5%	2,327	23.51
U-BIOPRED	148	93	73.1%	4,348	46.75
eTOX	97	92	69.6%	4,184	45.48
Pharma-Cog	94	88	44.3%	3,196	36.32
MARCAR	61	60	73.3%	1,518	25.30
PROACTIVE	34	29	89.7%	1,520	52.41
SAFE-T	23	21	38.1%	637	30.33

Table 5.2.3 SUMMARY CITATION INDICATORS FOR IMI1 PROJECTS IN CALL 1, 2010-2021

PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
NEWMEDS	214	2.06	1.10	31.78	25.7%
EUROPAIN	181	2.55	1.37	24.99	36.5%
IMIDIA	141	1.65	1.03	31.43	19.9%
SUMMIT	136	1.42	0.90	38.19	16.2%
PROTECT	99	1.03	0.91	40.28	8.1%
U-BIOPRED	93	2.49	1.33	22.09	37.6%
eTOX	92	1.76	1.24	30.89	20.7%
Pharma-Cog	88	1.13	0.84	44.54	14.8%
MARCAR	60	1.04	0.76	42.33	10.0%
PROACTIVE	29	2.22	1.57	26.19	27.6%
SAFE-T	21	1.73	1.06	33.06	14.3%
<b>Overall (IMI projects)</b>	<b>7,856</b>	<b>2.03</b>	<b>1.23</b>	<b>34.98</b>	<b>25.3%</b>

- Of the projects in call 1, NEWMEDS had the highest number of publications (220) and PROACTIVE had the highest percentage of open access papers (89.7%).

### 5.3 Summary bibliometric analyses for IMI 1 projects – call 2

Figure 5.3.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 call 2 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers. The same data is shown in Figure 5.3.1 and Figure 5.3.2, however Figure 5.3.1 has a smaller x-axis range that excludes BTCure so that the other projects are less clustered.

Figure 5.3.5 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALL 2, 2010-2021

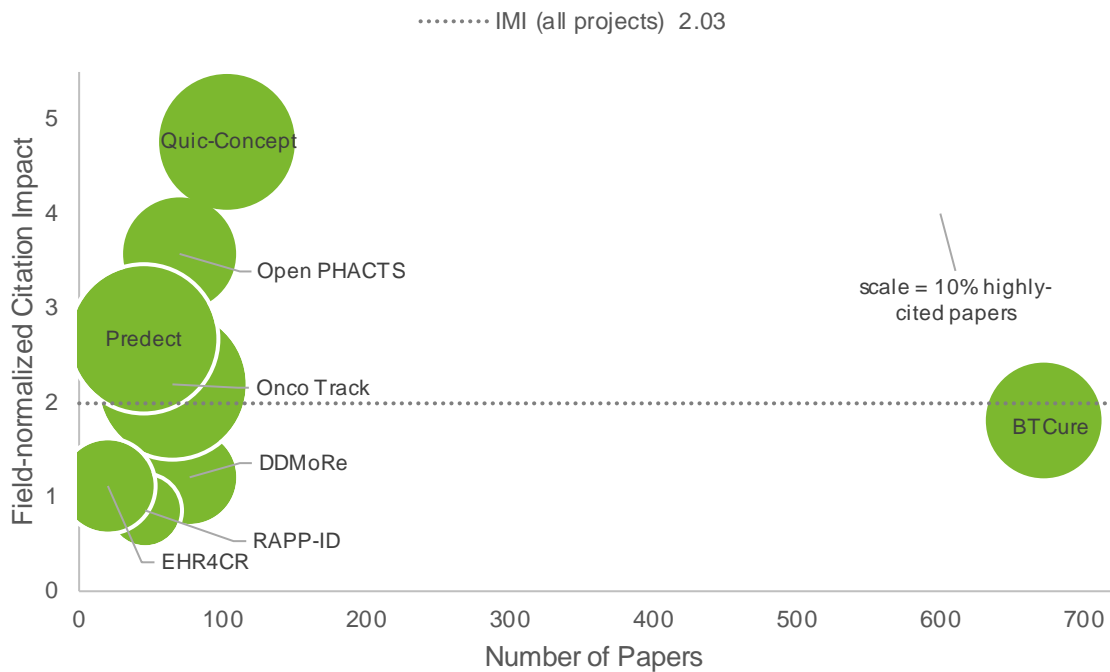
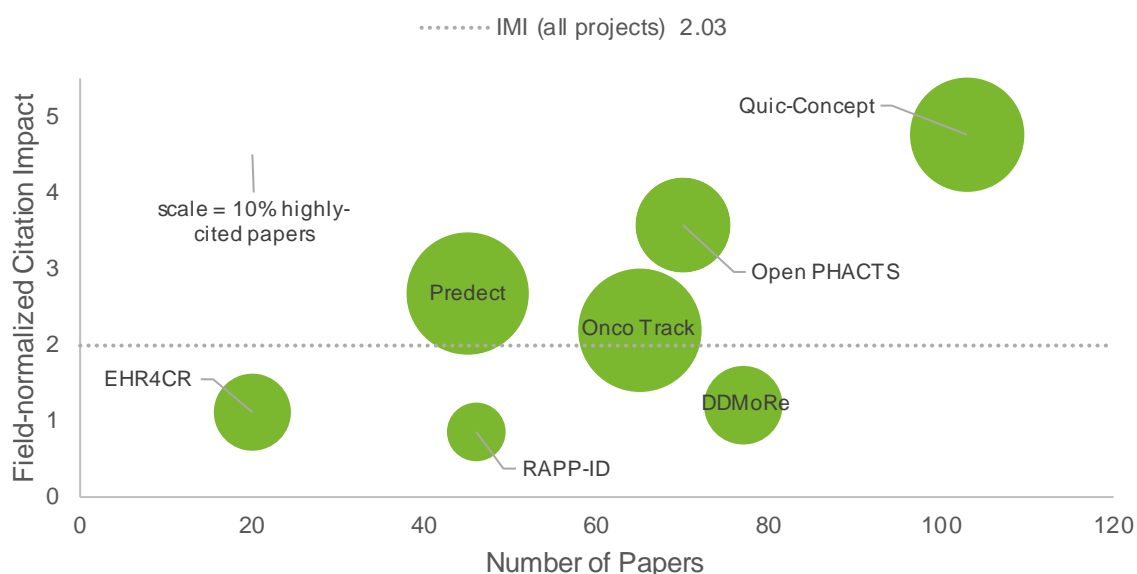


Figure 5.3.6 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALL, 2010-2021. SAME GRAPH AS FIGURE 5.3.1 BUT WITH A SMALLER X-AXIS RANGE



The data in Figure 5.3.1 and Figure 5.3.2 shows that:

- The average field-normalised citation impact of most IMI 1 call 2 projects was above world average apart from RAPP-ID which had the lowest citation impact (0.86). Similarly, all except RAPP-ID had a higher percentage of highly cited papers than the world average (10%).
- BTCURE remains the most prolific IMI 1 call 2 project with 672 papers and a citation impact of 1.81, which is lower than the citation impact of all IMI project papers (2.03).
- QUIC-CONCEPT is the most highly cited project with a citation impact more than four times the world average (4.76).
- Open Phacts, OncoTrack and Predect are also well cited with a citation impact of 3.57, 2.19 and 2.67, respectively.
- Half of the projects in this call had an average citation impact greater than the average citation impact of all IMI project papers (2.03).

Table 5.3.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 2 publications. Table 5.3.2 shows the normalised citation impact (normalised against world average values) of IMI 1 call 2 projects and is an expansion of the data shown in Figure 5.3.1 and Figure 5.3.2.

Table 5.3.4 BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALL 2, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
BTCure	719	672	72.5%	28,560	42.50
Quic-Concept	104	103	85.4%	10,696	103.84
DDMoRe	82	77	70.1%	1,617	21.00
Open PHACTS	73	70	90.0%	5,819	83.13
Onco Track	69	65	70.8%	4,321	66.48
RAPP-ID	47	46	71.7%	1,038	22.57
Predict	49	45	84.4%	2,988	66.40
EHR4CR	23	20	80.0%	436	21.80

Table 5.3.5 SUMMARY CITATION INDICATORS FOR IMI 1 PROJECTS IN CALL 2, 2010-2021

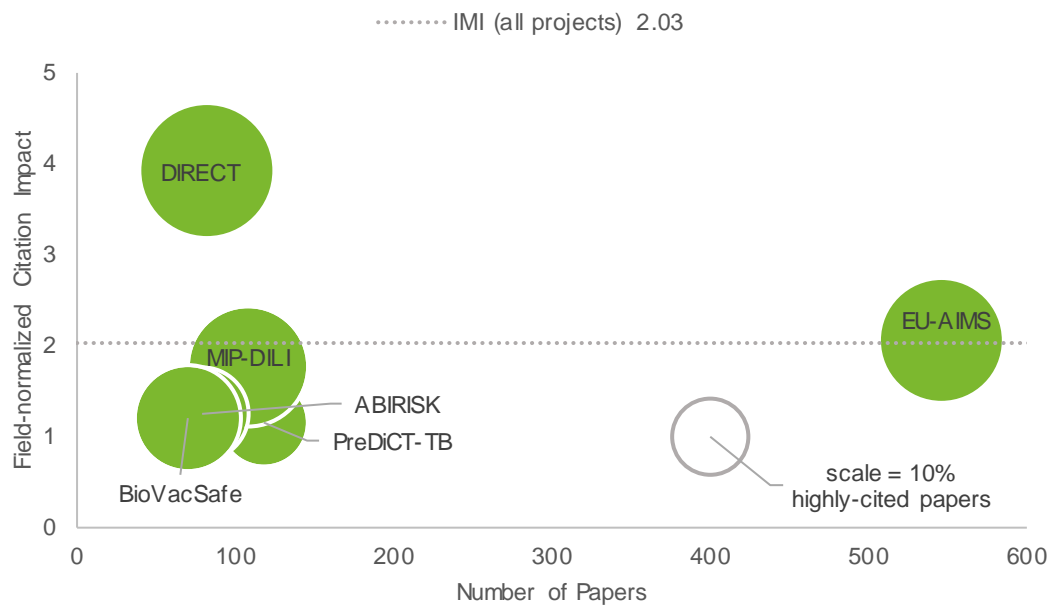
PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
BTCure	672	1.81	0.99	30.43	24.0%
Quic-Concept	103	4.76	2.19	31.90	33.0%
DDMoRe	77	1.21	1.07	46.27	15.6%
Open PHACTS	70	3.57	1.84	36.53	22.9%
Onco Track	65	2.19	1.19	29.00	38.5%
RAPP-ID	46	0.86	0.77	41.68	8.7%
Predict	45	2.67	1.49	34.99	37.8%
EHR4CR	20	1.11	1.10	40.75	15.0%
<b>Overall (IMI projects)</b>	<b>7,856</b>	<b>2.03</b>	<b>1.23</b>	<b>34.98</b>	<b>25.3%</b>

- Among IMI 1 call 2 projects Open PHACTS has the highest percentage of open access papers (90%).
- OncoTrack has the highest percentage of highly cited papers (38.5%)

## 5.4 Summary bibliometric analyses for IMI 1 projects – call 3

Figure 5.4.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 call 3 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.4.7 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALL 3, 2010-2021, 2010-2021



The data in Figure 5.4.1 shows that:

- The average citation impact and percentage of highly cited papers for all projects in this call was above the world average.
- EU-AIMS was by far the most prolific IMI 1, call 3 project with 546 papers. The field-normalised citation impact of this research was twice the world average (2.06) and above average for all IMI research (2.03).
- Research associated with DIRECT was very well-cited with a field-normalised citation impact of almost four times (3.93) the world average and nearly a third (32.9%) of its papers were highly cited.

Table 5.4.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 3 publications. Table 5.4.2 shows the normalised citation impact (normalised against world average values) of IMI 1 call 3 projects and is an expansion of the data shown in Figure 5.4.1.

Table 5.4.6 SUMMARY BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALL 3, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
EU-AIMS	565	546	83.0%	19,198	35.16
PreDiCT-TB	124	118	92.4%	2,545	21.57
MIP-DILI	116	108	64.8%	3,278	30.35
DIRECT	109	82	86.6%	3,735	45.55
ABIRISK	100	79	64.6%	2,076	26.28
BioVacSafe	73	70	80.0%	2,179	31.13

Table 5.4.7 SUMMARY CITATION INDICATORS FOR IMI 1 PROJECTS IN CALL 3, 2010-2021

PROJECT	NUMBER OF PAPERS	CITATION IMPACT			% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL	AVERAGE PERCENTILE	
EU-AIMS	546	2.06	1.12	29.94	28.0%
PreDiCT-TB	118	1.16	0.79	42.81	13.6%
MIP-DILI	108	1.77	1.35	34.44	25.9%
DIRECT	82	3.93	0.99	32.64	32.9%
ABIRISK	79	1.25	0.86	43.41	15.2%
BioVacSafe	70	1.20	0.94	33.23	20.0%
Overall (IMI projects)	7,856	2.03	1.23	34.98	25.3%

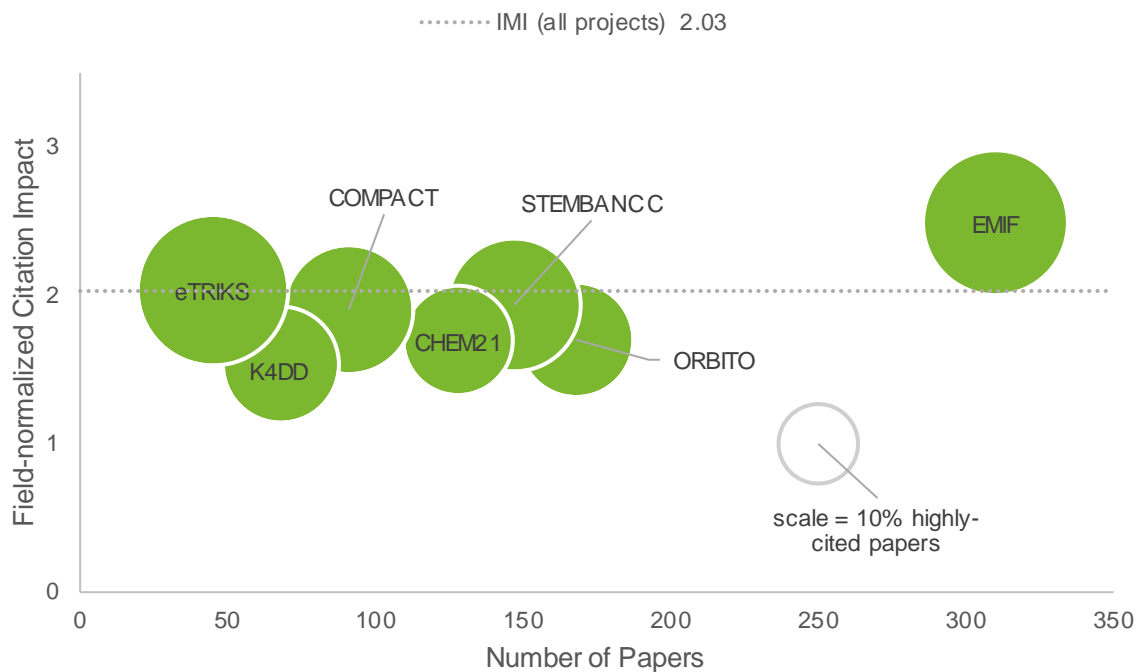
- PreDiCT-TB had the highest percentage of open access papers (92.4%).



## 5.5 Summary bibliometric analyses for IMI 1 projects – call 4

Figure 5.5.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 call 4 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.5.8 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALL 4, 2010-2021



The data in Figure 5.5.1 shows that:

- The average field-normalised citation impact of all projects in this call is above world average.
- EMIF produced the highest number of papers in call 4, with 310 papers published by the end of 2021 and has a field-normalised citation impact two and half times the world average (2.49).
- eTRIKS has the highest percentage of highly cited papers (37.8%)
- Two-of-the-seven projects in this call had an average field-normalised citation impact greater than the average citation impact for all IMI project research (2.03).

Table 5.5.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 4 publications. Table 5.5.2 shows the normalised citation impact (normalised against world average values) of IMI 1 call 4 projects and is an expansion of the data shown in Figure 5.5.1.

Table 5.5.8 BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALL 4, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
EMIF	331	310	84.2%	12,728	41.06
ORBITO	171	168	36.3%	4,475	26.64
STEMBANCC	153	147	81.6%	4,905	33.37
CHEM21	131	128	50.0%	5,499	42.96
COMPACT	91	91	53.8%	4,174	45.87
K4DD	70	68	75.0%	1,795	26.40
eTRIKS	56	45	95.6%	1,660	36.89

Table 5.5.9 SUMMARY CITATION INDICATORS FOR IMI 1 PROJECTS IN CALL 4, 2010-2021

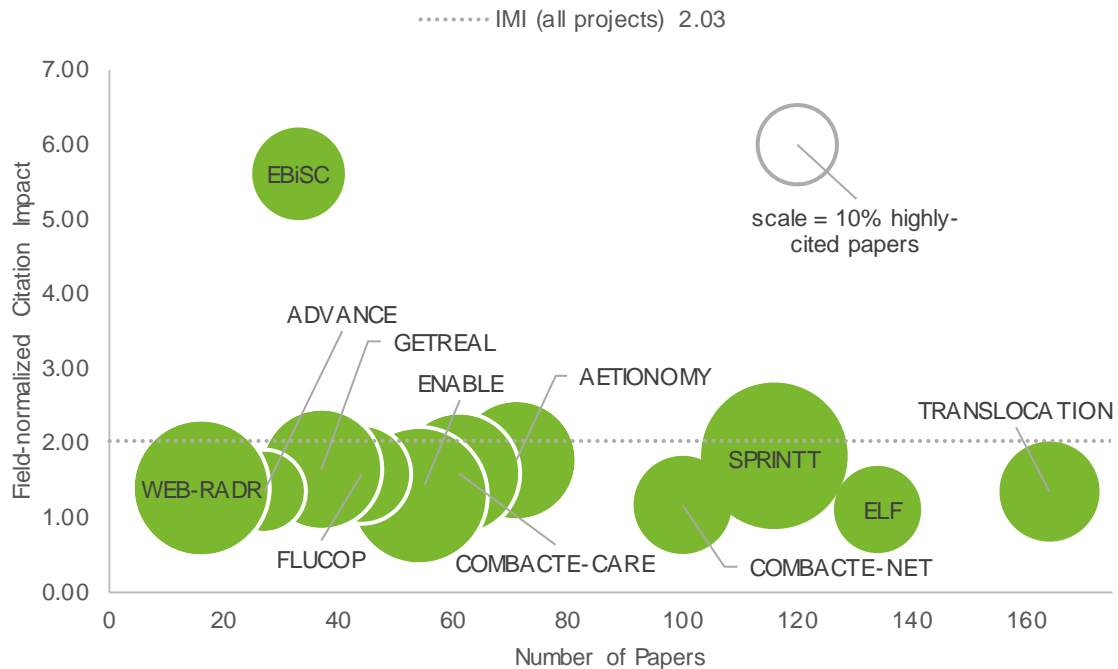
PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
EMIF	310	2.49	1.26	29.10	34.8%
ORBITO	168	1.70	1.27	29.23	21.4%
STEMBANCC	147	1.94	1.28	31.38	29.3%
CHEM21	128	1.70	1.27	36.72	19.5%
COMPACT	91	1.90	1.42	29.94	27.5%
K4DD	68	1.53	1.20	33.04	22.1%
eTRIKS	45	2.04	1.22	30.24	37.8%
Overall (IMI projects)	7,856	2.03	1.23	34.98	25.3%

- eTRIKS has the highest percentage of open access papers (95.6%).

## 5.6 Summary bibliometric analyses for IMI 1 projects – calls 5-10

Figure 5.6.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 calls 5-10 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.6.9 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALLS 5-10, 2010-2021



The data in Figure 5.6.1 shows that:

- Research associated with EBiSC was very well cited with an astounding field-normalised citation impact of more than five times the world average (5.61). However, the total number of EBiSC papers is still relatively low (33), so it is possible that only a few highly cited papers has inflated the citation impact.
- SPRINTT has the highest percentage of highly cited papers (37.9%)
- TRANSLOCATION produced the most papers (164) likely due to it being one of the longest running projects from IMI 1 calls 5-10.
- All the projects in calls 5-10 have a field-normalised citation impact greater than the world average but below average for all IMI project research (2.03), apart from EBiSC.

Table 5.6.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 5-10 publications. Table 5.6.2 shows the normalised citation impact (normalised against world average values) of IMI 1 calls 5-10 projects and is an expansion of the data shown in Figure 5.6.1.

Table 5.6.10 BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALLs 5-10, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
TRANSLOCATION	164	164	67.1%	4,123	25.14
ELF	135	134	80.6%	2,495	18.62
SPRINTT	123	116	57.8%	3,611	31.13
COMBACTE-NET	109	100	85.0%	1,370	13.70
AETIONOMY	74	71	81.7%	1,866	26.28
COMBACTE-CARE	66	61	86.9%	1,446	23.70
ENABLE	56	55	85.5%	1,104	20.07
PRECISESADS	74	54	63.0%	1,027	19.02
DRIVE-AB	60	54	83.3%	1,303	24.13
FLUCOP	45	44	84.1%	752	17.09
GETREAL	43	37	81.1%	858	23.19
EBISC	36	33	93.9%	2,694	81.64
ADVANCE	28	27	92.6%	397	14.70
WEB-RADR	17	16	87.5%	283	17.69

Table 5.6.11 SUMMARY CITATION INDICATORS FOR IMI 1 PROJECTS IN CALLS 5-10, 2010-2021

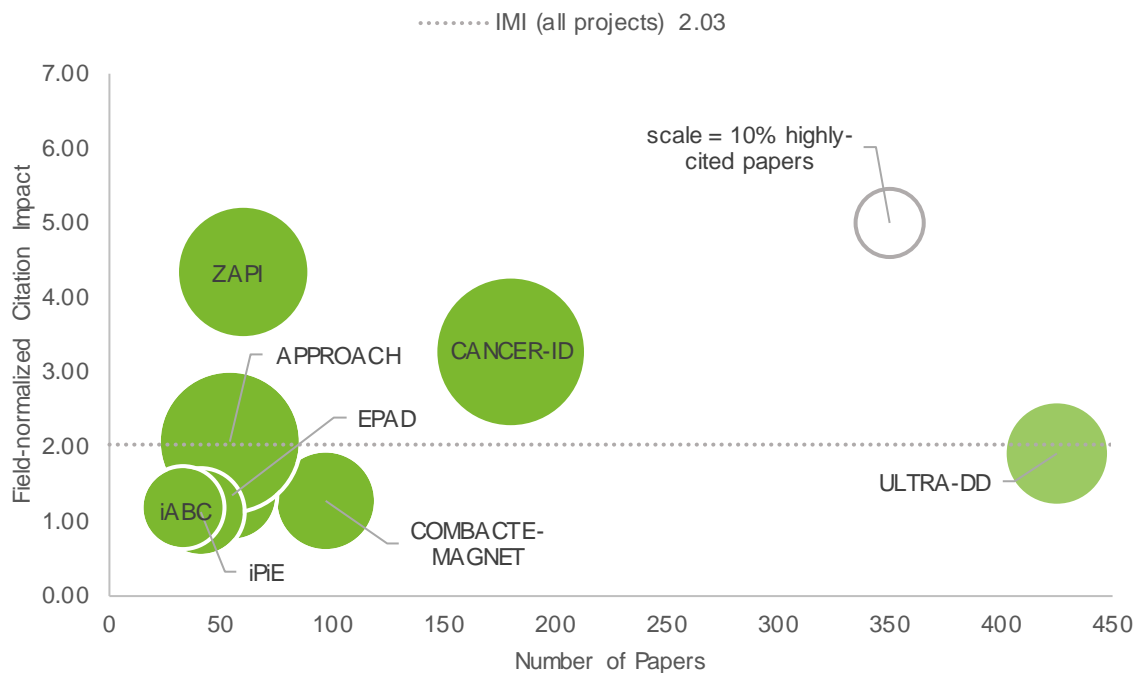
PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
TRANSLOCATION	164	1.35	0.98	36.32	17.7%
ELF	134	1.11	1.01	41.96	13.4%
SPRINTT	116	1.83	1.87	26.97	37.9%
COMBACTE-NET	100	1.17	0.91	37.63	17.0%
AETIONOMY	71	1.77	1.24	34.72	23.9%
COMBACTE-CARE	61	1.59	0.94	35.90	24.6%
ENABLE	55	1.45	1.10	34.83	21.8%
PRECISESADS	54	1.38	0.89	35.25	16.7%
DRIVE-AB	54	1.30	0.99	32.62	31.5%
FLUCOP	44	1.58	0.95	44.12	15.9%
GETREAL	37	1.65	1.13	36.77	24.3%
EBiSC	33	5.61	2.80	33.52	15.2%
ADVANCE	27	1.36	1.48	32.70	11.1%
WEB-RADR	16	1.40	1.32	33.20	31.3%
Overall (IMI projects)	7,856	2.03	1.23	34.98	25.3%

- EBiSC has the highest percentage (93.9%) of open access papers.

## 5.7 Summary bibliometric analyses for IMI 1 projects – call 11

Figure 5.7.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers for IMI 1 call 11 projects. Only projects with at least 10 papers and one highly cited paper over the time period (2010-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.7.10 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 1 PROJECTS – CALL 11, 2010-2021



The data in Figure 5.7.1 shows that:

- ULTRA-DD produced by far the most papers (425).
- All the projects performed above world average for percentage of highly cited papers and field-normalised citation impact.
- Research papers associated with APPROACH, CANCER-ID and ZAPI were very well-cited with field-normalised citation impacts of two (2.07), three (3.27), and four (4.34) times the world average, respectively. They are also the only projects in this call that are higher than the average for all IMI projects (2.03).
- Over half of CANCER-ID papers are highly cited (52.2%).

Table 5.7.1 shows raw citation impact and the percentage of open access papers by project for IMI 1 call 11 publications. Table 5.7.2 shows the normalised citation impact (normalised against world average values) of IMI 1 call 11 projects and is an expansion of the data shown in Figure 5.7.1.

Table 5.7.12 BIBLIOMETRIC INDICATORS FOR IMI 1 PROJECTS IN CALL 11, 2010-2021

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
ULTRA-DD	434	425	85.2%	9,417	22.16
CANCER-ID	208	180	73.3%	9,700	53.89
COMBACTE-MAGNET	108	97	84.5%	1,583	16.32
ZAPI	63	60	95.0%	2,847	47.45
EPAD	59	55	87.3%	817	14.85
APPROACH	67	54	72.2%	1,887	34.94
iPiE	42	41	70.7%	741	18.07
iABC	51	33	75.8%	365	11.06

Table 5.7.13 SUMMARY CITATION INDICATORS FOR IMI 1 PROJECTS IN CALL 11, 2010-2021

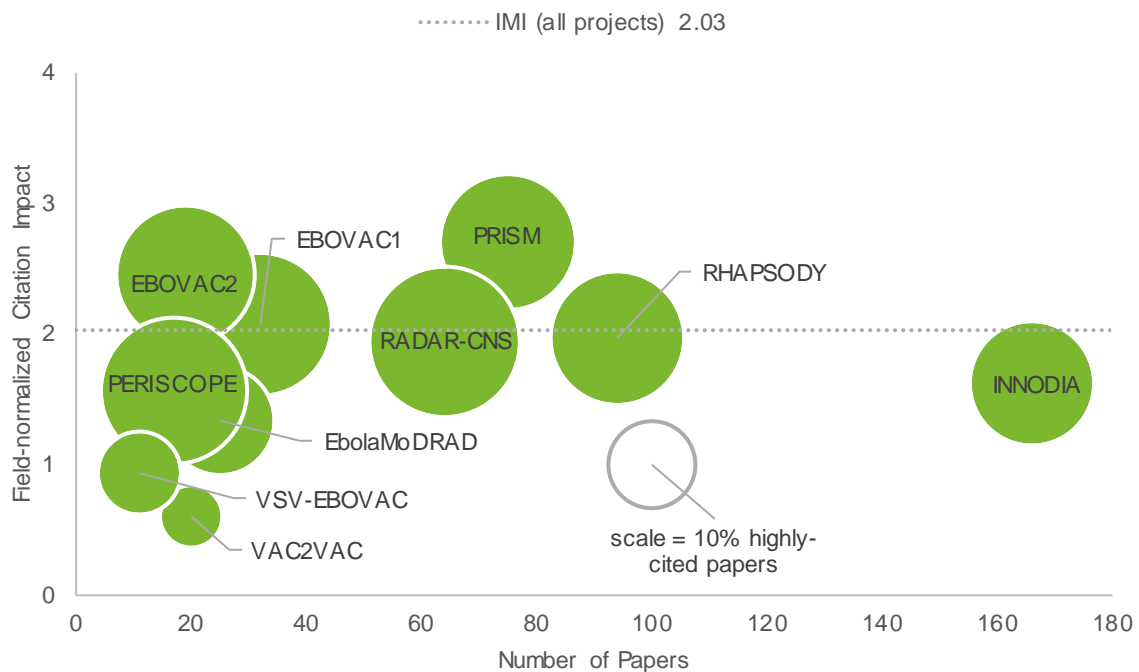
PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
ULTRA-DD	425	1.91	1.07	32.74	24.5%
CANCER-ID	180	3.27	1.58	18.26	52.2%
COMBACTE-MAGNET	97	1.28	0.88	38.61	22.7%
ZAPI	60	4.34	2.34	29.74	40.0%
EPAD	55	1.35	1.01	37.91	18.2%
APPROACH	54	2.07	1.72	30.39	46.3%
iPiE	41	1.13	0.93	37.71	17.1%
iABC	33	1.19	0.91	47.82	15.2%
<b>Overall (IMI projects)</b>	<b>7,856</b>	<b>2.03</b>	<b>1.23</b>	<b>34.98</b>	<b>25.3%</b>

- ZAPI has the highest percentage (95%) of open access papers.

## 5.8 Summary bibliometric analyses for IMI 2 calls 1-4 projects

Figure 5.8.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers from IMI 2 projects from calls 1-4. Only projects with at least 10 papers and one highly cited paper over the time period (2015-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.8.11 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 2 PROJECTS – CALLS 1-4, 2015-2021



The data in Figure 5.8.1 shows that:

- INNODIA was the most productive project, publishing 166 papers.
- PERISCOPE and VAC2VAC have published enough papers to be included in this year's analysis.
- Apart from VAC2VAC and VSV-EBOVAC, all the projects meet or exceed the world average (10%) for highly cited papers.
- PRISM and EBOVAC2 are the most impactful projects with a field-normalized citation impact of more than two times the world average, 2.70 and 2.45, respectively.
- Three-out-of-ten projects performed above the average field-normalized citation impact for all IMI projects (2.03)



Table 5.8.1 shows raw citation impact and percentage of open access papers by project for IMI 2 calls 1-4 publications and Table 5.8.2 shows indicators for IMI 2 calls 1-4 project research where citation impact has been normalised against world average values.

Table 5.8.14 BIBLIOMETRIC INDICATORS FOR IMI 2 CALLS 1-4 PROJECTS, 2015-2021<sup>10</sup>

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
INNODIA	203	166	77.3%	2,684	16.17
RHAPSODY	115	94	78.3%	1,576	16.77
PRISM	87	75	78.2%	1,625	21.67
RADAR-CNS	87	64	80.5%	901	14.08
EBOVAC1	34	32	100.0%	808	25.25
EbolaMoDRAD	26	25	73.1%	355	14.20
VAC2VAC	20	20	95.0%	61	3.05
EBOVAC2	19	19	100.0%	384	20.21
PERISCOPE	18	17	100.0%	103	6.06
VSV-EBOVAC	12	11	75.0%	206	18.73

- All the EBOVAC1, EBOVAC2, and PERISCOPE project papers are open access.
- EbolaMoDRAD has the lowest percentage of open access papers (73.1%).

<sup>10</sup> Note that IMI 2 funded researchers are contractually obliged to make their scientific articles open access through Green or Gold routes. However, for some of other document types, such as editorials, reviews or conference proceedings open access publication is strongly encouraged but not mandatory. Nevertheless, it is obvious that fewer than all of IMI's papers are classified as open access in this analysis, and this is likely to be due to ancillary factors. See footnote 6 for further explanations.

Table 5.8.15 SUMMARY CITATION INDICATORS FOR IMI 2 CALLS 1-4 PROJECTS, 2015-2021

PROJECT	CITATION IMPACT				% OF HIGHLY CITED PAPERS
	NUMBER OF PAPERS	NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL	AVERAGE PERCENTILE	
INNODIA	166	1.62	1.15	38.30	21.1%
RHAPSODY	94	1.97	1.05	37.51	24.5%
PRISM	75	2.70	1.10	41.21	25.3%
RADAR-CNS	64	1.94	1.66	36.67	31.3%
EBOVAC1	32	2.07	1.42	26.71	28.1%
EbolaMoDRAD	25	1.34	0.95	40.02	16.0%
VAC2VAC	20	0.60	0.66	62.12	5.0%
EBOVAC2	19	2.45	1.38	31.09	26.3%
PERISCOPE	17	1.57	1.64	42.36	29.4%
VSV-EBOVAC	11	0.94	0.62	26.76	9.1%
Overall (IMI projects)	7,856	2.03	1.23	34.98	25.3%

## 5.9 Summary bibliometric analyses for IMI 2 calls 5-10 projects

Figure 5.9.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers from IMI 2 projects from calls 5-10. Only projects with at least 10 papers and one highly cited paper over the time period (2017-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers. The same data is shown in Figure 5.9.1 and Figure 5.9.2, however Figure 5.9.2 has a smaller x-axis range in order to get a better view of the clustered projects in the bottom left corner of Figure 5.9.1.

Figure 5.9.1 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 2 PROJECTS – CALLS 5-10, 2017-2021

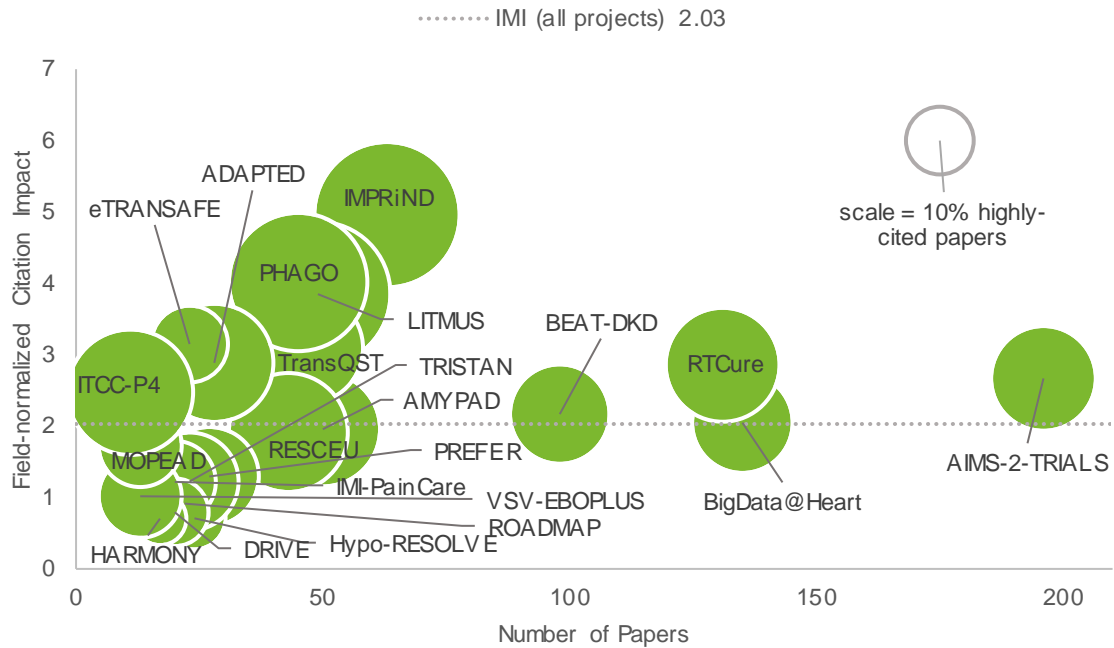
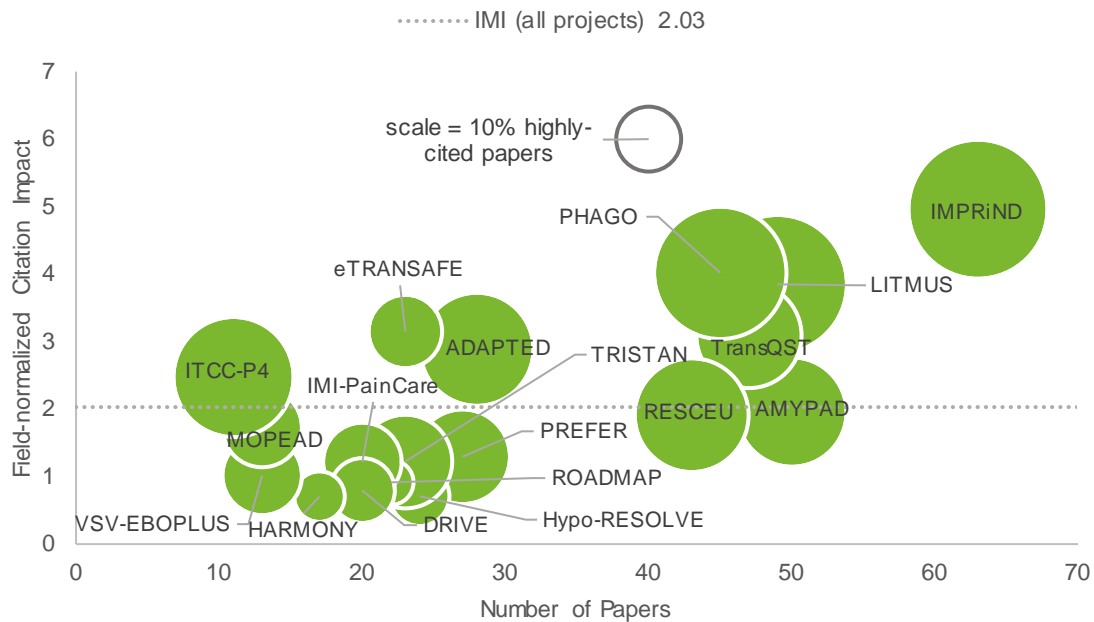


Figure 5.9.2 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 2 PROJECTS – CALLS 5-10, 2017-2020. SMALLER AXIS RANGE.



The data in Figure 5.9.1 and Figure 5.9.2 shows that:

- The AIMS-2-Trials project published the most papers, 196 papers and had a field-normalized citation impact of 2.67, more than 2.5 times higher than the world average (1). IMPRIND remains the top project in terms of field-normalized citation impact with a citation impact of nearly 5 times (4.96) the world average (1). It has (49.2%) of its papers that are highly cited.

Table 5.9.1 shows raw citation impact and percentage of open access papers by project for IMI 2 calls 5-10 publications and Table 5.9.2 shows indicators for IMI 2 calls 5-10 project research where citation impact has been normalised against world average values.

Table 5.9.16 BIBLIOMETRIC INDICATORS FOR IMI 2 CALLS 5-10 PROJECTS, 2017-2021<sup>11</sup>

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
AIMS-2-TRIALS	210	196	92.9%	2,331	11.89
BigData@Heart	157	135	92.6%	1,215	9.00
RTCure	146	131	85.5%	2,718	20.75
BEAT-DKD	106	98	87.8%	1,369	13.97
IMPRiND	66	63	88.9%	3,077	48.84
AMYPAD	56	50	94.0%	584	11.68
LITMUS	59	49	81.6%	786	16.04
TransQST	52	47	83.0%	898	19.11
PHAGO	46	45	100.0%	1,268	28.18
RESCEU	45	43	100.0%	602	14.00
ADAPTED	30	28	92.9%	533	19.04
PREFER	41	27	100.0%	247	9.15
Hypo-RESOLVE	33	24	87.5%	66	2.75
TRISTAN	23	23	91.3%	331	14.39
eTRANSafe	30	23	95.7%	483	21.00
ROADMAP	28	22	100.0%	190	8.64
IMI-PainCare	27	20	90.0%	97	4.85
DRIVE	21	20	70.0%	114	5.70
HARMONY	30	17	88.2%	263	15.47
VSV-EBOPUS	14	13	84.6%	177	13.62
MOPEAD	13	13	100.0%	63	4.85
ITCC-P4	11	11	81.8%	133	12.09

- Most of the projects in IMI 2 Calls 5-10 have more than 80% of their papers as open access, except for DRIVE which only 70% of the projects are open access.

<sup>11</sup> Note that IMI 2 funded researchers are contractually obliged to make their scientific articles open access through Green or Gold routes. However, for some of other document types, such as editorials, reviews or conference proceedings open access publication is strongly encouraged but not mandatory. Nevertheless, it is obvious that fewer than all of IMI's papers are classified as open access in this analysis, and this is likely to be due to ancillary factors. See footnote 6 for further explanations.

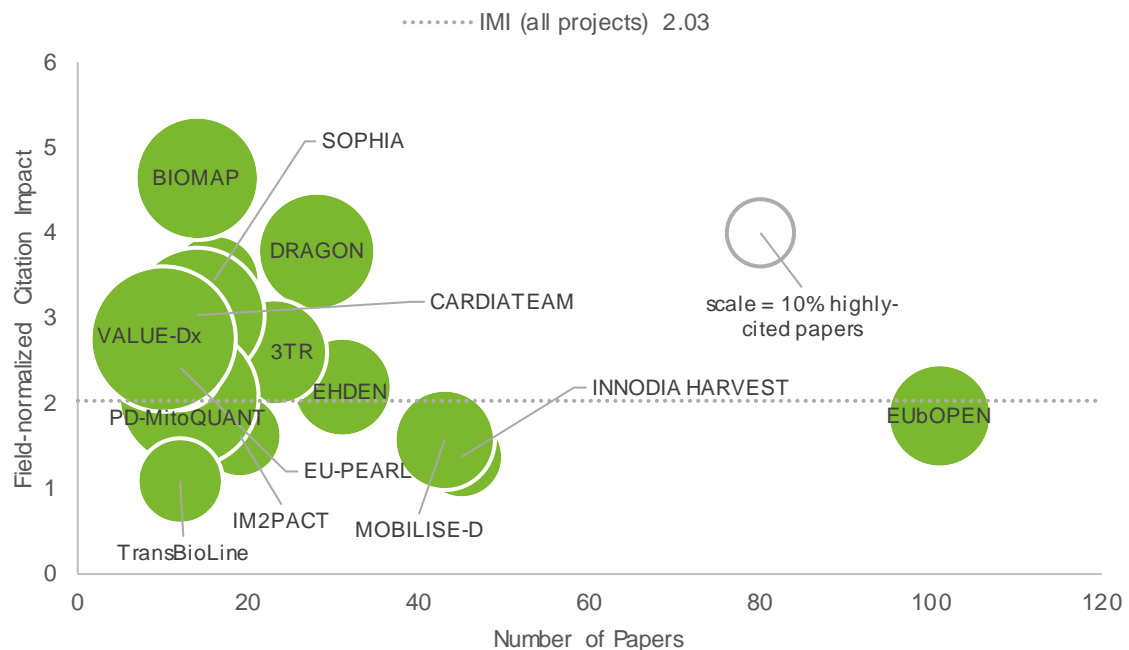
Table 5.9.17 SUMMARY CITATION INDICATORS FOR IMI 2 CALLS 5-10 PROJECTS, 2017-2020

PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
AIMS-2-TRIALS	196	2.67	1.28	42.98	25.0%
BigData@Heart	135	2.05	1.32	45.04	23.0%
RTCure	131	2.86	1.45	33.65	29.8%
BEAT-DKD	98	2.17	0.97	39.22	22.4%
IMPRiND	63	4.96	1.73	22.03	49.2%
AMYPAD	50	1.96	1.00	35.43	30.0%
LITMUS	49	3.85	1.54	34.72	49.0%
TransQST	47	3.09	1.98	37.38	27.7%
PHAGO	45	4.02	1.92	27.88	44.4%
RESCEU	43	1.91	1.17	30.67	32.6%
ADAPTED	28	2.89	1.57	45.15	32.1%
PREFER	27	1.29	1.46	41.48	22.2%
Hypo-RESOLVE	24	0.71	0.54	62.34	8.3%
TRISTAN	23	1.22	0.89	39.48	21.7%
eTRANSafe	23	3.15	1.81	42.05	13.0%
ROADMAP	22	0.91	0.53	47.84	4.5%
IMI-PainCare	20	1.22	1.06	54.06	15.0%
DRIVE	20	0.79	0.80	57.89	10.0%
HARMONY	17	0.70	0.44	55.42	5.9%
VSV-EBOPUS	13	1.02	1.08	27.59	15.4%
MOPEAD	13	1.73	0.58	64.56	15.4%
ITCC-P4	11	2.48	1.20	26.50	36.4%
<b>Overall (IMI projects)</b>	<b>7,856</b>	<b>2.03</b>	<b>1.23</b>	<b>34.98</b>	<b>25.3%</b>

## 5.10 Summary bibliometric analyses for IMI 2 calls 11-23 projects

Figure 5.10.1 compares the number of papers, average field-normalised citation impact and share of highly cited papers from IMI 2 projects from calls 11-23. Only projects with at least 10 papers and one highly cited paper over the time period (2019-2021) are shown. The area of the 'bubble' is proportional to the share of highly cited papers. The dotted horizontal line indicates the average field-normalised citation impact for all IMI project papers.

Figure 5.10.1 PAPER NUMBERS, AVERAGE FIELD-NORMALISED CITATION IMPACT AND SHARE OF HIGHLY CITED RESEARCH FOR SELECTED IMI 2 PROJECTS – CALLS 11-23, 2019-2021



The data in Figure 5.10.1 and Figure shows that:

- Publication rates in calls 11-23 are growing, with EUbOPEN still being the most prolific with 101 papers. Many of these projects are still quite new, and the oldest publication was published in 2019.
- BIOMAP had the highest average field-normalised citation impact which was more than 4 times (4.64) higher than the world average (1). However, the number of papers (14) is still quite low so the field-normalised citation impact should be considered with caution since one highly cited paper can inflate the metric.
- Half (5 out of 10) of Value-Dx papers were highly cited making it the project with the highest percentage of highly cited papers.

Table 5.10.1 shows raw citation impact and percentage of open access papers by project for IMI 2 calls 11-23 publications and Table 5.9.2 shows indicators for IMI 2 calls 11-23 project research where citation impact has been normalised against world average values.

Table 5.10.18 BIBLIOMETRIC INDICATORS FOR IMI 2 CALLS 11-23 PROJECTS, 2019-2021<sup>13</sup>

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	CITATIONS	RAW CITATION IMPACT
EUbOPEN	102	101	77.2%	397	3.93
INNODIA HARVEST	53	45	97.8%	215	4.78
MOBILISE-D	48	43	88.4%	212	4.93
EHDEN	41	31	93.5%	152	4.90
DRAGON	32	28	92.9%	194	6.93
3TR	25	23	78.3%	172	7.48
IM2PACT	19	19	84.2%	45	2.37
SOPHIA	17	16	87.5%	130	8.13
BIOMAP	18	14	78.6%	313	22.36
CARDIATEAM	16	14	100.0%	228	16.29
PD-MitoQUANT	13	13	92.3%	192	14.77
TransBioLine	12	12	91.7%	28	2.33
EU-PEARL	14	12	66.7%	41	3.42
ConcePTION	12	11	81.8%	18	1.64
VALUE-Dx	10	10	100.0%	57	5.70

- Nearly half of the included projects have at least 90% of their papers as open access.
- EU-PEARL and EUbOPEN have the lowest percentage of open access papers, 66.7% and 77.2% respectively.

<sup>13</sup> Note that IMI 2 funded researchers are contractually obliged to make their scientific articles open access through Green or Gold routes. However, for some of other document types, such as editorials, reviews or conference proceedings open access publication is strongly encouraged but not mandatory. Nevertheless, it is obvious that fewer than all of IMI's papers are classified as open access in this analysis, and this is likely to be due to ancillary factors. See footnote 6 for further explanations.

Table 5.10.19 SUMMARY CITATION INDICATORS FOR IMI 2 CALLS 11-23 PROJECTS, 2019-2021

PROJECT	NUMBER OF PAPERS	CITATION IMPACT		AVERAGE PERCENTILE	% OF HIGHLY CITED PAPERS
		NORMALISED AT FIELD LEVEL	NORMALISED AT JOURNAL LEVEL		
EUbOPEN	101	1.85	1.48	53.46	24.8%
INNODIA HARVEST	45	1.37	1.35	47.28	15.6%
MOBILISE-D	43	1.57	1.36	48.50	23.3%
EHDEN	31	2.19	1.58	51.80	22.6%
DRAGON	28	3.78	3.58	40.19	32.1%
3TR	23	2.60	1.26	46.69	26.1%
IM2PACT	19	1.62	1.45	60.16	15.8%
SOPHIA	16	3.45	1.23	69.50	18.8%
BIOMAP	14	4.64	0.91	20.29	35.7%
CARDIATEAM	14	3.03	1.19	33.06	42.9%
PD-MitoQUANT	13	2.08	1.84	28.08	46.2%
EU-PEARL	12	2.42	3.02	61.77	16.7%
TransBioLine	12	1.09	2.52	54.02	16.7%
VALUE-Dx	10	2.75	1.95	39.27	50.0%
Overall (IMI projects)	7,856	2.03	1.23	34.98	25.3%



## 6 GEOGRAPHIC CLUSTERING ANALYSIS

### 6.1 Locations where IMI-funded research takes place

This section of the report analyses geographic clusters where IMI research occurs, the citation impact of research published by these clusters and the clusters' constituent institutions.

Substantial clusters of research activity were identified in Europe and North America. While IMI project research also involves institutions in other parts of the world, publication rates for other geographies were low. This analysis, therefore, focuses on Europe and North America and we have identified the 36 and 18 geographic clusters respectively with the highest output.

Clusters have a 20km radius and the clusters in Europe and North America tend to focus on major cities with an existing strong academic research base. The largest European clusters are London (1,800 publications), Amsterdam (1,515 publications), Stockholm (843 publications), Paris (758 publications) and Oxford (757 publications). The largest clusters in North America are Boston (392 publications), Toronto (368 publications), New York (257 publications), Bethesda (173 publications), and Montreal (138 publications).

IMI research performs well above the national averages for citation impact for all the European and North American clusters. The highest European clusters for citation impact are Maastricht (3.83) and Zurich (3.65) both more than three times their national averages of 1.71 for both.

A relatively high percentage of IMI research is open access, with the Lyon, France cluster being among the highest with 98.8% of its IMI project research as open access papers and Rome being the lowest with over two-thirds (67.9%) of its publications being open access. The USA cluster with the highest percentage of IMI research was Seattle with 94.4 % of its publication being open access.

Around 40% of all EU-28 biomedical research involves international co-authorship while in comparison rates of international collaboration for IMI project research are very high for most clusters, especially in North America where most clusters have around 90% international collaboration which is expected as IMI is European funding organisation that primarily funds researchers working in EU-28. The European cluster with the highest rate of internationally collaborative papers was Basel with 94.5% of its research involving international co-authorship. While the European cluster, Rome, had the lowest at 75.3% international collaboration.

The clusters are visualised on maps in FIGURE 6.1.1 and Figure 6.1.2. Both maps are scaled separately so that the most intensive areas of output are shaded red and the areas of lowest output are blue. This means that the same colour shading is not comparable between maps. Table 6.1.1 to Table 6.1.4 show the research publication outputs of the individual clusters along with bibliometric indicators of their research performance. The citation metrics in Table 6.1.2 and Table 6.1.4 are shaded green when the performance of a cluster of IMI-supported research outperforms the national average performance for biomedical research.<sup>15</sup>

The institutions that constitute the top five clusters within the European and North American regions are shown in Table 6.1.5 and Table 6.1.6 respectively. The five journal subject categories in which the

---

<sup>15</sup> Web of Science journal categories which capture biomedically related publications used to calculate the national baselines are listed in [Annex 2](#).

top five clusters published most frequently within the European and North American regions are shown in Table 6.1.7 and TABLE 6.1.8 respectively.

FIGURE 6.1.1 MAP SHOWING EUROPEAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH, 2010-2021

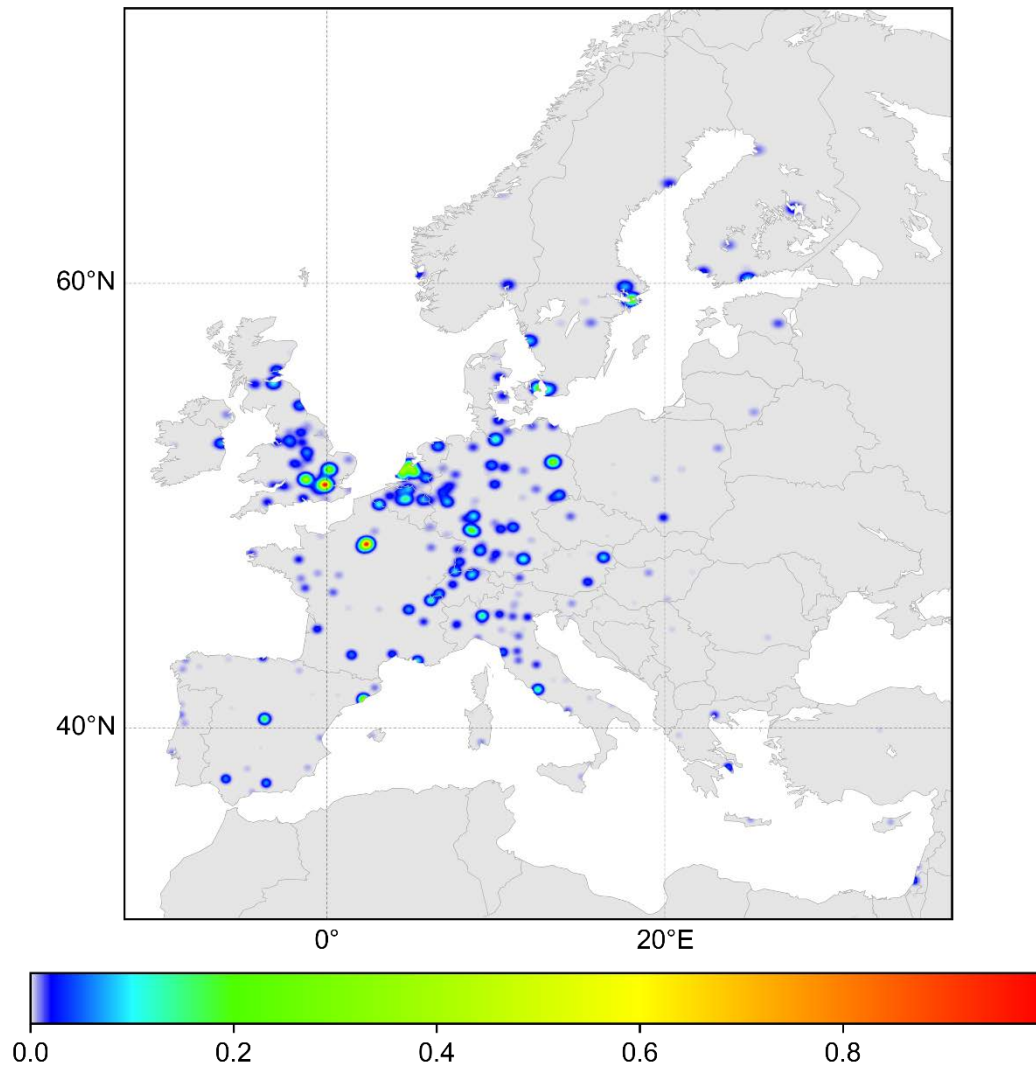


Figure 6.1.2 MAP SHOWING NORTH AMERICAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH, 2010-2021

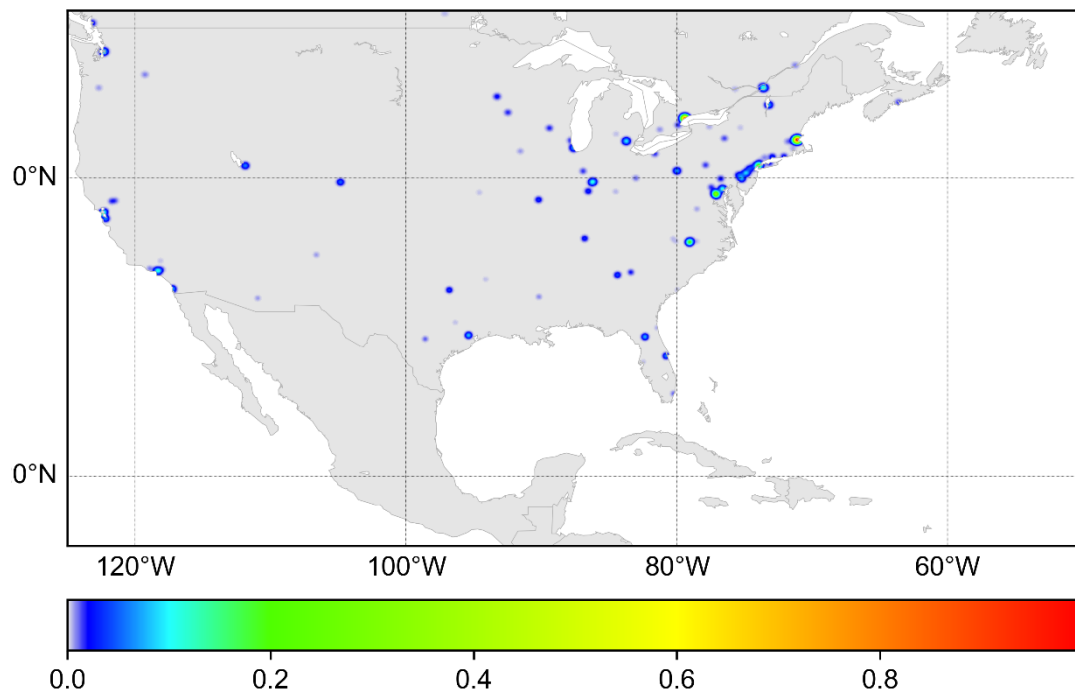


Table 6.1.1 OUTPUT AND RESEARCH PERFORMANCE OF EUROPEAN GEOGRAPHIC CLUSTERS OF IMI  
IMI PROJECT RESEARCH, 2010-2021

CLUSTER	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	RAW CITATION IMPACT	% OF INTERNATIONALLY COLLABORATIVE PUBLICATIONS
London (UK)	1,800	1,634	87.9%	35.60	83.2%
Amsterdam (Netherlands)	1,515	1,371	81.8%	35.79	82.0%
Stockholm (Sweden)	843	781	79.1%	36.92	80.0%
Paris (France)	758	710	81.5%	39.73	84.9%
Oxford (UK)	757	723	93.5%	32.56	83.7%
Cambridge (UK)	666	619	90.0%	45.30	81.6%
Copenhagen (Denmark)	646	596	75.0%	31.59	84.2%
Barcelona (Spain)	522	474	84.2%	33.95	80.6%
Berlin (Germany)	432	406	85.0%	37.11	82.3%
Mannheim (Germany)	422	408	79.9%	45.03	86.5%
Leuven (Belgium)	401	360	87.5%	34.90	90.8%
Madrid (Spain)	368	336	86.3%	27.68	78.9%
Basel (Switzerland)	361	327	78.9%	31.93	94.5%
Uppsala (Sweden)	330	312	78.5%	26.05	75.3%
Nijmegen (Netherlands)	320	300	84.3%	37.41	83.0%
Rome (Italy)	316	287	67.9%	38.67	75.3%
Frankfurt (Germany)	310	292	74.0%	23.83	84.9%
Vienna (Austria)	307	282	81.6%	25.41	84.0%
Milan (Italy)	306	262	77.1%	38.65	84.4%
Groningen (Netherlands)	305	287	92.3%	33.64	82.2%
Hamburg (Germany)	289	262	83.2%	36.33	80.5%
Gothenburg (Sweden)	288	269	80.3%	39.92	90.0%
Geneva (Switzerland)	277	254	86.6%	43.94	87.4%
Munich (Germany)	276	248	75.4%	37.05	83.5%
Maastricht (Netherlands)	268	257	93.8%	66.65	93.0%
Edinburgh (UK)	230	207	93.7%	42.10	82.1%
Helsinki (Finland)	199	191	88.0%	42.61	88.0%
Zurich (Switzerland)	187	173	89.6%	50.49	87.3%
Bonn (Germany)	178	166	89.8%	32.47	79.5%
Lausanne (Switzerland)	167	152	89.5%	37.75	86.2%

CLUSTER	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	RAW CITATION IMPACT	% OF INTERNATIONALLY COLLABORATIVE PUBLICATIONS
Beerse (Belgium)	167	158	76.6%	25.18	91.8%
Dresden (Germany)	165	149	87.9%	32.14	92.6%
Tubingen (Germany)	163	154	77.3%	28.92	76.6%
Marseille (France)	128	116	76.7%	39.62	82.8%
Lille (France)	91	85	74.1%	29.72	88.2%
Lyon (France)	88	80	98.8%	44.41	90.0%

Table 6.1.2 research performance of European geographic clusters of IMI PROJECT RESEARCH COMPARED TO NATIONAL BENCHMARKS, 2010-2021

CLUSTER	FIELD-NORMALISED CITATION IMPACT		JOURNAL-NORMALISED CITATION IMPACT		% OF HIGHLY CITED PAPERS	
	CLUSTER	NATIONAL	CLUSTER	NATIONAL	CLUSTER	NATIONAL
London (UK)	2.58	1.53	1.37	1.10	31.5%	17.2%
Amsterdam (Netherlands)	2.41	1.71	1.32	1.14	28.7%	19.4%
Stockholm (Sweden)	2.26	1.62	1.29	1.13	27.8%	17.8%
Oxford (UK)	2.54	1.53	1.36	1.10	32.4%	17.2%
Paris (France)	2.83	1.48	1.33	1.08	32.3%	15.5%
Cambridge (UK)	3.33	1.53	1.48	1.10	35.9%	17.2%
Copenhagen (Denmark)	2.28	1.71	1.16	1.15	26.2%	18.4%
Barcelona (Spain)	2.86	1.36	1.43	1.09	29.7%	14.2%
Mannheim (Germany)	2.92	1.34	1.31	1.08	33.3%	14.8%
Berlin (Germany)	2.87	1.34	1.43	1.08	29.6%	14.8%
Leuven (Belgium)	2.45	1.80	1.48	1.22	30.3%	19.7%
Madrid (Spain)	2.59	1.36	1.45	1.09	26.2%	14.2%
Basel (Switzerland)	2.03	1.71	1.42	1.17	28.1%	19.1%
Uppsala (Sweden)	1.98	1.62	1.17	1.13	23.1%	17.8%
Nijmegen (Netherlands)	2.55	1.71	1.33	1.14	32.7%	19.4%
Frankfurt (Germany)	2.04	1.34	1.24	1.08	27.4%	14.8%
Groningen (Netherlands)	2.38	1.71	1.13	1.14	24.4%	19.4%
Rome (Italy)	2.69	1.40	1.81	1.19	35.2%	15.3%
Vienna (Austria)	2.17	1.59	1.30	1.16	25.9%	17.4%
Gothenburg (Sweden)	2.88	1.62	1.59	1.13	36.1%	17.8%
Hamburg (Germany)	2.61	1.34	1.16	1.08	30.9%	14.8%
Milan (Italy)	2.80	1.40	1.42	1.19	36.6%	15.3%
Maastricht (Netherlands)	3.83	1.71	1.84	1.14	34.2%	19.4%
Geneva (Switzerland)	2.93	1.71	1.06	1.17	33.9%	19.1%
Munich (Germany)	3.04	1.34	1.37	1.08	33.1%	14.8%
Edinburgh (UK)	2.69	1.53	1.40	1.10	33.3%	17.2%
Helsinki (Finland)	3.01	1.60	1.38	1.10	39.8%	16.9%
Zurich (Switzerland)	3.65	1.71	1.51	1.17	38.7%	19.1%
Bonn (Germany)	2.42	1.34	1.47	1.08	25.3%	14.8%
Beerse (Belgium)	1.98	1.80	1.31	1.22	23.4%	19.7%
Tubingen (Germany)	3.03	1.34	1.24	1.08	31.8%	14.8%

CLUSTER	FIELD-NORMALISED CITATION IMPACT		JOURNAL-NORMALISED CITATION IMPACT		% OF HIGHLY CITED PAPERS	
	CLUSTER	NATIONAL	CLUSTER	NATIONAL	CLUSTER	NATIONAL
Lausanne (Switzerland)	2.92	1.71	1.17	1.17	27.0%	19.1%
Dresden (Germany)	2.72	1.34	0.98	1.08	25.5%	14.8%
Marseille (France)	2.72	1.48	1.45	1.08	36.2%	15.5%
Lille (France)	1.87	1.48	0.93	1.08	29.4%	15.5%
Lyon (France)	3.05	1.48	1.32	1.08	38.8%	15.5%

Table 6.1.3 OUTPUT AND RESEARCH PERFORMANCE OF NORTH AMERICAN GEORGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH, 2010-2021

CLUSTER	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	% OF OPEN ACCESS PAPERS	RAW CITATION IMPACT	% OF INTERNATIONALLY COLLABORATIVE PAPERS
Boston (USA)	392	376	87.2%	57.18	98.4%
Toronto (Canada)	368	360	87.5%	43.93	90.3%
New York (USA)	257	252	84.1%	54.67	99.6%
Bethesda (USA)	173	166	80.7%	56.70	98.2%
Montreal (Canada)	138	137	86.9%	48.02	98.5%
Chapel Hill (USA)	138	135	93.3%	35.96	88.1%
Indianapolis (USA)	125	116	78.4%	45.19	98.3%
San Francisco (USA)	107	103	86.4%	88.06	100.0%
Burlington (USA)	91	89	88.8%	28.81	100.0%
Baltimore (USA)	88	85	92.9%	79.05	100.0%
Los Angeles (USA)	86	85	85.9%	62.93	98.8%
Titusville (USA)	76	69	73.9%	16.04	97.1%
Seattle (USA)	72	71	94.4%	67.28	98.6%
Philadelphia (USA)	68	65	92.3%	57.75	98.5%
Ann Arbor (USA)	64	62	90.3%	75.16	98.4%
La Jolla (USA)	63	62	91.9%	68.39	100.0%
Houston (USA)	56	54	90.7%	55.37	100.0%
Gainesville (USA)	48	44	68.2%	38.16	97.7%

Table 6.1.4 RESEARCH PERFORMANCE OF NORTH AMERICAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH COMPARED TO NATIONAL BENCHMARKS, 2010-2021

CLUSTER	FIELD-NORMALISED CITATION IMPACT		JOURNAL-NORMALISED CITATION IMPACT		% OF HIGHLY CITED PAPERS	
	CLUSTER	NATIONAL	CLUSTER	NATIONAL	CLUSTER	NATIONAL
Boston (USA)	3.87	1.32	1.43	1.03	38.8%	15.4%
Toronto (Canada)	2.93	1.49	1.42	1.08	34.2%	16.1%
New York (USA)	4.53	1.32	1.50	1.03	34.9%	15.4%
Bethesda (USA)	3.52	1.32	1.51	1.03	45.2%	15.4%
Montreal (Canada)	3.32	1.49	1.17	1.08	33.6%	16.1%
Chapel Hill (USA)	3.16	1.32	1.26	1.03	31.9%	15.4%
Indianapolis (USA)	3.64	1.32	1.44	1.03	33.6%	15.4%
San Francisco (USA)	6.57	1.32	2.00	1.03	52.4%	15.4%
Burlington (USA)	1.79	1.32	0.79	1.03	21.3%	15.4%
Baltimore (USA)	6.33	1.32	1.60	1.03	50.6%	15.4%
Los Angeles (USA)	5.43	1.32	1.63	1.03	44.7%	15.4%
Seattle (USA)	5.14	1.32	2.00	1.03	46.5%	15.4%
Titusville (USA)	1.89	1.32	1.30	1.03	27.5%	15.4%
Philadelphia (USA)	6.55	1.32	2.00	1.03	43.1%	15.4%
La Jolla (USA)	5.68	1.32	1.45	1.03	50.0%	15.4%
Ann Arbor (USA)	6.86	1.32	2.13	1.03	58.1%	15.4%
Houston (USA)	4.52	1.32	1.91	1.03	55.6%	15.4%
Gainesville (USA)	2.14	1.32	1.65	1.03	45.5%	15.4%

Table 6.1.5 INSTITUTIONS CONSTITUTING TOP-FIVE, BY NUMBER OF PUBLICATIONS, EUROPEAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH, 2010-2021

CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
London	UK	King's College London	723
		Imperial College London	492
		University College London	454
		GlaxoSmithKline	116
		South London & Maudsley NHS Trust	89
		Birkbeck University London	84



CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		London School of Hygiene & Tropical Medicine	77
		Guy's & St Thomas' NHS Foundation Trust	73
		Royal Brompton Hospital	71
		University College London Hospitals NHS Foundation Trust	58
		Queen Mary University London	53
		St Georges University London	46
		Royal Brompton & Harefield NHS Foundation Trust	41
		The Medicines & Healthcare Products Regulatory Agency	27
		UK Research and Innovation, India	24
		Medical Research Council UK (MRC)	24
		Francis Crick Institute	24
		Royal Marsden NHS Foundation Trust	22
		Alan Turing Inst	22
		UCB Pharma SA	20
		Institute of Cancer Research - UK	20
		European Med Agcy	18
		London School Economics & Political Science	16
		National Institute for Health & Care Excellence	15
		Hlth Data Res UK	14
		South London & Maudsley NHS Fdn	14
		University of Westminster	13
		UCL Medical School	13
		National Institute for Biological Standards & Control	12
		HEPTARES THERAPEUT LTD	11
		King's College Hospital	11
		EMA	10
		Moorfields Eye Hospital NHS Foundation Trust	10
		Public Health England	9
		University of London Royal Veterinary College	9
		Royal Coll Gen Practitioners	9
		South London & Maudsley NHS Fdn Trust Slam	9
		Royal London Hospital	9
		Takeda Pharmaceutical Company Ltd	9
		TAKEDA DEV CTR EUROPE LTD	9

CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		Cancer Research UK	9
		King's College Hospital NHS Foundation Trust	9
		City University London	8
		Celltech Grp	8
		MRC Social Genet & Dev Psychiat SGDP Ctr	8
		St Georges Univ Hosp NHS Fdn Trust	7
		University of London	7
		Inst Psychiat Psychol & Neurosci	7
		AMGEN LTD	7
		Amgen	7
		Royal Brompton NIHR Biomed Res Unit	7
		South London & Maudsley Fdn NHS Trust	6
		UK Dementia Res Inst	6
		Genet Alliance UK	5
		Barts Health NHS Trust	5
		UK Research & Innovation (UKRI)	3
<b>Amsterdam</b>	<b>Netherlands</b>	Leiden University	409
		Utrecht University Medical Center	351
		Vrije Universiteit Amsterdam	328
		Erasmus MC	270
		Academic Medical Center Amsterdam	239
		University of Amsterdam	204
		Utrecht University	149
		VU UNIVERSITY MEDICAL CENTER	106
		Netherlands National Institute for Public Health & the Environment	55
		Erasmus University Rotterdam	35
		Delft University of Technology	16
		Wilhelmina Kinderziekenhuis	14
		Emma Children's Hospital	14
		Netherlands Cancer Institute	13
		Janssen Vaccines & Prevent BV	11
		ICIN Netherlands Heart Inst	10
		ReSViNET Fdn	10
		Netherlands Heart Inst	10

CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		Lygature	9
		Med Evaluat Board	9
		Netherlands Institute for Health Services Research	9
		Leiden Univ Med Ctr	8
		GGz inGeest	7
		Jan van Breemen Res Inst Reade	6
		St. Antonius Hospital Utrecht	6
		Amsterdam Univ Med Ctr	6
		Weibel Consulting	5
		Erasmus MC - Sophia Children's Hospital	2
		Leiden Univ Excl LUMC	1
<b>Stockholm</b>	<b>Sweden</b>	Karolinska Institutet	641
		Karolinska University Hospital	292
		Royal Institute of Technology	76
		Stockholm University	55
		Stockholm County Council	55
		Stockholm Hlth Care Serv	12
		Danderyds Hospital	10
		Publ Hlth Agcy Sweden	9
		AstraZeneca	8
		SciLifeLab	7
<b>Paris</b>	<b>France</b>	Universite de Paris	432
		Institut National de la Sante et de la Recherche Medicale (Inserm)	413
		UDICE French Res Univ	314
		Sorbonne Universite	205
		CEA	149
		Hopital Universitaire Cochin - APHP	115
		CNRS - National Institute for Biology (INSB)	103
		Hopital Universitaire Pitie-Salpetriere - APHP	101
		Centre National de la Recherche Scientifique (CNRS)	94
		Institut Pasteur Paris	72
		Sanofi France	52
		Institut de Recherches Internationales Servier	36
		Hopital Universitaire Bichat-Claude Bernard - APHP	32

CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		Assistance Publique Hopitaux Paris (APHP)	29
		Institut Curie	24
		Hopital Universitaire Bicetre - APHP	22
		Hopital Universitaire Saint-Louis - APHP	22
		Orsay Hosp	21
		Gustave Roussy	20
		Hopital Universitaire Necker-Enfants Malades - APHP	17
		Hopital Universitaire Europeen Georges-Pompidou - APHP	17
		CNRS - Institute of Chemistry (INC)	14
		Hopital Universitaire Saint-Antoine - APHP	14
		Universite Grenoble Alpes (UGA)	14
		Hopital Universitaire Beaujon - APHP	13
		Assistance Publique-Hopitaux de Marseille	12
		Hopital Universitaire Henri-Mondor - APHP	12
		Hopital Universitaire Paul-Brousse - APHP	12
		Hopital Universitaire Robert-Debre - APHP	11
		Museum National d'Histoire Naturelle (MNHN)	10
		CNRS - Institute of Ecology & Environment (INEE)	9
		Servier	7
		Hopital Universitaire Ambroise-Pare - APHP	6
		SOLEIL Synchrotron	6
		Univ Paris Est ComUE	6
		EURORDIS Rare Dis Europe	5
		Universite Paris 13	5
		Vaccine Res Inst	5
		Universite Paris Saclay	4
		Sanofi-Aventis	2
		Universite de Versailles Saint-Quentin-En-Yvelines	1
		Aix-Marseille Universite	1
<b>Oxford</b>	<b>UK</b>	University of Oxford	694
		Wellcome Centre for Human Genetics	100
		Oxford University Hospitals NHS Foundation Trust	30
		Diamond Light Source	26
		Ludwig Institute for Cancer Research	13



CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		Tufts University	6
		Northeastern University	5
<b>Toronto</b>	<b>Canada</b>	University of Toronto	211
		Structural Genomics Consortium	163
		Princess Margaret Cancer Centre	81
		Hospital for Sick Children (SickKids)	81
		Baycrest	67
		Centre for Addiction & Mental Health - Canada	32
		Ontario Institute for Cancer Research	26
		Holland Bloorview Kids Rehabilitation Hospital	20
		University Health Network Toronto	19
		Lunenfeld Tanenbaum Research Institute	17
		Toronto General Hospital	8
		Saint Michaels Hospital Toronto	6
<b>New York</b>	<b>USA</b>	Icahn School of Medicine at Mount Sinai	79
		Columbia University	63
		Pfizer	46
		New York University	32
		Northwell Health	19
		Memorial Sloan Kettering Cancer Center	18
		Albert Einstein College of Medicine	17
		NewYork-Presbyterian Hospital	14
		NYU Langone Medical Center	8
		Yeshiva University	6
		Rutgers State University New Brunswick	4
<b>Bethesda</b>	<b>USA</b>	AstraZeneca	24
		NIH National Heart Lung & Blood Institute (NHLBI)	22
		National Institutes of Health (NIH) - USA	22
		NIH National Institute of Mental Health (NIMH)	17
		NIH National Institute of Allergy & Infectious Diseases (NIAID)	15
		NIH National Cancer Institute (NCI)	13
		US Food & Drug Administration (FDA)	12
		NIH National Institute on Aging (NIA)	11

CLUSTER	COUNTRY	INSTITUTIONS	NUMBER OF PUBLICATIONS
		NIH National Human Genome Research Institute (NHGRI)	10
		Medimmune	9
		NIH National Institute of Arthritis & Musculoskeletal & Skin Diseases (NIAMS)	8
		NIH National Institute of Neurological Disorders & Stroke (NINDS)	8
		NIH National Institute of Diabetes & Digestive & Kidney Diseases (NIDDK)	7
		George Washington University	6
		Naval Research Laboratory	5
		GlaxoSmithKline	5
<b>Montreal</b>	<b>Canada</b>	University of Montreal	92
		McGill University	81
		CHU St Justine	17

Table 6.1.7 FIVE JOURNAL SUBJECT CATEGORIES IN WHICH TOP-FIVE, BY NUMBER OF PUBLICATIONS, EUROPEAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH PUBLISHED MOST FREQUENTLY, 2010-2021

CLUSTER	COUNTRY	JOURNAL SUBJECT CATEGORY	NUMBER OF PUBLICATIONS
<b>London</b>	<b>United Kingdom</b>	Neurosciences	370
		Psychiatry	203
		Clinical Neurology	190
		Pharmacology & Pharmacy	158
		Immunology	130
<b>Amsterdam</b>	<b>Netherlands</b>	Pharmacology & Pharmacy	186
		Rheumatology	180
		Neurosciences	173
		Immunology	159
		Clinical Neurology	113
<b>Stockholm</b>	<b>Sweden</b>	Rheumatology	130
		Immunology	101
		Neurosciences	97
		Clinical Neurology	82
		Biochemistry & Molecular Biology	64

CLUSTER	COUNTRY	JOURNAL SUBJECT CATEGORY	NUMBER OF PUBLICATIONS
Paris	France	Neurosciences	145
		Psychiatry	70
		Pharmacology & Pharmacy	61
		Biochemistry & Molecular Biology	60
		Endocrinology & Metabolism	60
Oxford	UK	Biochemistry & Molecular Biology	145
		Neurosciences	104
		Endocrinology & Metabolism	68
		Cell Biology	64
		Chemistry, Medicinal	62

TABLE 6.1.8 FIVE JOURNAL SUBJECT CATEGORIES IN WHICH TOP-FIVE, BY NUMBER OF PUBLICATIONS, NORTH AMERICAN GEOGRAPHIC CLUSTERS OF IMI PROJECT RESEARCH PUBLISHED MOST FREQUENTLY, 2010-2021

CLUSTER	COUNTRY	JOURNAL SUBJECT CATEGORY	NUMBER OF PUBLICATIONS
Boston	USA	Neurosciences	52
		Genetics & Heredity	46
		Endocrinology & Metabolism	40
		Biochemistry & Molecular Biology	38
		Clinical Neurology	36
Toronto	Canada	Biochemistry & Molecular Biology	99
		Neurosciences	73
		Psychiatry	65
		Chemistry, Medicinal	41
		Cell Biology	38
New York	USA	Pharmacology & Pharmacy	44
		Neurosciences	39
		Psychiatry	37
		Genetics & Heredity	25
		Immunology	20



CLUSTER	COUNTRY	JOURNAL SUBJECT CATEGORY	NUMBER OF PUBLICATIONS
Bethesda	USA	Pharmacology & Pharmacy	32
		Immunology	26
		Neurosciences	23
		Psychiatry	21
		Public, Environmental & Occupational Health	21
Montreal	Canada	Neurosciences	46
		Psychiatry	44
		Biochemistry & Molecular Biology	19
		Psychology, Developmental	13
		Genetics & Heredity	13

## 7 COLLABORATION ANALYSIS FOR IMI RESEARCH

### 7.1 Collaboration analysis for IMI research

International research collaboration is increasing<sup>16</sup> and although the reasons for this have not been fully clarified they are likely to include increasing access to facilities, resources, knowledge, people and expertise. In addition, international collaboration has been shown to be associated with an increase in the number of citations received by research papers, although this does depend upon the partner countries involved.<sup>17</sup> Co-authorship is likely to be a good indicator of collaboration, although there will be research collaborations that do not result in co-authored papers, and co-authored papers which may have required limited collaboration. Alternative data-based approaches, for example using information about co-funding or international exchanges, have limitations in terms of both comprehensiveness and validity.

In this report, co-authorship of papers<sup>18</sup> is used as an indicator of collaboration between different sectors, institutions and countries.

In this analysis, different institutions/organisation are assigned to sectors with the following definitions:

<sup>16</sup> Adams J (2013) Collaborations: the fourth age of research. *Nature*, **497**, 557-560.

<sup>17</sup> Adams, J., Gurney, K., & Marshall, S. (2007). Patterns of international collaboration for the UK and leading partners. A report by Evidence Ltd to the UK Office of Science and Innovation. 27pp.

<sup>18</sup> In the collaboration analysis papers rather than publications are analysed as some publications, such as editorials do not communicate novel research finding so cannot be considered a product of research collaboration.

- **Medical:** Organisations with the primary function of providing patient care. Typical these are public, private and university hospitals, though we have included in this sector Chinese medicine hospitals and umbrella organisations such as hospital systems (e.g., Mt Sinai) or UK National Health Services Healthcare Trusts.
- **Corporate:** Private or public companies or enterprises that operate for-profit. For IMI projects most corporate organisations are pharmaceuticals, others manufacture medical devices or provide information technology services. Included in this sector are any organisation with a suffix indicating limited liability (e.g., AB, LTD, GmbH, SA, LLC, INC and AG). Other organisations were identified as corporate from their website. It can be challenging to assign smaller organisations, potential small and medium sized enterprises (SMEs) to this category as they may have a limited online presence and if a SME has spun out from a university it can be difficult to ascertain the current relationship between the spin out and academic institution.
- **Academic:** Public and private universities and university departments. This includes research institutes, that may not have a teaching remit but have a clear affiliation to one or more universities and programs of research spanning multiple academic institutions.
- **Government:** Includes state, regional or federally funded research institutions, laboratories and facilities such as NIH or the World Health Organization (WHO); country or regional funders that disperse public money to research (e.g., BBSRC in the UK); government departments and agencies.
- **Other:** Organisation that do not fit in any other sector but have a role in the healthcare or research infrastructure. For example, research institutions not attached to a government, university or hospital; non-governmental organisations like patient groups, advocacy groups, not-for profits and charities; professional associations for healthcare professionals; non-governmental funders; regulators and tissue sample banks.
- **Unknown:** If an organisation cannot be identified as belonging to any of the other sectors, then it is assigned as unknown.

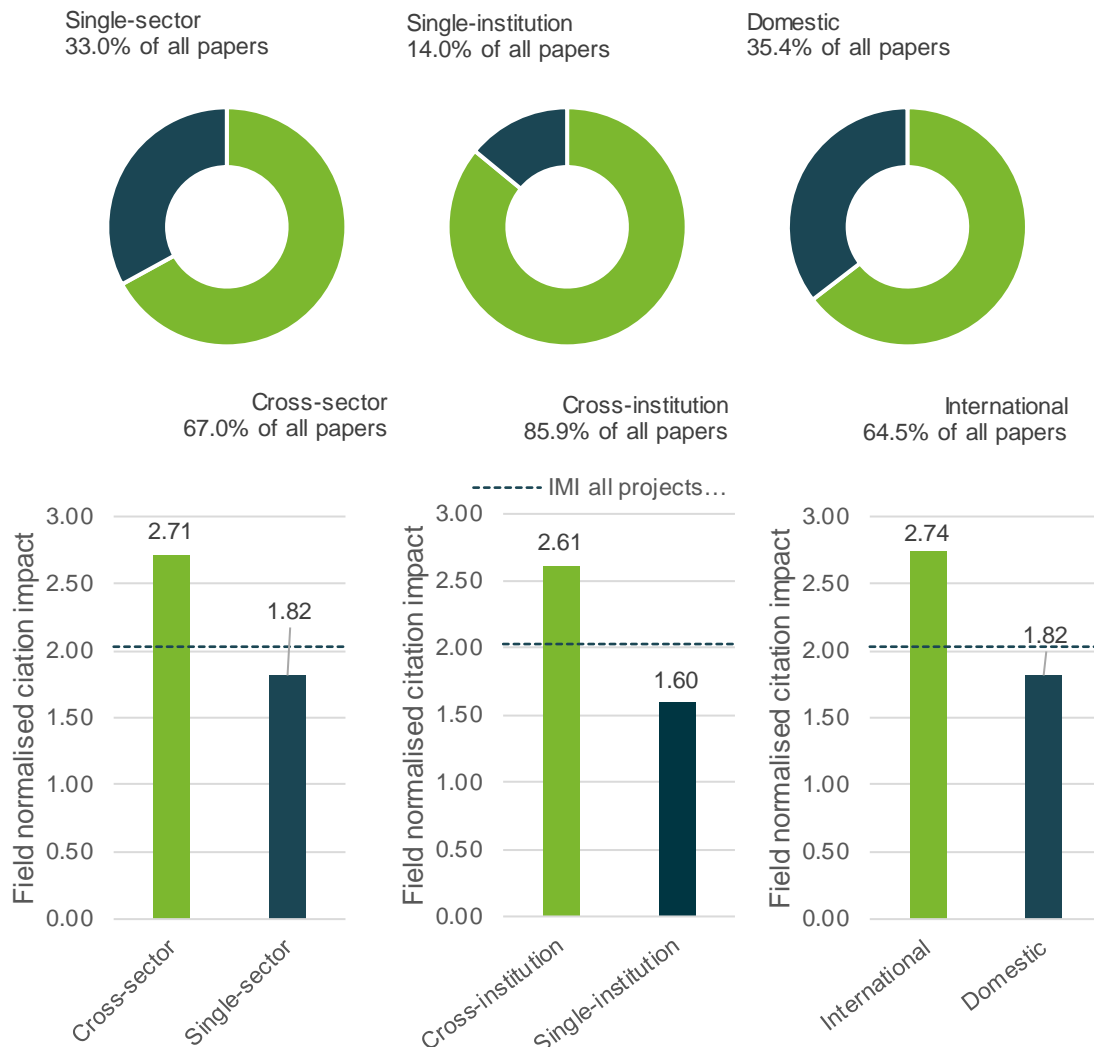
A paper is defined as cross-sector if the co-authors are affiliated to organisations that are assigned to different sectors. For example, if a paper has author addresses corresponding to the University of Copenhagen (academic) and the company Novartis (corporate), it would be classified as cross-sector. If a paper only has author addresses corresponding to the University of Cambridge (academic) and Utrecht University (academic), it would be classified as single-sector since both addresses are academic institutions, but it would be defined as cross-institution as more than one institution is listed in the addresses. A paper is defined as international if more than one country is listed in the addresses, or domestic if only a single country is listed.

The data in Table 7.1.1 compares the output and field-normalised citation impact of collaborative IMI project research with its non-collaborative research. Figure 7.1.1 presents the same data visually.

Table 7.1.1 CROSS-SECTOR, CROSS-INSTITUTION AND INTERNATIONAL OUTPUT AND FIELD-NORMALISED CITATION IMPACT OF IMI PROJECT RESEARCH, 2010-2021

	NUMBER OF PAPERS	% OF PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
Cross-sector	5,261	67.0%	2.71
Single-sector	2,589	33.0%	1.82
Cross-institution	6,749	85.9%	2.61
Single-institution	1,102	14.0%	1.60
International	5,069	64.5%	2.74
Domestic	2,782	35.4%	1.82

Figure 7.1.1 FIELD-NORMALISED CITATION IMPACT AND PERCENTAGE OF CROSS-SECTOR, CROSS-INSTITUTION AND INTERNATIONAL COLLABORATIVE PAPERS FROM IMI PROJECT RESEARCH, 2010-2021



- Nearly two-thirds of (67%) of all IMI project papers were published by co-authors working in different sectors.
- The majority (85.9%) of IMI project papers involved collaboration between different institutions.
- More than half (64.5%) of all IMI project papers involved international collaboration.
- Collaborative IMI project research was internationally influential with field-normalised citation impacts over 2.5-times the world average (1.00), regardless of the type of collaborations.
- IMI's collaborative research has an average field-normalised citation impact that is almost 50% higher than IMI's non-collaborative research and the non-collaborative research field-normalised citation impact was below average for IMI project research (2.03).

## 7.2 Collaboration analysis by IMI project

This section analyses the collaboration of IMI research at the individual project level.

Table 7.2.1 shows the number, percentage, and field-normalised citation impact of IMI research papers with co-authors from more than one country. Table 7.2.2 shows number, percentage, and field-normalised citation impact of IMI research papers with co-authors from more than one institution. Table 7.2.3 shows number, percentage, and field-normalised citation impact of IMI research papers with co-authors from more than one sector.

Figure 7.2.1 to Figure 7.2.5 are maps showing international collaboration for the five IMI projects with the highest number of papers: BTCURE, EU-AIMS, ULTRA-DD, EMIF, and NEWMEDS. The countries with the most frequent collaboration are the darkest shade of blue and gradually gets lighter the less collaboration there is.

It should be noted that the last column in Table 7.2.1 to Table 7.2.3 shows the field-normalised citation impact of those papers involving collaboration of the type being analysed, rather than for all papers belonging to a project. Therefore, in Table 7.2.1, the last column contains the field-normalised citation impact of only the internationally collaborative papers for each project. Similarly, the last column in Table 7.2.2 contains only the field-normalised citation impact of the papers with co-authors from more than one institution, and in Table 7.2.3, the last column contains only the field-normalised citation impact of cross-sector papers.

The key findings of Section 7.2 are:

- BTCURE had the highest number of papers with co-authors from more than one country, institution and sector (Table 7.1.1-Table 7.2.3). This may be due to BTCURE having the highest overall number of papers.
- EU-AIMS had the second highest number of papers with authors from more than one country, institution and sector (Table 7.1.1-Table 7.2.3). Again, this also may be due to EU-AIMS having the second highest overall number of papers
- For those projects with at least 100 papers, BigData@Heart has the highest percentage of its papers that are co-authors from more than one country (76.3%), institution (98.5%) and sector (90.4%).

- The majority of collaborative papers from the top five projects were co-authored with researchers from the United States (USA), Germany and the UK (Figure 7.2.1 to Figure 7.2.5).
- In general, there is a high level of collaboration within Europe for all of the top five projects. The most frequently collaborating European countries were the UK, Sweden, the Netherlands, France and Germany.
- EU-AIMS, NEWMEDS and ULTRA-DD had substantial input from Canadian researchers and ULTRA-DD had a noteworthy amount of collaboration from Chinese researchers (Figure 7.2.2- Figure 7.2.5).

Table 7.2.2 NUMBER, PERCENTAGE AND CITATION IMPACT<sup>19</sup> OF IMI-SUPPORTED RESEARCH PAPERS WITH AUTHORS FROM MORE THAN ONE COUNTRY, 2010-2021

PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
BTCure	672	398	59.2%	2.11
EU-AIMS	546	388	71.1%	2.44
ULTRA-DD	425	318	74.8%	2.22
EMIF	310	232	74.8%	2.94
NEWMEDS	214	137	64.0%	2.42
AIMS-2-TRIALS	196	145	74.0%	4.32
EUROPAIN	181	77	42.5%	3.42
CANCER-ID	180	92	51.1%	4.12
ORBITO	168	94	56.0%	1.78
INNODIA	166	121	72.9%	2.16
TRANSLOCATION	164	93	56.7%	1.68
STEMBANCC	147	83	56.5%	2.22
IMIDIA	141	81	57.4%	1.95
SUMMIT	136	93	68.4%	1.74
BigData@Heart	135	103	76.3%	2.74
ELF	134	76	56.7%	1.11
RTCure	131	70	53.4%	4.13
CHEM21	128	46	35.9%	2.33
PreDiCT-TB	118	69	58.5%	1.47
SPRINTT	116	73	62.9%	2.06

<sup>19</sup> The last column is the citation impact of only the internationally collaborative papers.

PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
MIP-DILI	108	57	52.8%	2.24
Quic-Concept	103	68	66.0%	6.74
EUbOPEN	101	64	63.4%	3.45
COMBACTE-NET	100	59	59.0%	1.30
PROTECT	99	71	71.7%	1.20
BEAT-DKD	98	74	75.5%	1.67
COMBACTE-MAGNET	97	65	67.0%	1.43
RHAPSODY	94	68	72.3%	2.79
U-BIOPRED	93	70	75.3%	3.03
eTOX	92	38	41.3%	1.91
COMPACT	91	50	54.9%	2.36
Pharma-Cog	88	71	80.7%	1.34
DIRECT	82	64	78.0%	4.66
ABIRISK	79	42	53.2%	1.30
DDMoRe	77	51	66.2%	1.27
PRISM	75	57	76.0%	4.26
AETIONOMY	71	37	52.1%	2.16
BioVacSafe	70	39	55.7%	1.39
Open PHACTS	70	43	61.4%	3.66
K4DD	68	40	58.8%	1.96
None	65	50	76.9%	3.87
Onco Track	65	32	49.2%	3.07
RADAR-CNS	64	50	78.1%	2.20
IMPRIND	63	42	66.7%	7.11
COMBACTE-CARE	61	43	70.5%	1.77
ZAPI	60	43	71.7%	5.80
MARCAR	60	30	50.0%	1.20
ENABLE	55	28	50.9%	1.28
EPAD	55	39	70.9%	1.68
DRIVE-AB	54	38	70.4%	1.39
APPROACH	54	46	85.2%	2.52
PRECISESADS	54	46	85.2%	1.54

PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
AMYPAD	50	41	82.0%	2.49
LITMUS	49	34	69.4%	5.44
TransQST	47	35	74.5%	2.79
RAPP-ID	46	25	54.3%	0.89
Prelect	45	33	73.3%	2.04
PHAGO	45	31	68.9%	3.71
INNODIA HARVEST	45	33	73.3%	2.14
eTRIKS	45	42	93.3%	2.37
FLUCOP	44	25	56.8%	1.29
RESCEU	43	33	76.7%	2.24
MOBILISE-D	43	29	67.4%	2.43
iPiE	41	13	31.7%	1.49
GETREAL	37	30	81.1%	1.55
iABC	33	24	72.7%	1.61
EBiSC	33	24	72.7%	1.65
EBOVAC1	32	22	68.8%	2.31
EHDEN	31	28	90.3%	2.72
PROACTIVE	29	25	86.2%	2.55
DRAGON	28	24	85.7%	4.88
ADAPTED	28	19	67.9%	3.93
ADVANCE	27	24	88.9%	1.30
PREFER	27	25	92.6%	1.64
EbolaMoDRAD	25	15	60.0%	1.43
Hypo-RESOLVE	24	20	83.3%	1.37
eTRANSafe	23	12	52.2%	1.12
TRISTAN	23	13	56.5%	1.48
3TR	23	12	52.2%	5.00
ROADMAP	22	17	77.3%	0.66
SAFE-T	21	12	57.1%	2.01
VAC2VAC	20	13	65.0%	0.82
DRIVE	20	7	35.0%	1.81
IMI-PainCare	20	12	60.0%	1.68

PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
EHR4CR	20	13	65.0%	1.40
EBOVAC2	19	11	57.9%	3.80
IM2PACT	19	10	52.6%	3.40
HARMONY	17	9	52.9%	0.99
PERISCOPE	17	7	41.2%	1.60
COMBACTE	16	2	12.5%	12.48
SOPHIA	16	10	62.5%	10.39
WEB-RADR	16	13	81.3%	1.50
BIOMAP	14	11	78.6%	4.75
CARDIATEAM	14	14	100.0%	3.54
VSV-EBOPLUS	13	11	84.6%	1.13
PD-MitoQUANT	13	7	53.8%	2.44
MOPEAD	13	11	84.6%	2.49
EU-PEARL	12	9	75.0%	1.58
TransBioLine	12	5	41.7%	2.00
ConcePTION	11	10	90.9%	1.01
VSV-EBOVAC	11	8	72.7%	1.06
ITCC-P4	11	9	81.8%	2.84
VALUE-Dx	10	9	90.0%	3.44
CARE	9	7	77.8%	9.23
VITAL	9	5	55.6%	0.32
c4c	9	9	100.0%	1.04
COMBACTE-CDI	9	9	100.0%	1.55
EQIPD	9	8	88.9%	3.04
MAD-CoV 2	8	8	100.0%	2.99
EBODAC	8	7	87.5%	2.39
ERA4TB	8	6	75.0%	2.78
RADAR-AD	8	4	50.0%	1.65
PARADIGM	7	6	85.7%	1.88
FAIRplus	7	2	28.6%	3.39
EUPATI	7	7	100.0%	0.75
EBOVAC3	7	7	100.0%	1.06



PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
ReSOLUTE	7	3	42.9%	0.91
KRONO	7	1	14.3%	0.53
NECESSITY	6	5	83.3%	2.66
EBISC2	6	6	100.0%	0.58
IDEA-FAST	6	6	100.0%	1.32
NeuroDeRisk	6	1	16.7%	0.24
MACUSTAR	6	4	66.7%	2.37
Immune-Image	5	5	100.0%	2.67
HIPPOCRATES	5	1	20.0%	11.00
iCONSENSUS	5	2	40.0%	1.25
EBOMAN	4	4	100.0%	3.88
MELLODDY	4	3	75.0%	0.57
DO->IT	4	4	100.0%	1.19
SafeSciMET	4	4	100.0%	0.89
ADAPT-SMART	4	2	50.0%	1.13
T2EVOLVE	3	1	33.3%	10.38
ND4BB	3	2	66.7%	1.65
IMMUCAN	3	2	66.7%	0.42
Eu2P	3	2	66.7%	0.06
imSAVAR	3	3	100.0%	5.01
ImmUniverse	3	2	66.7%	0.00
OPTIMA	2	1	50.0%	1.05
PIONEER	2	2	100.0%	1.26
COVID-RED	2	2	100.0%	0.44
STOPFOP	2	2	100.0%	2.17
NGN-PET	2	1	50.0%	1.05
Trials@Home	2	1	50.0%	2.41
Inno4Vac	2	1	50.0%	0.00
VHFMoDRAD	2	1	50.0%	0.38
PEVIA	2	2	100.0%	0.80
Impentri	1	1	100.0%	0.00
Pharmatrain	1	1	100.0%	0.10

PROJECT	NUMBER OF PAPERS	NUMBER OF INTERNATIONALLY COLLABORATIVE PAPERS	% OF INTERNATIONALLY COLLABORATIVE PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
EBOVAC	1	1	100.0%	3.27
EMTRAIN	1	1	100.0%	0.10
BIGPICTURE	1	1	100.0%	4.88
HARMONY PLUS	1	1	100.0%	11.67

Table 7.2.3 NUMBER, PERCENTAGE AND CITATION IMPACT<sup>20</sup> OF IMI-SUPPORTED RESEARCH PAPERS WITH AUTHORS FROM MORE THAN ONE INSTITUTION, 2010-2021

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
BTCure	672	550	81.8%	1.99
EU-AIMS	546	510	93.4%	2.32
ULTRA-DD	425	384	90.4%	2.14
EMIF	310	291	93.9%	2.75
NEWMEDS	214	177	82.7%	2.33
AIMS-2-TRIALS	196	187	95.4%	3.84
EUROPAIN	181	125	69.1%	2.90
CANCER-ID	180	154	85.6%	3.53
ORBITO	168	132	78.6%	1.83
INNODIA	166	155	93.4%	1.95
TRANSLOCATION	164	117	71.3%	1.63
STEMBANCC	147	118	80.3%	2.17
IMIDIA	141	118	83.7%	1.76
SUMMIT	136	121	89.0%	1.58
BigData@Heart	135	133	98.5%	2.96
ELF	134	99	73.9%	1.19
RTCure	131	122	93.1%	3.72
CHEM21	128	68	53.1%	2.05
PreDiCT-TB	118	99	83.9%	1.22

<sup>20</sup> The last column in is only the citation impact of the papers from more than one institution.

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
SPRINTT	116	100	86.2%	1.97
MIP-DILI	108	80	74.1%	1.97
Quic-Concept	103	98	95.1%	5.18
EUbOPEN	101	83	82.2%	3.17
COMBACTE-NET	100	90	90.0%	1.26
PROTECT	99	97	98.0%	1.05
BEAT-DKD	98	89	90.8%	2.55
COMBACTE-MAGNET	97	83	85.6%	1.46
RHAPSODY	94	81	86.2%	2.59
U-BIOPRED	93	83	89.2%	2.70
eTOX	92	52	56.5%	1.73
COMPACT	91	70	76.9%	2.05
Pharma-Cog	88	82	93.2%	1.26
DIRECT	82	79	96.3%	4.16
ABIRISK	79	68	86.1%	1.41
DDMoRe	77	64	83.1%	1.23
PRISM	75	70	93.3%	3.84
AETIONOMY	71	71	100.0%	1.83
BioVacSafe	70	43	61.4%	1.32
Open PHACTS	70	57	81.4%	3.67
K4DD	68	55	80.9%	1.77
None	65	60	92.3%	3.66
Onco Track	65	53	81.5%	2.36
RADAR-CNS	64	60	93.8%	2.43
IMPRiND	63	56	88.9%	5.84
COMBACTE-CARE	61	59	96.7%	1.72
ZAPI	60	49	81.7%	5.30
MARCAR	60	43	71.7%	1.19
ENABLE	55	50	90.9%	1.57
EPAD	55	47	85.5%	1.66
DRIVE-AB	54	48	88.9%	1.43
APPROACH	54	50	92.6%	2.40

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
PRECISESADS	54	51	94.4%	1.53
AMYPAD	50	49	98.0%	2.47
LITMUS	49	44	89.8%	5.49
TransQST	47	40	85.1%	2.55
RAPP-ID	46	37	80.4%	0.91
Prelect	45	36	80.0%	1.97
PHAGO	45	37	82.2%	4.98
INNODIA HARVEST	45	40	88.9%	2.11
eTRIKS	45	44	97.8%	2.32
FLUCOP	44	42	95.5%	1.89
RESCEU	43	40	93.0%	2.33
MOBILISE-D	43	42	97.7%	2.35
iPiE	41	34	82.9%	1.23
GETREAL	37	36	97.3%	1.71
iABC	33	29	87.9%	1.49
EBiSC	33	30	90.9%	5.16
EBOVAC1	32	25	78.1%	2.25
EHDEN	31	29	93.5%	2.72
PROACTIVE	29	29	100.0%	2.25
DRAGON	28	28	100.0%	5.29
ADAPTED	28	27	96.4%	3.67
ADVANCE	27	26	96.3%	1.43
PREFER	27	26	96.3%	1.62
EbolaMoDRAD	25	23	92.0%	1.39
Hypo-RESOLVE	24	21	87.5%	1.37
eTRANSafe	23	15	65.2%	1.12
TRISTAN	23	21	91.3%	1.50
3TR	23	21	91.3%	3.29
ROADMAP	22	20	90.9%	0.97
SAFE-T	21	20	95.2%	1.79
VAC2VAC	20	17	85.0%	0.95
DRIVE	20	19	95.0%	1.31

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
IMI-PainCare	20	18	90.0%	1.49
EHR4CR	20	19	95.0%	1.18
EBOVAC2	19	18	94.7%	2.57
IM2PACT	19	16	84.2%	3.31
HARMONY	17	14	82.4%	0.83
PERISCOPE	17	12	70.6%	1.35
COMBACTE	16	13	81.3%	3.31
SOPHIA	16	14	87.5%	8.89
WEB-RADR	16	14	87.5%	1.45
BIOMAP	14	13	92.9%	4.88
CARDIATEAM	14	14	100.0%	3.54
VSV-EBOPLUS	13	12	92.3%	1.06
PD-MitoQUANT	13	12	92.3%	2.18
MOPEAD	13	13	100.0%	2.81
EU-PEARL	12	11	91.7%	4.15
TransBioLine	12	12	100.0%	1.45
ConcePTION	11	11	100.0%	0.96
VSV-EBOVAC	11	9	81.8%	0.98
ITCC-P4	11	11	100.0%	2.48
VALUE-Dx	10	9	90.0%	3.44
CARE	9	8	88.9%	10.23
VITAL	9	7	77.8%	0.55
c4c	9	9	100.0%	1.04
COMBACTE-CDI	9	9	100.0%	1.55
EQIPD	9	8	88.9%	3.04
MAD-CoV 2	8	8	100.0%	2.99
EBODAC	8	8	100.0%	2.38
ERA4TB	8	7	87.5%	2.12
RADAR-AD	8	7	87.5%	1.20
PARADIGM	7	6	85.7%	1.88
FAIRplus	7	3	42.9%	2.93
EUPATI	7	7	100.0%	0.75

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
EBOVAC3	7	7	100.0%	1.06
ReSOLUTE	7	5	71.4%	1.21
KRONO	7	6	85.7%	2.94
NECESSITY	6	6	100.0%	2.53
EBiSC2	6	6	100.0%	0.58
IDEA-FAST	6	6	100.0%	1.32
NeuroDeRisk	6	3	50.0%	0.12
MACUSTAR	6	5	83.3%	2.23
Immune-Image	5	5	100.0%	2.67
HIPPOCRATES	5	4	80.0%	5.41
iCONSENSUS	5	4	80.0%	2.10
EBOMAN	4	4	100.0%	3.88
MELLODDY	4	3	75.0%	0.57
DO->IT	4	4	100.0%	1.19
SafeSciMET	4	4	100.0%	0.89
DECISION	4	4	100.0%	2.60
ADAPT-SMART	4	3	75.0%	0.77
T2EVOLVE	3	3	100.0%	3.66
ND4BB	3	3	100.0%	1.36
IMMUCAN	3	3	100.0%	0.87
Eu2P	3	3	100.0%	2.01
imSAVAR	3	3	100.0%	5.01
ImmUniverse	3	3	100.0%	0.00
OPTIMA	2	2	100.0%	0.75
PIONEER	2	2	100.0%	1.26
COVID-RED	2	2	100.0%	0.44
STOPFOP	2	2	100.0%	2.17
NGN-PET	2	1	50.0%	1.05
Trials@Home	2	1	50.0%	2.41
Inno4Vac	2	2	100.0%	0.00
VHFMoDRAD	2	2	100.0%	0.38
PEVIA	2	2	100.0%	0.80

PROJECT	NUMBER OF PAPERS	NUMBER OF PAPERS FROM MORE THAN ONE INSTITUTION	% OF PAPERS FROM MORE THAN ONE INSTITUTION	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
Impentri	1	1	100.0%	0.00
Pharmatrain	1	1	100.0%	0.10
EBOVAC	1	1	100.0%	3.27
EMTRAIN	1	1	100.0%	0.10
BIGPICTURE	1	1	100.0%	4.88
HARMONY PLUS	1	1	100.0%	11.67
COMBINE	1	1	100.0%	0.07
PERSIST-SEQ	1	1	100.0%	0.00

Table 7.2.4 NUMBER, PERCENTAGE AND CITATION IMPACT<sup>21</sup> OF IMI-SUPPORTED RESEARCH PAPERS WITH AUTHORS FROM MORE THAN ONE SECTOR, 2010-2021

PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
BTcure	672	434	64.6%	2.10
EU-AIMS	546	397	72.7%	2.41
ULTRA-DD	425	268	63.1%	2.49
EMIF	310	254	81.9%	2.61
NEWMEDS	214	133	62.1%	2.40
AIMS-2-TRIALS	196	140	71.4%	3.79
EUROPAIN	181	99	54.7%	3.06
CANCER-ID	180	135	75.0%	3.69
ORBITO	168	107	63.7%	1.95
INNODIA	166	134	80.7%	1.98
TRANSLOCATION	164	60	36.6%	1.77
STEMBANCC	147	74	50.3%	2.22
IMIDIA	141	75	53.2%	2.02
SUMMIT	136	102	75.0%	1.55
BigData@Heart	135	122	90.4%	3.10
ELF	134	46	34.3%	1.06

<sup>21</sup> The last column is only field-normalised citation impact for cross sector papers only.

PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
RTCure	131	106	80.9%	4.06
CHEM21	128	30	23.4%	2.25
PreDiCT-TB	118	67	56.8%	1.21
SPRINTT	116	84	72.4%	2.00
MIP-DILI	108	72	66.7%	1.91
Quic-Concept	103	74	71.8%	3.95
EUbOPEN	101	59	58.4%	3.33
COMBACTE-NET	100	79	79.0%	1.34
PROTECT	99	96	97.0%	1.06
BEAT-DKD	98	70	71.4%	2.87
COMBACTE-MAGNET	97	68	70.1%	1.44
RHAPSODY	94	56	59.6%	2.32
U-BIOPRED	93	77	82.8%	2.83
eTOX	92	28	30.4%	2.15
COMPACT	91	22	24.2%	3.38
Pharma-Cog	88	75	85.2%	1.31
DIRECT	82	62	75.6%	4.63
ABIRISK	79	60	75.9%	1.45
DDMoRe	77	50	64.9%	1.35
PRISM	75	60	80.0%	4.32
AETIONOMY	71	46	64.8%	2.23
BioVacSafe	70	32	45.7%	1.32
Open PHACTS	70	42	60.0%	4.62
K4DD	68	37	54.4%	1.72
None	65	49	75.4%	4.09
Onco Track	65	41	63.1%	2.38
RADAR-CNS	64	40	62.5%	2.87
IMPRiND	63	42	66.7%	4.21
COMBACTE-CARE	61	56	91.8%	1.75
ZAPI	60	40	66.7%	5.61
MARCAR	60	25	41.7%	1.22
ENABLE	55	32	58.2%	1.43



PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
EPAD	55	42	76.4%	1.76
DRIVE-AB	54	40	74.1%	1.39
APPROACH	54	44	81.5%	2.13
PRECISESADS	54	43	79.6%	1.68
AMYPAD	50	46	92.0%	2.51
LITMUS	49	41	83.7%	5.64
TransQST	47	28	59.6%	2.88
RAPP-ID	46	15	32.6%	1.09
Prelect	45	31	68.9%	1.89
PHAGO	45	31	68.9%	5.45
INNODIA HARVEST	45	35	77.8%	2.05
eTRIKS	45	37	82.2%	2.51
FLUCOP	44	40	90.9%	1.93
RESCEU	43	36	83.7%	2.44
MOBILISE-D	43	33	76.7%	2.16
iPiE	41	21	51.2%	1.16
GETREAL	37	31	83.8%	1.87
iABC	33	28	84.8%	1.52
EBiSC	33	23	69.7%	6.07
EBOVAC1	32	23	71.9%	2.24
EHDEN	31	25	80.6%	2.93
PROACTIVE	29	29	100.0%	2.25
DRAGON	28	25	89.3%	5.14
ADAPTED	28	26	92.9%	3.83
ADVANCE	27	24	88.9%	1.38
PREFER	27	25	92.6%	1.69
EbolaMoDRAD	25	17	68.0%	1.50
Hypo-RESOLVE	24	14	58.3%	1.14
eTRANSafe	23	11	47.8%	1.10
TRISTAN	23	19	82.6%	1.38
3TR	23	21	91.3%	3.29
ROADMAP	22	20	90.9%	0.97

PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
SAFE-T	21	20	95.2%	1.79
VAC2VAC	20	14	70.0%	1.00
DRIVE	20	17	85.0%	1.33
IMI-PainCare	20	15	75.0%	1.57
EHR4CR	20	17	85.0%	1.19
EBOVAC2	19	10	52.6%	3.92
IM2PACT	19	11	57.9%	2.61
HARMONY	17	14	82.4%	0.83
PERISCOPE	17	6	35.3%	1.35
COMBACTE	16	8	50.0%	4.04
SOPHIA	16	11	68.8%	10.62
WEB-RADR	16	12	75.0%	1.35
BIOMAP	14	12	85.7%	5.28
CARDIATEAM	14	14	100.0%	3.54
VSV-EBOPLUS	13	9	69.2%	1.19
PD-MitoQUANT	13	9	69.2%	2.54
MOPEAD	13	13	100.0%	2.81
EU-PEARL	12	10	83.3%	4.15
TransBioLine	12	11	91.7%	1.56
ConcePTION	11	11	100.0%	0.96
VSV-EBOVAC	11	6	54.5%	1.04
ITCC-P4	11	11	100.0%	2.48
VALUE-Dx	10	7	70.0%	3.87
CARE	9	5	55.6%	12.00
VITAL	9	5	55.6%	0.30
c4c	9	9	100.0%	1.04
COMBACTE-CDI	9	9	100.0%	1.55
EQIPD	9	5	55.6%	3.94
MAD-CoV 2	8	7	87.5%	2.98
EBODAC	8	7	87.5%	2.59
ERA4TB	8	6	75.0%	2.12
RADAR-AD	8	7	87.5%	1.20

PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
PARADIGM	7	6	85.7%	1.88
FAIRplus	7	1	14.3%	6.77
EUPATI	7	7	100.0%	0.75
EBOVAC3	7	4	57.1%	1.18
ReSOLUTE	7	4	57.1%	1.29
KRONO	7	4	57.1%	2.32
NECESSITY	6	6	100.0%	2.53
EBiSC2	6	6	100.0%	0.58
IDEA-FAST	6	2	33.3%	0.45
NeuroDeRisk	6	2	33.3%	0.12
MACUSTAR	6	5	83.3%	2.23
Immune-Image	5	5	100.0%	2.67
HIPPOCRATES	5	4	80.0%	5.41
iCONSENSUS	5	3	60.0%	1.25
EBOMAN	4	4	100.0%	3.88
MELLODDY	4	3	75.0%	0.57
DO->IT	4	3	75.0%	1.18
SafeSciMET	4	4	100.0%	0.89
DECISION	4	3	75.0%	2.91
ADAPT-SMART	4	3	75.0%	0.77
T2EVOLVE	3	2	66.7%	5.19
ND4BB	3	2	66.7%	1.32
IMMUCAN	3	2	66.7%	0.42
Eu2P	3	1	33.3%	0.00
imSAVAR	3	3	100.0%	5.01
ImmUniverse	3	3	100.0%	0.00
OPTIMA	2	2	100.0%	0.75
PIONEER	2	2	100.0%	1.26
COVID-RED	2	2	100.0%	0.44
STOPFOP	2	1	50.0%	3.81
NGN-PET	2	1	50.0%	1.05
Trials@Home	2	1	50.0%	2.41

PROJECT	NUMBER OF PAPERS	NUMBER OF CROSS SECTOR PAPERS	% OF CROSS SECTOR PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)
Inno4Vac	2	2	100.0%	0.00
VHFMoDRAD	2	2	100.0%	0.38
PEVIA	2	2	100.0%	0.80
Impentri	1	1	100.0%	0.00
Pharmatrain	1	1	100.0%	0.10
EBOVAC	1	1	100.0%	3.27
EMTRAIN	1	1	100.0%	0.10
BIGPICTURE	1	1	100.0%	4.88
HARMONY PLUS	1	1	100.0%	11.67
COMBINE	1	1	100.0%	0.07

Figure 7.2.2 INTERNATIONAL COLLABORATION BY COUNTRY, FOR IMI PROJECT: BTCURE, 2010-2021

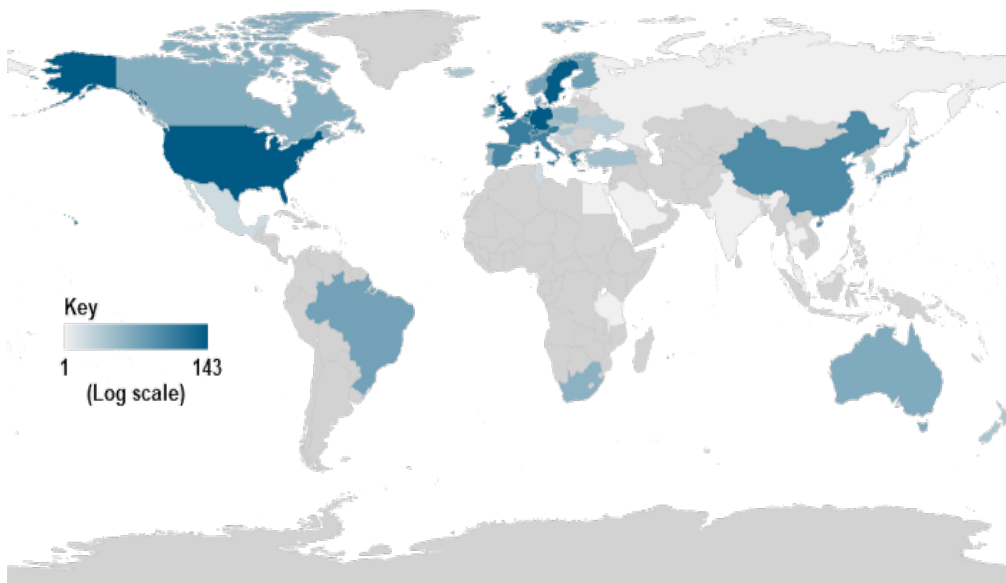


Figure 7.2.3 INTERNATIONAL COLLABORATION BY COUNTRY, FOR IMI PROJECT: EMIF, 2010-2021

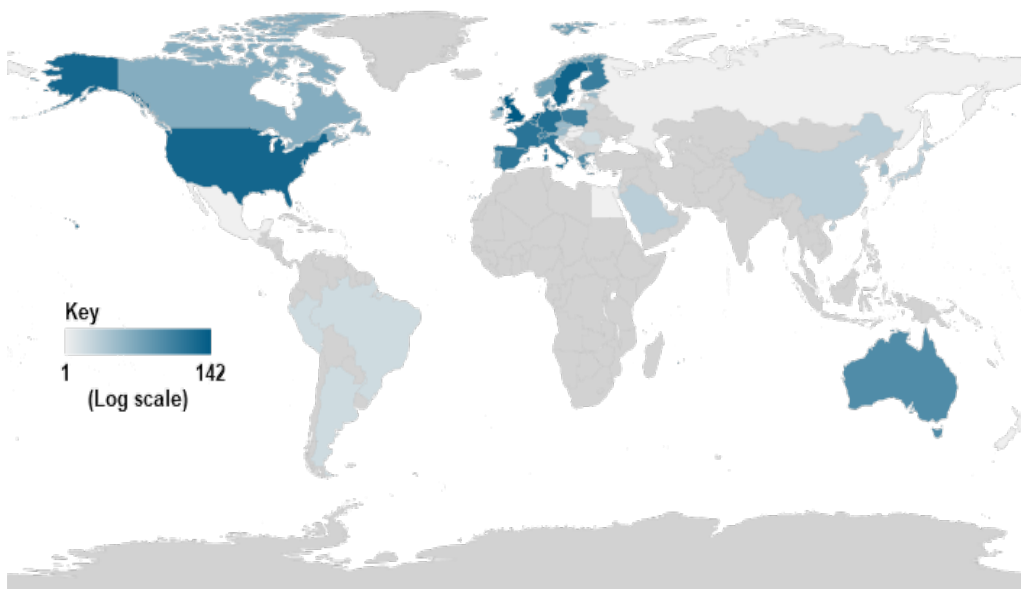


Figure 7.2.4 INTERNATIONAL COLLABORATION BY COUNTRY, FOR IMI PROJECT: EU-AIMS, 2010-2021

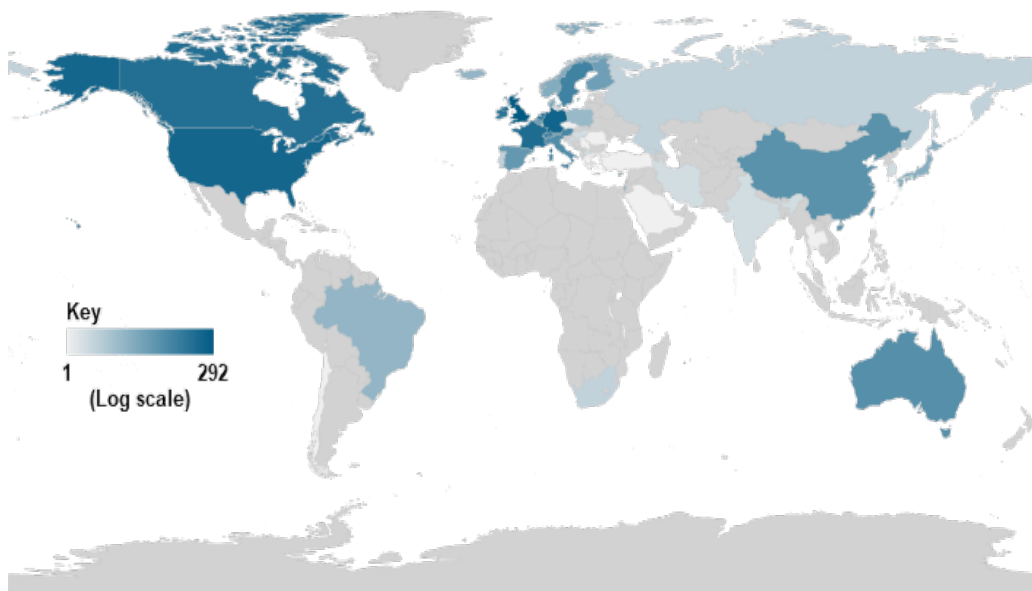


Figure 7.2.5 INTERNATIONAL COLLABORATION BY COUNTRY, FOR IMI PROJECT: NEWMEDS, 2010-2021

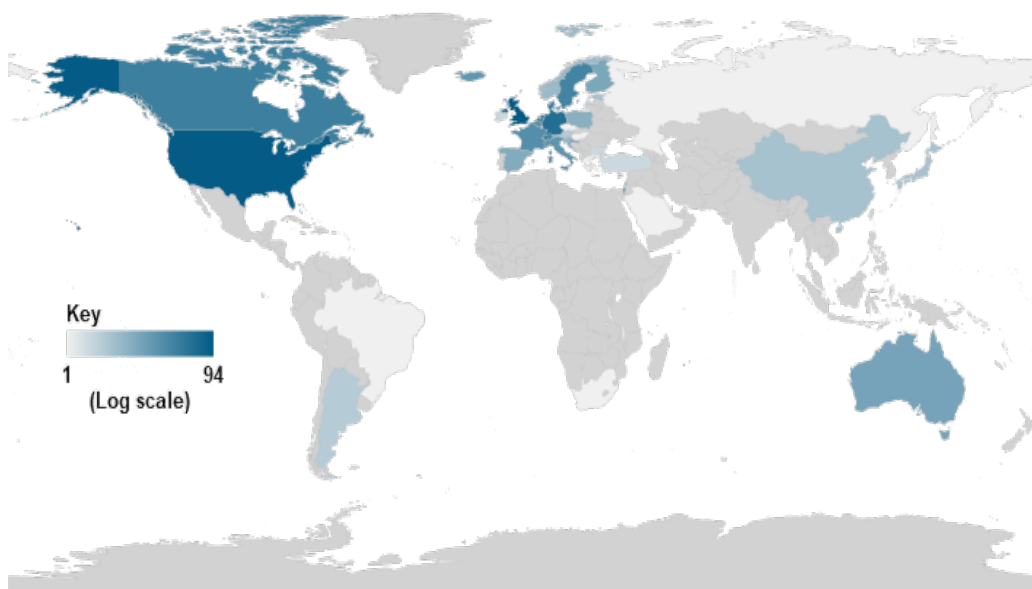
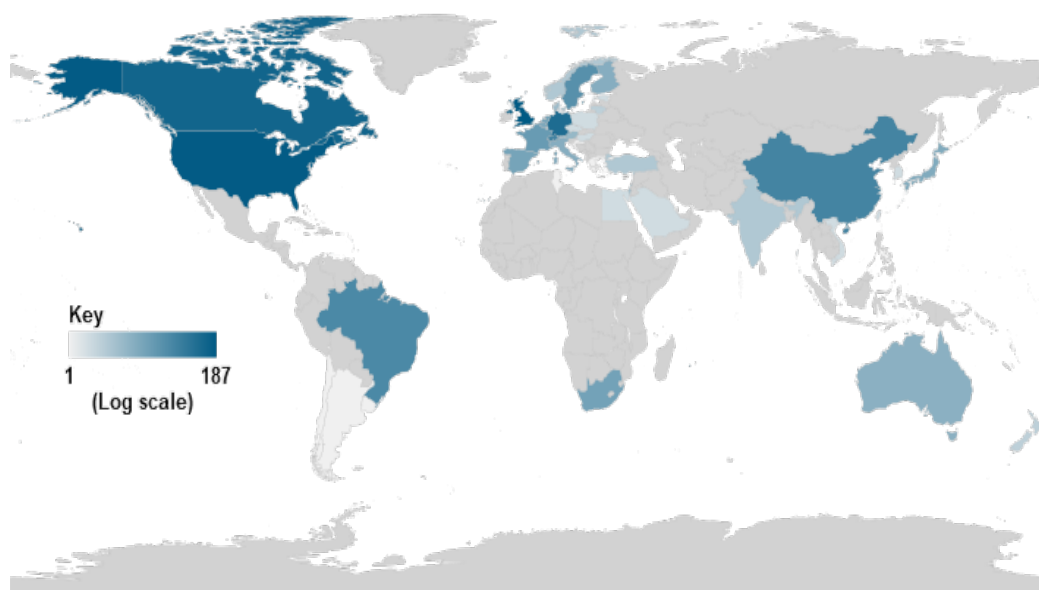


Figure 7.2.6 INTERNATIONAL COLLABORATION BY COUNTRY, FOR IMI PROJECT: ULTRA-DD, 2010-2021



### 7.3 Collaboration metrics for IMI research

This section of the report analyses the types of collaboration that occurred within each IMI project and examines the stability of institutional collaborations within each project.

In common with other metrics based on papers and citations, the indicators we present here work best with larger sample sizes. Indicators based on small numbers of papers will be less informative than those calculated for larger bodies of work. Therefore, the analysis presented in this section is for projects with at least 20 papers published between 2010 and 2021.

In the ninth (2018) and earlier versions of this report metric 3 indicated the intensity of international collaboration, in the tenth report (2019) it was updated to measure the stability of institutional collaborations which is what it shows in this report.

The results for all projects are shown in [Annex 5](#).

Three metrics were used to evaluate the collaborative nature of IMI projects:

- Metric 1 (Cross-sector Score) – Fraction of “cross-sector” papers with co-authors affiliated to institutions in different sectors (Figure 7.3.7.1). The institutions affiliated with each author on an IMI project paper were manually assigned by Clarivate to the relevant sector. Author affiliations were obtained through the Web of Science.
- Metric 2 (International Score) – Percentage of internationally collaborative papers. In calculating the international score for each project, greater weighting is given to papers with multilateral collaboration (co-authors from more than two countries), compared to bilateral collaboration (co-authors from two countries) (Figure 7.3.8.1). The country location of each author was determined using author addresses extracted in the Web of Science.

- Metric 3 (Stability Score) – Stability of institutional collaboration over the lifetime of the project. The collaboration stability for pairs of collaborating institutions was calculated following the method proposed by Y. Bu et al.<sup>22</sup> A stable institutional collaboration has a stable output, i.e. pairs of institutions co-publish a similar volume of papers in consecutive years for the duration of a project. The stability score for each project is the mean average stability of all the collaborating institutional pairs that have contributed to that IMI project research.

Each metric is calculated for an IMI project and can take a value between 0 and 1, with 1 indicating more collaborative activity. The collaboration index is a sum of all three metrics and the maximum possible value for a project is 3.

### 7.3.1 Metric 1 (cross-sector score): fraction of cross sector collaborative papers

Authors institutional affiliations, as they appear on IMI project research were assigned to sectors. Sector assignments were then used to classify each paper as “within one sector”, when all co-authors work within the same sector or “cross-sector” when co-authors work in two or more different sectors. The number and percentage of cross-sector papers for projects are presented in Table 7.2.3.

Figure 7.3.1.1 shows the total number of “within one sector” and “cross-sector” papers for each project. Projects are ordered by the number of cross-sector collaborative papers. The dark blue bars represent the number or fraction of “cross-sector” papers. The fraction of cross-sector papers in each project is referred to in the figure as “Cross-Sector Score”. Only projects with more than 20 associated papers are shown.

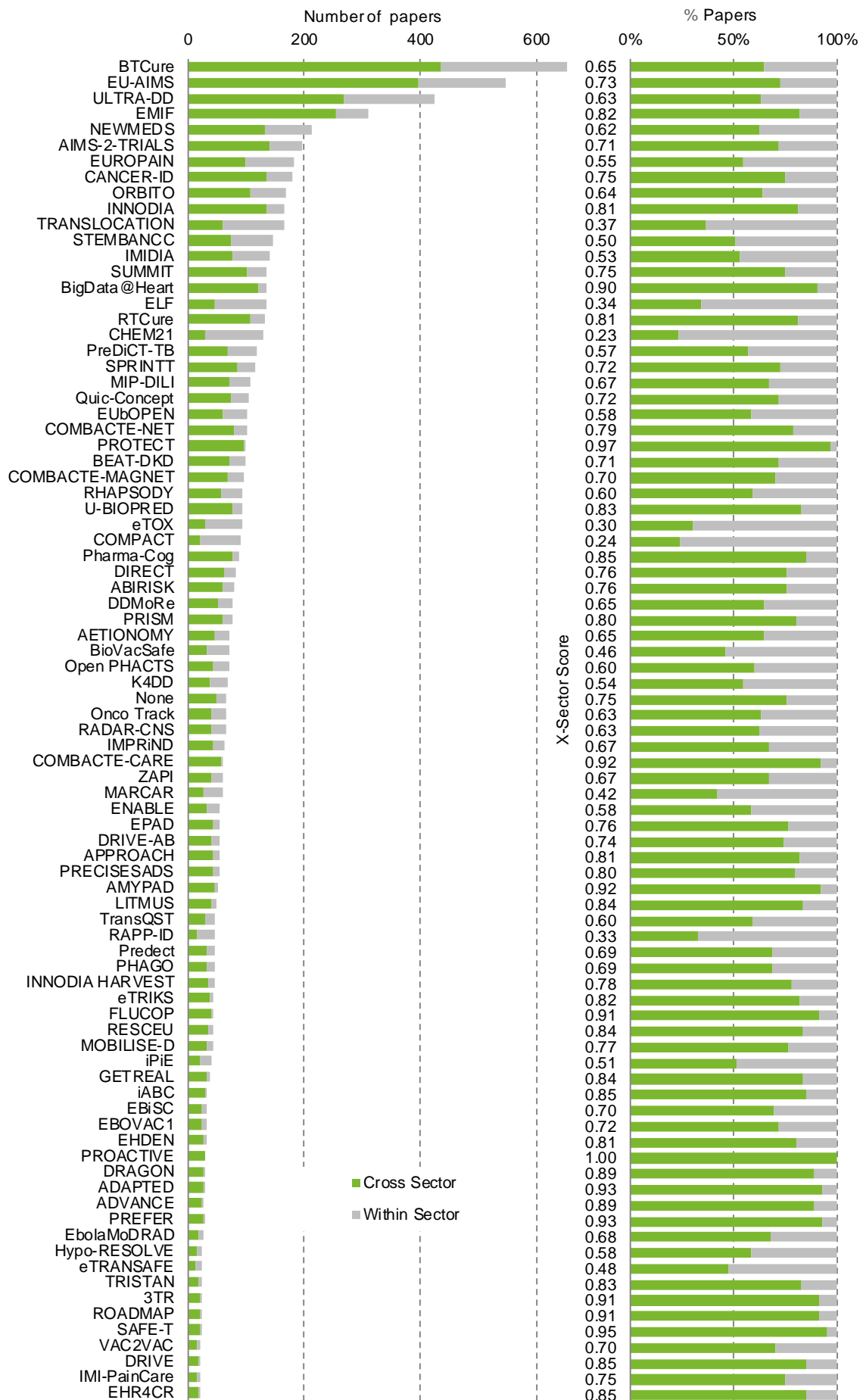
- BTCURE had the greatest number of cross-sector collaborative papers, 434 out of a total of 672. PROACTIVE, PROTECT and SAFE-T had the highest percentage of cross-sector collaborative papers (100%, 97% and 95%, respectively).

---

<sup>22</sup> Bu, Y., Murray, D.S., Ding, Y. et al. (2018) Measuring the stability of scientific collaboration. *Scientometrics*, **114**, 463.



FIGURE 7.3.1.1 NUMBER AND FRACTION OF CROSS-SECTOR COLLABORATIVE PAPERS BY PROJECT, 2010-2021. ORDERED BY NUMBER OF CROSS-SECTOR PAPERS



### 7.3.2 Metric 2 (international score): fraction of internationally collaborative papers

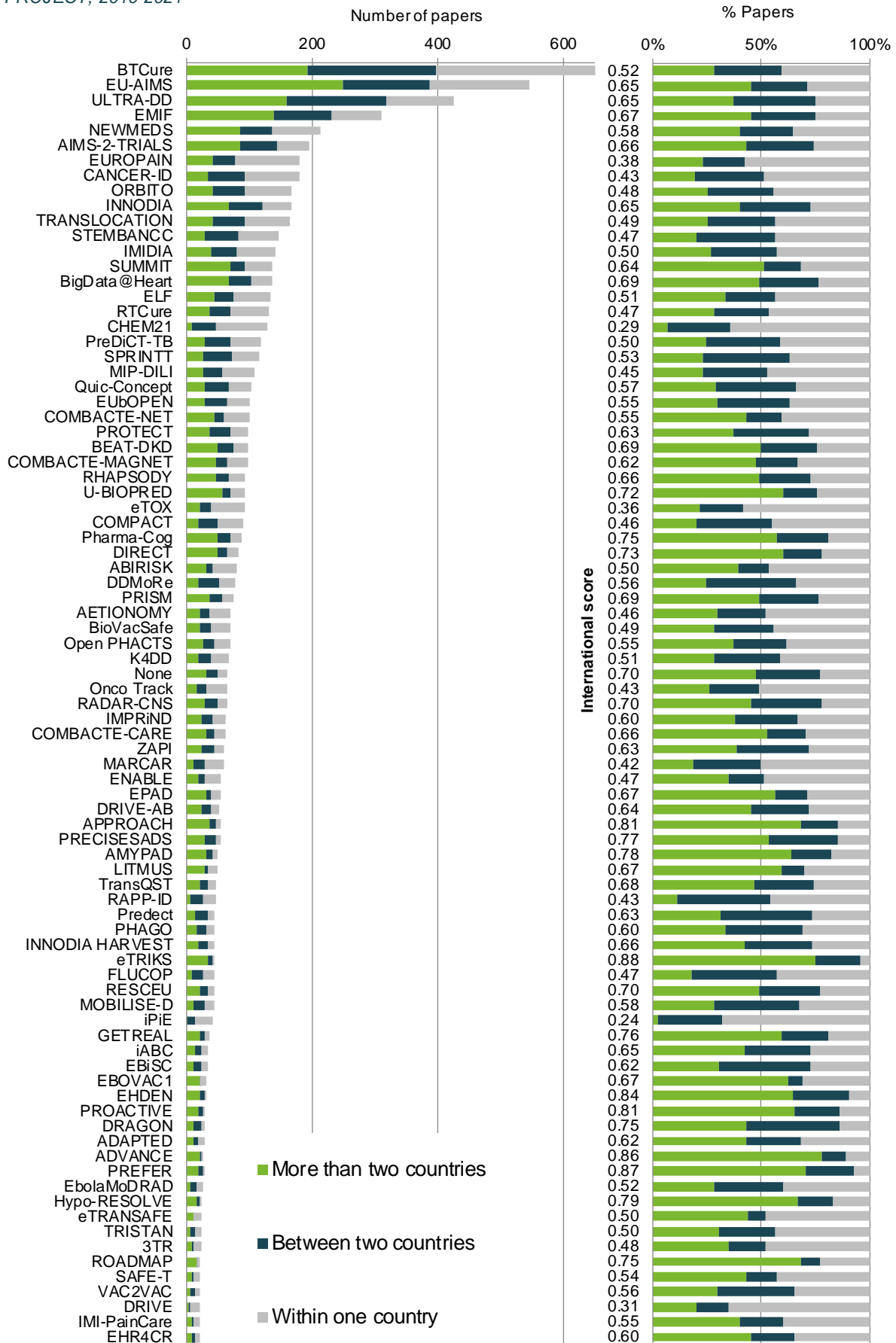
Author names and affiliations were extracted for all IMI project papers. The number of countries in the author affiliations for each paper was counted and used to classify the papers as “more than two countries”, “two countries” or “within one country” (same as domestic in the Section 7.1).

Figure 7.3.2.1 below shows the total number of papers for each project. Projects are ordered by the number of papers with author affiliations from more than one country. The bar colours reflect the fraction of papers that include international collaboration between “two countries” (bilateral) and “more than two countries” (multilateral). Only projects with more than 20 associated papers are shown.

The International Score was calculated by weighting each paper that involved only two countries by 0.75 and each paper that involved more than two countries by 1.00. The sum of the weighted papers was then divided by the total number of project papers. Total number of internationally collaborative papers for each project is shown in Table 7.2.1.

- BTCURE had the most internationally collaborative papers involving two or more countries (398 out of 672), with an International Score of 0.52.
- EU-AIMS had the most internationally collaborative papers involving more than two countries. (248 out of 546), with an international Score of 0.65.
- eTRICKS, PREFER, and ADVANCE had the highest International Scores (0.88, 0.87 and 0.86 respectively).

FIGURE 7.3.2.1 NUMBER AND FRACTION OF INTERNATIONALLY COLLABORATIVE PAPERS BY PROJECT, 2010-2021

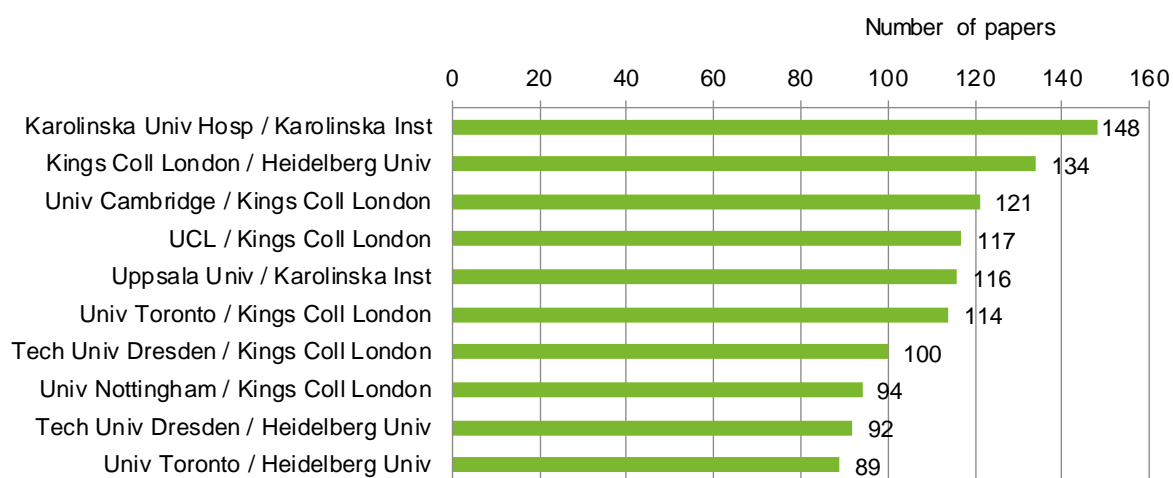


### 7.3.3 Metric 3 (stability score): stability of institutional collaboration

This section looks in depth at institutional collaboration activities in IMI funded research. Figure 7.3.3.1 shows the ten most productive, collaborating institution pairs, by total number of collaborative papers. Figure 7.3.3.2 shows the ten institutions that collaborate with the highest number of other institutions. Figure 7.3.3.3 shows the distribution of Metric 3 scores for IMI projects. Table 7.3.3.1 is an expansion of the data in Figure 7.3.3.3, showing the Metric 3 score for all projects with at least 20 papers and the number of collaborating institution pairs. The number and proportion of papers with authors from more than one institution for each project is shown in Table 7.2.2.

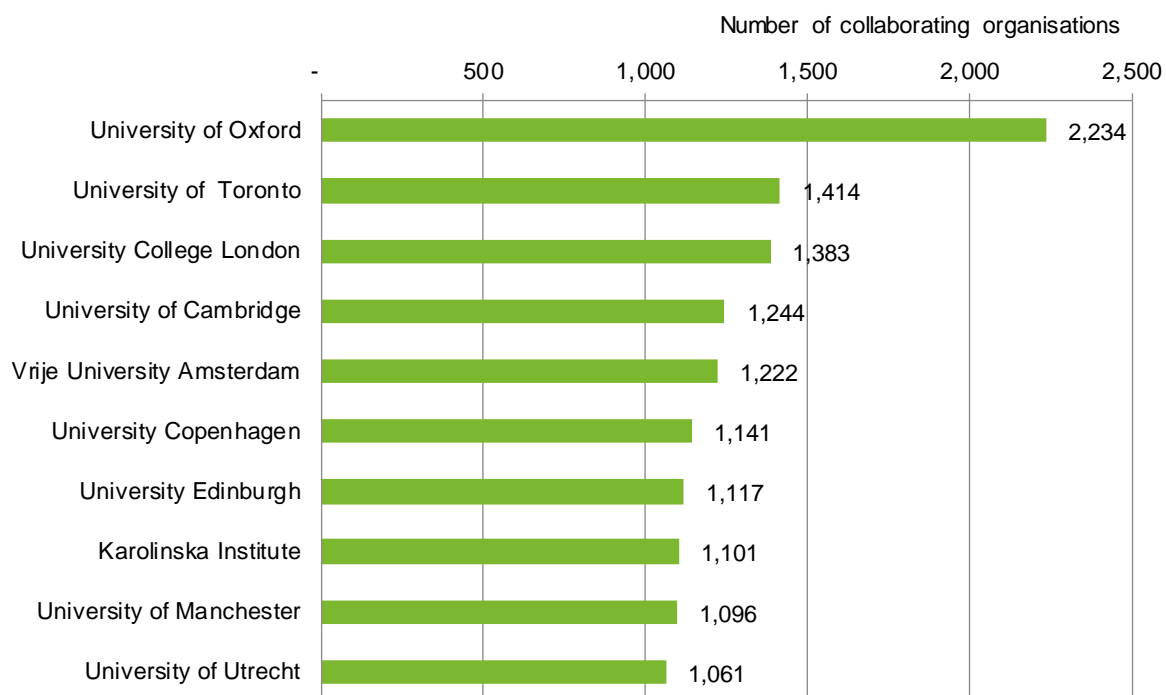
A project's Metric 3 is the mean average stability of collaborations between pairs of institutions that have co-authored papers that belong to that project. Pairs of institutions must have published two or more papers together as part of the same IMI project to be considered. A second requirement is that the IMI projects must have started in, or before, 2019. If a project started after 2019, too little time has elapsed for most pairs of institutions to have published more than one paper.

FIGURE 7.3.3.1 THE TEN MOST PRODUCTIVE PAIRS OF COLLABORATING INSTITUTIONS, 2010-2021



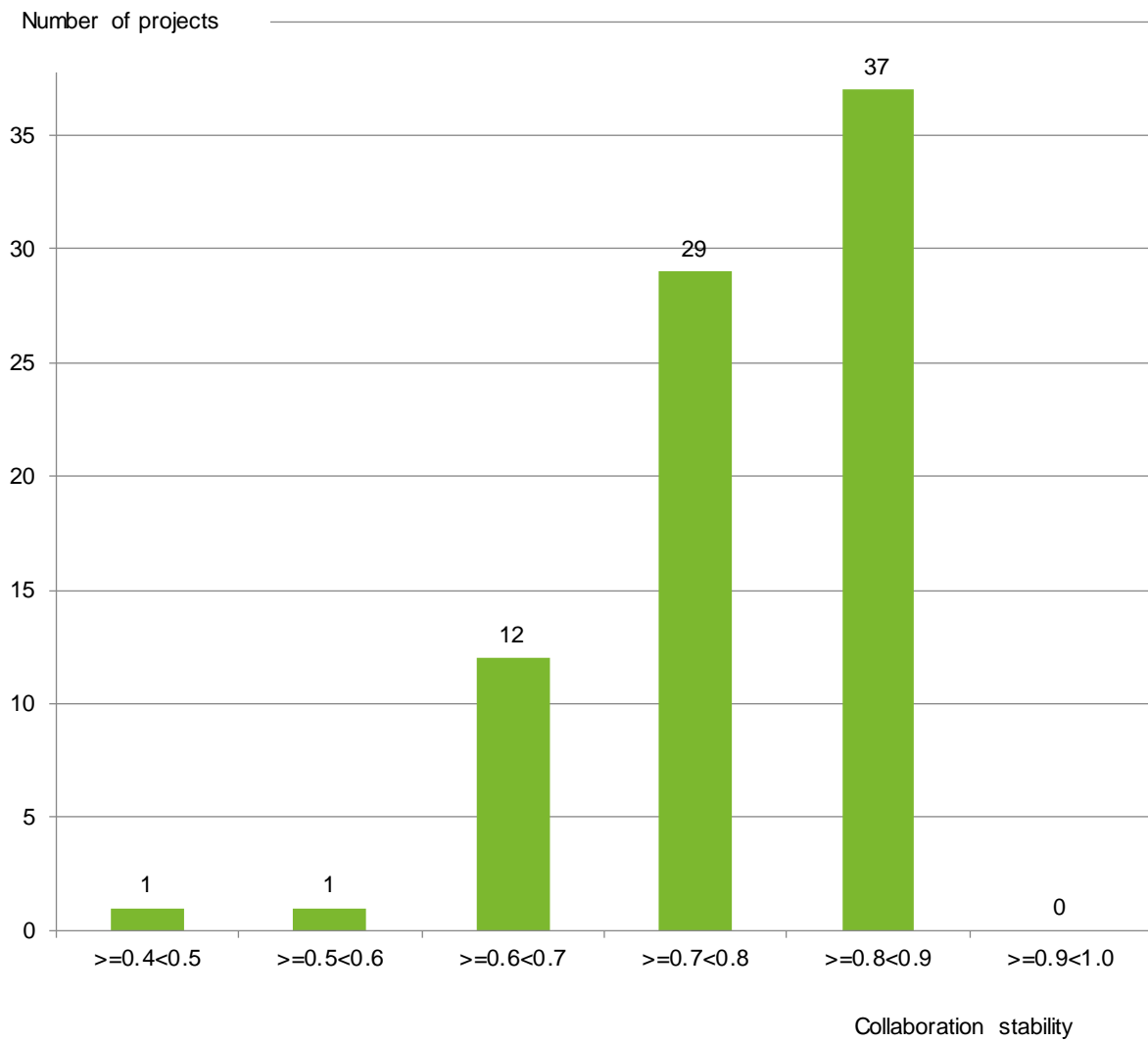
- The institutions that collaborated most frequently on IMI project papers were Karolinska University Hospital and Karolinska Institute, researchers at these institutions together, co-authored 148 publications. This is a change from last year's report where Kings College London and Heidelberg University were the top collaborating pair, now second.
- Kings College London is part of six out of ten pairs of the most productive collaborative institutions.

FIGURE 7.3.3.2 THE TEN INSTITUTIONS THAT HAVE COLLABORATED WITH THE GREATEST NUMBER OF OTHER INSTITUTIONS, 2010-2021



- The University of Oxford has collaborated with 2,234 other institutions on IMI project papers, the most of any of the other institutions.
- The University of Utrecht is new to the top 10 collaborating organisations, replacing the University of Amsterdam.
- Half of the ten most collaborative institutions are located in the United Kingdom.

FIGURE 7.3.3.3 METRIC 3: STABILITY SCORE DISTRIBUTION, 2010-2021



- Most IMI projects have a stability score of between 0.70 and 0.90 indicating that most of the collaboration between institutions is relatively stable.

Table 7.3.5 STABILITY SCORE FOR IMI PROJECTS, NUMBER OF COLLABORATING INSTITUTION PAIRS, TOTAL NUMBER OF PAPERS AND PROJECT START YEAR FOR ALL PROJECTS WITH AT LEAST 20 PAPERS THAT STARTED IN OR BEFORE 2019, 2010-2021

PROJECT	STABILITY SCORE (METRIC 3)	NUMBER OF COLLABORATING INSTITUTION PAIRS	NUMBER OF PAPERS	PROJECT START YEAR
BTCure	0.84	1,128	672	2011
EU-AIMS	0.82	3,763	546	2012
ULTRA-DD	0.77	410	425	2015
EMIF	0.84	2,933	310	2012
NEWMEDS	0.83	865	214	2010
AIMS-2-TRIALS	0.69	946	196	2018
EUROPAIN	0.85	345	181	2010
CANCER-ID	0.75	241	180	2015
ORBITO	0.76	351	168	2013
INNODIA	0.87	431	166	2010
TRANSLOCATION	0.81	78	164	2013
STEMBANCC	0.82	64	147	2013
IMIDIA	0.84	159	141	2010
SUMMIT	0.83	7,004	136	2011
BigData@Heart	0.69	4,090	135	2017
ELF	0.77	49	134	2014
RTCure	0.70	242	131	2017
CHEM21	0.80	22	128	2013
PreDiCT-TB	0.81	67	118	2013
SPRINTT	0.80	277	116	2014
MIP-DILI	0.82	145	108	2012
Quic-Concept	0.81	157	103	2012
COMBACTE-NET	0.89	672	100	2013
PROTECT	0.86	300	99	2010
BEAT-DKD	0.70	335	98	2017
COMBACTE-MAGNET	0.86	625	97	2015
RHAPSODY	0.81	428	94	2016
U-BIOPRED	0.86	1,019	93	2010
eTOX	0.86	126	92	2010
COMPACT	0.75	41	91	2014

PROJECT	STABILITY SCORE (METRIC 3)	NUMBER OF COLLABORATING INSTITUTION PAIRS	NUMBER OF PAPERS	PROJECT START YEAR
Pharma-Cog	0.85	1,066	88	2010
DIRECT	0.84	1,299	82	2012
ABIRISK	0.84	492	79	2012
DDMoRe	0.81	59	77	2012
PRISM	0.75	222	75	2017
AETIONOMY	0.80	96	71	2014
BioVacSafe	0.81	26	70	2012
Open PHACTS	0.76	63	70	2011
K4DD	0.80	38	68	2013
Onco Track	0.83	87	65	2011
RADAR-CNS	0.82	178	64	2016
IMPRiND	0.75	46	63	2017
COMBACTE-CARE	0.80	690	61	2015
MARCAR	0.84	39	60	2011
ZAPI	0.77	62	60	2015
ENABLE	0.83	46	55	2015
EPAD	0.81	217	55	2015
APPROACH	0.81	109	54	2015
DRIVE-AB	0.75	84	54	2015
PRECISESADS	0.75	226	54	2015
AMYPAD	0.77	349	50	2017
LITMUS	0.75	204	49	2018
TransQST	0.76	23	47	2017
RAPP-ID	0.85	14	46	2011
eTRIKS	0.75	714	45	2014
PHAGO	0.72	61	45	2017
Prelect	0.79	26	45	2012
FLUCOP	0.73	37	44	2015
RESCEU	0.69	491	43	2018
iPIE	0.76	22	41	2016
GETREAL	0.78	37	37	2015
EBiSC	0.77	17	33	2015
iABC	0.82	129	33	2015



PROJECT	STABILITY SCORE (METRIC 3)	NUMBER OF COLLABORATING INSTITUTION PAIRS	NUMBER OF PAPERS	PROJECT START YEAR
EBOVAC1	0.82	53	32	2015
EHDEN	0.63	896	31	2019
PROACTIVE	0.85	199	29	2011
ADAPTED	0.68	1,210	28	2017
ADVANCE	0.77	300	27	2015
PREFER	0.71	135	27	2017
EbolaMoDRAD	0.55	33	25	2016
Hypo-RESOLVE	0.64	56	24	2019
eTRANSafe	0.66	10	23	2018
TRISTAN	0.64	25	23	2017
ROADMAP	0.66	75	22	2017
SAFE-T	0.86	21	21	2011
DRIVE	0.79	8	20	2018
EHR4CR	0.78	51	20	2012
IMI-PainCare	0.67	50	20	2019
VAC2VAC	0.69	6	20	2018

- COMBACTE-NET has the highest stability score (0.89) while EbolaMoDRAD has the lowest (0.55).
- There is considerable variation in the number collaborating institution pairs that does not appear to be proportional to the number of project papers or dependent on the project start year. For example, BTCure started in 2011 and has the highest output of papers (672), only has 1,128 institution pairs compared with SUMMIT that started in the same year, has only produced 136 papers but has 7,004 institution pairs. This suggests that SUMMIT publishes papers with many authors from multiple institutions. In fact, one of SUMMIT's papers has 267 affiliations.

## 7.4 Collaboration index

The cross-sector score (Metrics 1) and international score (Metric 2) (described above) measure different types of collaboration. The first measures the fraction of papers that involve cross-sector collaborations, and the second reflects the fraction of papers that involve multilateral and bilateral international collaborations. Metric 3 or stability score is based on the collaboration stability of pairs of institutional collaborators that contribute to IMI project research. We compute a “collaboration index” across IMI projects as the sum of all three of the metrics. These data are shown in Table 7.4.1 for all IMI projects with 20 or more papers.

This year's collaboration index is not comparable with the collaboration index in the ninth (2018) and earlier versions of this report as Metric 3 was updated in the tenth report (2019) to indicate the stability of institutional collaboration rather than intensity.

- PROACTIVE had the highest overall collaboration index score (2.62) followed by ADVANCE (2.58).

Table 7.4.6 SUMMARY SCORE FOR COLLABORATION METRICS, TOTAL NUMBER OF PAPERS AND FIELD-NORMALISED CITATION IMPACT FOR IMI PROJECTS WITH AT LEAST 20 PAPERS, 2010-2021

PROJECT	CROSS-SECTOR SCORE (METRIC 1)	INTERNATIONAL SCORE (METRIC 2)	STABILITY SCORE (METRIC 3) <sup>23</sup>	COLLABORATION INDEX	NUMBER OF PAPERS	CITATION IMPACT (FIELD-NORMALISED)
BTCure	0.65	0.52	0.84	2.00	672	1.81
EU-AIMS	0.73	0.65	0.82	2.19	546	2.06
ULTRA-DD	0.63	0.65	0.77	2.06	425	1.91
EMIF	0.82	0.67	0.84	2.34	310	2.49
NEWMEDS	0.62	0.58	0.83	2.03	214	2.06
AIMS-2-TRIALS	0.71	0.66	0.69	2.07	196	2.67
EUROPAIN	0.55	0.38	0.85	1.78	181	2.55
CANCER-ID	0.75	0.43	0.75	1.93	180	3.27
ORBITO	0.64	0.48	0.76	1.88	168	1.70
INNODIA	0.81	0.65	0.87	2.32	166	1.62
TRANSLOCATION	0.37	0.49	0.81	1.66	164	1.35
STEMBANCC	0.50	0.47	0.82	1.79	147	1.94
IMIDIA	0.53	0.50	0.84	1.87	141	1.65
SUMMIT	0.75	0.64	0.83	2.22	136	1.42
BigData@Heart	0.90	0.69	0.69	2.29	135	2.05
ELF	0.34	0.51	0.77	1.62	134	1.11
RTCure	0.81	0.47	0.70	1.98	131	2.86
CHEM21	0.23	0.29	0.80	1.33	128	1.70
PreDiCT-TB	0.57	0.50	0.81	1.88	118	1.16
SPRINTT	0.72	0.53	0.80	2.05	116	1.83
MIP-DILI	0.67	0.45	0.82	1.94	108	1.77
Quic-Concept	0.72	0.57	0.81	2.09	103	4.76
EUbOPEN	0.58	0.55	n/a	n/a	101	1.85

<sup>23</sup> Some projects do not have a Stability score due to the project not being active for at least 3 years. The Collaboration Index was not calculated for projects with no Stability Score.

PROJECT	CROSS-SECTOR SCORE (METRIC 1)	INTERNATIONAL SCORE (METRIC 2)	STABILITY SCORE (METRIC 3) <sup>23</sup>	COLLABORATION INDEX	NUMBER OF PAPERS	CITATION IMPACT (FIELD-NORMALISED)
COMBACTE-NET	0.79	0.55	0.89	2.23	100	1.17
PROTECT	0.97	0.63	0.86	2.46	99	1.03
BEAT-DKD	0.71	0.69	0.70	2.11	98	2.17
COMBACTE-MAGNET	0.70	0.62	0.86	2.18	97	1.28
RHAPSODY	0.60	0.66	0.81	2.07	94	1.97
U-BIOPRED	0.83	0.72	0.86	2.41	93	2.49
eTOX	0.30	0.36	0.86	1.53	92	1.76
COMPACT	0.24	0.46	0.75	1.46	91	1.90
Pharma-Cog	0.85	0.75	0.85	2.45	88	1.13
DIRECT	0.76	0.73	0.84	2.33	82	3.93
ABIRISK	0.76	0.50	0.84	2.10	79	1.25
DDMoRe	0.65	0.56	0.81	2.02	77	1.21
PRISM	0.80	0.69	0.75	2.25	75	2.70
AETIONOMY	0.65	0.46	0.80	1.91	71	1.77
Open PHACTS	0.60	0.55	0.76	1.92	70	3.57
BioVacSafe	0.46	0.49	0.81	1.76	70	1.20
K4DD	0.54	0.51	0.80	1.86	68	1.53
None	0.75	0.70	0.49	1.94	65	3.50
Onco Track	0.63	0.43	0.83	1.90	65	2.19
RADAR-CNS	0.63	0.70	0.82	2.15	64	1.94
IMPRiND	0.67	0.60	0.75	2.01	63	4.96
COMBACTE-CARE	0.92	0.66	0.80	2.37	61	1.59
ZAPI	0.67	0.63	0.77	2.07	60	4.34
MARCAR	0.42	0.42	0.84	1.68	60	1.04
EPAD	0.76	0.67	0.81	2.25	55	1.35
ENABLE	0.58	0.47	0.83	1.88	55	1.45
APPROACH	0.81	0.81	0.81	2.43	54	2.07
PRECISESADS	0.80	0.77	0.75	2.32	54	1.38
DRIVE-AB	0.74	0.64	0.75	2.13	54	1.30
AMYPAD	0.92	0.78	0.77	2.47	50	1.96
LITMUS	0.84	0.67	0.75	2.25	49	3.85
TransQST	0.60	0.68	0.76	2.03	47	3.09
RAPP-ID	0.33	0.43	0.85	1.61	46	0.86
eTRIKS	0.82	0.88	0.75	2.46	45	2.04

PROJECT	CROSS-SECTOR SCORE (METRIC 1)	INTERNATIONAL SCORE (METRIC 2)	STABILITY SCORE (METRIC 3) <sup>23</sup>	COLLABORATION INDEX	NUMBER OF PAPERS	CITATION IMPACT (FIELD-NORMALISED)
Predict	0.69	0.63	0.79	2.11	45	2.67
INNODIA HARVEST	0.78	0.66	n/a	n/a	45	1.37
PHAGO	0.69	0.60	0.72	2.01	45	4.02
FLUCOP	0.91	0.47	0.73	2.11	44	1.58
RESCEU	0.84	0.70	0.69	2.22	43	1.91
MOBILISE-D	0.77	0.58	n/a	n/a	43	1.57
iPiE	0.51	0.24	0.76	1.51	41	1.13
GETREAL	0.84	0.76	0.78	2.37	37	1.65
iABC	0.85	0.65	0.82	2.32	33	1.19
EBiSC	0.70	0.62	0.77	2.09	33	5.61
EBOVAC1	0.72	0.67	0.82	2.21	32	2.07
EHDEN	0.81	0.84	0.63	2.27	31	2.19
PROACTIVE	1.00	0.81	0.85	2.66	29	2.22
DRAGON	0.89	0.75	n/a	n/a	28	3.78
ADAPTED	0.93	0.62	0.68	2.23	28	2.89
PREFER	0.93	0.87	0.71	2.51	27	1.29
ADVANCE	0.89	0.86	0.77	2.52	27	1.36
EbolaMoDRAD	0.68	0.52	0.55	1.75	25	1.34
Hypo-RESOLVE	0.58	0.79	0.64	2.02	24	0.71
TRISTAN	0.83	0.50	0.64	1.97	23	1.22
eTRANSafe	0.48	0.50	0.66	1.64	23	3.15
3TR	0.91	0.48	n/a	n/a	23	2.60
ROADMAP	0.91	0.75	0.66	2.32	22	0.91
SAFE-T	0.95	0.54	0.86	2.35	21	1.73
VAC2VAC	0.70	0.56	0.69	1.96	20	0.60
EHR4CR	0.85	0.60	0.78	2.23	20	1.11
IMI-PainCare	0.75	0.55	0.67	1.97	20	1.22
DRIVE	0.85	0.31	0.79	1.95	20	0.79

## 8 BENCHMARKING ANALYSIS – IMI PROJECT RESEARCH AGAINST RESEARCH FROM SELECTED COMPARATORS

This section of the report analyses the output and citation impact of IMI project research benchmarked against research supported by other Public-Private Partnerships, and funders of biomedical research across Europe, Asia, Australia, and North America.

The publications funded by each comparator were identified using specific searches of the funding acknowledgment data provided by authors and extracted in Web of Science. This is the same process by which IMI project publications have been identified. Authors may not always acknowledge their sources of funding and may not always do so correctly. Therefore, the coverage of the datasets used in these analyses may not be complete and may not be entirely accurate; however, the sample represented by these datasets is sufficient to allow a comparison to be made.

### 8.1 Identifying comparators

The seven funders listed in Table 8.1.1 are used as comparators for IMI in this report. They are the same comparators as in the previous twelfth report produced in 2021. Each comparator had sufficient publications to allow a meaningful analysis.

Table 8.1.1 SUMMARY OF INFORMATION OF IMI-SELECTED COMPARATORS, 2010-2021

COMPARATOR	NUMBER OF PUBLICATIONS (2010-2021)	NUMBER OF PAPERS (2010-2021)	COUNTRY	REGION
Critical Path (C-Path)	568	528	USA	North America
Commonwealth Scientific and Industrial Research Organisation (CSIRO) <sup>24</sup>	1,036	1,003	Australia	Australia
Foundation for the National Institutes of Health (FNIH)	5,070	4,782	USA	North America
Grand Challenges in Global Health (GCGH)	890	889	USA	North America
Indian Council of Medical Research (ICMR)	17,852	17,323	India	Asia
Medical Research Council (MRC)	141,230	126,335	UK	Europe
Wellcome Trust (WT)	96,581	89,768	UK	Europe

<sup>24</sup> The dataset containing all publications attributed to CSIRO between 2010 and 2021 has been reduced to include only medically related publications for these analyses. A list of Web of Science journal categories which capture medically related publications is given in [Annex 2](#).

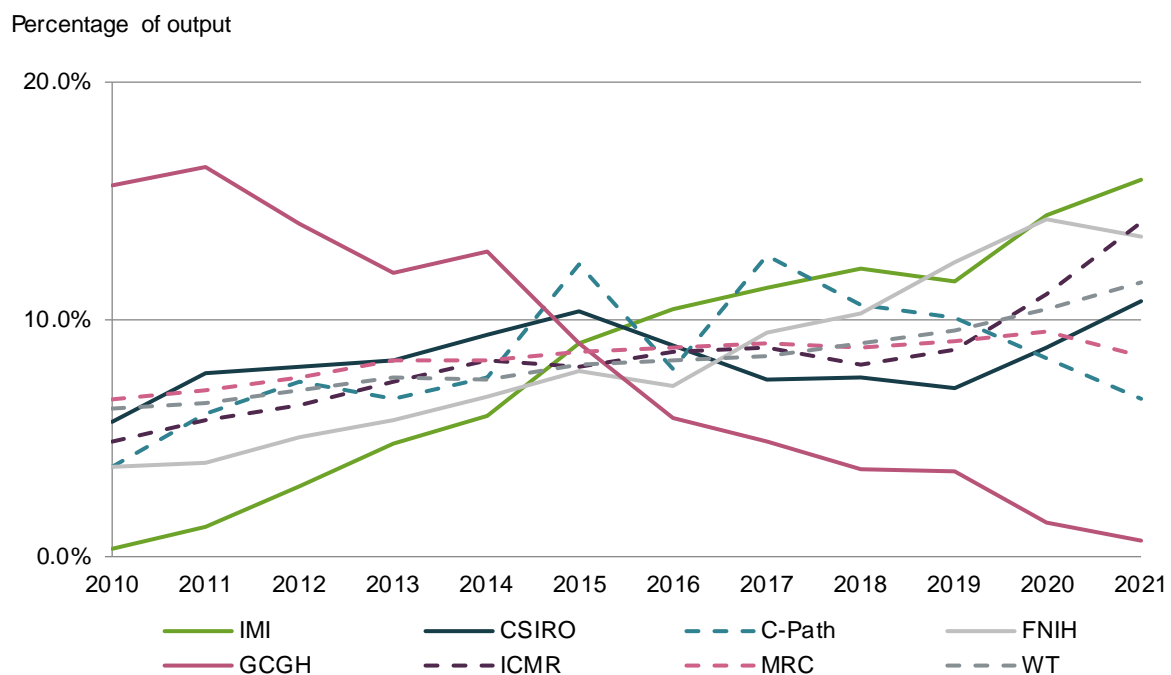
## 8.2 Trends in output: IMI project research compared with selected comparators

This section of the report analyses trends in the performance of IMI project research and the selected comparators.

### 8.2.1 Trends in output: IMI project research compared with selected comparators

The output of IMI and the comparators varies widely (some produced many papers and some relatively few), therefore a visual comparison of absolute paper counts would not provide an understanding of their growth relative to one another. To provide a more easily interpretable comparison, Figure 8.2.1.1 shows the percentage of each organisation's total paper count between 2010 and 2021 published in each year. Table 8.2.1.1 shows the same data as in Figure 8.2.1.1 and Table 8.2.1.2 show the number of papers per year for IMI and the selected comparators.

Figure 8.2.1.1 TRENDS IN OUTPUT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- Most of IMI's research output was published in the last four years 2018-2021, accounting for more than half of its paper output.
- IMI has experienced the most rapid increase in percentage of output, only in 2019 seeing a slight decrease.
- GCGH has sustained a decreasing percentage of output since 2011. Similarly, C-path has been on a downward trend since 2017.
- After peaking in 2015, CSIRO was on a steady decrease in percentage of output but has seen a shift upwards since 2019.

FIGURE 8.2.1.2 COMPARING PERCENTAGE OUTPUT IN THE FIRST FIVE YEARS (2010–2015) TO MOST RECENT FIVE YEARS (2016–2021) – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

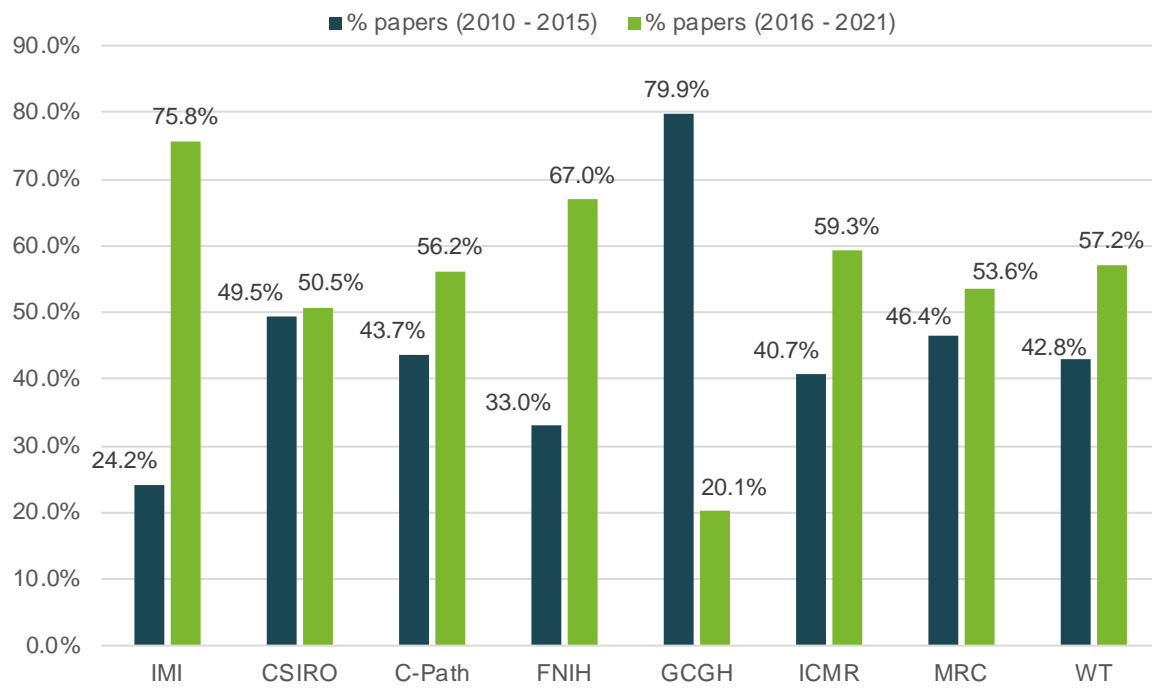


TABLE 8.2.1.1 SHARE OF OUTPUT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	0.3%	5.7%	3.8%	3.8%	15.6%	4.8%	6.6%	6.2%
2011	1.2%	7.8%	6.1%	3.9%	16.4%	5.7%	7.0%	6.4%
2012	3.0%	8.0%	7.4%	5.0%	14.1%	6.4%	7.6%	7.0%
2013	4.7%	8.3%	6.6%	5.7%	11.9%	7.4%	8.3%	7.5%
2014	6.0%	9.4%	7.6%	6.7%	12.8%	8.3%	8.3%	7.5%
2015	9.0%	10.4%	12.3%	7.8%	9.0%	8.0%	8.6%	8.1%
2016	10.4%	8.9%	8.0%	7.2%	5.8%	8.6%	8.8%	8.3%
2017	11.3%	7.5%	12.7%	9.4%	4.8%	8.8%	9.0%	8.4%
2018	12.2%	7.6%	10.6%	10.2%	3.7%	8.1%	8.8%	9.0%
2019	11.6%	7.1%	10.0%	12.4%	3.6%	8.7%	9.1%	9.5%
2020	14.4%	8.8%	8.3%	14.2%	1.5%	11.1%	9.5%	10.4%
2021	15.9%	10.8%	6.6%	13.4%	0.7%	14.1%	8.4%	11.6%

TABLE 8.2.1.2 NUMBER OF PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	26	57	20	181	139	840	8,375	5,603
2011	97	78	32	188	146	993	8,824	5,782
2012	235	80	39	240	125	1,101	9,566	6,302
2013	371	83	35	274	106	1,284	10,427	6,774
2014	468	94	40	321	114	1,432	10,466	6,732
2015	707	104	65	373	80	1,393	10,923	7,245
2016	816	89	42	346	52	1,494	11,102	7,424
2017	889	75	67	451	43	1,522	11,351	7,584
2018	957	76	56	490	33	1,402	11,167	8,051
2019	913	71	53	595	32	1,507	11,513	8,520
2020	1,129	88	44	680	13	1,915	11,985	9,374
2021	1,248	108	35	643	6	2,440	10,636	10,377
<b>Total</b>	<b>7,856</b>	<b>1,003</b>	<b>528</b>	<b>4,782</b>	<b>889</b>	<b>17,323</b>	<b>126,335</b>	<b>89,768</b>

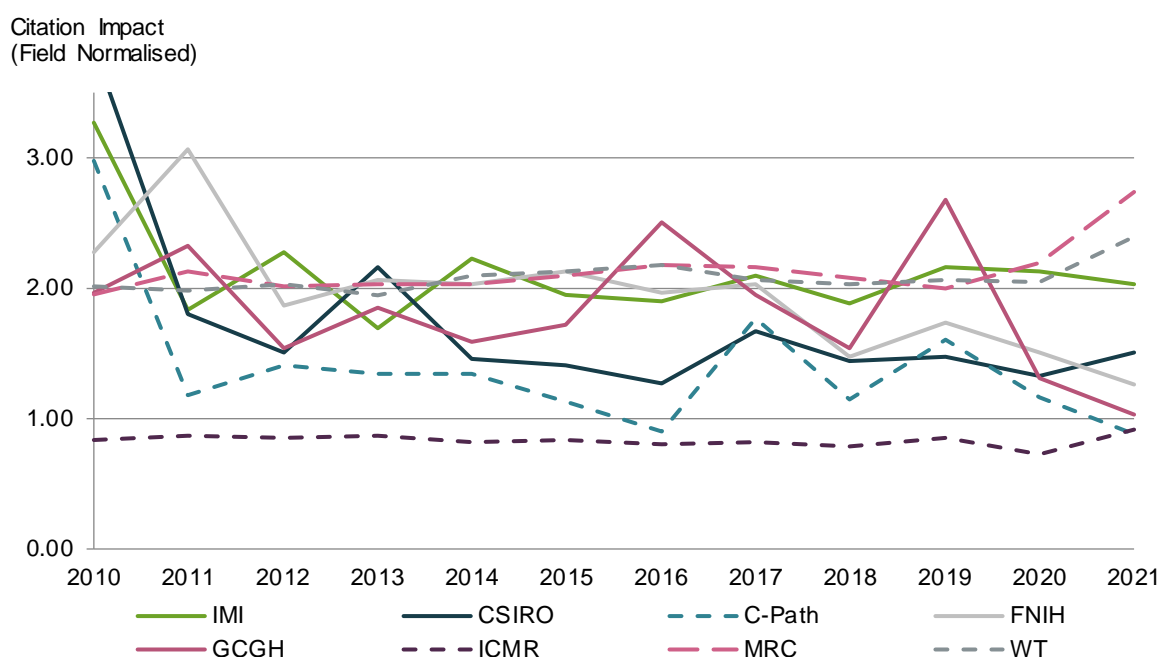


## 8.2.2 Trends in field-normalised citation impact: IMI project research compared with selected comparators

As discussed in Section 3, citations accumulate over time at a rate that is dependent upon the field of research. Therefore, it is standard bibliometric practice to normalise citation counts for these two factors. In this report, field-normalised citation impact has been calculated by dividing the citations received by each publication by the world average citations per publication for the relevant year and field.

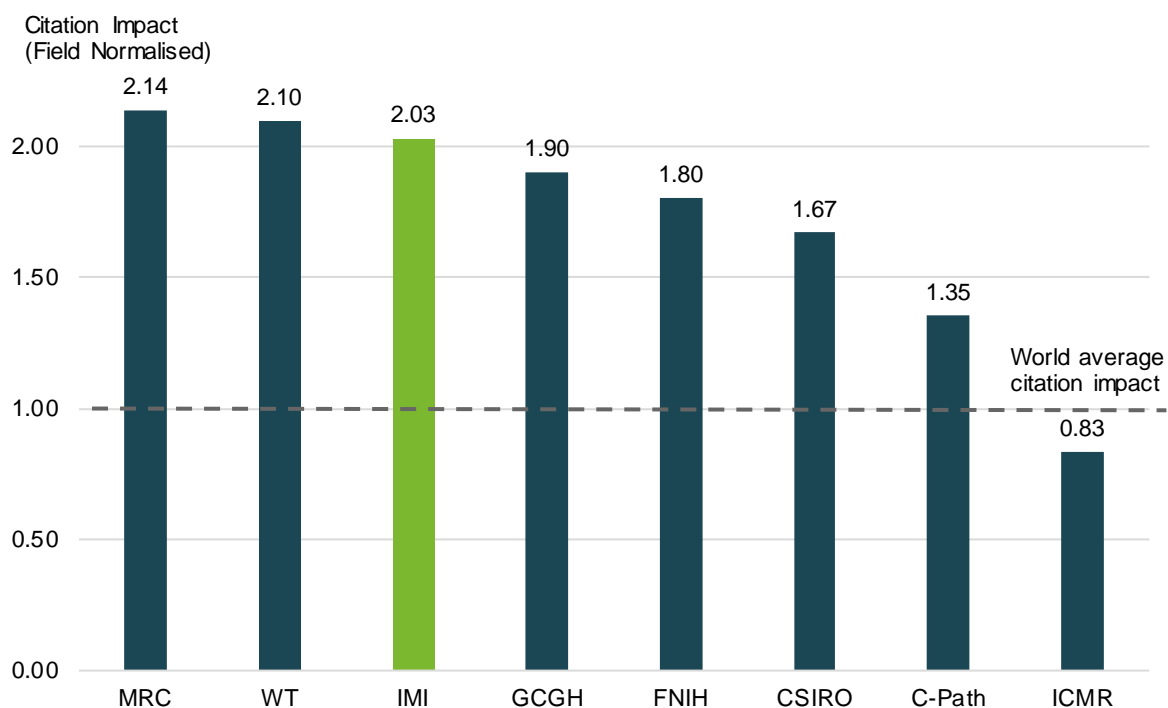
Figure 8.2.2.1 shows the annual trends in field-normalised citation impact of IMI and the comparators between 2010 and 2021 and Figure 8.2.2.2 shows the average field-normalised citation impact of IMI and the comparators between 2010 and 2021. Table 8.2.2.1 has the same data as in Figure 8.2.2.1 and Figure 8.2.2.2.

FIGURE 8.2.2.1 TRENDS IN FIELD-NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- The field-normalised citation impact of IMI, MRC and the WT were stable at close to twice the world average between 2010 and 2021, indicating highly cited, internationally significant research.
- The exceptionally high field-normalised citation impact of IMI, CSIRO, and C-Path project research in 2010 was driven by a small number of highly cited papers.
- ICMR has consistently underperformed in comparison to the world average between 2010-2021.
- In last year's report CSIRO's field normalized citation impact had dropped significantly in 2020, However, in this year's report we see that the field normalized citation impact for 2020 is similar to previous years supporting last year's conclusion that it was likely due to a significant amount of CSIRO's papers not having enough time to be cited. For 2021, CSIRO's citation impact (1.51) is slightly higher than 2020 (1.33).

Figure 8.2.2.2 AVERAGE FIELD-NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- The average field-normalised citation impact of IMI project research (2.03) between 2010 and 2021 was two times the world average and was comparable to MRC's and WT's citation impact and ahead of all other comparators.
- Only ICMR's average field-normalised citation impact (0.83) was below world average (1.00).

Table 8.2.2.1 FIELD-NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	3.27	3.84	2.98	2.28	1.96	0.84	1.95	2.02
2011	1.83	1.79	1.17	3.07	2.32	0.86	2.13	1.98
2012	2.27	1.51	1.41	1.87	1.54	0.85	2.01	2.03
2013	1.69	2.16	1.35	2.06	1.85	0.86	2.03	1.95
2014	2.22	1.46	1.35	2.04	1.59	0.82	2.02	2.10
2015	1.95	1.41	1.12	2.12	1.71	0.84	2.10	2.13
2016	1.89	1.27	0.90	1.96	2.51	0.80	2.17	2.17
2017	2.09	1.67	1.77	2.03	1.95	0.82	2.16	2.07
2018	1.89	1.43	1.15	1.47	1.54	0.79	2.08	2.02
2019	2.16	1.47	1.60	1.73	2.68	0.86	2.00	2.06
2020	2.13	1.33	1.17	1.50	1.31	0.73	2.20	2.05
2021	2.04	1.51	0.88	1.26	1.03	0.92	2.74	2.39
Average	2.03	1.67	1.35	1.80	1.90	0.83	2.14	2.10

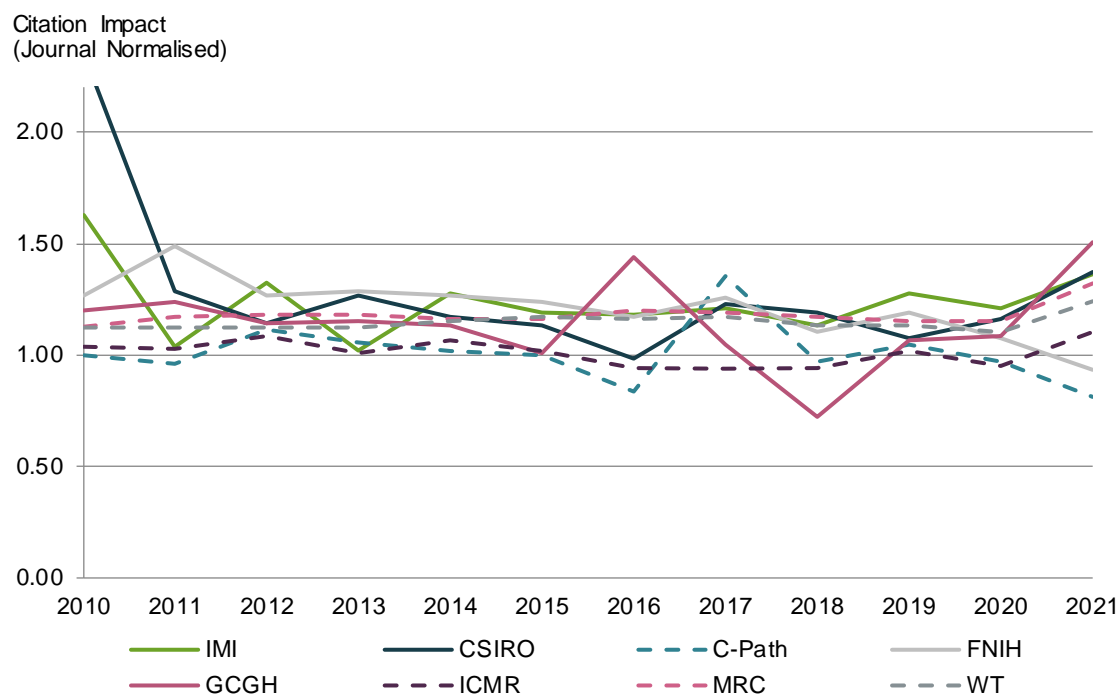
- In 2012 and 2014 IMI had the highest field-normalised citation impact (2.27 and 2.22 respectively) of the funding organisations analysed and since 2017 has remained in the Top 3.

### 8.2.3 Trends in journal-normalised citation impact: IMI project research compared with selected comparators

As discussed in Section 3, an alternative indicator to field-normalised citation impact is citation impact normalised at the journal level. The journal-normalised citation impact is calculated by dividing the number of citations a paper received by the average number of citations for the year and the journal in which the paper is published. As for the field-normalised citation impact, the world average for journal-normalised citation impact is 1.00.

Figure 8.2.3.1 shows the annual trends in journal-normalised citation impact of IMI and the comparators between 2010 and 2021. Figure 8.2.2.2 shows the average field-normalised citation impact of IMI and the comparators between 2010 and 2021. Table 8.2.3.1 shows the same data as in Figure 8.2.3.1 and Figure 8.2.3.2.

FIGURE 8.2.3.1 TRENDS IN JOURNAL NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- IMI project research has a journal-normalised citation impact that has remained above the world average between 2010 and 2021, indicating that IMI research performs well in the journals they are published.
- IMI projects had the highest journal normalised citation impact for 2012, 2014, 2019 and 2020.
- The journal-normalised citation impact of ICMR, MRC and WT remained relatively stable between 2010 and 2021, while that of the other comparators such as CSIRO and GCGH showed greater variability. This is to be expected given the smaller number of papers funded by CSIRO and GCGH relative to the output of research institutions like the MRC and WT.

FIGURE 8.2.3.2 AVERAGE JOURNAL-NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

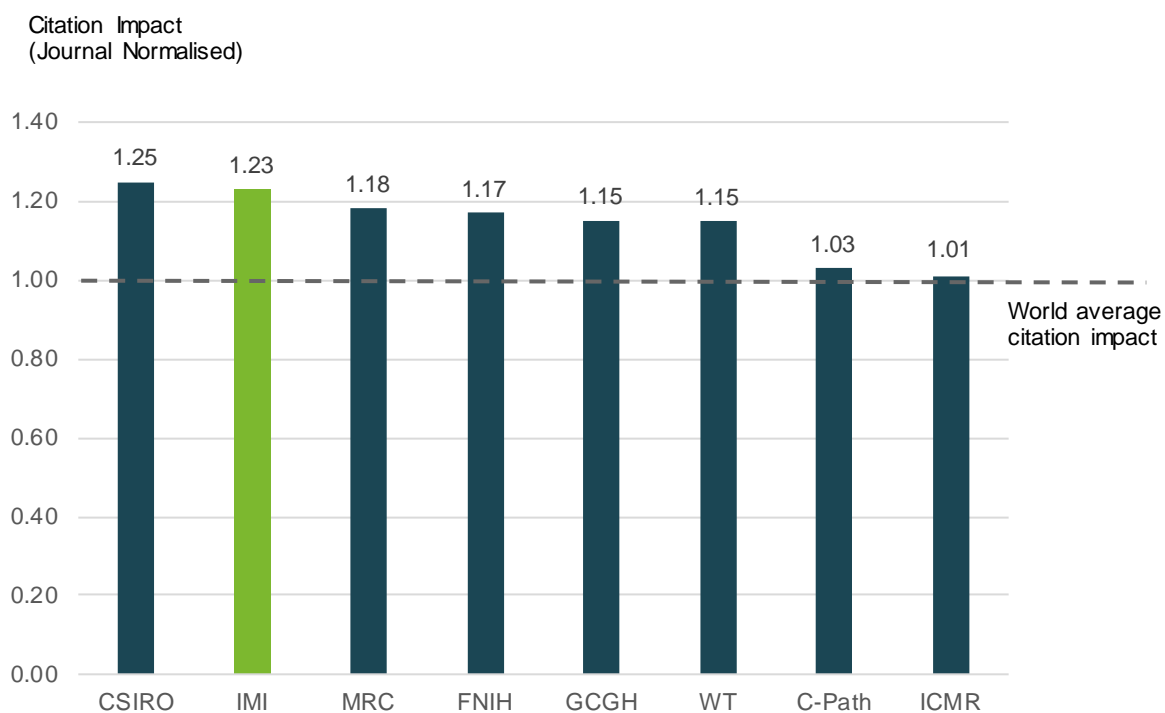


TABLE 8.2.3.1 JOURNAL-NORMALISED CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

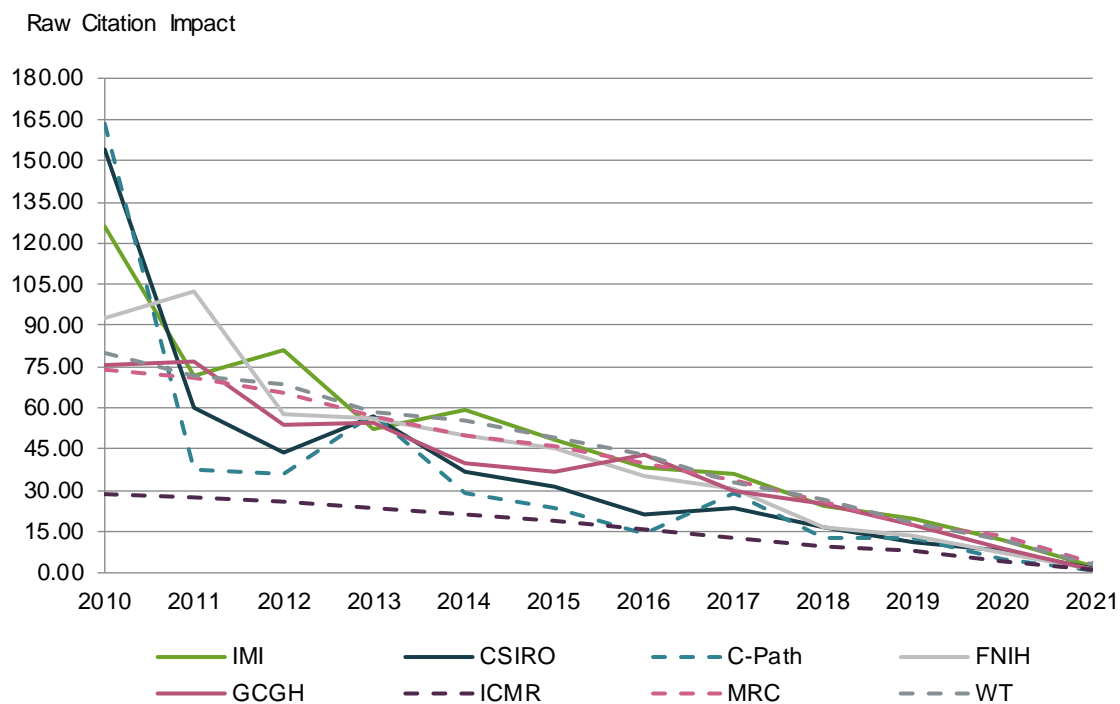
YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	1.63	2.35	1.00	1.26	1.20	1.04	1.13	1.12
2011	1.03	1.28	0.96	1.49	1.24	1.02	1.17	1.13
2012	1.32	1.14	1.11	1.26	1.14	1.08	1.18	1.12
2013	1.02	1.27	1.06	1.29	1.16	1.01	1.18	1.12
2014	1.28	1.17	1.02	1.27	1.14	1.06	1.16	1.15
2015	1.19	1.13	1.00	1.24	1.00	1.02	1.16	1.17
2016	1.18	0.98	0.84	1.17	1.43	0.95	1.20	1.16
2017	1.21	1.22	1.35	1.25	1.05	0.94	1.19	1.17
2018	1.13	1.19	0.97	1.10	0.72	0.94	1.17	1.13
2019	1.28	1.07	1.04	1.19	1.06	1.02	1.15	1.13
2020	1.21	1.16	0.97	1.08	1.08	0.95	1.15	1.10
2021	1.36	1.37	0.81	0.93	1.50	1.10	1.32	1.24
Average	1.23	1.25	1.03	1.17	1.15	1.01	1.18	1.15

## 8.2.4 Trends in raw citation impact: IMI project research compared with selected comparators

The raw (un-normalised) citation impact of a group of papers is calculated by dividing the sum of citations by the total number of papers published. As such it is the mean average number of citations to a paper. This indicator must be used with caution as it is not normalised to field or year.

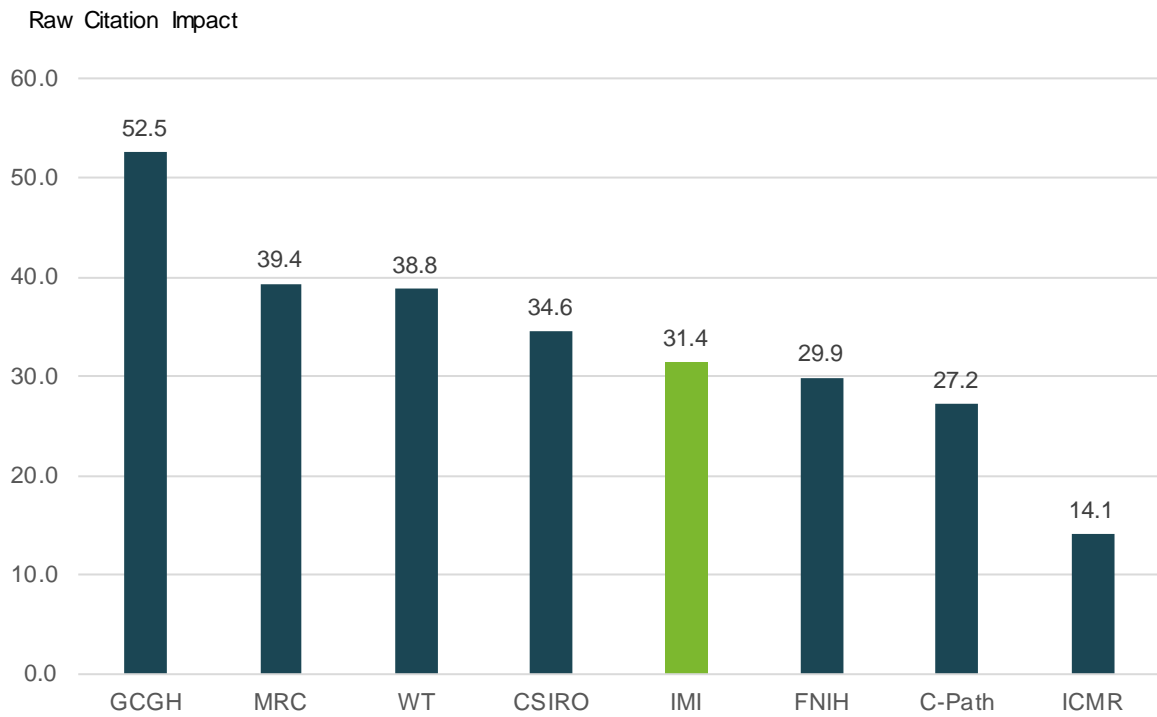
Figure 8.2.4.1 shows the annual trends in average raw citation impact of IMI and the comparators for papers published each year between 2010 and 2021. Figure 8.2.4.2 shows the average raw citation impact of IMI and the comparators for papers published between 2010 and 2021. Table 8.2.4.1 has the same data as in Figure 8.2.4.1 and Figure 8.2.4.2.

FIGURE 8.2.4.1 TRENDS IN RAW CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- The raw citation impact of all organisations in the most recent years between 2010 to 2021 are lower in comparison to previous years. This is expected as more recent publications have had less time to accumulate citations, and the raw citation impact is not normalised.
- IMI's 2021 raw citation impact (2.35) is higher than all comparators raw citation impacts except for WT and MRC

FIGURE 8.2.4.2 AVERAGE RAW CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- IMI's average raw citation impact between 2010 and 2021 (31.4) is higher than three out of the seven comparators (C-Path (27.2) ICMR (14.1) and FNIH (29.9)).
- IMI's Raw Citation impact increased the most (15%) from last year, relative to the comparators.
- GCGH had the highest raw citation impact (52.5).

TABLE 8.2.4.1 RAW CITATION IMPACT – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

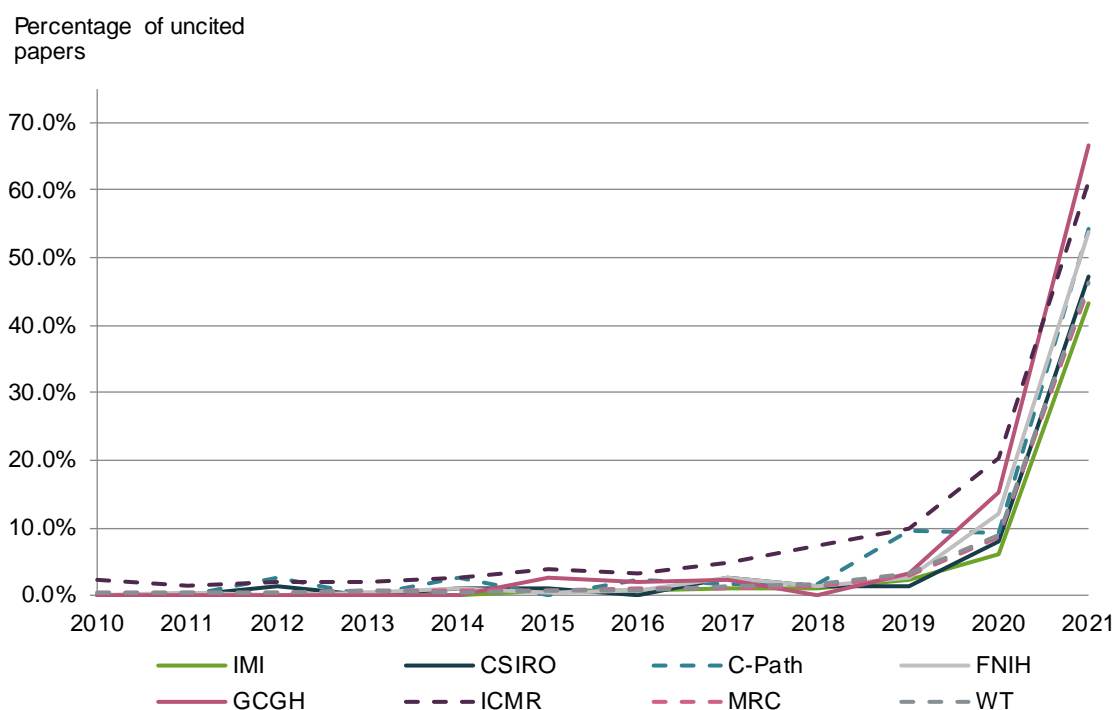
YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	126.04	154.09	163.48	92.82	75.41	28.64	73.92	80.03
2011	71.50	59.85	37.78	102.42	76.90	27.52	71.34	71.73
2012	80.79	43.63	36.25	57.78	53.75	25.79	65.37	68.87
2013	52.54	57.10	57.49	56.23	54.45	23.81	56.74	58.86
2014	59.05	36.82	28.95	50.28	40.18	21.58	50.33	55.43
2015	48.75	31.42	23.62	45.38	36.55	18.59	45.73	49.05
2016	38.29	21.40	13.93	34.89	42.60	15.44	39.57	42.82
2017	36.12	23.74	28.81	30.53	30.05	12.77	33.36	33.06
2018	24.16	16.46	12.78	16.75	24.88	9.71	25.68	26.56
2019	19.33	11.51	12.65	13.51	17.41	7.93	17.29	17.87
2020	12.24	7.68	4.63	7.15	8.77	4.48	13.11	12.29
2021	2.35	1.62	0.87	1.26	1.17	1.15	3.63	3.02
<b>Average</b>	<b>31.4</b>	<b>34.6</b>	<b>27.2</b>	<b>29.9</b>	<b>52.5</b>	<b>14.1</b>	<b>39.4</b>	<b>38.8</b>



## 8.2.5 Trends in uncited research: IMI project research compared with selected comparators

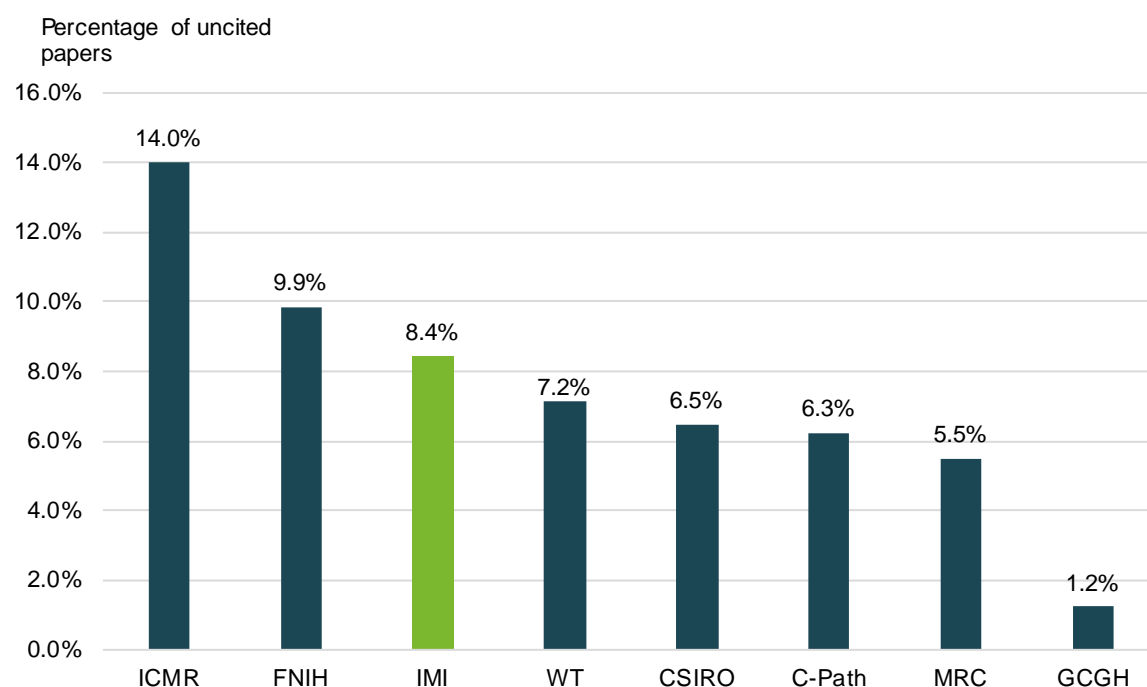
Most publication datasets will include papers which have no citations. Figure 8.2.5.1 shows the percentage of uncited papers between 2010 and 2021 for IMI and the selected comparators. Figure 8.2.5.1 shows the trend in average percentage of uncited papers between 2010 and 2021 for IMI and the selected comparators. Figure 8.2.5.2 shows the average percentage of uncited papers between 2010 and 2021 for IMI and the selected comparators. Table 8.2.5.1 has the same data as in Figure 8.2.5.1 and Figure 8.2.5.2.

FIGURE 8.2.5.1 TRENDS IN UNCITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- The similar trends in uncited papers indicate the similar citation life-cycle for biomedical research funded across all the benchmarking organisations. More recent publications are less likely to be cited than older publications. Therefore, the higher percentage of uncited papers in most recent years should not be taken as evidence that these articles are more likely to remain uncited.
- IMI has the lowest percentage of uncited papers in 2021. While ICMR has most often had one of the highest. This helps explain ICMR's lower than average citation impact.

FIGURE 8.2.5.1 AVERAGE PERCENTAGE OF UNCITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- Around 8% of IMI project papers remained uncited between 2010 and 2021
- C-Paths percentage of uncited papers has decreased by nearly 4% since last year's report. The largest change of all the comparators.
- GCGH has the lowest percentage of uncited papers, around 2% of its papers uncited

TABLE 8.2.5.1 PERCENTAGE OF UNCITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	0.0%	0.0%	0.0%	0.0%	0.0%	2.3%	0.4%	0.4%
2011	0.0%	0.0%	0.0%	0.5%	0.0%	1.4%	0.5%	0.4%
2012	0.0%	1.3%	2.6%	0.0%	0.0%	2.0%	0.4%	0.4%
2013	0.5%	0.0%	0.0%	0.4%	0.0%	2.1%	0.6%	0.6%
2014	0.0%	1.1%	2.5%	0.9%	0.0%	2.4%	0.7%	0.4%
2015	0.7%	1.0%	0.0%	0.3%	2.5%	3.9%	0.7%	0.8%
2016	0.6%	0.0%	2.4%	0.6%	1.9%	3.3%	1.0%	0.8%
2017	1.1%	2.7%	1.5%	2.7%	2.3%	4.8%	1.2%	1.3%
2018	0.9%	1.3%	1.8%	1.4%	0.0%	7.4%	1.5%	1.6%
2019	2.4%	1.4%	9.4%	2.7%	3.1%	9.9%	2.8%	3.3%
2020	6.1%	8.0%	9.1%	12.2%	15.4%	20.2%	8.7%	9.0%
2021	43.3%	47.2%	54.3%	53.8%	66.7%	61.3%	45.6%	46.3%
<b>Total</b>	<b>8.4%</b>	<b>6.5%</b>	<b>6.3%</b>	<b>9.9%</b>	<b>1.2%</b>	<b>14.0%</b>	<b>5.5%</b>	<b>7.2%</b>

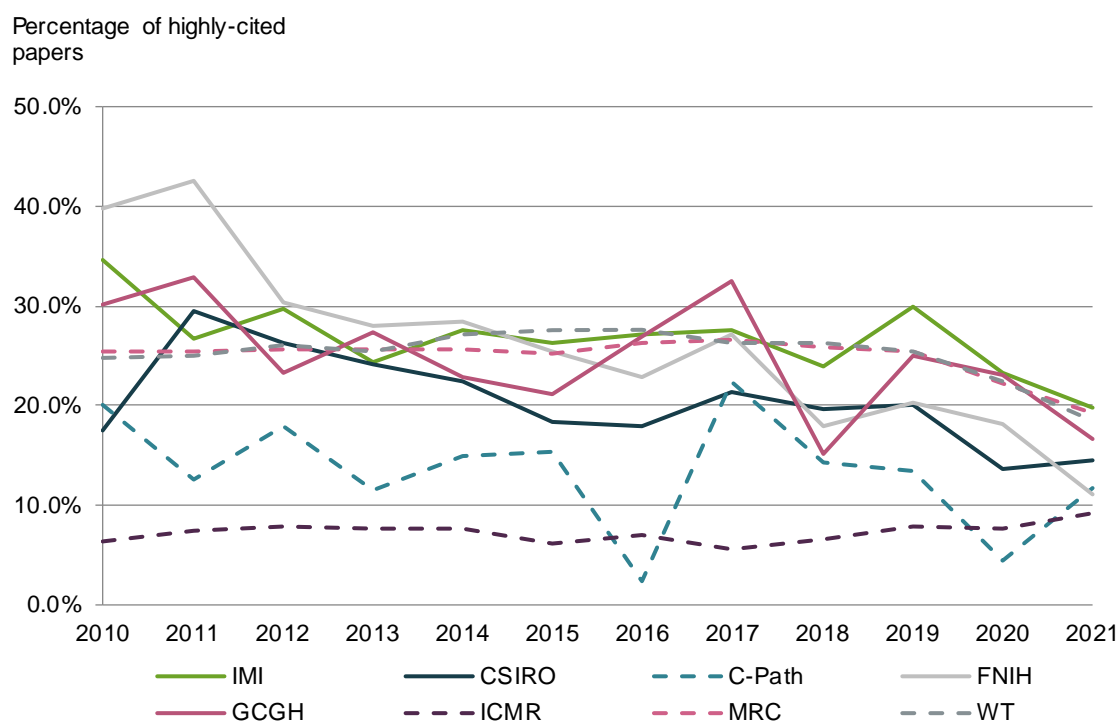
- No IMI project papers published between 2010 and 2012 or in 2014 are uncited.

### 8.2.6 Trends in highly cited research: IMI project research compared with selected comparators

As discussed in Section 3, highly cited work is recognised as having a greater impact, and citation counts have been correlated with other qualitative evaluations of research performance, such as peer review. For institutional research evaluation, we have found that the world’s top 10% of most highly cited papers is often a suitable definition of highly cited work. Therefore, if more than 10% of an entity’s publications are in the top 10% of the world’s most highly cited papers, then it has performed better than expected.

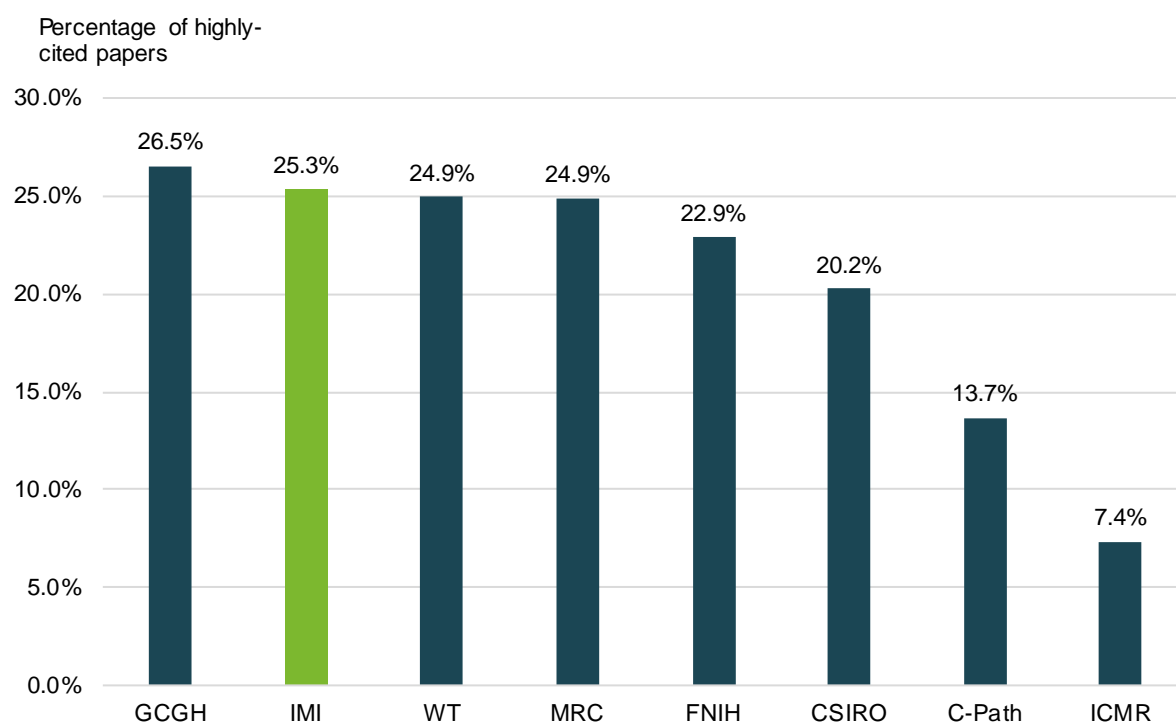
Figure 8.2.6.1 shows the annual trends in percentage of highly cited papers between 2010 and 2021 for IMI and the selected comparators. Figure 8.2.6.2 shows the total percentage of highly cited papers between 2010 and 2021 for IMI and the selected comparators. Table 8.2.6.1 has the same data as in Figure 8.2.6.1 and Figure 8.2.6.2.

FIGURE 8.2.6.1 TRENDS IN HIGHLY CITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- Between 2010 and 2021, IMI and most of the comparators had an above average percentage (10%) of highly cited papers the exceptions being ICMR, which was consistently below the world average, and C-Path, which was below average in 2016 and 2020.
- In most years, IMI is among the organisations with the highest percentage of highly cited papers. IMI has had the highest percentage of highly-cited papers since 2019.

FIGURE 8.2.6.2 PERCENTAGE OF HIGHLY CITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- IMI ranks second in comparison to the comparators for percentage of highly-cited papers, with only GCGH outperforming IMI.
- Around a quarter of papers published by IMI and the comparators between 2010 and 2021 were highly cited. C-Path had a comparatively lower proportions of highly cited papers (13.7%) while ICMR was well below world average performance (7.4%).

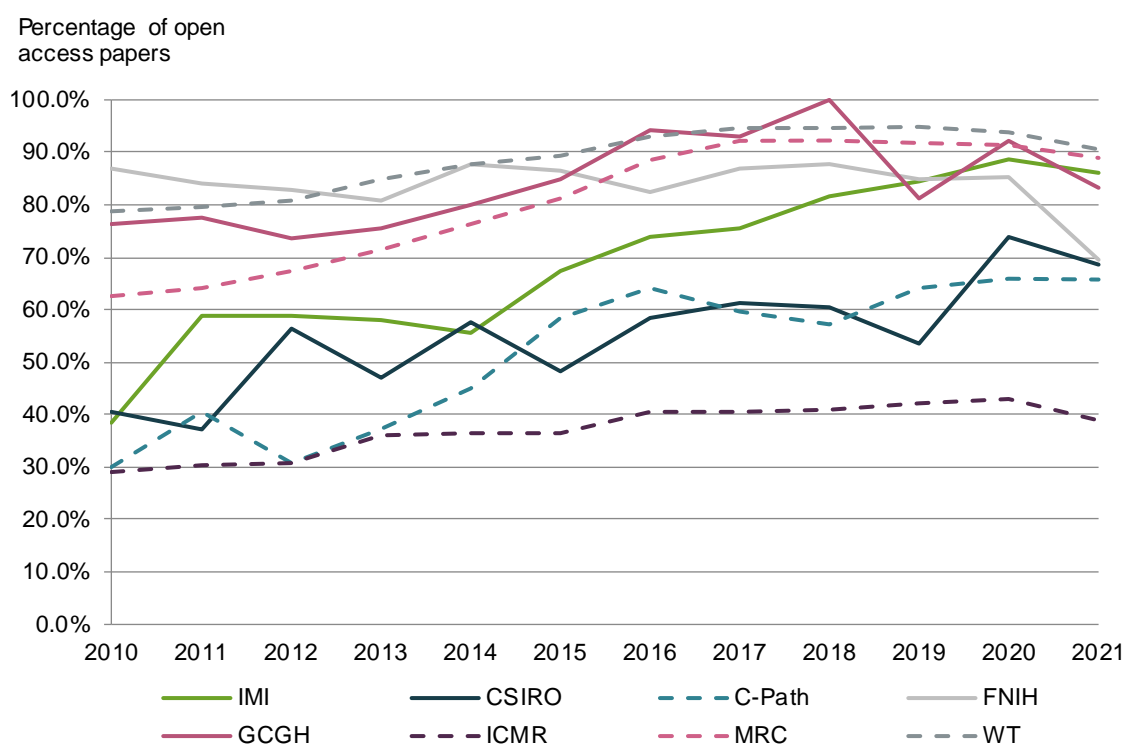
TABLE 8.2.6.1 PERCENTAGE OF HIGHLY CITED PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	34.6%	17.5%	20.0%	39.8%	30.2%	6.4%	25.4%	24.9%
2011	26.8%	29.5%	12.5%	42.6%	32.9%	7.4%	25.5%	25.0%
2012	29.8%	26.3%	17.9%	30.4%	23.2%	7.8%	25.7%	26.0%
2013	24.3%	24.1%	11.4%	28.1%	27.4%	7.6%	25.7%	25.4%
2014	27.6%	22.3%	15.0%	28.3%	22.8%	7.6%	25.7%	27.2%
2015	26.3%	18.3%	15.4%	25.5%	21.2%	6.2%	25.2%	27.5%
2016	27.2%	18.0%	2.4%	22.8%	26.9%	7.1%	26.3%	27.6%
2017	27.7%	21.3%	22.4%	27.1%	32.6%	5.6%	26.6%	26.3%
2018	24.0%	19.7%	14.3%	18.0%	15.2%	6.5%	25.9%	26.2%
2019	29.9%	20.0%	13.5%	20.4%	25.0%	7.9%	25.5%	25.4%
2020	23.2%	13.6%	4.5%	18.2%	23.1%	7.7%	22.2%	22.3%
2021	19.8%	14.4%	11.8%	11.1%	16.7%	9.2%	19.3%	18.5%
<b>Total</b>	<b>25.3%</b>	<b>20.2%</b>	<b>13.7%</b>	<b>22.9%</b>	<b>26.5%</b>	<b>7.4%</b>	<b>24.9%</b>	<b>24.9%</b>

## 8.2.7 Trends in open access research: IMI project research compared with selected comparators

Figure 8.2.7.1 shows annual trends in the percentage of open access papers between 2010 and 2021 for IMI and the selected comparators. Figure 8.2.7.2 shows the total percentage of open access papers between 2010 and 2021 for IMI and the selected comparators. Table 8.2.7.1 shows the same data as in Figure 8.2.7.1 and Figure 8.2.7.2.<sup>25</sup>

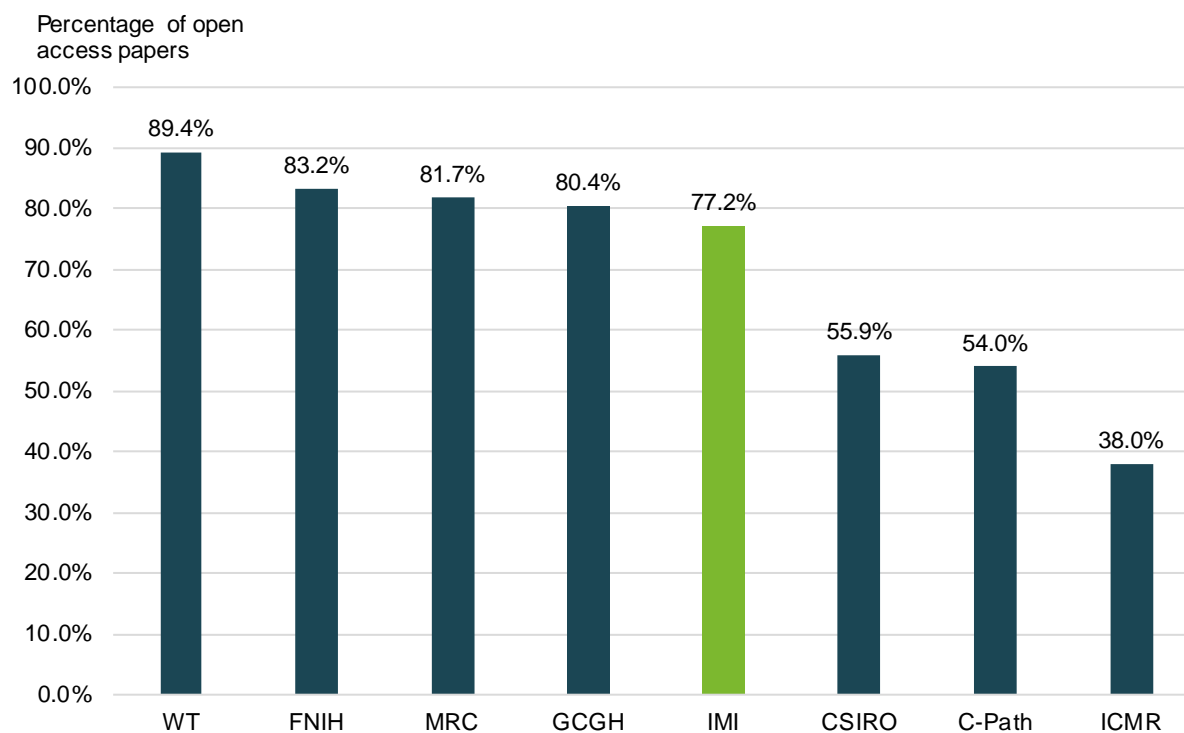
FIGURE 8.2.7.1 TRENDS IN OPEN ACCESS PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- IMI and most of the comparators have increased their output of open access papers between 2010 and 2021, except for FNIH which is trending downward.
- IMI increased its percentage of open access papers at a faster rate than any of the comparators.

<sup>25</sup> The Web of Science open access data come from the Directory of Open Access Journals (DOAJ) and collaborations with Impact Story and Our Research's Unpaywall services. The Web of Science therefore provides unrivalled coverage of open access publications that are published through DOAJ Gold, Other Gold, Green Published, Green Accepted or Bronze routes. It is also possible that some publishers make publications available without following a recognised open access route. In these cases publications will not be indexed as open access in the Web of Science or in this report.

FIGURE 8.2.7.2 TOTAL PERCENTAGE OF OPEN ACCESS PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021



- Most organisations, including IMI, have published more than half of their publications as open access. IMI had a lower share of open access papers compared to FNIH, GCGH, MRC, and WT.
- WT has the highest total percentage of open access papers (89.4%) between 2010 and 2021. In contrast ICMR, had the lowest percentage of open access papers (38.0%).

TABLE 8.2.7.1 PERCENTAGE OF OPEN ACCESS PAPERS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

YEAR	IMI	CSIRO	C-PATH	FNIH	GCGH	ICMR	MRC	WT
2010	38.5%	40.4%	30.0%	86.7%	76.3%	29.0%	62.6%	78.8%
2011	58.8%	37.2%	40.6%	84.0%	77.4%	30.2%	64.1%	79.4%
2012	58.7%	56.3%	30.8%	82.9%	73.6%	30.6%	67.5%	80.9%
2013	58.0%	47.0%	37.1%	80.7%	75.5%	35.9%	71.5%	84.8%
2014	55.6%	57.4%	45.0%	87.5%	79.8%	36.6%	76.3%	87.7%
2015	67.2%	48.1%	58.5%	86.3%	85.0%	36.3%	81.3%	89.4%
2016	73.9%	58.4%	64.3%	82.4%	94.2%	40.6%	88.7%	93.1%
2017	75.6%	61.3%	59.7%	86.9%	93.0%	40.5%	92.1%	94.6%
2018	81.8%	60.5%	57.1%	87.8%	100.0%	40.8%	92.3%	94.5%
2019	84.7%	53.5%	64.2%	84.9%	81.3%	42.3%	91.7%	94.9%
2020	88.7%	73.9%	65.9%	85.3%	92.3%	43.0%	91.5%	93.9%
2021	86.1%	68.5%	65.7%	69.5%	83.3%	38.9%	88.9%	90.7%
<b>Total</b>	<b>77.2%</b>	<b>55.9%</b>	<b>54.0%</b>	<b>83.2%</b>	<b>80.4%</b>	<b>38.0%</b>	<b>81.7%</b>	<b>89.4%</b>

### 8.3 Summary of bibliometric indicators: IMI project research compared with selected comparators

Although IMI has only been funding research for just over a decade, its performance is on par with well-established funding bodies that have been operating for much longer, like the MRC and the Wellcome Trust, as indicated by comparable citation impacts, and percentages of highly cited papers (Table 8.3.1).

Table 8.3.2 SUMMARY OF BIBLIOMETRIC INDICATORS – IMI PROJECT RESEARCH COMPARED WITH SELECTED COMPARATORS, 2010-2021

PROJECT	NUMBER OF PAPERS	CITATION IMPACT (NORMALISED AT FIELD LEVEL)	PERCENTAGE OF UNCITED PAPERS	PERCENTAGE OF HIGHLY CITED PAPERS
IMI	7,856	2.03	8.4%	25.3%
C-Path	528	1.35	6.3%	13.7%
CSIRO	1,003	1.67	6.5%	20.2%
FNIH	4,782	1.80	9.9%	22.9%
GCGH	889	1.90	1.2%	26.5%
ICMR	17,323	0.83	14.0%	7.4%
MRC	126,335	2.14	5.5%	24.9%
WT	89,768	2.10	7.2%	24.9%



## ANNEX 1: Bibliometrics and citation analysis

Bibliometrics are about publications and their citations. The academic field emerged from 'information science' and now usually refers to the methods used to study and index texts and information.

Publications cite other publications. These citation links grow into networks, and their numbers are likely to be related to the significance or impact of the publication. The meaning of the publication is determined from keywords and content. Citation analysis and content analysis have therefore become a common part of bibliometric methodology. Historically, bibliometric methods were used to trace relationships amongst academic journal citations. Now, bibliometrics are important in indexing research performance.

Bibliometric data have particular characteristics of which the user should be aware, and these are considered here.

Journal papers (publications, sources) report research work. Papers refer to or 'cite' earlier work relevant to the material being reported. New papers are cited in their turn. Papers that accumulate more citations are thought of as having greater 'impact', which is interpreted as significance or influence on their field. Citation counts are therefore recognised as a measure of impact, which can be used to index the excellence of the research from a particular group, institution or country.

The origins of citation analysis as a tool that could be applied to research performance can be traced to the mid-1950s, when Eugene Garfield proposed the concept of citation indexing and introduced the Science Citation Index, the Social Sciences Citation Index and the Arts & Humanities Citation Index, produced by the Institute of Scientific Information (now Clarivate).<sup>26</sup>

We can count citations, but they are only 'indicators' of impact or quality – not metrics. Most impact indicators use average citation counts from groups of papers, because some individual papers may have unusual or misleading citation profiles. These outliers are diluted in larger samples.

### **Data source**

The data we use come from the Clarivate Web of Science databases which give access not only to journals but also to conference proceedings, books, patents, websites, and chemical structures, compounds and reactions. It has a unified structure that integrates all data and search terms together and therefore provides a level of comparability not found in other databases. It is widely acknowledged to be the world's leading source of citation and bibliometric data. The Clarivate Web of Science Core Collection is part of the Web of Science and focuses on research published in journals and conferences in science, medicine, arts, humanities and social sciences.

The Web of Science was originally created as an awareness and information retrieval tool but it has acquired an important primary use as a tool for research evaluation, using citation analysis and bibliometrics. Data coverage is both current and retrospective in the sciences, social sciences, arts and humanities, in some cases back to 1900. Within the research community this data source was previously referred to by the acronym 'ISI'.

Unlike other databases, the Web of Science and underlying databases are selective, that is: the journals abstracted are selected using rigorous editorial and quality criteria. The authoritative,

---

<sup>26</sup> Garfield, E (1955) Citation Indexes for Science – New dimension in documentation through association of ideas. *Science*: 122, 108-111.

multidisciplinary content covers over 12,000 of the highest impact journals worldwide, including open access journals, and over 150,000 conference proceedings. The abstracted journals encompass the majority of significant, frequently cited scientific reports and, more importantly, an even greater proportion of the scientific research output which is cited. This selective process ensures that the citation counts remain relatively stable in given research fields and do not fluctuate unduly from year to year, which increases the usability of such data for performance evaluation.

Clarivate has extensive experience with databases on research inputs, activity and outputs and has developed innovative analytical approaches for benchmarking and interpreting international, national and institutional research impact.

### ***Database categories***

The source data can be grouped in various classification systems. Most of these are based on groups of journals that have a relatively high cross-citation linkage and naturally cluster together. Custom classifications use subject maps in third-party data such as the OECD categories set out in the Frascati manual.

Clarivate frequently uses the broader field categories in the InCites: Essential Science Indicators™ and the finer journal categories in the Web of Science. There are 22 fields in Essential Science Indicators and 254 fields in Web of Science. In either case, our bibliometric analyses draw on the full range of data available in the underlying database, so analyses in our reports will differ slightly from anything created ‘on the fly’ from data in the web interface.

The lists of journal categories in these systems are attached at the end of this document.

Most analyses start with an overall view across the data, then move to a view across broad categories and only then focus in at a finer level in the areas of greatest interest to policy, programme or organisational purpose.

### ***Assigning papers to addresses***

A paper is assigned to each country and each organisation whose address appears at least once for any author on that paper. One paper counts once and only once for each assignment, however many address variants occur for the country or organisation. No weighting is applied.

For example, a paper has five authors, thus:

<b>AUTHOR</b>	<b>ORGANISATION</b>	<b>COUNTRY</b>		
<b>Gurney, KA</b>	Univ Leeds	UK	<b>Counts for Univ Leeds</b>	<b>Counts for UK</b>
<b>Adams, J</b>	Univ Leeds	UK	No gain for Univ Leeds	No gain for UK
<b>Kochalko, D</b>	Univ C San Diego	USA	<b>Counts for UCSD</b>	<b>Counts for USA</b>
<b>Munshi, S</b>	Gujarat Univ	India	<b>Counts for Gujarat Univ</b>	<b>Counts for India</b>
<b>Pendlebury, D</b>	Univ Oregon	USA	<b>Counts for Univ Oregon</b>	No gain for USA

So this one paper with five authors would be included once in the tallies for each of four universities and once in the tallies for each of three countries.

Work carried out within Clarivate, and research published elsewhere, indicates that fractional weighting based on the balance of authors by organisation and country makes little difference to the conclusions of an analysis at an aggregate level. Such fractional analysis can introduce unforeseen errors in the attempt to create a detailed but uncertain assignment. Partitioning credit would make a greater difference at a detailed, group level but the analysis can then be manually validated.

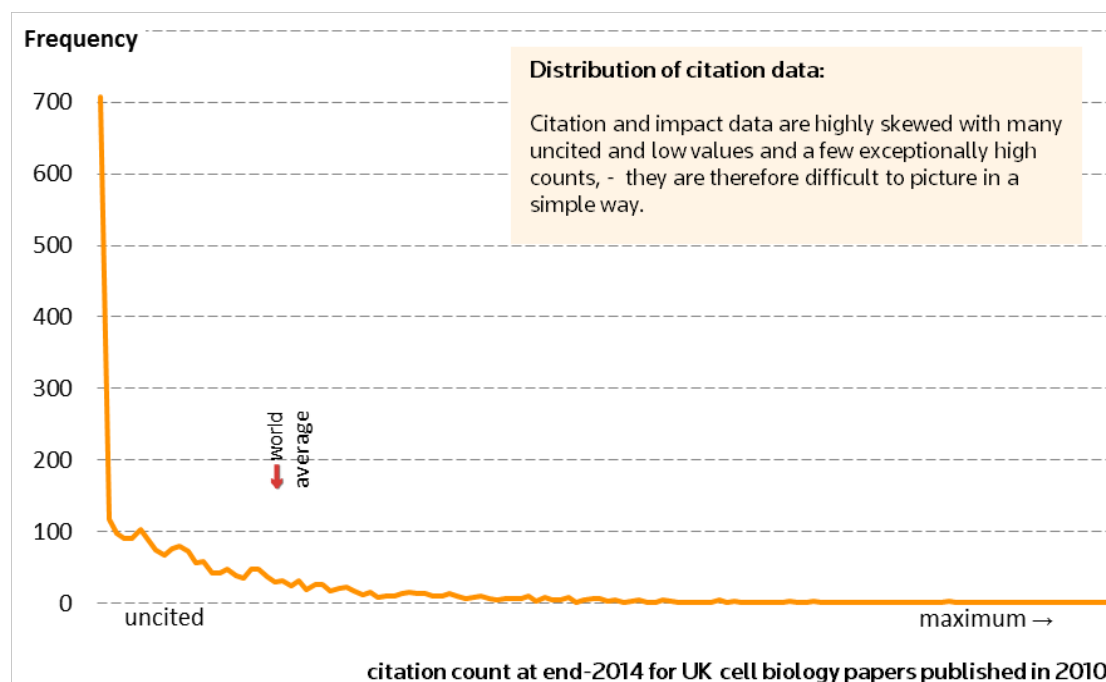
### Citation counts

A publication accumulates citation counts when it is referred to by more recent publications. Some papers get cited frequently and many get cited rarely or never, so the distribution of citations is highly skewed.

Why are many papers never cited? Certainly some papers remain uncited because their content is of little or no impact, but that is not the only reason. It might be because they have been published in a journal not read by researchers to whom the paper might be interesting. It might be that they represent important but 'negative' work reporting a blind alley to be avoided by others. The publication may be a commentary in an editorial, rather than a normal journal article and thus of general rather than research interest. Or it might be that the work is a 'sleeping beauty' that has yet to be recognised for its significance.

Other papers can be very highly cited: hundreds, even thousands of times. Again, there are multiple reasons for this. Most frequently cited work is being recognised for its innovative significance and impact on the research field of which it speaks. Impact here is a good reflection of quality: it is an indicator of excellence. But there are other papers which are frequently cited because their significance is slightly different: they describe key methodology; they are a thoughtful and wide-ranging review of a field; or they represent contentious views which others seek to refute.

Citation analysis cannot make value judgments about why an article is uncited nor about why it is highly cited. The analysis can only report the citation impact that the publication has achieved. We normally assume, based on many other studies linking bibliometric and peer judgments, that high citation counts correlate on average with the quality of the research.



The figure shows the skewed distribution of more or less frequently cited papers from a sample of UK authored publications in cell biology. The skew in the distribution varies from field to field. It is to compensate for such factors that actual citation counts must be normalised, or rebased, against a world baseline.

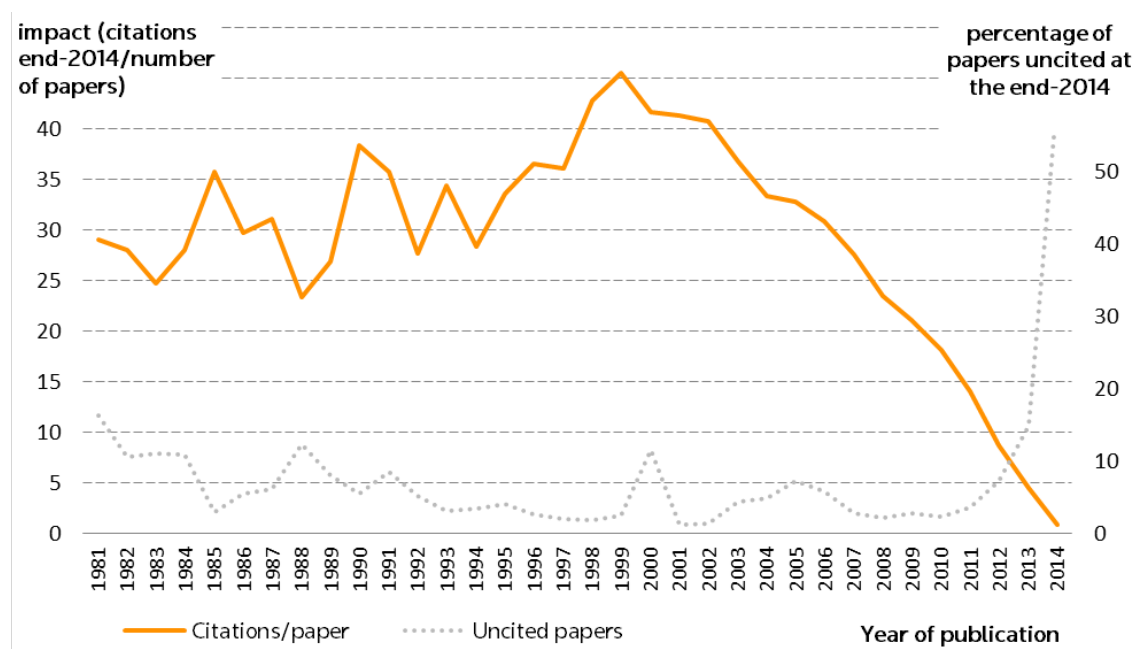
We do not seek to account separately for the effect of self-citation. If the citation count is significantly affected by self-citation then the paper is likely to have been infrequently cited. This is therefore only of consequence for low impact activity. Studies show that for large samples at national and organisational level the effect of self-citation has little or no effect on the analytical outcomes and would not alter interpretation of the results.

### Time factors

Citations accumulate over time. Older papers therefore have, on average, more citations than more recent work. The graph below shows the pattern of citation accumulation for a set of 33 journals in the journal category **Materials Science, Biomaterials**. Papers less than eight years old are, on average, still accumulating additional citations. The citation count goes on to reach a plateau for older sources.

The graph shows that the percentage of papers that have never been cited drops over about five years. Beyond five years, between 5% and 10% or more of papers remain uncited.

Account must be taken of these time factors in comparing current research with historical patterns. For these reasons, it is sometimes more appropriate to use a fixed five-year window of papers and citations to compare two periods than to look at the longer term profile of citations and of uncitedness for a recent year and an historical year.



### Discipline factors

Citation rates vary between disciplines and fields. For the UK science base as a whole, ten years produces a general plateau beyond which few additional citations would be expected. On the whole, citations accumulate more rapidly and plateau at a higher level in biological sciences than physical sciences, and natural sciences generally cite at a higher rate than social sciences.

Papers are assigned to disciplines (journal categories or research fields) by Clarivate, bringing cognate research areas together. The journal category classification scheme has been recently revised and updated. Before 2007, journals were assigned to the older, well established Current Contents categories which were informed by extensive work by Thomson and with the research community since the early 1960s. This scheme has been superseded by the 252 Web of Science journal categories which allow for greater disaggregation for the growing volume of research which is published and abstracted.

Papers are allocated according to the journal in which the paper is published. Some journals may be considered to be part of the publication record for more than one research field. As the example below illustrates, the journal *Acta Biomaterialia* is assigned to two journal categories: **Materials Science, Biomaterials and Engineering, Biomedical**.

Very few papers are not assigned to any research field and as such will not be included in specific analyses using normalised citation impact data. The journals included in the Clarivate databases and how they are selected are detailed here: [mjl.clarivate.com/](http://mjl.clarivate.com/).

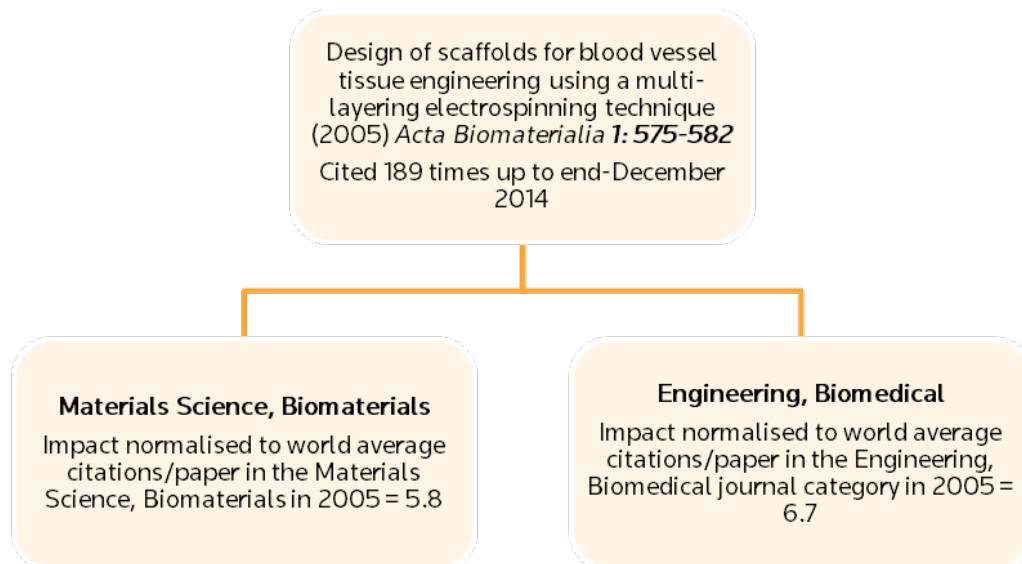
Some journals with a very diverse content, including the prestigious journals *Nature and Science* were classified as **Multidisciplinary** in databases created prior to 2007. The papers from these **Multidisciplinary** journals are now re-assigned to more specific research fields using an algorithm based on the research area(s) of the references cited by the article.

### **Normalised citation impact**

Because citations accumulate over time at a rate that is dependent upon the field of research, all analyses must take both field and year into account. In other words, because the absolute citation count for a specific article is influenced by its field and by the year it was published, we can only make comparisons of indexed data after normalising with reference to these two variables.

We only use citation counts for reviews and articles in calculations of impact, because document type influences the citation count. For example, a review will often be cited more frequently than an article in the same field, but editorials and meeting abstracts are rarely cited and citation rates for conference proceedings are extremely variable. The most common normalisation factors are the average citations per paper for (1) the year and (2) either the field or the journal in which the paper was published. This normalisation is also referred to as 'rebasings' the citation count.

Impact is therefore most commonly analysed in terms of 'normalised impact', or NCI. The following schematic illustrates how the normalised citation impact is calculated at paper level and journal category level.



This article in the journal *Acta Biomaterialia* is assigned to two journal categories: **Materials Science, Biomaterials** and **Engineering, Biomedical**. The world average baselines for, as an example, **Materials science, Biomaterials** are calculated by summing the citations to all the articles and reviews published worldwide in the journal *Acta Biomaterialia* and the other 32 journals assigned to this category for each year and dividing this by the total number of articles and reviews published in the journal category. This gives the category-specific normalised citation impact (in the above example the category-specific field-normalised citation impact for **Materials Science, Biomaterials** is 5.8 and the category-specific field-normalised citation impact for **Engineering, Biomedical** is higher at 6.7). Most papers (nearly two-thirds) are assigned to a single journal category whilst a minority are assigned to more than 5.

Citation data provided by Clarivate are assigned on an annual census date referred to as the Article Time Period. For the majority of publications, the Article Time Period is the same as the year of publication, but for a few publications (especially those published at the end of the calendar year in less main-stream journals) the Article Time Period may vary from the actual year of publication.

World average impact data are sourced from the Clarivate National Science Indicators baseline data for 2016.

### Mean normalised citation impact

Research performance has historically been indexed by using average citation impact, usually compared to a world average that accounts for time and discipline. As noted, however, the distribution of citations amongst papers is highly skewed because many papers are never cited while a few papers accumulate very large citation counts. That means that an average may be misleading if assumptions are made about the distribution of the underlying data.

In fact, almost all research activity metrics are skewed: for research income, PhD numbers and publications there are many low activity values and a few exceptionally high values. In reality, therefore, the skewed distribution means that average impact tends to be greater than and often significantly different from either the median or mode in the distribution. This should be borne in mind when reviewing analytical outcomes.

The average (normalised) citation impact can be calculated at an individual paper level where it can be associated with more than one journal category. It can also be calculated for a set of papers at any level from a single country to an individual researcher's output. In the example above, the average citation impact of the Acta Biomaterialia paper can be expressed as  $((5.8 + 6.7)/2) = 6.3$ .

### **What are uncited papers?**

It may be a surprise that some journal papers are never subsequently cited after publication, even by their authors. This accounts for about half the total global output for a typical, recent 10-year period. We cannot tell why papers are not cited. It is likely that a significant proportion of papers remain uncited because they are reporting negative results which are an essential matter of record in their field but make the content less likely to be referenced in other papers. Inevitably, other papers are uncited because their content is trivial or marginal to the mainstream. However, it should not be assumed that this is the case for all such papers.

There is variation in non-citation between countries and between fields. For example, relatively more engineering papers tend to remain uncited than papers in other sciences, indicative of a disciplinary factor but not a quality factor. While there is also an obvious increase in the likelihood of citation over time, most papers that are going to be cited will be cited within a few years of publication.

## Journal category systems used in our analyses.

### WEB OF SCIENCE

Acoustics	Classics	Engineering, multidisciplinary
Agricultural economics & policy	Clinical neurology	Engineering, ocean
Agricultural engineering	Communication	Engineering, petroleum
Agriculture, dairy & animal science	Computer science, artificial intelligence	Entomology
Agriculture, multidisciplinary	Computer science, cybernetics	Environmental sciences
Agriculture, soil science	Computer science, hardware & architecture	Environmental studies
Agronomy	Computer science, information systems	Ergonomics
Allergy	Computer science, interdisciplinary applications	Ethics
Anatomy & morphology	Computer science, software engineering	Ethnic studies
Andrology	Computer science, theory & methods	Evolutionary biology
Anesthesiology	Construction & building technology	Family studies
Anthropology	Criminology & penology	Film, radio, television
Applied linguistics	Critical care medicine	Fisheries
Archaeology	Crystallography	Folklore
Architecture	Dance	Food science & technology
Area studies	Demography	Forestry
Art	Dentistry, oral surgery & medicine	Gastroenterology & hepatology
Asian studies	Dermatology	Genetics & heredity
Astronomy & astrophysics	Developmental biology	Geochemistry & geophysics
Automation & control systems	Ecology	Geography
Behavioral sciences	Economics	Geography, physical
Biochemical research methods	Education & educational research	Geology
Biochemistry & molecular biology	Education, scientific disciplines	Geosciences, multidisciplinary
Biodiversity conservation	Education, special	Geriatrics & gerontology
Biology	Electrochemistry	Health care sciences & services
Biology, miscellaneous	Emergency medicine	Health policy & services
Biophysics	Endocrinology & metabolism	Hematology
Biotechnology & applied microbiology	Energy & fuels	History
Business	Engineering, aerospace	History & philosophy of science
Business, finance	Engineering, biomedical	History of social sciences
Cardiac & cardiovascular systems	Engineering, chemical	Horticulture
Cell biology	Engineering, civil	Humanities, multidisciplinary
Chemistry, analytical	Engineering, electrical & electronic	Imaging science & photographic technology
Chemistry, applied	Engineering, environmental	Immunology
Chemistry, inorganic & nuclear	Engineering, geological	Industrial relations & labor
Chemistry, medicinal	Engineering, industrial	Infectious diseases
Chemistry, multidisciplinary	Engineering, manufacturing	Information & library science
Chemistry, organic	Engineering, marine	Instruments & instrumentation
Chemistry, physical	Engineering, mechanical	Integrative & complementary medicine
International relations	Mining & mineral processing	Psychology
Language & linguistics	Multidisciplinary sciences	Psychology, applied



Language & linguistics theory	Music	Psychology, biological
Law	Mycology	Psychology, clinical
Limnology	Nanoscience & nanotechnology	Psychology, developmental
Linguistics	Neuroimaging	Psychology, educational
Literary reviews	Neurosciences	Psychology, experimental
Literary theory & criticism		Psychology, mathematical
Literature	Nuclear science & technology	Psychology, multidisciplinary
Literature, African, Australian, Canadian	Nursing	Psychology, psychoanalysis
Literature, American	Nutrition & dietetics	Psychology, social
Literature, British Isles	Obstetrics & gynecology	Public administration
Literature, German, Dutch, Scandinavian	Oceanography	Public, environmental & occupational health
Literature, romance	Oncology	Radiology, nuclear medicine & medical imaging
Literature, Slavic	Operations research & management science	Rehabilitation
Management	Ophthalmology	Religion
Marine & freshwater biology	Optics	Remote sensing
Materials science, biomaterials	Ornithology	Reproductive biology
Materials science, ceramics	Orthopedics	Respiratory system
Materials science, characterization & testing	Otorhinolaryngology	Rheumatology
Materials science, coatings & films	Paleontology	Robotics
Materials science, composites	Parasitology	Social issues
Materials science, multidisciplinary	Pathology	Social sciences, biomedical
Materials science, paper & wood	Pediatrics	Social sci, interdisciplinary
Materials science, textiles	Peripheral vascular disease	Social sci, mathematical methods
Math & computational biology	Pharmacology & pharmacy	Social work
Mathematics	Philosophy	Sociology
Mathematics, applied	Physics, applied	Soil science
Mathematics, interdisciplinary applications	Physics, atomic, molecular & chemical	Spectroscopy
Mechanics	Physics, condensed matter	Sport sciences
Medical ethics	Physics, fluids & plasmas	Statistics & probability
Medical informatics	Physics, mathematical	Substance abuse
Medical laboratory technology	Physics, multidisciplinary	Surgery
Medicine, general & internal	Physics, nuclear	Telecommunications
Medicine, legal	Physics, particles & fields	Theater
Medicine, research & experimental	Physiology	Thermodynamics
Medieval & renaissance studies	Planning & development	Toxicology
Metallurgy & metallurgical engineering	Plant sciences	Transplantation
Meteorology & atmospheric sci	Poetry	Transportation
Microbiology	Political science	Transportation science & technology
Microscopy	Polymer science	Tropical medicine
Mineralogy	Psychiatry	
Urban studies		
Urology & nephrology		
Veterinary		
Veterinary sciences		
Virology		
Water resources		

Women's studies  
Zoology

#### ESSENTIAL SCIENCE INDICATORS

Agricultural Sciences	Geosciences	Pharmacology
Biology & Biochemistry	Immunology	Physics
Chemistry	Law	Plant & Animal Science
Clinical Medicine	Materials Science	Psychology/Psychiatry
Computer Science	Mathematics	Social Sciences, general
Ecology/Environment	Microbiology	Space Science
Economics & Business	Molecular Biology & Genetics	
Education	Multidisciplinary	
Engineering	Neurosciences & Behaviour	

## ANNEX 2: Biomedically related journal categories

This Annex lists the Web of Science journal categories which capture biomedically related publications.

Allergy	Physiology
Anaesthesiology	Primary Health Care
Anatomy & Morphology	Psychiatry
Andrology	Psychology
Audiology & Speech-Language Pathology	Psychology, Applied
Behavioural Sciences	Psychology, Biological
Cardiac & Cardiovascular Systems	Psychology, Clinical
Cell & Tissue Engineering	Psychology, Developmental
Clinical Neurology	Psychology, Educational
Critical Care Medicine	Psychology, Experimental
Dentistry, Oral Surgery & Medicine	Psychology, Mathematical
Dermatology	Psychology, Psychoanalysis
Emergency Medicine	Psychology, Social
Endocrinology & Metabolism	Public, Environmental & Occupational Health
Ergonomics	Radiology, Nuclear Medicine & Medical Imaging
Gastroenterology & Hepatology	Rehabilitation
Geriatrics & Gerontology	Reproductive Biology
Gerontology	Respiratory System
Haematology	Rheumatology
Health Care Sciences & Services	Substance Abuse
Health Policy & Services	Surgery
Immunology	Transplantation
Infectious Diseases	Tropical Medicine
Integrative & Complementary Medicine	Urology & Nephrology
Medical Ethics	Virology
Medical Informatics	
Medical Laboratory Technology	
Medicine, General & Internal	
Medicine, Legal	
Medicine, Research & Experimental	
Neuroimaging	
Neurosciences	
Nursing	
Nutrition & Dietetics	
Obstetrics & Gynaecology	
Oncology	
Ophthalmology	
Orthopaedics	
Otorhinolaryngology	
Paediatrics	
Pathology	
Peripheral Vascular Disease	
Pharmacology & Pharmacy	

## ANNEX 3: Total number of Web of Science Publications from IMI projects between 2010 and 2021 by country

COUNTRY	NUMBER OF PUBLICATIONS
UK	3,684
Germany	2,756
Netherlands	2,108
USA	2,066
Sweden	1,418
France	1,382
Italy	1,235
Spain	1,061
Switzerland	1,049
Belgium	867
Denmark	633
Canada	568
Austria	508
Finland	402
Australia	313
Peoples R China	286
Greece	249
Norway	225
Ireland	203
Poland	172
Japan	155
Portugal	150
Brazil	137
Israel	120
Singapore	91
Hungary	89
South Africa	80
Estonia	70
Czech Republic	70
Luxembourg	70
India	61

COUNTRY	NUMBER OF PUBLICATIONS
South Korea	52
Saudi Arabia	50
Iceland	48
Turkey	46
Taiwan	46
Lithuania	41
Slovenia	38
New Zealand	38
Cyprus	35
Egypt	34
Croatia	32
Argentina	28
Romania	26
Russia	23
Serbia	21
Thailand	20
Qatar	18
Kenya	17
Iran	16
Chile	16
Latvia	15
Palestine	11
Vietnam	10
Lebanon	10
Mexico	9
Tanzania	9
Bulgaria	9
Colombia	8
Ukraine	8
Malta	7
Uganda	7
Sierra Leone	7
Pakistan	6
Georgia	6

COUNTRY	NUMBER OF PUBLICATIONS
Liechtenstein	6
Tunisia	6
Uruguay	6
Nigeria	6
U Arab Emirates	5
Kuwait	5
Slovakia	5
Peru	5
Jordan	5
DEM REP CONGO	5
Gabon	5
Philippines	5
Iraq	5
Guinea	5
Mozambique	4
Malaysia	4
Mali	4
Gambia	4
Burkina Faso	3
Bangladesh	3
BELARUS	3
Malawi	3
Senegal	3
Oman	2
Nepal	2
Monaco	2
Moldova	2
Macedonia	2
Liberia	2
Guatemala	2
Ghana	2
Sri Lanka	2
Ethiopia	2
Cote Ivoire	2

COUNTRY	NUMBER OF PUBLICATIONS
Bosnia & Herceg	2
Kosovo	1
Kazakhstan	1
Indonesia	1
Benin	1
Armenia	1
Uzbekistan	1
Algeria	1
Ecuador	1
Zambia	1
Cook Islands	1
Cameroon	1
Cambodia	1
Niger	1
Burundi	1
Morocco	1
Mongolia	1
Albania	1
Rwanda	1
Botswana	1
Zimbabwe	1
North Macedonia	1

## ANNEX 4: Total number of Web of Science Publications, papers and open access papers between 2010 and 2021 by Project

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
BTCure	719	672	487	72.5%
EU-AIMS	565	546	453	83.0%
ULTRA-DD	434	425	362	85.2%
EMIF	331	310	261	84.2%
NEWMEDS	220	214	124	57.9%
AIMS-2-TRIALS	210	196	182	92.9%
CANCER-ID	208	180	132	73.3%
INNODIA	203	166	146	88.0%
EUROPAIN	183	181	73	40.3%
ORBITO	171	168	61	36.3%
TRANSLOCATION	164	164	110	67.1%
BigData@Heart	157	135	125	92.6%
STEMBANCC	153	147	120	81.6%
IMIDIA	151	141	121	85.8%
U-BIOPRED	148	93	68	73.1%
RTCure	146	131	112	85.5%
SUMMIT	141	136	106	77.9%
ELF	135	134	108	80.6%
CHEM21	131	128	64	50.0%
PreDiCT-TB	124	118	109	92.4%
SPRINTT	123	116	67	57.8%
MIP-DILI	116	108	70	64.8%
RHAPSODY	115	94	87	92.6%
COMBACTE-NET	109	100	85	85.0%
DIRECT	109	82	71	86.6%
COMBACTE-MAGNET	108	97	82	84.5%
BEAT-DKD	106	98	86	87.8%
Quic-Concept	104	103	88	85.4%
EUbOPEN	102	101	78	77.2%



PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
PROTECT	101	99	46	46.5%
ABIRISK	100	79	51	64.6%
eTOX	97	92	64	69.6%
Pharma-Cog	94	88	39	44.3%
COMPACT	91	91	49	53.8%
None	89	65	55	84.6%
PRISM	87	75	62	82.7%
RADAR-CNS	87	64	55	85.9%
DDMoRe	82	77	54	70.1%
AETIONOMY	74	71	58	81.7%
PRECISESADS	74	54	34	63.0%
Open PHACTS	73	70	63	90.0%
BioVacSafe	73	70	56	80.0%
K4DD	70	68	51	75.0%
Onco Track	69	65	46	70.8%
APPROACH	67	54	39	72.2%
COMBACTE-CARE	66	61	53	86.9%
IMPRIND	66	63	56	88.9%
ZAPI	63	60	57	95.0%
MARCAR	61	60	44	73.3%
DRIVE-AB	60	54	45	83.3%
EPAD	59	55	48	87.3%
LITMUS	59	49	40	81.6%
ENABLE	56	55	47	85.5%
AMYPAD	56	50	47	94.0%
eTRIKS	56	45	43	95.6%
INNODIA HARVEST	53	45	44	97.8%
TransQST	52	47	39	83.0%
iABC	51	33	25	75.8%
Prelect	49	45	38	84.4%
MOBILISE-D	48	43	38	88.4%
RAPP-ID	47	46	33	71.7%
PHAGO	46	45	45	100.0%

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
FLUCOP	45	44	37	84.1%
RESCEU	45	43	43	100.0%
GETREAL	43	37	30	81.1%
iPiE	42	41	29	70.7%
EHDEN	41	31	29	93.5%
PREFER	41	27	27	100.0%
EBiSC	36	33	31	93.9%
EBOVAC1	34	32	32	100.0%
PROACTIVE	34	29	26	89.7%
Hypo-RESOLVE	33	24	21	87.5%
DRAGON	32	28	26	92.9%
ADAPTED	30	28	26	92.9%
eTRANSafe	30	23	21	91.3%
HARMONY	30	17	15	88.2%
ROADMAP	28	22	22	100.0%
ADVANCE	28	27	25	92.6%
IMI-PainCare	27	20	14	70.0%
EbolaMoDRAD	26	25	18	72.0%
3TR	25	23	18	78.3%
EHR4CR	23	20	16	80.0%
SAFE-T	23	21	8	38.1%
TRISTAN	23	23	22	95.7%
DRIVE	21	20	18	90.0%
VAC2VAC	20	20	19	95.0%
IM2PACT	19	19	16	84.2%
EBOVAC2	19	19	19	100.0%
BIOMAP	18	14	11	78.6%
PERISCOPE	18	17	17	100.0%
COMBACTE	17	16	10	62.5%
IDEA-FAST	17	6	5	83.3%
SOPHIA	17	16	14	87.5%
WEB-RADR	17	16	14	87.5%
CARDIATEAM	16	14	14	100.0%

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
EQIPD	15	9	8	88.9%
VSV-EBOPLUS	14	13	11	84.6%
EU-PEARL	14	12	8	66.7%
MOPEAD	13	13	13	100.0%
PD-MitoQUANT	13	13	12	92.3%
RADAR-AD	12	8	8	100.0%
ConcePTION	12	11	9	81.8%
TransBioLine	12	12	11	91.7%
MACUSTAR	12	6	6	100.0%
VSV-EBOVAC	12	11	8	72.7%
ITCC-P4	11	11	9	81.8%
c4c	11	9	6	66.7%
COMBACTE-CDI	11	9	7	77.8%
CARE	11	9	9	100.0%
VALUE-Dx	10	10	10	100.0%
KRONO	10	7	7	100.0%
MAD-CoV 2	9	8	8	100.0%
VITAL	9	9	9	100.0%
ReSOLUTE	9	7	7	100.0%
ERA4TB	8	8	8	100.0%
EBODAC	8	8	8	100.0%
FAIRplus	8	7	7	100.0%
EUPATI	8	7	7	100.0%
NeuroDeRisk	7	6	5	83.3%
EBOVAC3	7	7	7	100.0%
NECESSITY	7	6	5	83.3%
PARADIGM	7	7	7	100.0%
HIPPOCRATES	6	5	5	100.0%
EBiSC2	6	6	6	100.0%
DECISION	6	4	4	100.0%
MELLODDY	6	4	4	100.0%
iCONSENSUS	5	5	4	80.0%
SafeSciMET	5	4	2	50.0%

PROJECT	NUMBER OF PUBLICATIONS	NUMBER OF PAPERS	NUMBER OF OPEN ACCESS PAPERS	% OF OPEN ACCESS PAPERS
Immune-Image	5	5	4	80.0%
imSAVAR	4	3	3	100.0%
ADAPT-SMART	4	4	2	50.0%
DO->IT	4	4	4	100.0%
EBOMAN	4	4	4	100.0%
T2EVOLVE	3	3	2	66.7%
ND4BB	3	3	3	100.0%
ImmUniverse	3	3	3	100.0%
IMMUCAN	3	3	3	100.0%
PEVIA	3	2	2	100.0%
PIONEER	3	2	2	100.0%
Eu2P	3	3	2	66.7%
COVID-RED	3	2	2	100.0%
Trials@Home	3	2	2	100.0%
HARMONY PLUS	2	1	1	100.0%
VHFMoDRAD	2	2	2	100.0%
EMTRAIN	2	1	0	0.0%
OPTIMA	2	2	2	100.0%
Inno4Vac	2	2	2	100.0%
NGN-PET	2	2	1	50.0%
STOPFOP	2	2	2	100.0%
BIGPICTURE	1	1	1	100.0%
Pharmatrain	1	1	1	100.0%
PERSIST-SEQ	1	1	0	0.0%
Impentri	1	1	1	100.0%
UNITE4TB	1	0	0	0.0%
RespiriTB	1	1	1	100.0%
RespiriNTM	1	1	1	100.0%
COMBINE	1	1	1	100.0%
EBOVAC	1	1	1	100.0%
FILODIAG	1	0	0	0.0%
Screen4Care	1	1	1	100.0%

## ANNEX 5: Collaboration index for all IMI supported research projects

This Annex provides the calculation of the collaboration indicators for all IMI supported research projects with at least one paper. Collaboration index only calculated for projects with a Stability score and at least 20 papers.

PROJECT	CROSS-SECTOR SCORE	INTERNATIONAL SCORE	STABILITY SCORE	COLLABORATION INDEX	TOTAL PAPERS	CITATION IMPACT (FIELD-NORMALISED)
BTCure	0.65	0.52	0.84	2.00	672	1.81
EU-AIMS	0.73	0.65	0.82	2.19	546	2.06
ULTRA-DD	0.63	0.65	0.77	2.06	425	1.91
EMIF	0.82	0.67	0.84	2.34	310	2.49
NEWMEDS	0.62	0.58	0.83	2.03	214	2.06
AIMS-2-TRIALS	0.71	0.66	0.69	2.07	196	2.67
EUROPAIN	0.55	0.38	0.85	1.78	181	2.55
CANCER-ID	0.75	0.43	0.75	1.93	180	3.27
ORBITO	0.64	0.48	0.76	1.88	168	1.70
INNODIA	0.81	0.65	0.87	2.32	166	1.62
TRANSLOCATION	0.37	0.49	0.81	1.66	164	1.35
STEMBANCC	0.50	0.47	0.82	1.79	147	1.94
IMIDIA	0.53	0.50	0.84	1.87	141	1.65
SUMMIT	0.75	0.64	0.83	2.22	136	1.42
BigData@Heart	0.90	0.69	0.69	2.29	135	2.05
ELF	0.34	0.51	0.77	1.62	134	1.11
RTCure	0.81	0.47	0.70	1.98	131	2.86
CHEM21	0.23	0.29	0.80	1.33	128	1.70
PreDiCT-TB	0.57	0.50	0.81	1.88	118	1.16
SPRINTT	0.72	0.53	0.80	2.05	116	1.83
MIP-DILI	0.67	0.45	0.82	1.94	108	1.77
Quic-Concept	0.72	0.57	0.81	2.09	103	4.76
EUbOPEN	0.58	0.55	0.00	1.13	101	1.85
COMBACTE-NET	0.79	0.55	0.89	2.23	100	1.17
PROTECT	0.97	0.63	0.86	2.46	99	1.03
BEAT-DKD	0.71	0.69	0.70	2.11	98	2.17
COMBACTE-MAGNET	0.70	0.62	0.86	2.18	97	1.28
RHAPSODY	0.60	0.66	0.81	2.07	94	1.97
U-BIOPRED	0.83	0.72	0.86	2.41	93	2.49
eTOX	0.30	0.36	0.86	1.53	92	1.76

PROJECT	CROSS-SECTOR SCORE	INTERNATIONAL SCORE	STABILITY SCORE	COLLABORATION INDEX	TOTAL PAPERS	CITATION IMPACT (FIELD-NORMALISED)
COMPACT	0.24	0.46	0.75	1.46	91	1.90
Pharma-Cog	0.85	0.75	0.85	2.45	88	1.13
DIRECT	0.76	0.73	0.84	2.33	82	3.93
ABIRISK	0.76	0.50	0.84	2.10	79	1.25
DDMoRe	0.65	0.56	0.81	2.02	77	1.21
PRISM	0.80	0.69	0.75	2.25	75	2.70
AETIONOMY	0.65	0.46	0.80	1.91	71	1.77
Open PHACTS	0.60	0.55	0.76	1.92	70	3.57
BioVacSafe	0.46	0.49	0.81	1.76	70	1.20
K4DD	0.54	0.51	0.80	1.86	68	1.53
None	0.75	0.70	0.49	1.94	65	3.50
Onco Track	0.63	0.43	0.83	1.90	65	2.19
RADAR-CNS	0.63	0.70	0.82	2.15	64	1.94
IMPRiND	0.67	0.60	0.75	2.01	63	4.96
COMBACTE-CARE	0.92	0.66	0.80	2.37	61	1.59
ZAPI	0.67	0.63	0.77	2.07	60	4.34
MARCAR	0.42	0.42	0.84	1.68	60	1.04
ENABLE	0.58	0.47	0.83	1.88	55	1.45
EPAD	0.76	0.67	0.81	2.25	55	1.35
DRIVE-AB	0.74	0.64	0.75	2.13	54	1.30
PRECISESADS	0.80	0.77	0.75	2.32	54	1.38
APPROACH	0.81	0.81	0.81	2.43	54	2.07
AMYPAD	0.92	0.78	0.77	2.47	50	1.96
LITMUS	0.84	0.67	0.75	2.25	49	3.85
TransQST	0.60	0.68	0.76	2.03	47	3.09
RAPP-ID	0.33	0.43	0.85	1.61	46	0.86
Predict	0.69	0.63	0.79	2.11	45	2.67
INNODIA HARVEST	0.78	0.66	0.00	1.43	45	1.37
eTRIKS	0.82	0.88	0.75	2.46	45	2.04
PHAGO	0.69	0.60	0.72	2.01	45	4.02
FLUCOP	0.91	0.47	0.73	2.11	44	1.58
MOBILISE-D	0.77	0.58	0.00	1.34	43	1.57
RESCEU	0.84	0.70	0.69	2.22	43	1.91
iPiE	0.51	0.24	0.76	1.51	41	1.13
GETREAL	0.84	0.76	0.78	2.37	37	1.65

PROJECT	CROSS-SECTOR SCORE	INTERNATIONAL SCORE	STABILITY SCORE	COLLABORATION INDEX	TOTAL PAPERS	CITATION IMPACT (FIELD-NORMALISED)
iABC	0.85	0.65	0.82	2.32	33	1.19
EBiSC	0.70	0.62	0.77	2.09	33	5.61
EBOVAC1	0.72	0.67	0.82	2.21	32	2.07
EHDEN	0.81	0.84	0.63	2.27	31	2.19
PROACTIVE	1.00	0.81	0.85	2.66	29	2.22
DRAGON	0.89	0.75	#N/A	#N/A	28	3.78
ADAPTED	0.93	0.62	#N/A	#N/A	28	2.89
PREFER	0.93	0.87	#N/A	#N/A	27	1.29
ADVANCE	0.89	0.86	#N/A	#N/A	27	1.36
EbolaMoDRAD	0.68	0.52	#N/A	#N/A	25	1.34
Hypo-RESOLVE	0.58	0.79	#N/A	#N/A	24	0.71
3TR	0.91	0.48	#N/A	#N/A	23	2.60
eTRANSafe	0.48	0.50	#N/A	#N/A	23	3.15
TRISTAN	0.83	0.50	#N/A	#N/A	23	1.22
ROADMAP	0.91	0.75	#N/A	#N/A	22	0.91
SAFE-T	0.95	0.54	#N/A	#N/A	21	1.73
VAC2VAC	0.70	0.56	#N/A	#N/A	20	0.60
DRIVE	0.85	0.31	#N/A	#N/A	20	0.79
IMI-PainCare	0.75	0.55	#N/A	#N/A	20	1.22
EHR4CR	0.85	0.60	#N/A	#N/A	20	1.11
EBOVAC2	0.53	0.51	#N/A	#N/A	19	2.45
IM2PACT	0.58	0.43	#N/A	#N/A	19	1.62
PERISCOPE	0.35	0.38	#N/A	#N/A	17	1.57
HARMONY	0.82	0.47	#N/A	#N/A	17	0.70
COMBACTE	0.50	0.11	#N/A	#N/A	16	3.01
WEB-RADR	0.75	0.75	#N/A	#N/A	16	1.40
SOPHIA	0.69	0.55	#N/A	#N/A	16	3.45
BIOMAP	0.86	0.73	#N/A	#N/A	14	4.64
CARDIATEAM	1.00	0.95	#N/A	#N/A	14	3.03
PD-MitoQUANT	0.69	0.46	#N/A	#N/A	13	2.08
MOPEAD	1.00	0.81	#N/A	#N/A	13	1.73
VSV-EBOPLUS	0.69	0.77	#N/A	#N/A	13	1.02
EU-PEARL	0.83	0.73	#N/A	#N/A	12	2.42
TransBioLine	0.92	0.40	#N/A	#N/A	12	1.09
ConcePTION	1.00	0.91	#N/A	#N/A	11	0.79
ITCC-P4	1.00	0.75	#N/A	#N/A	11	2.48

PROJECT	CROSS-SECTOR SCORE	INTERNATIONAL SCORE	STABILITY SCORE	COLLABORATION INDEX	TOTAL PAPERS	CITATION IMPACT (FIELD-NORMALISED)
VSV-EBOVAC	0.55	0.64	#N/A	#N/A	11	0.94
VALUE-Dx	0.70	0.82	#N/A	#N/A	10	2.75
c4c	1.00	0.92	#N/A	#N/A	9	0.69
COMBACTE-CDI	1.00	0.94	#N/A	#N/A	9	1.38
CARE	0.56	0.64	#N/A	#N/A	9	9.09
VITAL	0.56	0.53	#N/A	#N/A	9	0.36
EQIPD	0.56	0.78	#N/A	#N/A	9	2.38
RADAR-AD	0.88	0.50	#N/A	#N/A	8	1.51
MAD-CoV 2	0.88	0.88	#N/A	#N/A	8	2.99
ERA4TB	0.75	0.66	#N/A	#N/A	8	1.11
EBODAC	0.88	0.81	#N/A	#N/A	8	2.38
EUPATI	1.00	0.96	#N/A	#N/A	7	0.75
FAIRplus	0.14	0.25	#N/A	#N/A	7	1.45
PARADIGM	0.86	0.82	#N/A	#N/A	7	1.61
EBOVAC3	0.57	0.89	#N/A	#N/A	7	0.60
ReSOLUTE	0.57	0.36	#N/A	#N/A	7	1.00
KRONO	0.57	0.14	#N/A	#N/A	7	2.52
IDEA-FAST	0.33	0.79	#N/A	#N/A	6	1.10
NeuroDeRisk	0.33	0.17	#N/A	#N/A	6	0.12
EBiSC2	1.00	0.92	#N/A	#N/A	6	0.52
NECESSITY	1.00	0.83	#N/A	#N/A	6	2.53
MACUSTAR	0.83	0.67	#N/A	#N/A	6	1.98
HIPPOCRATES	0.80	0.15	#N/A	#N/A	5	4.32
Immune-Image	1.00	0.90	#N/A	#N/A	5	2.67
iCONSENSUS	0.60	0.35	#N/A	#N/A	5	1.26
MELLODDY	0.75	0.75	#N/A	#N/A	4	0.62
EBOMAN	1.00	0.94	#N/A	#N/A	4	3.88
DO->IT	0.75	0.81	#N/A	#N/A	4	1.19
DECISION	0.75	0.00	#N/A	#N/A	4	2.60
SafeSciMET	1.00	1.00	#N/A	#N/A	4	0.89
ADAPT-SMART	0.75	0.50	#N/A	#N/A	4	0.58
Eu2P	0.33	0.67	#N/A	#N/A	3	2.01
ND4BB	0.67	0.58	#N/A	#N/A	3	1.36
imSAVAR	1.00	0.83	#N/A	#N/A	3	5.01
T2EVOLVE	0.67	0.25	#N/A	#N/A	3	3.66
ImmUniverse	1.00	0.58	#N/A	#N/A	3	0.00



PROJECT	CROSS-SECTOR SCORE	INTERNATIONAL SCORE	STABILITY SCORE	COLLABORATION INDEX	TOTAL PAPERS	CITATION IMPACT (FIELD-NORMALISED)
IMMUCAN	0.67	0.58	#N/A	#N/A	3	0.87
COVID-RED	1.00	1.00	#N/A	#N/A	2	0.44
PIONEER	1.00	1.00	#N/A	#N/A	2	1.26
Inno4Vac	1.00	0.38	#N/A	#N/A	2	0.00
STOPFOP	0.50	0.88	#N/A	#N/A	2	2.17
PEVIA	1.00	0.88	#N/A	#N/A	2	0.80
Trials@Home	0.50	0.38	#N/A	#N/A	2	1.20
VHFMoDRAD	1.00	0.38	#N/A	#N/A	2	0.38
OPTIMA	1.00	0.50	#N/A	#N/A	2	0.75
NGN-PET	0.50	0.50	#N/A	#N/A	2	1.38
HARMONY PLUS	1.00	1.00	#N/A	#N/A	1	11.67
Impentri	1.00	1.00	#N/A	#N/A	1	0.00
BIGPICTURE	1.00	1.00	#N/A	#N/A	1	4.88
EBOVAC	1.00	1.00	#N/A	#N/A	1	3.27
PERSIST-SEQ	0.00	0.00	#N/A	#N/A	1	0.00
RespiriNTM	0.00	0.00	#N/A	#N/A	1	4.39
Pharmatrain	1.00	1.00	#N/A	#N/A	1	0.10
EMTRAIN	1.00	1.00	#N/A	#N/A	1	0.10
Screen4Care	0.00	0.00	#N/A	#N/A	1	0.00
RespiriTB	0.00	0.00	#N/A	#N/A	1	4.39
COMBINE	1.00	0.00	#N/A	#N/A	1	0.07

## ANNEX 6: Bibliography of hot papers and highly cited papers

This Annex provides bibliographic data for hot and highly cited papers. Hot papers are papers that receive citations soon after publication, relative to other papers of the same field and age. For the purpose of this report, highly cited papers have been defined as those articles and reviews which belong to the world's top decile of papers in that journal category and year of publication, when ranked by number of citations received. A percentage that is above 10 indicates above-average performance.

Papers are listed in ascending alphabetical order (project, first author) and unassigned papers, are listed at the end of each section.

This section lists papers that have been identified as current hot papers or that have been identified as highly cited in the IMI project publication dataset.

### Hot papers associated with IMI projects

AIMS-2-TRIALS: Moreno, Carmen et al. How mental health care should change as a consequence of the COVID-19 pandemic, *LANCET PSYCHIAT* 7: 813-824

DIRECT: Aguet, Francois et al. The GTEx Consortium atlas of genetic regulatory effects across human tissues, *SCIENCE* 369: 1318-1330

EuOPEN: Attwood, Misty M. et al. Trends in kinase drug discovery: targets, indications and inhibitor design, *NAT REV DRUG DISCOV* 20: 839-861

EUROPAIN: Kosek, Eva et al. Chronic nociplastic pain affecting the musculoskeletal system: clinical criteria and grading system, *PAIN* 162: 2629-2634

IMI-PainCare: Kosek, Eva et al. Chronic nociplastic pain affecting the musculoskeletal system: clinical criteria and grading system, *PAIN* 162: 2629-2634

IMPRiND: Shi, Yang et al. Structure-based classification of tauopathies, *NATURE* 598: 359-+

PHAGO: Meinhardt, Jenny et al. Olfactory transmucosal SARS-CoV-2 invasion as a port of central nervous system entry in individuals with COVID-19, *NAT NEUROSCI* 24: 168-175

PRISM: Moreno, Carmen et al. How mental health care should change as a consequence of the COVID-19 pandemic, *LANCET PSYCHIAT* 7: 813-824

Quic-Concept: Zwanenburg, Alex et al. The Image Biomarker Standardization Initiative: Standardized Quantitative Radiomics for High-Throughput Image-based Phenotyping, *RADIOLOGY* 295: 328-338

RTCure: Simon, David et al. SARS-CoV-2 vaccination responses in untreated, conventionally treated and anticytokine-treated patients with immune-mediated inflammatory diseases, *ANN RHEUM DIS* 80: 1312-1316

RTCure: Haberman, Rebecca H. et al. Methotrexate hampers immunogenicity to BNT162b2 mRNA COVID-19 vaccine in immune-mediated inflammatory disease, *ANN RHEUM DIS* 80: 1339-1344

SOPHIA: Stefan, Norbert et al. Global pandemics interconnected - obesity, impaired metabolic health and COVID-19, *NAT REV ENDOCRINOL* 17: 135-149

## Highly cited papers associated with IMI projects

This section lists papers that perform above average as defined by citation counts in the 10<sup>th</sup> percentile.

3TR: Stengel, Stephanie T. et al. Activating Transcription Factor 6 Mediates Inflammatory Signals in Intestinal Epithelial Cells Upon Endoplasmic Reticulum Stress, *GASTROENTEROLOGY* 159: 1357-+

3TR: Bernardes, Joana P. et al. Longitudinal Multi-omics Analyses Identify Responses of Megakaryocytes, Erythroid Cells, and Plasmablasts as Hallmarks of Severe COVID-19, *IMMUNITY* 53: 1296-+

3TR: Kolmert, Johan et al. Urinary Leukotriene E-4 and Prostaglandin D-2 Metabolites Increase in Adult and Childhood Severe Asthma Characterized by Type 2 Inflammation A Clinical Observational Study, *AM J RESP CRIT CARE* 203: 37-53

3TR: Yordanova, Iveta A. et al. The Worm-Specific Immune Response in Multiple Sclerosis Patients Receiving Controlled *Trichuris suis* Ova Immunotherapy, *LIFE-BASEL* 11:

3TR: Schreiber, Stefan et al. Therapeutic Interleukin-6 Trans-signaling Inhibition by Olamkicept (sgp130Fc) in Patients With Active Inflammatory Bowel Disease, *GASTROENTEROLOGY* 160: 2354-+

3TR: Hoepel, Willianne et al. High titers and low fucosylation of early human anti-SARS-CoV-2 IgG promote inflammation by alveolar macrophages, *SCI TRANSL MED* 13:

ABIRISK: Kieseier, Bernd C. et al. Disease Amelioration With Tocilizumab in a Treatment-Resistant Patient With Neuromyelitis Optica Implication for Cellular Immune Responses, *JAMA NEUROL* 70: 390-393

ABIRISK: Wenniger, Lucas J. Maillette de Buy et al. Immunoglobulin G4+clones identified by next-generation sequencing dominate the B cell receptor repertoire in immunoglobulin G4 associated cholangitis, *HEPATOLOGY* 57: 2390-2398

ABIRISK: Warnke, Clemens et al. Changes to anti-JCV antibody levels in a Swedish national MS cohort, *J NEUROL NEUROSUR PS* 84: 1199-1205

ABIRISK: Shankar, G. et al. Assessment and Reporting of the Clinical Immunogenicity of Therapeutic Proteins and Peptides-Harmonized Terminology and Tactical Recommendations, *AAPS J* 16: 658-673

ABIRISK: Ungar, Bella et al. The temporal evolution of antidrug antibodies in patients with inflammatory bowel disease treated with infliximab, *GUT* 63: 1258-1264

ABIRISK: Warnke, Clemens et al. Cerebrospinal Fluid JC Virus Antibody Index for Diagnosis of Natalizumab-Associated Progressive Multifocal Leukoencephalopathy, *ANN NEUROL* 76: 792-801

ABIRISK: Hemmer, Bernhard et al. Role of the innate and adaptive immune responses in the course of multiple sclerosis, *LANCET NEUROL* 14: 406-419

ABIRISK: Warnke, Clemens et al. Natalizumab exerts a suppressive effect on surrogates of B cell function in blood and CSF, *MULT SCLER J* 21: 1036-1044

ABIRISK: Ringelstein, Marius et al. Long-term Therapy With Interleukin 6 Receptor Blockade in Highly Active Neuromyelitis Optica Spectrum Disorder, *JAMA NEUROL* 72: 756-763

ABIRISK: Diebold, Martin et al. Dimethyl fumarate influences innate and adaptive immunity in multiple sclerosis, *J AUTOIMMUN* 86: 39-50

ABIRISK: Quistebert, Jocelyn et al. Incidence and risk factors for adalimumab and infliximab anti-drug antibodies in rheumatoid arthritis: A European retrospective multicohort analysis, *SEMIN ARTHRITIS RHEU* 48: 967-975

ABIRISK: Cassotta, Antonino et al. A single T cell epitope drives the neutralizing anti-drug antibody response to natalizumab in multiple sclerosis patients, *NAT MED* 25: 1402-+

ADAPTED: van der Lee, Sven J. et al. The effect of *ITAPOE* and other common genetic variants on the onset of Alzheimers disease and dementia: a community-based cohort study, *LANCET NEUROL* 17: 434-444

ADAPTED: van der Lee, Sven J. et al. Circulating metabolites and general cognitive ability and dementia: Evidence from 11 cohort studies, *ALZHEIMERS DEMENT* 14: 707-722

ADAPTED: Tynkkynen, Juho et al. Association of branched-chain amino acids and other circulating metabolites with risk of incident dementia and Alzheimers disease: A prospective study in eight cohorts, *ALZHEIMERS DEMENT* 14: 723-733

ADAPTED: Wevers, Nienke R. et al. A perfused human blood-brain barrier on-a-chip for high-throughput assessment of barrier function and antibody transport, *FLUIDS BARRIERS CNS* 15:

ADAPTED: van der Lee, Sven J. et al. A nonsynonymous mutation in *PLCG2* reduces the risk of Alzheimers disease, dementia with Lewy bodies and frontotemporal dementia, and increases the likelihood of longevity, *ACTA NEUROPATHOL* 138: 237-250

ADAPTED: Moreno-Grau, Sonia et al. Genome-wide association analysis of dementia and its clinical endophenotypes reveal novel loci associated with Alzheimers disease and three causality networks: The GR@ACE project, *ALZHEIMERS DEMENT* 15: 1333-1347

ADAPTED: Cenini, Giovanna et al. Dissecting Alzheimers disease pathogenesis in human 2D and 3D models, *MOL CELL NEUROSCI* 110:

ADAPTED: Roberto, Natalia et al. Neuropsychiatric profiles and conversion to dementia in mild cognitive impairment, a latent class analysis, *SCI REP-UK* 11:

ADAPTED: de Rojas, Itziar et al. Common variants in Alzheimers disease and risk stratification by polygenic risk scores, *NAT COMMUN* 12:

ADVANCE: Pebody, R. et al. Effectiveness of seasonal influenza vaccine for adults and children in preventing laboratory-confirmed influenza in primary care in the United Kingdom: 2015/16 end-of-season results, *EUROSURVEILLANCE* 21: 41-51

ADVANCE: Karafillakis, Emilie et al. The benefit of the doubt or doubts over benefits? A systematic literature review of perceived risks of vaccines in European populations, *VACCINE* 35: 4840-4850

ADVANCE: Willame, Corinne et al. Incidence Rates of Autoimmune Diseases in European Healthcare Databases: A Contribution of the ADVANCE Project, DRUG SAFETY 44: 383-395

AETIONOMY: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

AETIONOMY: Domingo Gispert, Juan et al. Cerebrospinal fluid sTREM2 levels are associated with gray matter volume increases and reduced diffusivity in early Alzheimers disease, ALZHEIMERS DEMENT 12: 1259-1272

AETIONOMY: Gautier, Clement A. et al. The endoplasmic reticulum-mitochondria interface is perturbed in PARK2 knockout mice and patients with PARK2 mutations, HUM MOL GENET 25: 2972-2984

AETIONOMY: Bedarf, J. R. et al. Functional implications of microbial and viral gut metagenome changes in early stage L-DOPA-naive Parkinsons disease patients, GENOME MED 9:

AETIONOMY: Mouton-Ligeri, Francois et al. PINK1/Parkin-Dependent Mitochondrial Surveillance: From Pleiotropy to Parkinsons Disease, FRONT MOL NEUROSCI 10:

AETIONOMY: Crous-Bou, Marta et al. Alzheimers disease prevention: from risk factors to early intervention, ALZHEIMERS RES THER 9:

AETIONOMY: Hoyt, Charles Tapley et al. PyBEL: a computational framework for Biological Expression Language, BIOINFORMATICS 34: 703-704

AETIONOMY: Brosseron, Frederic et al. Characterization and clinical use of inflammatory cerebrospinal fluid protein markers in Alzheimers disease, ALZHEIMERS RES THER 10:

AETIONOMY: Mouton-Liger, Francois et al. Parkin deficiency modulates NLRP3 inflammasome activation by attenuating an A20-dependent negative feedback loop, GLIA 66: 1736-1751

AETIONOMY: Froehlich, Holger et al. From hype to reality: data science enabling personalized medicine, BMC MED 16:

AETIONOMY: McWilliams, Thomas G. et al. Phosphorylation of Parkin at serine 65 is essential for its activation in vivo, OPEN BIOL 8:

AETIONOMY: Bonello, Fiona et al. LRRK2 impairs PINK1/Parkin-dependent mitophagy via its kinase activity: pathologic insights into Parkinsons disease, HUM MOL GENET 28: 1645-1660

AETIONOMY: Mubeen, Sarah et al. The Impact of Pathway Database Choice on Statistical Enrichment Analysis and Predictive Modeling, FRONT GENET 10:

AETIONOMY: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), GENOME MED 12:

AETIONOMY: Corti, Olga et al. Autophagy in neurodegeneration: New insights underpinning therapy for neurological diseases, J NEUROCHEM 154: 354-371

AETIONOMY: Antonell, Anna et al. Synaptic, axonal damage and inflammatory cerebrospinal fluid biomarkers in neurodegenerative dementias, ALZHEIMERS DEMENT 16: 262-272

AETIONOMY: Brosseron, Frederic et al. Multicenter Alzheimers and Parkinsons disease immune biomarker verification study, ALZHEIMERS DEMENT 16: 292-304

AIMS-2-TRIALS: Greenberg, David M. et al. Testing the Empathizing-Systemizing theory of sex differences and the Extreme Male Brain theory of autism in half a million people, P NATL ACAD SCI USA 115: 12152-12157

AIMS-2-TRIALS: Holiga, Stefan et al. Patients with autism spectrum disorders display reproducible functional connectivity alterations, SCI TRANSL MED 11:

AIMS-2-TRIALS: Bolte, Sven et al. The contribution of environmental exposure to the etiology of autism spectrum disorder, CELL MOL LIFE SCI 76: 1275-1297

AIMS-2-TRIALS: Greven, Corina U. et al. Sensory Processing Sensitivity in the context of Environmental Sensitivity: A critical review and development of research agenda, NEUROSCI BIOBEHAV R 98: 287-305

AIMS-2-TRIALS: Pretzsch, Charlotte Marie et al. Effects of cannabidiol on brain excitation and inhibition systems, a randomised placebo-controlled single dose trial during magnetic resonance spectroscopy in adults with and without autism spectrum disorder, NEUROPSYCHOPHARMACOL 44: 1398-1405

AIMS-2-TRIALS: Warriar, Varun et al. Social and non-social autism symptoms and trait domains are genetically dissociable, COMMUN BIOL 2:

AIMS-2-TRIALS: Wolfers, Thomas et al. From pattern classification to stratification: towards conceptualizing the heterogeneity of Autism Spectrum Disorder, NEUROSCI BIOBEHAV R 104: 240-254

AIMS-2-TRIALS: Lombardo, Michael, V et al. Big data approaches to decomposing heterogeneity across the autism spectrum, MOL PSYCHIATR 24: 1435-1450

AIMS-2-TRIALS: Nystrom, Par et al. Joint Attention in Infancy and the Emergence of Autism, BIOL PSYCHIAT 86: 631-638

AIMS-2-TRIALS: Postema, Merel C. et al. Altered structural brain asymmetry in autism spectrum disorder in a study of 54 datasets, NAT COMMUN 10:

AIMS-2-TRIALS: Oldehinkel, Marianne et al. Altered Connectivity Between Cerebellum, Visual, and Sensory-Motor Networks in Autism Spectrum Disorder: Results from the EU-AIMS Longitudinal European Autism Project, BIOL PSYCHIAT-COGN N 4: 260-270

AIMS-2-TRIALS: Fraguas, David et al. Dietary Interventions for Autism Spectrum Disorder: A Meta-analysis, PEDIATRICS 144:

AIMS-2-TRIALS: Lord, Catherine et al. Autism spectrum disorder, NAT REV DIS PRIMERS 6:

AIMS-2-TRIALS: Pohl, A. L. et al. A comparative study of autistic and non-autistic womens experience of motherhood, MOL AUTISM 11:

AIMS-2-TRIALS: Moessnang, Carolin et al. Social brain activation during mentalizing in a large autism cohort: the Longitudinal European Autism Project, MOL AUTISM 11:

AIMS-2-TRIALS: Pelton, Mirabel K. et al. Understanding Suicide Risk in Autistic Adults: Comparing the Interpersonal Theory of Suicide in Autistic and Non-autistic Samples, *J AUTISM DEV DISORD* 50: 3620-3637

AIMS-2-TRIALS: Martinez, Kenia et al. Sensory-to-Cognitive Systems Integration Is Associated With Clinical Severity in Autism Spectrum Disorder, *J AM ACAD CHILD PSY* 59: 422-433

AIMS-2-TRIALS: Begum Ali, Jannath et al. Early Motor Differences in Infants at Elevated Likelihood of Autism Spectrum Disorder and/or Attention Deficit Hyperactivity Disorder, *J AUTISM DEV DISORD* 50: 4367-4384

AIMS-2-TRIALS: Hoogman, Martine et al. Consortium neuroscience of attention deficit/hyperactivity disorder and autism spectrum disorder: The ENIGMA adventure, *HUM BRAIN MAPP* 43: 37-55

AIMS-2-TRIALS: Dohmatob, Elvis et al. Dark control: The default mode network as a reinforcement learning agent, *HUM BRAIN MAPP* 41: 3318-3341

AIMS-2-TRIALS: Ching, Christopher R. K. et al. Mapping Subcortical Brain Alterations in 22q11.2 Deletion Syndrome: Effects of Deletion Size and Convergence With Idiopathic Neuropsychiatric Illness, *AM J PSYCHIAT* 177: 589-600

AIMS-2-TRIALS: de Pablo, Gonzalo Salazar et al. Impact of coronavirus syndromes on physical and mental health of health care workers: Systematic review and meta-analysis, *J AFFECT DISORDERS* 275: 48-57

AIMS-2-TRIALS: Hornberg, Hanna et al. Rescue of oxytocin response and social behaviour in a mouse model of autism, *NATURE* 584: 252-+

AIMS-2-TRIALS: Warriar, Varun et al. Elevated rates of autism, other neurodevelopmental and psychiatric diagnoses, and autistic traits in transgender and gender-diverse individuals, *NAT COMMUN* 11:

AIMS-2-TRIALS: Moreno, Carmen et al. How mental health care should change as a consequence of the COVID-19 pandemic, *LANCET PSYCHIAT* 7: 813-824

AIMS-2-TRIALS: van Kessel, Robin et al. Autism and education-Teacher policy in Europe: Policy mapping of Austria, Hungary, Slovakia and Czech Republic, *RES DEV DISABIL* 105:

AIMS-2-TRIALS: Oakley, Bethany F. M. et al. How do core autism traits and associated symptoms relate to quality of life? Findings from the Longitudinal European Autism Project, *AUTISM* 25: 389-404

AIMS-2-TRIALS: Davies, Robert W. et al. Using common genetic variation to examine phenotypic expression and risk prediction in 22q11.2 deletion syndrome, *NAT MED* 26:

AIMS-2-TRIALS: Fraguas, David et al. Assessment of School Anti-Bullying Interventions A Meta-analysis of Randomized Clinical Trials, *JAMA PEDIATR* 175: 44-55

AIMS-2-TRIALS: Piccardi, Elena Serena et al. Behavioural and neural markers of tactile sensory processing in infants at elevated likelihood of autism spectrum disorder and/or attention deficit hyperactivity disorder, *J NEURODEV DISORD* 13:

AIMS-2-TRIALS: Mossa, Adele et al. Developmental impaired Akt signaling in the Shank1 and Shank3 double knock-out mice, *MOL PSYCHIATR* 26: 1928-1944

AIMS-2-TRIALS: Gomez, Andrea M. et al. Neurexins: molecular codes for shaping neuronal synapses, *NAT REV NEUROSCI* 22: 137-151

AIMS-2-TRIALS: Grabrucker, Stefanie et al. Activation of the medial preoptic area (MPOA) ameliorates loss of maternal behavior in a Shank2 mouse model for autism, *EMBO J* 40:

AIMS-2-TRIALS: Peng, Han et al. Accurate brain age prediction with lightweight deep neural networks, *MED IMAGE ANAL* 68:

AIMS-2-TRIALS: van Kessel, Robin et al. Inclusive education in the European Union: A fuzzy-set qualitative comparative analysis of education policy for autism, *SOC WORK PUBLIC HLTH* 36: 286-299

AIMS-2-TRIALS: Adhya, Dwaipayan et al. Atypical Neurogenesis in Induced Pluripotent Stem Cells From Autistic Individuals, *BIOL PSYCHIAT* 89: 486-496

AIMS-2-TRIALS: Hull, Laura et al. Is social camouflaging associated with anxiety and depression in autistic adults?, *MOL AUTISM* 12:

AIMS-2-TRIALS: Floris, Dorothea L. et al. Towards robust and replicable sex differences in the intrinsic brain function of autism, *MOL AUTISM* 12:

AIMS-2-TRIALS: Dumas, Guillaume et al. Systematic detection of brain protein-coding genes under positive selection during primate evolution and their roles in cognition, *GENOME RES* 31: 484-496

AIMS-2-TRIALS: Gui, Anna et al. Attentive brain states in infants with and without later autism, *TRANSL PSYCHIAT* 11:

AIMS-2-TRIALS: Roman-Urrestarazu, Andres et al. Association of Race/Ethnicity and Social Disadvantage With Autism Prevalence in 7 Million School Children in England, *JAMA PEDIATR* 175:

AIMS-2-TRIALS: Maria, Yanez Lopez et al. Simultaneous quantification of GABA, Glx and GSH in the neonatal human brain using magnetic resonance spectroscopy, *NEUROIMAGE* 233:

AIMS-2-TRIALS: Fusar-Poli, Paolo et al. Preventive psychiatry: a blueprint for improving the mental health of young people, *WORLD PSYCHIATRY* 20: 200-221

AIMS-2-TRIALS: Oakley, Bethany et al. COVID-19 health and social care access for autistic people: European policy review, *BMJ OPEN* 11:

AIMS-2-TRIALS: Constantino, John N. et al. Clinical and Translational Implications of an Emerging Developmental Substructure for Autism, *ANNU REV CLIN PSYCHO* 17: 365-389

AIMS-2-TRIALS: Persico, Antonio M. et al. The pediatric psychopharmacology of autism spectrum disorder: A systematic review - Part I: The past and the present, *PROG NEURO-PSYCHOPH* 110:

AIMS-2-TRIALS: Floris, Dorothea L. et al. Atypical Brain Asymmetry in Autism-A Candidate for Clinically Meaningful Stratification, *BIOL PSYCHIAT-COGN* 6: 802-812



AIMS-2-TRIALS: Del Bianco, Teresa et al. Temporal Profiles of Social Attention Are Different Across Development in Autistic and Neurotypical People, *BIOL PSYCHIAT-COGN N* 6: 813-824

AIMS-2-TRIALS: Eyre, Michael et al. The Developing Human Connectome Project: typical and disrupted perinatal functional connectivity, *BRAIN* 144: 2199-2213

AMYPAD: Crous-Bou, Marta et al. Alzheimers disease prevention: from risk factors to early intervention, *ALZHEIMERS RES THER* 9:

AMYPAD: Tur, Carmen et al. Assessing treatment outcomes in multiple sclerosis trials and in the clinical setting, *NAT REV NEUROL* 14: 75-93

AMYPAD: Collij, Lyduine E. et al. Assessing Amyloid Pathology in Cognitively Normal Subjects Using F-18-Flutemetamol PET: Comparing Visual Reads and Quantitative Methods, *J NUCL MED* 60: 541-547

AMYPAD: Fantoni, Enrico et al. The Spatial-Temporal Ordering of Amyloid Pathology and Opportunities for PET Imaging, *J NUCL MED* 61: 166-171

AMYPAD: Arabi, Hossein et al. Deep learning-guided joint attenuation and scatter correction in multitracer neuroimaging studies, *HUM BRAIN MAPP* 41: 3667-3679

AMYPAD: Mutsaerts, Henk J. M. M. et al. EXploreASL: An image processing pipeline for multi-center ASL perfusion MRI studies, *NEUROIMAGE* 219:

AMYPAD: Collij, Lyduine E. et al. Multitracer model for staging cortical amyloid deposition using PET imaging, *NEUROLOGY* 95: E1538-E1553

AMYPAD: Chetelat, Gael et al. Amyloid-PET and F-18-FDG-PET in the diagnostic investigation of Alzheimers disease and other dementias, *LANCET NEUROL* 19: 951-962

AMYPAD: Tsvetanov, Kamen A. et al. The effects of age on resting-state BOLD signal variability is explained by cardiovascular and cerebrovascular factors, *PSYCHOPHYSIOLOGY* 58:

AMYPAD: Villemagne, Victor L. et al. Molecular Imaging Approaches in Dementia, *RADIOLOGY* 298: 517-530

AMYPAD: Boccardi, Marina et al. The strategic biomarker roadmap for the validation of Alzheimers diagnostic biomarkers: methodological update, *EUR J NUCL MED MOL I* 48: 2070-2085

AMYPAD: Solomon, Alina et al. Multidomain interventions: state-of-the-art and future directions for protocols to implement precision dementia risk reduction. A user manual for Brain Health Services-part 4 of 6, *ALZHEIMERS RES THER* 13:

AMYPAD: Ranson, Janice M. et al. Modifiable risk factors for dementia and dementia risk profiling. A user manual for Brain Health Services-part 2 of 6, *ALZHEIMERS RES THER* 13:

AMYPAD: Visser, Leonie N. C. et al. Dementia risk communication. A user manual for Brain Health Services-part 3 of 6, *ALZHEIMERS RES THER* 13:

AMYPAD: Milne, Richard et al. Societal and equity challenges for Brain Health Services. A user manual for Brain Health Services-part 6 of 6, *ALZHEIMERS RES THER* 13:

APPROACH: Musumeci, Giuseppe et al. Biomarkers of Chondrocyte Apoptosis and Autophagy in Osteoarthritis, INT J MOL SCI 16: 20560-20575

APPROACH: Rahmati, Maryam et al. Inflammatory mediators in osteoarthritis: A critical review of the state-of-the-art, current prospects, and future challenges, BONE 85: 81-90

APPROACH: Richardson, Stephen M. et al. Mesenchymal stem cells in regenerative medicine: Focus on articular cartilage and intervertebral disc regeneration, METHODS 99: 69-80

APPROACH: Mobasheri, A. et al. Osteoarthritis Year in Review 2016: biomarkers (biochemical markers), OSTEOARTHR CARTILAGE 25: 199-208

APPROACH: Mobasheri, Ali et al. The role of metabolism in the pathogenesis of osteoarthritis, NAT REV RHEUMATOL 13: 302-311

APPROACH: Fellows, Christopher R. et al. Adipose, Bone Marrow and Synovial Joint-Derived Mesenchymal Stem Cells for Cartilage Repair, FRONT GENET 7:

APPROACH: Sanchez, C. et al. Chondrocyte secretome: a source of novel insights and exploratory biomarkers of osteoarthritis, OSTEOARTHR CARTILAGE 25: 1199-1209

APPROACH: Luo, Yunyun et al. The minor collagens in articular cartilage, PROTEIN CELL 8: 560-572

APPROACH: Rahmati, Maryam et al. Aging and osteoarthritis: Central role of the extracellular matrix, AGEING RES REV 40: 20-30

APPROACH: Henrotin, Y. et al. Osteoarthritis biomarkers derived from cartilage extracellular matrix: Current status and future perspectives, ANN PHYS REHABIL MED 59: 145-148

APPROACH: Mobasheri, Ali et al. An update on the pathophysiology of osteoarthritis, ANN PHYS REHABIL MED 59: 333-339

APPROACH: Francisco, Vera et al. Adipokines and inflammation: is it a question of weight?, BRIT J PHARMACOL 175: 1569-1579

APPROACH: Thomas, Sally et al. What is the evidence for a role for diet and nutrition in osteoarthritis?, RHEUMATOLOGY 57: 61-74

APPROACH: Mobasheri, Ali et al. The chondrocyte channelome: A narrative review, JOINT BONE SPINE 86: 29-35

APPROACH: Guan, Vivienne X. et al. A systematic review of osteoarthritis prevention and management with dietary phytochemicals from foods, MATURITAS 122: 35-43

APPROACH: Lewis, Rebecca et al. Strategies for optimising musculoskeletal health in the 21st century, BMC MUSCULOSKEL DIS 20:

APPROACH: Mobasheri, Ali et al. Molecular taxonomy of osteoarthritis for patient stratification, disease management and drug development: biochemical markers associated with emerging clinical phenotypes and molecular endotypes, CURR OPIN RHEUMATOL 31: 80-89

APPROACH: Loef, Marieke et al. Fatty acids and osteoarthritis: different types, different effects, JOINT BONE SPINE 86: 451-458

APPROACH: Van Spil, Willem Evert et al. Osteoarthritis phenotypes and novel therapeutic targets, BIOCHEM PHARMACOL 165: 41-48

APPROACH: Loef, M. et al. The association of plasma fatty acids with hand and knee osteoarthritis: the NEO study, OSTEOARTHR CARTILAGE 28: 223-230

APPROACH: Widera, Pawel et al. Multi-classifier prediction of knee osteoarthritis progression from incomplete imbalanced longitudinal data, SCI REP-UK 10:

APPROACH: Gielis, Willem Paul et al. Scoring Osteoarthritis Reliably in Large Joints and the Spine Using Whole-Body CT: OsteoArthritis Computed Tomography-Score (OACT-Score), J PERS MED 11:

APPROACH: Saberi, Morteza et al. Targeting mitochondrial dysfunction with small molecules in intervertebral disc aging and degeneration, GEROSCIENCE 43: 517-537

APPROACH: Zheng, Linli et al. The role of metabolism in chondrocyte dysfunction and the progression of osteoarthritis, AGEING RES REV 66:

APPROACH: Cordero-Barreal, Alfonso et al. An Update on the Role of Leptin in the Immuno-Metabolism of Cartilage, INT J MOL SCI 22:

BEAT-DKD: Zschiedrich, Stefan et al. Targeting mTOR Signaling Can Prevent the Progression of FSGS, J AM SOC NEPHROL 28: 2144-2157

BEAT-DKD: Schell, Christoph et al. The Evolving Complexity of the Podocyte Cytoskeleton, J AM SOC NEPHROL 28: 3166-3174

BEAT-DKD: Henique, Carole et al. Genetic and pharmacological inhibition of microRNA-92a maintains podocyte cell cycle quiescence and limits crescentic glomerulonephritis, NAT COMMUN 8:

BEAT-DKD: Anders, Hans-Joachim et al. CKD in diabetes: diabetic kidney disease versus nondiabetic kidney disease, NAT REV NEPHROL 14: 361-377

BEAT-DKD: Hoehne, Martin et al. Single-nephron proteomes connect morphology and function in proteinuric kidney disease, KIDNEY INT 93: 1308-1319

BEAT-DKD: Viau, Amandine et al. Cilia-localized LKB1 regulates chemokine signaling, macrophage recruitment, and tissue homeostasis in the kidney, EMBO J 37:

BEAT-DKD: Selby, Nicholas M. et al. Magnetic resonance imaging biomarkers for chronic kidney disease: a position paper from the European Cooperation in Science and Technology Action PARENCHIMA, NEPHROL DIAL TRANSPL 33: II4-II14

BEAT-DKD: Wolf, Marcos et al. Magnetic resonance imaging T-1- and T-2-mapping to assess renal structure and function: a systematic review and statement paper, NEPHROL DIAL TRANSPL 33: II41-II50

BEAT-DKD: Wanner, Nicola et al. DNA Methyltransferase 1 Controls Nephron Progenitor Cell Renewal and Differentiation, J AM SOC NEPHROL 30: 63-78

BEAT-DKD: Brinkkoetter, Paul T. et al. Anaerobic Glycolysis Maintains the Glomerular Filtration Barrier Independent of Mitochondrial Metabolism and Dynamics, CELL REP 27: 1551-+

BEAT-DKD: Conserva, Francesca et al. Urinary miRNA-27b-3p and miRNA-1228-3p correlate with the progression of Kidney Fibrosis in Diabetic Nephropathy, SCI REP-UK 9:

BEAT-DKD: Mendichovszky, Iosif et al. Technical recommendations for clinical translation of renal MRI: a consensus project of the Cooperation in Science and Technology Action PARENCHIMA, MAGN RESON MATER PHY 33: 131-140

BEAT-DKD: Ljimini, Alexandra et al. Consensus-based technical recommendations for clinical translation of renal diffusion-weighted MRI, MAGN RESON MATER PHY 33: 177-195

BEAT-DKD: Buhl, Eva M. et al. Dysregulated mesenchymal PDGFR-beta drives kidney fibrosis, EMBO MOL MED 12:

BEAT-DKD: Meyer-Schwesinger, Catherine et al. A novel mouse model of phospholipase A2 receptor 1-associated membranous nephropathy mimics podocyte injury in patients, KIDNEY INT 97: 913-919

BEAT-DKD: Ramnath, Raina D. et al. Blocking matrix metalloproteinase-mediated syndecan-4 shedding restores the endothelial glycocalyx and glomerular filtration barrier function in early diabetic kidney disease, KIDNEY INT 97: 951-965

BEAT-DKD: Hapca, Simona et al. The Relationship between AKI and CKD in Patients with Type 2 Diabetes: An Observational Cohort Study, J AM SOC NEPHROL 32: 138-150

BEAT-DKD: Fazzini, F. et al. Association of mitochondrial DNA copy number with metabolic syndrome and type 2 diabetes in 14 176 individuals, J INTERN MED 290: 190-202

BEAT-DKD: Zaibi, Nawel et al. Protective effects of dapagliflozin against oxidative stress-induced cell injury in human proximal tubular cells, PLOS ONE 16:

BEAT-DKD: Rogg, Manuel et al. SRGAP1 Controls Small Rho GTPases To Regulate Podocyte Foot Process Maintenance, J AM SOC NEPHROL 32: 563-579

BEAT-DKD: Zimmermann, Marina et al. Deep learning-based molecular morphometrics for kidney biopsies, JCI INSIGHT 6:

BEAT-DKD: Lee, Matthew M. Y. et al. Effect of Empagliflozin on Left Ventricular Volumes in Patients With Type 2 Diabetes, or Prediabetes, and Heart Failure With Reduced Ejection Fraction (SUGAR-DM-HF), CIRCULATION 143: 516-525

BigData@Heart: Hemingway, Harry et al. Big data from electronic health records for early and late translational cardiovascular research: challenges and potential, EUR HEART J 39: 1481-+

BigData@Heart: DeSalvo, Daniel J. et al. Continuous glucose monitoring and glycemic control among youth with type 1 diabetes: International comparison from the T1D Exchange and DPV Initiative, PEDIATR DIABETES 19: 1271-1275

BigData@Heart: Schrage, Benedikt et al. Association Between Use of Primary-Prevention Implantable Cardioverter-Defibrillators and Mortality in Patients With Heart Failure A Prospective

Propensity Score-Matched Analysis From the Swedish Heart Failure Registry, *CIRCULATION* 140: 1530-1539

BigData@Heart: Jagodzinski, Annika et al. Rationale and Design of the Hamburg City Health Study, *EUR J EPIDEMIOL* 35: 169-181

BigData@Heart: Bunting, Karina V. et al. A Practical Guide to Assess the Reproducibility of Echocardiographic Measurements, *J AM SOC ECHOCARDIOG* 32: 1505-1515

BigData@Heart: Li, Qianrui et al. Diagnosis and treatment for hyperuricemia and gout: a systematic review of clinical practice guidelines and consensus statements, *BMJ OPEN* 9:

BigData@Heart: Willems, Stephan et al. Cabins, castles, and constant hearts: rhythm control therapy in patients with atrial fibrillation, *EUR HEART J* 40: 3793-+

BigData@Heart: Allara, Elias et al. Genetic Determinants of Lipids and Cardiovascular Disease Outcomes A Wide-Angled Mendelian Randomization Investigation, *CIRC-GENOM PRECIS ME* 12: 543-551

BigData@Heart: Denaxas, Spiros et al. UK phenomics platform for developing and validating electronic health record phenotypes: CALIBER, *J AM MED INFORM ASSN* 26: 1545-1559

BigData@Heart: Seligman, William H. et al. Development of an international standard set of outcome measures for patients with atrial fibrillation: a report of the International Consortium for Health Outcomes Measurement (ICHOM) atrial fibrillation working group, *EUR HEART J* 41: 1132-1140

BigData@Heart: Vollmer, Sebastian et al. Machine learning and artificial intelligence research for patient benefit: 20 critical questions on transparency, replicability, ethics, and effectiveness, *BMJ-BRIT MED J* 368:

BigData@Heart: van Ouwkerk, Antoinette F. et al. Epigenetic and Transcriptional Networks Underlying Atrial Fibrillation, *CIRC RES* 127: 34-50

BigData@Heart: Schmidt, Amand F. et al. Genetic drug target validation using Mendelian randomisation, *NAT COMMUN* 11:

BigData@Heart: Bean, Daniel M. et al. Angiotensin-converting enzyme inhibitors and angiotensin II receptor blockers are not associated with severe COVID-19 infection in a multi-site UK acute hospital trust, *EUR J HEART FAIL* 22: 967-974

BigData@Heart: Dretzke, Janine et al. Predicting recurrent atrial fibrillation after catheter ablation: a systematic review of prognostic models, *EUROPACE* 22: 748-760

BigData@Heart: Meyer, Hannah V. et al. Genetic and functional insights into the fractal structure of the heart, *NATURE* 584: 589-+

BigData@Heart: Lai, Alvina G. et al. Estimated impact of the COVID-19 pandemic on cancer services and excess 1-year mortality in people with cancer and multimorbidity: near real-time data on cancer care, cancer deaths and a population-based cohort study, *BMJ OPEN* 10:

BigData@Heart: Savarese, Gianluigi et al. Association between renin-angiotensin-aldosterone system inhibitor use and COVID-19 hospitalization and death: a 1.4 million patient nationwide registry analysis, *EUR J HEART FAIL* 23: 476-485

BigData@Heart: Schrage, Benedikt et al. Lower socioeconomic status predicts higher mortality and morbidity in patients with heart failure, *HEART* 107: 229-236

BigData@Heart: Sun, Luanluan et al. Polygenic risk scores in cardiovascular risk prediction: A cohort study and modelling analyses, *PLOS MED* 18:

BigData@Heart: Carr, Ewan et al. Evaluation and improvement of the National Early Warning Score (NEWS2) for COVID-19: a multi-hospital study, *BMC MED* 19:

BigData@Heart: Becher, Peter M. et al. Use of sodium-glucose co-transporter 2 inhibitors in patients with heart failure and type 2 diabetes mellitus: data from the Swedish Heart Failure Registry, *EUR J HEART FAIL* 23: 1012-1022

BigData@Heart: Katsoulis, M. et al. Obesity during the COVID-19 pandemic: both cause of high risk and potential effect of lockdown? A population-based electronic health record study, *PUBLIC HEALTH* 191: 41-47

BigData@Heart: Antoniadou, Charalambos et al. The year in cardiovascular medicine 2020: digital health and innovation, *EUR HEART J* 42: 732-739

BigData@Heart: Gaziano, Liam et al. Actionable druggable genome-wide Mendelian randomization identifies repurposing opportunities for COVID-19, *NAT MED* 27:

BigData@Heart: Wood, Angela et al. Linked electronic health records for research on a nationwide cohort of more than 54 million people in England: data resource, *BMJ-BRIT MED J* 372:

BigData@Heart: Yuan, Shuai et al. Homocysteine, B vitamins, and cardiovascular disease: a Mendelian randomization study, *BMC MED* 19:

BigData@Heart: Uijl, Alicia et al. Identification of distinct phenotypic clusters in heart failure with preserved ejection fraction, *EUR J HEART FAIL* 23: 973-982

BigData@Heart: Jordan, Elizabeth et al. Evidence-Based Assessment of Genes in Dilated Cardiomyopathy, *CIRCULATION* 144: 7-19

BigData@Heart: Idris, Iskandar et al. Lower risk of hospitalization for heart failure, kidney disease and death with sodium-glucose co-transporter-2 inhibitors compared with dipeptidyl peptidase-4 inhibitors in type 2 diabetes regardless of prior cardiovascular or kidney disease: A retrospective, *DIABETES OBES METAB* 23: 2207-2214

BigData@Heart: Banerjee, Amitava et al. Excess deaths in people with cardiovascular diseases during the COVID-19 pandemic, *EUR J PREV CARDIOL* 28: 1599-1608

BIGPICTURE: Moulin, Pierre et al. IMI-Bigpicture: A Central Repository for Digital Pathology, *TOXICOL PATHOL* 49: 711-713

BIOMAP: Fyhrquist, Nanna et al. Microbe-host interplay in atopic dermatitis and psoriasis, *NAT COMMUN* 10:

BIOMAP: Tsoi, Lam C. et al. Progression of acute-to-chronic atopic dermatitis is associated with quantitative rather than qualitative changes in cytokine responses, *J ALLERGY CLIN IMMUN* 145: 1406-1415

BIOMAP: Langan, Sinead M. et al. Atopic dermatitis, *LANCET* 396: 345-360

BIOMAP: Mobus, Lena et al. Atopic dermatitis displays stable and dynamic skin transcriptome signatures, *J ALLERGY CLIN IMMUN* 147: 213-223

BIOMAP: Marrs, Tom et al. Gut microbiota development during infancy: Impact of introducing allergenic foods, *J ALLERGY CLIN IMMUN* 147: 613-+

BioVacSafe: Kaufmann, Stefan H. E. et al. Tuberculosis vaccines: Time to think about the next generation, *SEMIN IMMUNOL* 25: 172-181

BioVacSafe: Weiner, J., III et al. Recent advances towards tuberculosis control: vaccines and biomarkers, *J INTERN MED* 275: 467-480

BioVacSafe: Kaufmann, Stefan H. E. et al. Progress in tuberculosis vaccine development and host-directed therapies-a state of the art review, *LANCET RESP MED* 2: 301-320

BioVacSafe: Andersen, Peter et al. Novel Vaccination Strategies against Tuberculosis, *CSH PERSPECT MED* 4:

BioVacSafe: Andersen, Peter et al. Tuberculosis vaccines - rethinking the current paradigm, *TRENDS IMMUNOL* 35: 387-395

BioVacSafe: Rappuoli, Rino et al. Vaccines, new opportunities for a new society, *P NATL ACAD SCI USA* 111: 12288-12293

BioVacSafe: Cliff, Jacqueline M. et al. The human immune response to tuberculosis and its treatment: a view from the blood, *IMMUNOL REV* 264: 88-102

BioVacSafe: Tricot, Sabine et al. Evaluating the Efficiency of Isotope Transmission for Improved Panel Design and a Comparison of the Detection Sensitivities of Mass Cytometer Instruments, *CYTOM PART A* 87A: 357-368

BioVacSafe: Olafsdottir, Thorunn et al. Molecular signatures of vaccine adjuvants, *VACCINE* 33: 5302-5307

BioVacSafe: Kaufmann, Stefan H. E. et al. Molecular Determinants in Phagocyte-Bacteria Interactions, *IMMUNITY* 44: 476-491

BioVacSafe: Kaufmann, Stefan H. E. et al. Host-directed therapies for bacterial and viral infections, *NAT REV DRUG DISCOV* 17: 35-56

BioVacSafe: Gao, Yong et al. Advances in HIV-1 Vaccine Development, *VIRUSES-BASEL* 10:

BioVacSafe: Zyla, Joanna et al. Gene set enrichment for reproducible science: comparison of CERNO and eight other algorithms, *BIOINFORMATICS* 35: 5146-5154

BioVacSafe: Pei, Gang et al. Cellular stress promotes NOD1/2-dependent inflammation via the endogenous metabolite sphingosine-1-phosphate, *EMBO J* 40:

BTCure: Cope, Andrew et al. The Th1 life cycle: molecular control of IFN-gamma to IL-10 switching, *TRENDS IMMUNOL* 32: 278-286

BTCure: Finzel, Stephanie et al. Repair of bone erosions in rheumatoid arthritis treated with tumour necrosis factor inhibitors is based on bone apposition at the base of the erosion, *ANN RHEUM DIS* 70: 1587-1593

BTCure: Shi, Jing et al. Autoantibodies recognizing carbamylated proteins are present in sera of patients with rheumatoid arthritis and predict joint damage, *P NATL ACAD SCI USA* 108: 17372-17377

BTCure: Heiland, Gisela Ruiz et al. High level of functional dickkopf-1 predicts protection from syndesmophyte formation in patients with ankylosing spondylitis, *ANN RHEUM DIS* 71: 572-574

BTCure: Akhmetshina, Alfiya et al. Activation of canonical Wnt signalling is required for TGF-beta-mediated fibrosis, *NAT COMMUN* 3:

BTCure: Gerlag, Danielle M. et al. EULAR recommendations for terminology and research in individuals at risk of rheumatoid arthritis: report from the Study Group for Risk Factors for Rheumatoid Arthritis, *ANN RHEUM DIS* 71: 638-641

BTCure: Suwannalai, P. et al. Avidity maturation of anti-citrullinated protein antibodies in rheumatoid arthritis, *ARTHRITIS RHEUM-US* 64: 1323-1328

BTCure: Harre, Ulrike et al. Induction of osteoclastogenesis and bone loss by human autoantibodies against citrullinated vimentin, *J CLIN INVEST* 122: 1791-1802

BTCure: Nikitopoulou, Ioanna et al. Autotaxin expression from synovial fibroblasts is essential for the pathogenesis of modeled arthritis, *J EXP MED* 209: 923-931

BTCure: Klarenbeek, P. L. et al. Inflamed target tissue provides a specific niche for highly expanded T-cell clones in early human autoimmune disease, *ANN RHEUM DIS* 71: 1088-1093

BTCure: Uderhardt, Stefan et al. 12/15-Lipoxygenase Orchestrates the Clearance of Apoptotic Cells and Maintains Immunologic Tolerance, *IMMUNITY* 36: 834-846

BTCure: Pandis, Ioannis et al. Identification of microRNA-221/222 and microRNA-323-3p association with rheumatoid arthritis via predictions using the human tumour necrosis factor transgenic mouse model, *ANN RHEUM DIS* 71: 1716-1723

BTCure: Giera, Martin et al. Lipid and lipid mediator profiling of human synovial fluid in rheumatoid arthritis patients by means of LC-MS/MS, *BBA-MOL CELL BIOL L* 1821: 1415-1424

BTCure: Schett, Georg et al. Bone erosion in rheumatoid arthritis: mechanisms, diagnosis and treatment, *NAT REV RHEUMATOL* 8: 656-664

BTCure: Le Friec, Gaele et al. The CD46-Jagged1 interaction is critical for human T(H)1 immunity, *NAT IMMUNOL* 13: 1213-+



BTCure: Wesley, Annmarie et al. Association between body mass index and anti-citrullinated protein antibody-positive and anti-citrullinated protein antibody-negative rheumatoid arthritis: Results from a population-based case-control study, *ARTHRIT CARE RES* 65: 107-112

BTCure: Trouw, Leendert A. et al. Closing the serological gap: promising novel biomarkers for the early diagnosis of rheumatoid arthritis, *AUTOIMMUN REV* 12: 318-322

BTCure: Schett, Georg et al. Diabetes Is an Independent Predictor for Severe Osteoarthritis Results from a longitudinal cohort study, *DIABETES CARE* 36: 403-409

BTCure: Finzel, Stephanie et al. Interleukin-6 receptor blockade induces limited repair of bone erosions in rheumatoid arthritis: a micro CT study, *ANN RHEUM DIS* 72: 396-400

BTCure: Amara, Khaled et al. Monoclonal IgG antibodies generated from joint-derived B cells of RA patients have a strong bias toward citrullinated autoantigen recognition, *J EXP MED* 210: 445-455

BTCure: Kiechl, Stefan et al. Blockade of receptor activator of nuclear factor-kappa B (RANKL) signaling improves hepatic insulin resistance and prevents development of diabetes mellitus, *NAT MED* 19: 358-363

BTCure: Cui, Jing et al. Genome-Wide Association Study and Gene Expression Analysis Identifies CD84 as a Predictor of Response to Etanercept Therapy in Rheumatoid Arthritis, *PLOS GENET* 9:

BTCure: Brink, Mikael et al. Multiplex Analyses of Antibodies Against Citrullinated Peptides in Individuals Prior to Development of Rheumatoid Arthritis, *ARTHRITIS RHEUM-US* 65: 899-910

BTCure: Shi, Jing et al. Brief Report: AntiCarbamylated Protein Antibodies Are Present in Arthralgia Patients and Predict the Development of Rheumatoid Arthritis, *ARTHRITIS RHEUM-US* 65: 911-915

BTCure: Trenkmann, Michelle et al. Tumor Necrosis Factor alpha-Induced MicroRNA-18a Activates Rheumatoid Arthritis Synovial Fibroblasts Through a Feedback Loop in NF-kappa B Signaling, *ARTHRITIS RHEUM-US* 65: 916-927

BTCure: Lundberg, Karin et al. Genetic and environmental determinants for disease risk in subsets of rheumatoid arthritis defined by the anticitrullinated protein/peptide antibody fine specificity profile, *ANN RHEUM DIS* 72: 652-658

BTCure: Lin, Neng-Yu et al. Autophagy regulates TNF alpha-mediated joint destruction in experimental arthritis, *ANN RHEUM DIS* 72: 761-768

BTCure: Guenther, Claudia et al. Apoptosis, necrosis and necroptosis: cell death regulation in the intestinal epithelium, *GUT* 62: 1062-1071

BTCure: Wenniger, Lucas J. Maillette de Buy et al. Immunoglobulin G4+clones identified by next-generation sequencing dominate the B cell receptor repertoire in immunoglobulin G4 associated cholangitis, *HEPATOLOGY* 57: 2390-2398

BTCure: Frey, Silke et al. The novel cytokine interleukin-36 alpha is expressed in psoriatic and rheumatoid arthritis synovium, *ANN RHEUM DIS* 72: 1569-1574

BTCure: Rose, Thomas et al. IFN and its response proteins, IP-10 and SIGLEC-1, are biomarkers of disease activity in systemic lupus erythematosus, *ANN RHEUM DIS* 72: 1639-1645

BTCure: de Hair, Maria J. H. et al. Smoking and overweight determine the likelihood of developing rheumatoid arthritis, ANN RHEUM DIS 72: 1654-1658

BTCure: Maresz, Katarzyna J. et al. Porphyromonas gingivalis Facilitates the Development and Progression of Destructive Arthritis through Its Unique Bacterial Peptidylarginine Deiminase (PAD), PLOS PATHOG 9:

BTCure: Frisell, Thomas et al. Familial Risks and Heritability of Rheumatoid Arthritis Role of Rheumatoid Factor/Anti-Citrullinated Protein Antibody Status, Number and Type of Affected Relatives, Sex, and Age, ARTHRITIS RHEUM-US 65: 2773-2782

BTCure: Quirke, Anne-Marie et al. Heightened immune response to autocitrullinated Porphyromonas gingivalis peptidylarginine deiminase: a potential mechanism for breaching immunologic tolerance in rheumatoid arthritis, ANN RHEUM DIS 73: 263-269

BTCure: Shi, Jing et al. Carbamylation and antibodies against carbamylated proteins in autoimmunity and other pathologies, AUTOIMMUN REV 13: 225-230

BTCure: Kumari, Snehlata et al. Tumor Necrosis Factor Receptor Signaling in Keratinocytes Triggers Interleukin-24-Dependent Psoriasis-like Skin Inflammation in Mice, IMMUNITY 39: 899-911

BTCure: Liszewski, M. Kathryn et al. Intracellular Complement Activation Sustains T Cell Homeostasis and Mediates Effector Differentiation, IMMUNITY 39: 1143-1157

BTCure: Okada, Yukinori et al. Genetics of rheumatoid arthritis contributes to biology and drug discovery, NATURE 506: 376-+

BTCure: Doorenspleet, M. E. et al. Rheumatoid arthritis synovial tissue harbours dominant B-cell and plasma-cell clones associated with autoreactivity, ANN RHEUM DIS 73: 756-762

BTCure: Shi, Jing et al. Anti-carbamylated protein (anti-CarP) antibodies precede the onset of rheumatoid arthritis, ANN RHEUM DIS 73: 780-783

BTCure: Evans, Hayley G. et al. TNF-alpha blockade induces IL-10 expression in human CD4+T cells, NAT COMMUN 5:

BTCure: Burska, Agata et al. Cytokines as Biomarkers in Rheumatoid Arthritis, MEDIAT INFLAMM 2014:

BTCure: Han, Buhm et al. Fine Mapping Seronegative and Seropositive Rheumatoid Arthritis to Shared and Distinct HLA Alleles by Adjusting for the Effects of Heterogeneity, AM J HUM GENET 94: 522-532

BTCure: Kleyer, Arnd et al. Bone loss before the clinical onset of rheumatoid arthritis in subjects with anticitrullinated protein antibodies, ANN RHEUM DIS 73: 854-860

BTCure: van Nies, J. A. B. et al. What is the evidence for the presence of a therapeutic window of opportunity in rheumatoid arthritis? A systematic literature review, ANN RHEUM DIS 73: 861-870

BTCure: de Aquino, Sabrina G. et al. Periodontal Pathogens Directly Promote Autoimmune Experimental Arthritis by Inducing a TLR2-and IL-1-Driven Th17 Response, J IMMUNOL 192: 4103-4111

BTCure: Bozec, Aline et al. T Cell Costimulation Molecules CD80/86 Inhibit Osteoclast Differentiation by Inducing the IDO/Tryptophan Pathway, SCI TRANSL MED 6:

BTCure: Reynisdottir, Gudrun et al. Structural Changes and Antibody Enrichment in the Lungs Are Early Features of Anti-Citrullinated Protein Antibody-Positive Rheumatoid Arthritis, ARTHRITIS RHEUMATOL 66: 31-39

BTCure: Kato, Masaru et al. Dual Role of Autophagy in Stress-Induced Cell Death in Rheumatoid Arthritis Synovial Fibroblasts, ARTHRITIS RHEUMATOL 66: 40-48

BTCure: de Hair, M. J. H. et al. Features of the Synovium of Individuals at Risk of Developing Rheumatoid Arthritis, ARTHRITIS RHEUMATOL 66: 513-522

BTCure: Menon, Bina et al. Interleukin-17+CD8+T Cells Are Enriched in the Joints of Patients With Psoriatic Arthritis and Correlate With Disease Activity and Joint Damage Progression, ARTHRITIS RHEUMATOL 66: 1272-1281

BTCure: James, Eddie A. et al. Citrulline-Specific Th1 Cells Are Increased in Rheumatoid Arthritis and Their Frequency Is Influenced by Disease Duration and Therapy, ARTHRITIS RHEUMATOL 66: 1712-1722

BTCure: Liu, Bi-Sheng et al. TLR-mediated STAT3 and ERK activation controls IL-10 secretion by human B cells, EUR J IMMUNOL 44: 2121-2129

BTCure: DAlessio, Silvia et al. VEGF-C-dependent stimulation of lymphatic function ameliorates experimental inflammatory bowel disease, J CLIN INVEST 124: 3863-3878

BTCure: Khmaladze, Ia et al. Mannan induces ROS-regulated, IL-17A-dependent psoriasis arthritis-like disease in mice, P NATL ACAD SCI USA 111: E3669-E3678

BTCure: Leppkes, Moritz et al. Pleiotropic functions of TNF-alpha in the regulation of the intestinal epithelial response to inflammation, INT IMMUNOL 26: 509-515

BTCure: Catrina, Anca I. et al. Lungs, joints and immunity against citrullinated proteins in rheumatoid arthritis, NAT REV RHEUMATOL 10: 645-653

BTCure: Klein, Kerstin et al. Epigenetics in rheumatoid arthritis, CURR OPIN RHEUMATOL 27: 76-82

BTCure: Rombouts, Yoann et al. Anti-citrullinated protein antibodies acquire a pro-inflammatory Fc glycosylation phenotype prior to the onset of rheumatoid arthritis, ANN RHEUM DIS 74: 234-241

BTCure: van Baarsen, Lisa G. M. et al. Heterogeneous expression pattern of interleukin 17A (IL-17A), IL-17F and their receptors in synovium of rheumatoid arthritis, psoriatic arthritis and osteoarthritis: possible explanation for nonresponse to anti-IL-17 therapy?, ARTHRITIS RES THER 16:

BTCure: Hensvold, Aase Haj et al. Environmental and genetic factors in the development of anticitrullinated protein antibodies (ACPAs) and ACPA-positive rheumatoid arthritis: an epidemiological investigation in twins, ANN RHEUM DIS 74: 375-380

BTCure: Palumbo-Zerr, Katrin et al. Orphan nuclear receptor NR4A1 regulates transforming growth factor-beta signaling and fibrosis, NAT MED 21: 150-158

BTCure: Choi, Ivy Y. et al. MRP8/14 serum levels as a strong predictor of response to biological treatments in patients with rheumatoid arthritis, ANN RHEUM DIS 74: 499-505

BTCure: Kelkka, Tiina et al. Reactive Oxygen Species Deficiency Induces Autoimmunity with Type 1 Interferon Signature, ANTIOXID REDOX SIGN 21: 2231-2245

BTCure: Guenther, Claudia et al. Caspase-8 controls the gut response to microbial challenges by Tnf-alpha-dependent and independent pathways, GUT 64: 601-U1111

BTCure: Pieters, Bartijn C. H. et al. Commercial Cow Milk Contains Physically Stable Extracellular Vesicles Expressing Immunoregulatory TGF-beta, PLOS ONE 10:

BTCure: Harre, Ulrike et al. Glycosylation of immunoglobulin G determines osteoclast differentiation and bone loss, NAT COMMUN 6:

BTCure: Koenders, Marije I. et al. Novel therapeutic targets in rheumatoid arthritis, TRENDS PHARMACOL SCI 36: 189-195

BTCure: Viatte, Sebastien et al. Association of HLA-DRB1 Haplotypes With Rheumatoid Arthritis Severity, Mortality, and Treatment Response, JAMA-J AM MED ASSOC 313: 1645-1656

BTCure: Gan, Ryan W. et al. Anti-carbamylated Protein Antibodies Are Present Prior to Rheumatoid Arthritis and Are Associated with Its Future Diagnosis, J RHEUMATOL 42: 572-579

BTCure: van Steenberg, Hanna W. et al. Characterising arthralgia in the preclinical phase of rheumatoid arthritis using MRI, ANN RHEUM DIS 74: 1225-1232

BTCure: Gao, Wei et al. Hypoxia and STAT3 signalling interactions regulate pro-inflammatory pathways in rheumatoid arthritis, ANN RHEUM DIS 74: 1275-1283

BTCure: Mascalzoni, Deborah et al. International Charter of principles for sharing bio-specimens and data, EUR J HUM GENET 23: 721-728

BTCure: Tacconi, Carlotta et al. Vascular Endothelial Growth Factor C Disrupts the Endothelial Lymphatic Barrier to Promote Colorectal Cancer Invasion, GASTROENTEROLOGY 148: 1438+

BTCure: Kolev, Martin et al. Complement Regulates Nutrient Influx and Metabolic Reprogramming during Th1 Cell Responses, IMMUNITY 42: 1033-1047

BTCure: Ytterberg, A. Jimmy et al. Shared immunological targets in the lungs and joints of patients with rheumatoid arthritis: identification and validation, ANN RHEUM DIS 74: 1772-1777

BTCure: Lenz, Tobias L. et al. Widespread non-additive and interaction effects within HLA loci modulate the risk of autoimmune diseases, NAT GENET 47: 1085+

BTCure: Arntz, Onno J. et al. Oral administration of bovine milk derived extracellular vesicles attenuates arthritis in two mouse models, MOL NUTR FOOD RES 59: 1701-1712

BTCure: Luo, Yubin et al. Microbiota from Obese Mice Regulate Hematopoietic Stem Cell Differentiation by Altering the Bone Niche, CELL METAB 22: 886-894

BTCure: Hecht, Carolin et al. Additive effect of anti-citrullinated protein antibodies and rheumatoid factor on bone erosions in patients with RA, ANN RHEUM DIS 74: 2151-2156

BTCure: Martin, Paul et al. Capture Hi-C reveals novel candidate genes and complex long-range interactions with related autoimmune risk loci, NAT COMMUN 6:

BTCure: Gao, W. et al. Tofacitinib regulates synovial inflammation in psoriatic arthritis, inhibiting STAT activation and induction of negative feedback inhibitors, ANN RHEUM DIS 75: 311-315

BTCure: Haschka, Judith et al. Relapse rates in patients with rheumatoid arthritis in stable remission tapering or stopping antirheumatic therapy: interim results from the prospective randomised controlled RETRO study, ANN RHEUM DIS 75: 45-51

BTCure: Catrina, Anca I. et al. Mechanisms involved in triggering rheumatoid arthritis, IMMUNOL REV 269: 162-174

BTCure: Holmdahl, Rikard et al. Ncf1 polymorphism reveals oxidative regulation of autoimmune chronic inflammation, IMMUNOL REV 269: 228-247

BTCure: Koliaraki, Vasiliki et al. IKK beta in intestinal mesenchymal cells promotes initiation of colitis-associated cancer, J EXP MED 212: 2235-2251

BTCure: Klein, Kerstin et al. The bromodomain protein inhibitor I-BET151 suppresses expression of inflammatory genes and matrix degrading enzymes in rheumatoid arthritis synovial fibroblasts, ANN RHEUM DIS 75: 422-429

BTCure: Raaschou, Pauline et al. Rheumatoid arthritis, anti-tumour necrosis factor treatment, and risk of squamous cell and basal cell skin cancer: cohort study based on nationwide prospectively recorded data from Sweden, BMJ-BRIT MED J 352:

BTCure: van de Bovenkamp, Fleur S. et al. The Emerging Importance of IgG Fab Glycosylation in Immunity, J IMMUNOL 196: 1435-1441

BTCure: Mantel, Angla et al. Rheumatoid arthritis is associated with a more severe presentation of acute coronary syndrome and worse short-term outcome, EUR HEART J 36: 3413-3422

BTCure: Rombouts, Yoann et al. Extensive glycosylation of ACPA-IgG variable domains modulates binding to citrullinated antigens in rheumatoid arthritis, ANN RHEUM DIS 75: 578-585

BTCure: Uluckan, Oezge et al. Chronic skin inflammation leads to bone loss by IL-17-mediated inhibition of Wnt signaling in osteoblasts, SCI TRANSL MED 8:

BTCure: de Lange-Brokaar, B. J. E. et al. Characterization of synovial mast cells in knee osteoarthritis: association with clinical parameters, OSTEOARTH R CARTILAGE 24: 664-671

BTCure: Vicente, Rita et al. Deregulation and therapeutic potential of microRNAs in arthritic diseases, NAT REV RHEUMATOL 12: 211-220

BTCure: Simon, David et al. Analysis of periarticular bone changes in patients with cutaneous psoriasis without associated psoriatic arthritis, ANN RHEUM DIS 75: 660-666

BTCure: Krishnamurthy, Akilan et al. Identification of a novel chemokine-dependent molecular mechanism underlying rheumatoid arthritis-associated autoantibody-mediated bone loss, ANN RHEUM DIS 75: 721-729

BTCure: Wigerblad, Gustaf et al. Autoantibodies to citrullinated proteins induce joint pain independent of inflammation via a chemokine-dependent mechanism, ANN RHEUM DIS 75: 730-738

BTCure: Gerlag, Danielle M. et al. Towards prevention of autoantibody-positive rheumatoid arthritis: from lifestyle modification to preventive treatment, RHEUMATOLOGY 55: 607-614

BTCure: Klocke, Katrin et al. Induction of autoimmune disease by deletion of CTLA-4 in mice in adulthood, P NATL ACAD SCI USA 113: E2383-E2392

BTCure: Vicente, Rita et al. Cellular senescence impact on immune cell fate and function, AGING CELL 15: 400-406

BTCure: Kawalkowska, Joanna et al. Abrogation of collagen-induced arthritis by a peptidyl arginine deiminase inhibitor is associated with modulation of T cell-mediated immune responses, SCI REP-UK 6:

BTCure: Kerkman, Priscilla F. et al. Identification and characterisation of citrullinated antigen-specific B cells in peripheral blood of patients with rheumatoid arthritis, ANN RHEUM DIS 75: 1170-1176

BTCure: Danks, Lynett et al. RANKL expressed on synovial fibroblasts is primarily responsible for bone erosions during joint inflammation, ANN RHEUM DIS 75: 1187-1195

BTCure: Chen, Zhu et al. Th2 and eosinophil responses suppress inflammatory arthritis, NAT COMMUN 7:

BTCure: Arbore, Giuseppina et al. A novel complement-metabolism-inflammasome axis as a key regulator of immune cell effector function, EUR J IMMUNOL 46: 1563-1573

BTCure: Udalova, Irina A. et al. Macrophage heterogeneity in the context of rheumatoid arthritis, NAT REV RHEUMATOL 12: 472-485

BTCure: Hess, Christoph et al. Complement-Mediated Regulation of Metabolism and Basic Cellular Processes, IMMUNITY 45: 240-254

BTCure: Rech, Juergen et al. Prediction of disease relapses by multibiomarker disease activity and autoantibody status in patients with rheumatoid arthritis on tapering DMARD treatment, ANN RHEUM DIS 75: 1637-1644

BTCure: Reynisdottir, Gudrun et al. Signs of immune activation and local inflammation are present in the bronchial tissue of patients with untreated early rheumatoid arthritis, ANN RHEUM DIS 75: 1722-1727

BTCure: Campbell, T. Mark et al. Mesenchymal Stem Cell Alterations in Bone Marrow Lesions in Patients With Hip Osteoarthritis, ARTHRITIS RHEUMATOL 68: 1648-1659

BTCure: Altawil, Reem et al. Remaining Pain in Early Rheumatoid Arthritis Patients Treated With Methotrexate, ARTHRIT CARE RES 68: 1061-1068

BTCure: Munoz, Luis E. et al. Nanoparticles size-dependently initiate self-limiting NETosis-driven inflammation, P NATL ACAD SCI USA 113: E5856-E5865

BTCure: Freeley, Simon et al. The ins and outs of complement-driven immune responses, IMMUNOL REV 274: 16-32

BTCure: Lopez-Mejias, Raquel et al. Cardiovascular risk assessment in patients with rheumatoid arthritis: The relevance of clinical, genetic and serological markers, AUTOIMMUN REV 15: 1013-1030

BTCure: Scher, Jose U. et al. The lung microbiota in early rheumatoid arthritis and autoimmunity, MICROBIOME 4:

BTCure: Faustini, Francesca et al. Subclinical joint inflammation in patients with psoriasis without concomitant psoriatic arthritis: a cross-sectional and longitudinal analysis, ANN RHEUM DIS 75: 2068-2074

BTCure: Pfeifle, Rene et al. Regulation of autoantibody activity by the IL-23-T(H)17 axis determines the onset of autoimmune disease, NAT IMMUNOL 18: 104-113

BTCure: Malmstrom, Vivianne et al. The immunopathogenesis of seropositive rheumatoid arthritis: from triggering to targeting, NAT REV IMMUNOL 17: 60-75

BTCure: Kolev, Martin et al. Keeping it All Going - Complement Meets Metabolism, FRONT IMMUNOL 8:

BTCure: Ajeganova, S. et al. The association between anti-carbamylated protein (anti-CarP) antibodies and radiographic progression in early rheumatoid arthritis: a study exploring replication and the added value to ACPA and rheumatoid factor, ANN RHEUM DIS 76: 112-118

BTCure: Budin-Ljosne, Isabelle et al. Dynamic Consent: a potential solution to some of the challenges of modern biomedical research, BMC MED ETHICS 18:

BTCure: Catrina, Anca I. et al. Mechanisms leading from systemic autoimmunity to joint-specific disease in rheumatoid arthritis, NAT REV RHEUMATOL 13: 79-86

BTCure: Hafkenscheid, Lise et al. Structural Analysis of Variable Domain Glycosylation of Anti-Citrullinated Protein Antibodies in Rheumatoid Arthritis Reveals the Presence of Highly Sialylated Glycans, MOL CELL PROTEOMICS 16: 278-287

BTCure: Frank-Bertoncelj, Mojca et al. Epigenetically-driven anatomical diversity of synovial fibroblasts guides joint-specific fibroblast functions, NAT COMMUN 8:

BTCure: Koliaraki, Vasiliki et al. Mesenchymal Cells in Colon Cancer, GASTROENTEROLOGY 152: 964-979

BTCure: Hellgren, K. et al. Rheumatoid Arthritis and Risk of Malignant Lymphoma, ARTHRITIS RHEUMATOL 69: 700-708

BTCure: Lubbers, R. et al. Production of complement components by cells of the immune system, CLIN EXP IMMUNOL 188: 183-194

BTCure: Trouw, Leendert A. et al. Beyond citrullination: other post-translational protein modifications in rheumatoid arthritis, *NAT REV RHEUMATOL* 13: 331-339

BTCure: Melagraki, Georgia et al. Cheminformatics-aided discovery of small-molecule Protein-Protein Interaction (PPI) dual inhibitors of Tumor Necrosis Factor (TNF) and Receptor Activator of NF-kappa B Ligand (RANKL), *PLOS COMPUT BIOL* 13:

BTCure: Harre, Ulrike et al. Cellular and molecular pathways of structural damage in rheumatoid arthritis, *SEMIN IMMUNOPATHOL* 39: 355-363

BTCure: Alissafi, Themis et al. Tregs restrain dendritic cell autophagy to ameliorate autoimmunity, *J CLIN INVEST* 127: 2789-2804

BTCure: Jonasdottir, H. S. et al. Targeted lipidomics reveals activation of resolution pathways in knee osteoarthritis in humans, *OSTEOARTH R CARTILAGE* 25: 1150-1160

BTCure: Lie, Elisabeth et al. Tumour necrosis factor inhibitor treatment and occurrence of anterior uveitis in ankylosing spondylitis: results from the Swedish biologics register, *ANN RHEUM DIS* 76: 1515-1521

BTCure: Olsson, Lina M. et al. A single nucleotide polymorphism in the NCF1 gene leading to reduced oxidative burst is associated with systemic lupus erythematosus, *ANN RHEUM DIS* 76: 1607-1613

BTCure: Arbore, Giuseppina et al. Intracellular complement - the complosome - in immune cell regulation, *MOL IMMUNOL* 89: 2-9

BTCure: Ganguly, Payal et al. Age-related Changes in Bone Marrow Mesenchymal Stromal Cells: A Potential Impact on Osteoporosis and Osteoarthritis Development, *CELL TRANSPLANT* 26: 1520-1529

BTCure: Rogier, Rebecca et al. Alteration of the intestinal microbiome characterizes preclinical inflammatory arthritis in mice and its modulation attenuates established arthritis, *SCI REP-UK* 7:

BTCure: Schoenau, Verena et al. The value of F-18-FDG-PET/CT in identifying the cause of fever of unknown origin (FUO) and inflammation of unknown origin (IUO): data from a prospective study, *ANN RHEUM DIS* 77: 70-77

BTCure: Scherer, Hans Ulrich et al. The B cell response to citrullinated antigens in the development of rheumatoid arthritis, *NAT REV RHEUMATOL* 14: 157-169

BTCure: Karouzakis, Emmanuel et al. Analysis of early changes in DNA methylation in synovial fibroblasts of RA patients before diagnosis, *SCI REP-UK* 8:

BTCure: Hedstrom, Anna Karin et al. Smoking and susceptibility to rheumatoid arthritis in a Swedish population-based case-control study, *EUR J EPIDEMIOL* 33: 415-423

BTCure: Webster, Amy P. et al. Increased DNA methylation variability in rheumatoid arthritis-discordant monozygotic twins, *GENOME MED* 10:



BTCure: Lloyd, Katy A. et al. Differential ACPA Binding to Nuclear Antigens Reveals a PAD-Independent Pathway and a Distinct Subset of Acetylation Cross-Reactive Autoantibodies in Rheumatoid Arthritis, FRONT IMMUNOL 9:

BTCure: Perucha, Esperanza et al. The cholesterol biosynthesis pathway regulates IL-10 expression in human Th1 cells, NAT COMMUN 10:

BTCure: Steen, Johanna et al. Recognition of Amino Acid Motifs, Rather Than Specific Proteins, by Human Plasma Cell-Derived Monoclonal Antibodies to Posttranslationally Modified Proteins in Rheumatoid Arthritis, ARTHRITIS RHEUMATOL 71: 196-209

BTCure: Ge, Changrong et al. Structural Basis of Cross-Reactivity of Anti-Citrullinated Protein Antibodies, ARTHRITIS RHEUMATOL 71: 210-221

BTCure: Gerlag, Danielle M. et al. Effects of B-cell directed therapy on the preclinical stage of rheumatoid arthritis: the PRAIRI study, ANN RHEUM DIS 78: 179-185

BTCure: Frangou, Eleni et al. REDD1/autophagy pathway promotes thromboinflammation and fibrosis in human systemic lupus erythematosus (SLE) through NETs decorated with tissue factor (TF) and interleukin-17A (IL-17A), ANN RHEUM DIS 78: 238-248

BTCure: Burja, Blaz et al. Olive Leaf Extract Attenuates Inflammatory Activation and DNA Damage in Human Arterial Endothelial Cells, FRONT CARDIOVASC MED 6:

BTCure: Ramwadhoebe, Tamara H. et al. Effect of rituximab treatment on T and B cell subsets in lymph node biopsies of patients with rheumatoid arthritis, RHEUMATOLOGY 58: 1075-1085

BTCure: Hedstrom, Anna Karin et al. Complex Relationships of Smoking, HLA-DRB1 Genes, and Serologic Profiles in Patients With Early Rheumatoid Arthritis: Update From a Swedish Population-Based Case-Control Study, ARTHRITIS RHEUMATOL 71: 1504-1511

BTCure: Hafkenscheid, Lise et al. N-Linked Glycans in the Variable Domain of IgG Anti-Citrullinated Protein Antibodies Predict the Development of Rheumatoid Arthritis, ARTHRITIS RHEUMATOL 71: 1626-1633

BTCure: Sokolova, Maria, V et al. A set of serum markers detecting systemic inflammation in psoriatic skin, entheses, and joint disease in the absence of C-reactive protein and its link to clinical disease manifestations, ARTHRITIS RES THER 22:

BTCure: Kolev, Martin et al. Diapedesis-Induced Integrin Signaling via LFA-1 Facilitates Tissue Immunity by Inducing Intrinsic Complement C3 Expression in Immune Cells, IMMUNITY 52: 513-+

BTCure: Klareskog, L. et al. The importance of differences, On environment and its interactions with genes and immunity in the causation of rheumatoid arthritis, J INTERN MED 287: 514-533

BTCure: Scherer, Hans Ulrich et al. The etiology of rheumatoid arthritis, J AUTOIMMUN 110:

BTCure: Reed, Evan et al. Presence of autoantibodies in seronegative rheumatoid arthritis associates with classical risk factors and high disease activity, ARTHRITIS RES THER 22:

BTCure: Lubbers, R. et al. Complement component C1q is produced by isolated articular chondrocytes, OSTEOARTH CARTILAGE 28: 675-684

BTCure: Sahlstroem, Peter et al. Different Hierarchies of Anti-Modified Protein Autoantibody Reactivities in Rheumatoid Arthritis, *ARTHRITIS RHEUMATOL* 72: 1643-1657

BTCure: Kristyanto, Hendy et al. Persistently activated, proliferative memory autoreactive B cells promote inflammation in rheumatoid arthritis, *SCI TRANSL MED* 12:

CANCER-ID: Barault, L. et al. Digital PCR quantification of MGMT methylation refines prediction of clinical benefit from alkylating agents in glioblastoma and metastatic colorectal cancer, *ANN ONCOL* 26: 1994-1999

CANCER-ID: Misale, Sandra et al. Vertical suppression of the EGFR pathway prevents onset of resistance in colorectal cancers, *NAT COMMUN* 6:

CANCER-ID: Chudziak, Jakub et al. Clinical evaluation of a novel microfluidic device for epitope-independent enrichment of circulating tumour cells in patients with small cell lung cancer, *ANALYST* 141: 669-678

CANCER-ID: Arena, Sabrina et al. MM-151 overcomes acquired resistance to cetuximab and panitumumab in colorectal cancers harboring EGFR extracellular domain mutations, *SCI TRANSL MED* 8:

CANCER-ID: Pantel, K. et al. The biology of circulating tumor cells, *ONCOGENE* 35: 1216-1224

CANCER-ID: Russo, Mariangela et al. Acquired Resistance to the TRK Inhibitor Entrectinib in Colorectal Cancer, *CANCER DISCOV* 6: 36-44

CANCER-ID: Russo, Mariangela et al. Tumor Heterogeneity and Lesion-Specific Response to Targeted Therapy in Colorectal Cancer, *CANCER DISCOV* 6: 147-153

CANCER-ID: Andree, Kiki C. et al. Challenges in circulating tumor cell detection by the CellSearch system, *MOL ONCOL* 10: 395-407

CANCER-ID: Bidard, Francois-Clement et al. Circulating tumor cells in breast cancer, *MOL ONCOL* 10: 418-430

CANCER-ID: Heitzer, Ellen et al. Non-invasive detection of genome-wide somatic copy number alterations by liquid biopsies, *MOL ONCOL* 10: 494-502

CANCER-ID: Hvichia, G. E. et al. A novel microfluidic platform for size and deformability based separation and the subsequent molecular characterization of viable circulating tumor cells, *INT J CANCER* 138: 2894-2904

CANCER-ID: Gorges, Tobias M. et al. Enumeration and Molecular Characterization of Tumor Cells in Lung Cancer Patients Using a Novel In Vivo Device for Capturing Circulating Tumor Cells, *CLIN CANCER RES* 22: 2197-2206

CANCER-ID: Alix-Panabieres, Catherine et al. Clinical Applications of Circulating Tumor Cells and Circulating Tumor DNA as Liquid Biopsy, *CANCER DISCOV* 6: 479-491

CANCER-ID: Gorges, Tobias M. et al. Heterogeneous PSMA expression on circulating tumor cells - a potential basis for stratification and monitoring of PSMA-directed therapies in prostate cancer, *ONCOTARGET* 7: 34930-34941

CANCER-ID: Stoecklein, Nikolas H. et al. Challenges for CTC-based liquid biopsies: low CTC frequency and diagnostic leukapheresis as a potential solution, EXPERT REV MOL DIAGN 16: 147-164

CANCER-ID: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

CANCER-ID: Ulz, Peter et al. Inferring expressed genes by whole-genome sequencing of plasma DNA, NAT GENET 48: 1273-1278

CANCER-ID: Hanssen, Annkathrin et al. Characterization of different CTC subpopulations in non-small cell lung cancer, SCI REP-UK 6:

CANCER-ID: Gorges, Tobias M. et al. Accession of Tumor Heterogeneity by Multiplex Transcriptome Profiling of Single Circulating Tumor Cells, CLIN CHEM 62: 1504-1515

CANCER-ID: Swennenhuis, J. F. et al. Improving the CellSearch (R) system, EXPERT REV MOL DIAGN 16: 1291-1305

CANCER-ID: van Emburgh, Beth O. et al. Acquired RAS or EGFR mutations and duration of response to EGFR blockade in colorectal cancer, NAT COMMUN 7:

CANCER-ID: Kuskel, Andra et al. Improved detection of circulating tumor cells in non-metastatic high-risk prostate cancer patients, SCI REP-UK 6:

CANCER-ID: Wang, Hongxia et al. Circulating and disseminated tumor cells: diagnostic tools and therapeutic targets in motion, ONCOTARGET 8: 1884-1912

CANCER-ID: Alix-Panabieres, Catherine et al. Epithelial-mesenchymal plasticity in circulating tumor cells, J MOL MED 95: 133-142

CANCER-ID: Bardelli, Alberto et al. Liquid Biopsies, What We Do Not Know (Yet), CANCER CELL 31: 172-179

CANCER-ID: Picco, Gabriele et al. Loss of AXIN1 drives acquired resistance to WNT pathway blockade in colorectal cancer cells carrying RSPO3 fusions, EMBO MOL MED 9: 293-303

CANCER-ID: Perakis, Samantha et al. Emerging concepts in liquid biopsies, BMC MED 15:

CANCER-ID: Zeune, Leonie et al. Multiscale Segmentation via Bregman Distances and Nonlinear Spectral Analysis, SIAM J IMAGING SCI 10: 111-146

CANCER-ID: Pailler, Emma et al. Circulating Tumor Cells with Aberrant ALK Copy Number Predict Progression-Free Survival during Crizotinib Treatment in ALK-Rearranged Non-Small Cell Lung Cancer Patients, CANCER RES 77: 2222-2230

CANCER-ID: Pietrantonio, Filippo et al. Heterogeneity of Acquired Resistance to Anti-EGFR Monoclonal Antibodies in Patients with Metastatic Colorectal Cancer, CLIN CANCER RES 23: 2414-2422

CANCER-ID: Cabel, L. et al. Circulating tumor cells and circulating tumor DNA: What surgical oncologists need to know?, EJSO-EUR J SURG ONC 43: 949-962

CANCER-ID: Cabel, Luc et al. Circulating tumor cells: clinical validity and utility, INT J CLIN ONCOL 22: 421-430

CANCER-ID: Pixberg, C. F. et al. Analysis of DNA methylation in single circulating tumor cells, ONCOGENE 36: 3223-3231

CANCER-ID: Lindsay, C. R. et al. A prospective examination of circulating tumor cell profiles in non-small-cell lung cancer molecular subgroups, ANN ONCOL 28: 1523-1531

CANCER-ID: Amirouchene-Angelozzi, Nabil et al. Tumor Evolution as a Therapeutic Target, CANCER DISCOV 7: 805-817

CANCER-ID: Siravegna, Giulia et al. Integrating liquid biopsies into the management of cancer, NAT REV CLIN ONCOL 14: 531-548

CANCER-ID: Siena, S. et al. Dynamic molecular analysis and clinical correlates of tumor evolution within a phase II trial of panitumumab-based therapy in metastatic colorectal cancer, ANN ONCOL 29: 119-126

CANCER-ID: Mastoraki, Sophia et al. ESR1 Methylation: A Liquid Biopsy-Based Epigenetic Assay for the Follow-up of Patients with Metastatic Breast Cancer Receiving Endocrine Treatment, CLIN CANCER RES 24: 1500-1510

CANCER-ID: Poudineh, Mahla et al. Profiling circulating tumour cells and other biomarkers of invasive cancers, NAT BIOMED ENG 2: 72-84

CANCER-ID: Riethdorf, Sabine et al. Clinical applications of the CellSearch platform in cancer patients, ADV DRUG DELIVER REV 125: 102-121

CANCER-ID: Mainardi, Sara et al. SHP2 is required for growth of KRAS-mutant non-small-cell lung cancer in vivo, NAT MED 24: 961-+

CANCER-ID: Siravegna, Giulia et al. Radiologic and Genomic Evolution of Individual Metastases during HER2 Blockade in Colorectal Cancer, CANCER CELL 34: 148-+

CANCER-ID: Anfossi, Simone et al. Clinical utility of circulating non-coding RNAs - an update, NAT REV CLIN ONCOL 15: 541-563

CANCER-ID: Bidard, Francois-Clement et al. Circulating Tumor Cells in Breast Cancer Patients Treated by Neoadjuvant Chemotherapy: A Meta-analysis, JNCI-J NATL CANCER I 110: 560-567

CANCER-ID: Zavridou, Martha et al. Evaluation of Preanalytical Conditions and Implementation of Quality Control Steps for Reliable Gene Expression and DNA Methylation Analyses in Liquid Biopsies, CLIN CHEM 64: 1522-1533

CANCER-ID: Barault, Ludovic et al. Discovery of methylated circulating DNA biomarkers for comprehensive non-invasive monitoring of treatment response in metastatic colorectal cancer, GUT 67: 1995-2005

CANCER-ID: Andree, Kiki C. et al. Toward a real liquid biopsy in metastatic breast and prostate cancer: Diagnostic LeukApheresis increases CTC yields in a European prospective multicenter study (CTCTrap), INT J CANCER 143: 2584-2591

CANCER-ID: Germano, Giovanni et al. The Clinical Impact of the Genomic Landscape of Mismatch Repair-Deficient Cancers, *CANCER DISCOV* 8: 1518-1528

CANCER-ID: Neumann, Martin H. D. et al. ctDNA and CTCs in Liquid Biopsy - Current Status and Where We Need to Progress, *COMPUT STRUCT BIOTEC* 16: 190-195

CANCER-ID: Fehm, Tanja N. et al. Diagnostic leukapheresis for CTC analysis in breast cancer patients: CTC frequency, clinical experiences and recommendations for standardized reporting, *CYTOM PART A* 93A: 1213-1219

CANCER-ID: Pantel, Klaus et al. Circulating Tumor Cells in Prostate Cancer: From Discovery to Clinical Utility, *CLIN CHEM* 65: 87-99

CANCER-ID: Heitzer, Ellen et al. Current and future perspectives of liquid biopsies in genomics-driven oncology, *NAT REV GENET* 20: 71-88

CANCER-ID: Lianidou, Evi et al. Liquid biopsies, *GENE CHROMOSOME CANC* 58: 219-232

CANCER-ID: Reimers, Natalie et al. Liquid biopsy: novel technologies and clinical applications, *CLIN CHEM LAB MED* 57: 312-316

CANCER-ID: de Wit, Sanne et al. Single tube liquid biopsy for advanced non-small cell lung cancer, *INT J CANCER* 144: 3127-3137

CANCER-ID: Siravegna, Giulia et al. Plasma HER2 (ERBB2) Copy Number Predicts Response to HER2-targeted Therapy in Metastatic Colorectal Cancer, *CLIN CANCER RES* 25: 3046-3053

CANCER-ID: Rothwell, Dominic G. et al. Utility of ctDNA to support patient selection for early phase clinical trials: the TARGET study, *NAT MED* 25: 738-+

CANCER-ID: Pantel, Klaus et al. Liquid biopsy and minimal residual disease - latest advances and implications for cure, *NAT REV CLIN ONCOL* 16: 409-424

CANCER-ID: Janning, Melanie et al. Determination of PD-L1 Expression in Circulating Tumor Cells of NSCLC Patients and Correlation with Response to PD-1/PD-L1 Inhibitors, *CANCERS* 11:

CANCER-ID: Lindsay, C. R. et al. EPAC-lung: pooled analysis of circulating tumour cells in advanced non-small cell lung cancer, *EUR J CANCER* 117: 60-68

CANCER-ID: Tamminga, Menno et al. Circulating tumor cells in advanced non-small cell lung cancer patients are associated with worse tumor response to checkpoint inhibitors, *J IMMUNOTHER CANCER* 7:

CANCER-ID: Klotten, Vera et al. Multicenter Evaluation of Circulating Plasma MicroRNA Extraction Technologies for the Development of Clinically Feasible Reverse Transcription Quantitative PCR and Next-Generation Sequencing Analytical Work Flows, *CLIN CHEM* 65: 1132-1140

CANCER-ID: Klotten, Vera et al. Circulating Tumor Cell PD-L1 Expression as Biomarker for Therapeutic Efficacy of Immune Checkpoint Inhibition in NSCLC, *CELLS-BASEL* 8:

CANCER-ID: Parikh, Aparna R. et al. Liquid versus tissue biopsy for detecting acquired resistance and tumor heterogeneity in gastrointestinal cancers, *NAT MED* 25: 1415-+

CANCER-ID: Ulz, Peter et al. Inference of transcription factor binding from cell-free DNA enables tumor subtype prediction and early detection, NAT COMMUN 10:

CANCER-ID: Keller, Laura et al. Unravelling tumour heterogeneity by single-cell profiling of circulating tumour cells, NAT REV CANCER 19: 553-567

CANCER-ID: Hofman, P. et al. Liquid biopsy in the era of immuno-oncology: is it ready for prime-time use for cancer patients?, ANN ONCOL 30: 1448-1459

CANCER-ID: Lazzari, Luca et al. Patient-Derived Xenografts and Matched Cell Lines Identify Pharmacogenomic Vulnerabilities in Colorectal Cancer, CLIN CANCER RES 25: 6243-6259

CANCER-ID: Pailler, Emma et al. Acquired Resistance Mutations to ALK Inhibitors Identified by Single Circulating Tumor Cell Sequencing in ALK-Rearranged Non-Small-Cell Lung Cancer, CLIN CANCER RES 25: 6671-6682

CANCER-ID: Tayoun, Tala et al. CTC-Derived Models: A Window into the Seeding Capacity of Circulating Tumor Cells (CTCs), CELLS-BASEL 8:

CANCER-ID: Siravegna, G. et al. How liquid biopsies can change clinical practice in oncology, ANN ONCOL 30: 1580-1590

CANCER-ID: Nanou, Afroditi et al. Tumour-derived extracellular vesicles in blood of metastatic cancer patients associate with overall survival, BRIT J CANCER 122: 801-811

CANCER-ID: Jeannot, Emmanuelle et al. A single droplet digital PCR for ESR1 activating mutations detection in plasma, ONCOGENE 39: 2987-2995

CANCER-ID: Cieslikowski, Wojciech A. et al. Circulating Tumor Cells as a Marker of Disseminated Disease in Patients with Newly Diagnosed High-Risk Prostate Cancer, CANCERS 12:

CANCER-ID: Zhou, Qing et al. Cell-free DNA analysis reveals POLR1D-mediated resistance to bevacizumab in colorectal cancer, GENOME MED 12:

CANCER-ID: Magri, Alessandro et al. High-dose vitamin C enhances cancer immunotherapy, SCI TRANSL MED 12:

CANCER-ID: Silveira, Amanda Bortolini et al. High-Accuracy Determination of Microsatellite Instability Compatible with Liquid Biopsies, CLIN CHEM 66: 606-613

CANCER-ID: Arena, Sabrina et al. A Subset of Colorectal Cancers with Cross-Sensitivity to Olaparib and Oxaliplatin, CLIN CANCER RES 26: 1372-1384

CANCER-ID: Heitzer, Ellen et al. Cell-Free DNA and Apoptosis: How Dead Cells Inform About the Living, TRENDS MOL MED 26: 519-528

CANCER-ID: Koch, Claudia et al. Characterization of circulating breast cancer cells with tumorigenic and metastatic capacity, EMBO MOL MED 12:

CANCER-ID: Weber, Sabrina et al. Technical Evaluation of Commercial Mutation Analysis Platforms and Reference Materials for Liquid Biopsy Profiling, CANCERS 12:

CANCER-ID: Cortes-Hernandez, Luis Enrique et al. Circulating tumor cell as the functional aspect of liquid biopsy to understand the metastatic cascade in solid cancer, MOL ASPECTS MED 72:

CANCER-ID: Valihrach, Lukas et al. Circulating miRNA analysis for cancer diagnostics and therapy, MOL ASPECTS MED 72:

CANCER-ID: Huggett, Jim F. et al. The Digital MIQE Guidelines Update: Minimum Information for Publication of Quantitative Digital PCR Experiments for 2020, CLIN CHEM 66: 1012-1029

CANCER-ID: Mauri, G. et al. The DNA damage response pathway as a land of therapeutic opportunities for colorectal cancer, ANN ONCOL 31: 1135-1147

CANCER-ID: Heidrich, Isabel et al. Liquid biopsies: Potential and challenges, INT J CANCER 148: 528-545

CANCER-ID: Keller, Laura et al. Clinical relevance of blood-based ctDNA analysis: mutation detection and beyond, BRIT J CANCER 124: 345-358

CANCER-ID: Drula, Rares et al. MicroRNAs from Liquid Biopsy Derived Extracellular Vesicles: Recent Advances in Detection and Characterization Methods, CANCERS 12:

CANCER-ID: Keller, Laura et al. Biology and clinical relevance of EpCAM, CELL STRESS 3: 165-180

CANCER-ID: Zavridou, Martha et al. Prognostic Significance of Gene Expression and DNA Methylation Markers in Circulating Tumor Cells and Paired Plasma Derived Exosomes in Metastatic Castration Resistant Prostate Cancer, CANCERS 13:

CANCER-ID: Neves, Rui P. L. et al. Proficiency Testing to Assess Technical Performance for CTC-Processing and Detection Methods in CANCER-ID, CLIN CHEM 67: 631-641

CANCER-ID: Alix-Panabieres, Catherine et al. Liquid Biopsy: From Discovery to Clinical Application, CANCER DISCOV 11: 858-873

CANCER-ID: Hofbauer, Lorenz C. et al. Novel approaches to target the microenvironment of bone metastasis, NAT REV CLIN ONCOL 18: 488-505

CANCER-ID: Smit, Daniel J. et al. Circulating tumor cells as a promising target for individualized drug susceptibility tests in cancer therapy, BIOCHEM PHARMACOL 188:

CARDIATEAM: Simmonds, Steven J. et al. Cellular and Molecular Differences between HFpEF and HFrEF: A Step Ahead in an Improved Pathological Understanding, CELLS-BASEL 9:

CARDIATEAM: Tschoepe, Carsten et al. Myocarditis and inflammatory cardiomyopathy: current evidence and future directions, NAT REV CARDIOL 18: 169-193

CARDIATEAM: de Boer, Rudolf A. et al. Common mechanistic pathways in cancer and heart failure. A scientific roadmap on behalf of the Translational Research Committee of the Heart Failure Association (HFA) of the European Society of Cardiology (ESC), EUR J HEART FAIL 22: 2272-2289

CARDIATEAM: Dia, Maya et al. Reduced reticulum-mitochondria Ca<sup>2+</sup> transfer is an early and reversible trigger of mitochondrial dysfunctions in diabetic cardiomyopathy, BASIC RES CARDIOL 115:

CARDIATEAM: Hazebroek, Mark R. et al. Intravenous immunoglobulin therapy in adult patients with idiopathic chronic cardiomyopathy and cardiac parvovirus B19 persistence: a prospective, double-blind, randomized, placebo-controlled clinical trial, EUR J HEART FAIL 23: 302-309

CARDIATEAM: Verdonschot, Job A. J. et al. Phenotypic clustering of dilated cardiomyopathy patients highlights important pathophysiological differences, EUR HEART J 42:

CARE: Maisonnasse, Pauline et al. Hydroxychloroquine use against SARS-CoV-2 infection in non-human primates, NATURE 585: 584+

CARE: Fenwick, Craig et al. Changes in SARS-CoV-2 Spike versus Nucleoprotein Antibody Responses Impact the Estimates of Infections in Population-Based Seroprevalence Studies, J VIROL 95:

CARE: Laporte, Manon et al. The SARS-CoV-2 and other human coronavirus spike proteins are fine-tuned towards temperature and proteases of the human airways, PLOS PATHOG 17:

CARE: Stevaert, Annelies et al. Betulonic Acid Derivatives Interfering with Human Coronavirus 229E Replication via the nsp15 Endoribonuclease, J MED CHEM 64: 5632-5644

CARE: Lo, Ho Sing et al. Simeprevir Potently Suppresses SARS-CoV-2 Replication and Synergizes with Remdesivir, ACS CENTRAL SCI 7: 792-802

CARE: Fenwick, Craig et al. A high-throughput cell- and virus-free assay shows reduced neutralization of SARS-CoV-2 variants by COVID-19 convalescent plasma, SCI TRANSL MED 13:

CHEM21: Windle, Claire L. et al. Engineering aldolases as biocatalysts, CURR OPIN CHEM BIOL 19: 25-33

CHEM21: Cioc, Razvan C. et al. Multicomponent reactions: advanced tools for sustainable organic synthesis, GREEN CHEM 16: 2958-2975

CHEM21: Prat, Denis et al. A survey of solvent selection guides, GREEN CHEM 16: 4546-4551

CHEM21: Scheller, Philipp N. et al. Enzyme Toolbox: Novel Enantiocomplementary Imine Reductases, CHEMBIOCHEM 15: 2201-2204

CHEM21: Hussain, Shahed et al. An (R)-Imine Reductase Biocatalyst for the Asymmetric Reduction of Cyclic Imines, CHEMCATCHEM 7: 579-583

CHEM21: Harsanyi, Antal et al. Organofluorine chemistry: applications, sources and sustainability, GREEN CHEM 17: 2081-2086

CHEM21: McElroy, C. Robert et al. Towards a holistic approach to metrics for the 21st century pharmaceutical industry, GREEN CHEM 17: 3111-3121

CHEM21: McKenna, Shane M. et al. Enzyme cascade reactions: synthesis of furandicarboxylic acid (FDCA) and carboxylic acids using oxidases in tandem, GREEN CHEM 17: 3271-3275

CHEM21: Reay, Alan J. et al. Unified mild reaction conditions for C2-selective Pd-catalysed tryptophan arylation, including tryptophan-containing peptides, ORG BIOMOL CHEM 13: 8298-8309



CHEM21: Ashcroft, Christopher P. et al. Survey of Solvent Usage in Papers Published in Organic Process Research & Development 1997-2012, *ORG PROCESS RES DEV* 19: 740-747

CHEM21: Reay, Alan J. et al. Catalytic C-H bond functionalisation chemistry: the case for quasi-heterogeneous catalysis, *CHEM COMMUN* 51: 16289-16307

CHEM21: Prat, Denis et al. CHEM21 selection guide of classical- and less classical-solvents, *GREEN CHEM* 18: 288-296

CHEM21: Both, Peter et al. Whole-Cell Biocatalysts for Stereoselective C-H Amination Reactions, *ANGEW CHEM INT EDIT* 55: 1511-1513

CHEM21: Vogl, Thomas et al. A Toolbox of Diverse Promoters Related to Methanol Utilization: Functionally Verified Parts for Heterologous Pathway Expression in *Pichia pastoris*, *ACS SYNTH BIOL* 5: 172-186

CHEM21: van der Heijden, Gydo et al. 2-Bromo-6-isocyanopyridine as a Universal Convertible Isocyanide for Multicomponent Chemistry, *ORG LETT* 18: 984-987

CHEM21: Mampuys, Pieter et al. Iodide-Catalyzed Synthesis of Secondary Thiocarbamates from Isocyanides and Thiosulfonates, *ORG LETT* 18: 2808-2811

CHEM21: Weninger, Astrid et al. Combinatorial optimization of CRISPR/Cas9 expression enables precision genome engineering in the methylotrophic yeast *Pichia pastoris*, *J BIOTECHNOL* 235: 139-149

CHEM21: Harsanyi, Antal et al. One-Step Continuous Flow Synthesis of Antifungal WHO Essential Medicine Flucytosine Using Fluorine, *ORG PROCESS RES DEV* 21: 273-276

CHEM21: Reay, Alan J. et al. Mild and Regioselective Pd(OAc)<sub>2</sub>-Catalyzed C-H Arylation of Tryptophans by [ArN<sub>2</sub>]X, Promoted by Tonic Acid, *ACS CATAL* 7: 5174-5179

CHEM21: Chapman, Michael R. et al. Simple and Versatile Laboratory Scale CSTR for Multiphasic Continuous-Flow Chemistry and Long Residence Times, *ORG PROCESS RES DEV* 21: 1294-1301

CHEM21: Aleku, Godwin A. et al. A reductive aminase from *Aspergillus oryzae*, *NAT CHEM* 9: 961-969

CHEM21: Vogl, Thomas et al. Engineered bidirectional promoters enable rapid multi-gene co-expression optimization, *NAT COMMUN* 9:

CHEM21: Grundtvig, Ines P. Rosinha et al. Screening of organic solvents for bioprocesses using aqueous-organic two-phase systems, *BIOTECHNOL ADV* 36: 1801-1814

CHEM21: Adams, Joseph P. et al. Biocatalysis: A Pharma Perspective, *ADV SYNTH CATAL* 361: 2421-2432

CHEM21: Wiltschi, Birgit et al. Enzymes revolutionize the bioproduction of value-added compounds: From enzyme discovery to special applications, *BIOTECHNOL ADV* 40:

COMBACTE: Schechner, Vered et al. Epidemiological Interpretation of Studies Examining the Effect of Antibiotic Usage on Resistance, *CLIN MICROBIOL REV* 26: 289-307

COMBACTE: Sztajer, Helena et al. Cross-feeding and interkingdom communication in dual-species biofilms of *Streptococcus mutans* and *Candida albicans*, ISME J 8: 2256-2271

COMBACTE: Deng, Zhi-Luo et al. Dysbiosis in chronic periodontitis: Key microbial players and interactions with the human host, SCI REP-UK 7:

COMBACTE: Gottschick, Cornelia et al. The urinary microbiota of men and women and its changes in women during bacterial vaginosis and antibiotic treatment, MICROBIOME 5:

COMBACTE: Lee, Andie S. et al. Methicillin-resistant *Staphylococcus aureus*, NAT REV DIS PRIMERS 4:

COMBACTE: Heilbronner, Simon et al. The microbiome-shaping roles of bacteriocins, NAT REV MICROBIOL 19: 726-739

COMBACTE-CARE: Docobo-Perez, F. et al. Pharmacodynamics of Fosfomycin: Insights into Clinical Use for Antimicrobial Resistance, ANTIMICROB AGENTS CH 59: 5602-5610

COMBACTE-CARE: Gutierrez-Gutierrez, Belen et al. A Predictive Model of Mortality in Patients With Bloodstream Infections due to Carbapenemase-Producing Enterobacteriaceae, MAYO CLIN PROC 91: 1362-1371

COMBACTE-CARE: Raquel Palacios-Baena, Zaira et al. Development and validation of the INCREMENT-ESBL predictive score for mortality in patients with bloodstream infections due to extended-spectrum- beta-lactamase-producing Enterobacteriaceae, J ANTIMICROB CHEMOTH 72: 906-913

COMBACTE-CARE: Gutierrez-Gutierrez, Belen et al. Effect of appropriate combination therapy on mortality of patients with bloodstream infections due to carbapenemase-producing Enterobacteriaceae (INCREMENT): a retrospective cohort study, LANCET INFECT DIS 17: 726-734

COMBACTE-CARE: Grabein, B. et al. Intravenous fosfomycin-back to the future. Systematic review and meta-analysis of the clinical literature, CLIN MICROBIOL INFECTION 23: 363-372

COMBACTE-CARE: Harris, P. N. A. et al. Proposed primary endpoints for use in clinical trials that compare treatment options for bloodstream infection in adults: a consensus definition, CLIN MICROBIOL INFECTION 23: 533-541

COMBACTE-CARE: de Kraker, M. E. A. et al. Good epidemiological practice: a narrative review of appropriate scientific methods to evaluate the impact of antimicrobial stewardship interventions, CLIN MICROBIOL INFECTION 23: 819-825

COMBACTE-CARE: Bassetti, M. et al. Management of KPC-producing *Klebsiella pneumoniae* infections, CLIN MICROBIOL INFECTION 24: 133-144

COMBACTE-CARE: Rodriguez-Bano, Jesus et al. Treatment of Infections Caused by Extended-Spectrum-Beta-Lactamase-, AmpC-, and Carbapenemase-Producing Enterobacteriaceae, CLIN MICROBIOL REV 31:

COMBACTE-CARE: Bluhmki, Tobias et al. A wild bootstrap approach for the Aalen-Johansen estimator, BIOMETRICS 74: 977-985

COMBACTE-CARE: Eliakim-Raz, Noa et al. Risk Factors for Treatment Failure and Mortality Among Hospitalized Patients With Complicated Urinary Tract Infection: A Multicenter Retrospective Cohort Study (RESCUING Study Group), *CLIN INFECT DIS* 68: 29-36

COMBACTE-CARE: Troeman, D. P. R. et al. Antimicrobial approaches in the prevention of *Staphylococcus aureus* infections: a review, *J ANTIMICROB CHEMOTH* 74: 281-294

COMBACTE-CARE: Cornely, Oliver A. et al. Pharmacokinetics and safety of aztreonam/avibactam for the treatment of complicated intra-abdominal infections in hospitalized adults: results from the REJUVENATE study, *J ANTIMICROB CHEMOTH* 75: 618-627

COMBACTE-CARE: Kaier, Klaus et al. Mechanical ventilation and the daily cost of ICU care, *BMC HEALTH SERV RES* 20:

COMBACTE-CARE: Behnke, Michael et al. Information technology aspects of large-scale implementation of automated surveillance of healthcare-associated infections, *CLIN MICROBIOL INFEC* 27: S29-S39

COMBACTE-CDI: Behnke, Michael et al. Information technology aspects of large-scale implementation of automated surveillance of healthcare-associated infections, *CLIN MICROBIOL INFEC* 27: S29-S39

COMBACTE-CDI: Boekhoud, Ilse M. et al. Haem is crucial for medium-dependent metronidazole resistance in clinical isolates of *Clostridioides difficile*, *J ANTIMICROB CHEMOTH* 76: 1731-1740

COMBACTE-MAGNET: Docobo-Perez, F. et al. Pharmacodynamics of Fosfomycin: Insights into Clinical Use for Antimicrobial Resistance, *ANTIMICROB AGENTS CH* 59: 5602-5610

COMBACTE-MAGNET: Perner, Anders et al. Sepsis: frontiers in diagnosis, resuscitation and antibiotic therapy, *INTENS CARE MED* 42: 1958-1969

COMBACTE-MAGNET: Juan, Carlos et al. Host and Pathogen Biomarkers for Severe *Pseudomonas aeruginosa* Infections, *J INFECT DIS* 215: S44-S51

COMBACTE-MAGNET: Hotterbeekx, An et al. In vivo and In vitro Interactions between *Pseudomonas aeruginosa* and *Staphylococcus spp.*, *FRONT CELL INFECT MI* 7:

COMBACTE-MAGNET: Tschudin-Sutter, Sarah et al. Contact Precautions for Preventing Nosocomial Transmission of Extended-Spectrum beta Lactamase-Producing *Escherichia coli*: A Point/Counterpoint Review, *CLIN INFECT DIS* 65: 342-347

COMBACTE-MAGNET: Grabein, B. et al. Intravenous fosfomycin-back to the future. Systematic review and meta-analysis of the clinical literature, *CLIN MICROBIOL INFEC* 23: 363-372

COMBACTE-MAGNET: de Kraker, M. E. A. et al. Good epidemiological practice: a narrative review of appropriate scientific methods to evaluate the impact of antimicrobial stewardship interventions, *CLIN MICROBIOL INFEC* 23: 819-825

COMBACTE-MAGNET: Nunez-Nunez, M. et al. The methodology of surveillance for antimicrobial resistance and healthcare-associated infections in Europe (SUSPIRE): a systematic review of publicly available information, *CLIN MICROBIOL INFEC* 24: 105-109

COMBACTE-MAGNET: Tacconelli, Evelina et al. Surveillance for control of antimicrobial resistance, LANCET INFECT DIS 18: E99-E106

COMBACTE-MAGNET: Lopez-Causape, Carla et al. The Versatile Mutational Resistome of *Pseudomonas aeruginosa*, FRONT MICROBIOL 9:

COMBACTE-MAGNET: Schrijver, R. et al. Review of antimicrobial resistance surveillance programmes in livestock and meat in EU with focus on humans, CLIN MICROBIOL INFECTION 24: 577-590

COMBACTE-MAGNET: Bluhmki, Tobias et al. A wild bootstrap approach for the Aalen-Johansen estimator, BIOMETRICS 74: 977-985

COMBACTE-MAGNET: Eliakim-Raz, Noa et al. Risk Factors for Treatment Failure and Mortality Among Hospitalized Patients With Complicated Urinary Tract Infection: A Multicenter Retrospective Cohort Study (RESCUING Study Group), CLIN INFECT DIS 68: 29-36

COMBACTE-MAGNET: Troeman, D. P. R. et al. Antimicrobial approaches in the prevention of *Staphylococcus aureus* infections: a review, J ANTIMICROB CHEMOTH 74: 281-294

COMBACTE-MAGNET: Horcajada, Juan P. et al. Epidemiology and Treatment of Multidrug-Resistant and Extensively Drug-Resistant *Pseudomonas aeruginosa* Infections, CLIN MICROBIOL REV 32:

COMBACTE-MAGNET: Kaier, Klaus et al. Mechanical ventilation and the daily cost of ICU care, BMC HEALTH SERV RES 20:

COMBACTE-MAGNET: Biddle, Michele S. Y. et al. Attitudes and approaches to patient and public involvement across Europe: A systematic review, HEALTH SOC CARE COMM 29: 18-27

COMBACTE-MAGNET: del Barrio-Tofino, Ester et al. *Pseudomonas aeruginosa* epidemic high-risk clones and their association with horizontally-acquired beta-lactamases: 2020 update, INT J ANTIMICROB AG 56:

COMBACTE-MAGNET: Wheatley, Rachel et al. Rapid evolution and host immunity drive the rise and fall of carbapenem resistance during an acute *Pseudomonas aeruginosa* infection, NAT COMMUN 12:

COMBACTE-MAGNET: Rodriguez-Bano, Jesus et al. Antimicrobial resistance research in a post-pandemic world: Insights on antimicrobial resistance research in the COVID-19 pandemic, J GLOB ANTIMICROB RE 25: 5-7

COMBACTE-MAGNET: Behnke, Michael et al. Information technology aspects of large-scale implementation of automated surveillance of healthcare-associated infections, CLIN MICROBIOL INFECTION 27: S29-S39

COMBACTE-MAGNET: Rodriguez-Bano, Jesus et al. Key considerations on the potential impacts of the COVID-19 pandemic on antimicrobial resistance research and surveillance, TROP SOC TROP MED H 115: 1122-1129

COMBACTE-NET: Tacke, Daniela et al. Primary prophylaxis of invasive fungal infections in patients with haematologic malignancies. 2014 update of the recommendations of the Infectious Diseases Working Party of the German Society for Haematology and Oncology, ANN HEMATOL 93: 1449-1456

COMBACTE-NET: Barbier, Francois et al. Colonization and infection with extended-spectrum beta-lactamase-producing Enterobacteriaceae in ICU patients: what impact on outcomes and carbapenem exposure?, *J ANTIMICROB CHEMOTH* 71: 1088-1097

COMBACTE-NET: Passaro, Leonor et al. Prevention of hospital-acquired pneumonia in non-ventilated adult patients: a narrative review, *ANTIMICROB RESIST IN 5*:

COMBACTE-NET: Israel, Laura et al. Human Adaptive Immunity Rescues an Inborn Error of Innate Immunity, *CELL* 168: 789+

COMBACTE-NET: Grabein, B. et al. Intravenous fosfomycin-back to the future. Systematic review and meta-analysis of the clinical literature, *CLIN MICROBIOL INFEC* 23: 363-372

COMBACTE-NET: Gravestock, Isaac et al. Adaptive power priors with empirical Bayes for clinical trials, *PHARM STAT* 16: 349-360

COMBACTE-NET: de Kraker, M. E. A. et al. Good epidemiological practice: a narrative review of appropriate scientific methods to evaluate the impact of antimicrobial stewardship interventions, *CLIN MICROBIOL INFEC* 23: 819-825

COMBACTE-NET: Bluhmki, Tobias et al. A wild bootstrap approach for the Aalen-Johansen estimator, *BIOMETRICS* 74: 977-985

COMBACTE-NET: Eliakim-Raz, Noa et al. Risk Factors for Treatment Failure and Mortality Among Hospitalized Patients With Complicated Urinary Tract Infection: A Multicenter Retrospective Cohort Study (RESCUING Study Group), *CLIN INFECT DIS* 68: 29-36

COMBACTE-NET: Troeman, D. P. R. et al. Antimicrobial approaches in the prevention of *Staphylococcus aureus* infections: a review, *J ANTIMICROB CHEMOTH* 74: 281-294

COMBACTE-NET: Abbas, M. et al. Impact of participation in a surgical site infection surveillance network: results from a large international cohort study, *J HOSP INFECT* 102: 267-276

COMBACTE-NET: Kaier, Klaus et al. Mechanical ventilation and the daily cost of ICU care, *BMC HEALTH SERV RES* 20:

COMBACTE-NET: Clemen, Ramona et al. Gas Plasma Technology Augments Ovalbumin Immunogenicity and OT-II T Cell Activation Conferring Tumor Protection in Mice, *ADV SCI* 8:

COMBACTE-NET: Wheatley, Rachel et al. Rapid evolution and host immunity drive the rise and fall of carbapenem resistance during an acute *Pseudomonas aeruginosa* infection, *NAT COMMUN* 12:

COMBACTE-NET: Rodriguez-Bano, Jesus et al. Antimicrobial resistance research in a post-pandemic world: Insights on antimicrobial resistance research in the COVID-19 pandemic, *J GLOB ANTIMICROB RE* 25: 5-7

COMBACTE-NET: Behnke, Michael et al. Information technology aspects of large-scale implementation of automated surveillance of healthcare-associated infections, *CLIN MICROBIOL INFEC* 27: S29-S39

COMBACTE-NET: Rodriguez-Bano, Jesus et al. Key considerations on the potential impacts of the COVID-19 pandemic on antimicrobial resistance research and surveillance, T ROY SOC TROP MED H 115: 1122-1129

COMPACT: Carvalho, Cristiane de Souza et al. Carrier interactions with the biological barriers of the lung: Advanced in vitro models and challenges for pulmonary drug delivery, ADV DRUG DELIVER REV 75: 129-140

COMPACT: Verdurmen, Wouter P. R. et al. Efficient cell-specific uptake of binding proteins into the cytoplasm through engineered modular transport systems, J CONTROL RELEASE 200: 13-22

COMPACT: Colombo, Stefano et al. Mechanistic profiling of the siRNA delivery dynamics of lipid-polymer hybrid nanoparticles, J CONTROL RELEASE 201: 22-31

COMPACT: Lorenzer, Cornelia et al. Going beyond the liver: Progress and challenges of targeted delivery of siRNA therapeutics, J CONTROL RELEASE 203: 1-15

COMPACT: Nordin, Joel Z. et al. Ultrafiltration with size-exclusion liquid chromatography for high yield isolation of extracellular vesicles preserving intact biophysical and functional properties, NANOMED-NANOTECHNOL 11: 879-883

COMPACT: Heldring, Nina et al. Therapeutic Potential of Multipotent Mesenchymal Stromal Cells and Their Extracellular Vesicles, HUM GENE THER 26: 506-517

COMPACT: Laechelt, Ulrich et al. Nucleic Acid Therapeutics Using Polyplexes: A Journey of 50 Years (and Beyond), CHEM REV 115: 11043-11078

COMPACT: Wiklander, Oscar P. B. et al. Extracellular vesicle in vivo biodistribution is determined by cell source, route of administration and targeting, J EXTRACELL VESICLES 4:

COMPACT: Willms, Eduard et al. Cells release subpopulations of exosomes with distinct molecular and biological properties, SCI REP-UK 6:

COMPACT: Kristensen, Mie et al. Applications and Challenges for Use of Cell-Penetrating Peptides as Delivery Vectors for Peptide and Protein Cargos, INT J MOL SCI 17:

COMPACT: Kuehn, Anna et al. Human Alveolar Epithelial Cells Expressing Tight Junctions to Model the Air-Blood Barrier, ALTEX-ALTERN ANIM EX 33: 251-260

COMPACT: Mager, Imre et al. Targeting blood-brain-barrier transcytosis - perspectives for drug delivery, NEUROPHARMACOLOGY 120: 4-7

COMPACT: Oswald, Mira et al. Targeting the Central Nervous System (CNS): A Review of Rabies Virus-Targeting Strategies, MOL PHARMACEUT 14: 2177-2196

COMPACT: OLoughlin, Aisling J. et al. Functional Delivery of Lipid-Conjugated siRNA by Extracellular Vesicles, MOL THER 25: 1580-1587

COMPACT: Dowaidar, Moataz et al. Graphene oxide nanosheets in complex with cell penetrating peptides for oligonucleotides delivery, BBA-GEN SUBJECTS 1861: 2334-2341

COMPACT: Dowaidar, Moataz et al. Magnetic Nanoparticle Assisted Self-assembly of Cell Penetrating Peptides-Oligonucleotides Complexes for Gene Delivery, *SCI REP-UK* 7:

COMPACT: Birch, Ditlev et al. Fluorophore labeling of a cell-penetrating peptide induces differential effects on its cellular distribution and affects cell viability, *BBA-BIOMEMBRANES* 1859: 2483-2494

COMPACT: de Groot, Anne Marit et al. Hollow microneedle-mediated intradermal delivery of model vaccine antigen-loaded PLGA nanoparticles elicits protective T cell-mediated immunity to an intracellular bacterium, *J CONTROL RELEASE* 266: 27-35

COMPACT: Du, Guangsheng et al. Intradermal vaccination with hollow microneedles: A comparative study of various protein antigen and adjuvant encapsulated nanoparticles, *J CONTROL RELEASE* 266: 109-118

COMPACT: Vermeulen, Lotte M. P. et al. Endosomal Size and Membrane Leakiness Influence Proton Sponge-Based Rupture of Endosomal Vesicles, *ACS NANO* 12: 2332-2345

COMPACT: Kletting, Stephanie et al. Co-Culture of Human Alveolar Epithelial (hAELVi) and Macrophage (THP-1) Cell Lines, *ALTEX-ALTERN ANIM EX* 35: 211-222

COMPACT: Sanduleanu, Sebastian et al. Tracking tumor biology with radiomics: A systematic review utilizing a radiomics quality score, *RADIOTHER ONCOL* 127: 349-360

COMPACT: Monkare, Juha et al. Development of PLGA nanoparticle loaded dissolving microneedles and comparison with hollow microneedles in intradermal vaccine delivery, *EUR J PHARM BIOPHARM* 129: 111-121

COMPACT: Sork, Helena et al. Heterogeneity and interplay of the extracellular vesicle small RNA transcriptome and proteome, *SCI REP-UK* 8:

COMPACT: Abdelhamid, Hani Nasser et al. Gene delivery using cell penetrating peptides-zeolitic imidazolate frameworks, *MICROPOR MESOPOR MAT* 300:

COVID-RED: Schaffner, Anna et al. Characterization of a Pan-Immunoglobulin Assay Quantifying Antibodies Directed against the Receptor Binding Domain of the SARS-CoV-2 S1-Subunit of the Spike Protein: A Population-Based Study, *J CLIN MED* 9:

DDMoRe: Nielsen, Elisabet I. et al. Pharmacokinetic-Pharmacodynamic Modeling of Antibacterial Drugs, *PHARMACOL REV* 65: 1053-1090

DDMoRe: Buechel, Finja et al. Path2Models: large-scale generation of computational models from biochemical pathway maps, *BMC SYST BIOL* 7:

DDMoRe: Delattre, Maud et al. A note on BIC in mixed-effects models, *ELECTRON J STAT* 8: 456-475

DDMoRe: Bender, Brendan C. et al. Population pharmacokinetic-pharmacodynamic modelling in oncology: a tool for predicting clinical response, *BRIT J CLIN PHARMACO* 79: 56-71

DDMoRe: Bergmann, Frank T. et al. COMBINE archive and OMEX format: one file to share all information to reproduce a modeling project, *BMC BIOINFORMATICS* 15:

DDMoRe: Chelliah, Vijayalakshmi et al. BioModels: ten-year anniversary, NUCLEIC ACIDS RES 43: D542-D548

DDMoRe: Dosne, Anne-Gaelle et al. Improving the estimation of parameter uncertainty distributions in nonlinear mixed effects models using sampling importance resampling, J PHARMACOKINET PHAR 43: 583-596

DDMoRe: McMurry, Julie A. et al. Identifiers for the 21st century: How to design, provision, and reuse persistent identifiers to maximize utility and impact of life science data, PLOS BIOL 15:

DDMoRe: Dosne, Anne-Gaelle et al. An automated sampling importance resampling procedure for estimating parameter uncertainty, J PHARMACOKINET PHAR 44: 509-520

DDMoRe: Malik-Sheriff, Rahuman S. et al. BioModels-15 years of sharing computational models in life science, NUCLEIC ACIDS RES 48: D407-D415

DDMoRe: Keating, Sarah M. et al. SBML Level 3: an extensible format for the exchange and reuse of biological models, MOL SYST BIOL 16:

DDMoRe: Busse, David et al. Which Analysis Approach Is Adequate to Leverage Clinical Microdialysis Data? A Quantitative Comparison to Investigate Exposure and Reponse Exemplified by Levofloxacin, PHARM RES-DORDR 38: 381-395

DECISION: Colavita, Francesca et al. COVID-19 Rapid Antigen Test as Screening Strategy at Points of Entry: Experience in Lazio Region, Central Italy, August-October 2020, BIOMOLECULES 11:

DECISION: Amendola, Alessandra et al. Saliva Is a Valid Alternative to Nasopharyngeal Swab in Chemiluminescence-Based Assay for Detection of SARS-CoV-2 Antigen, J CLIN MED 10:

DIRECT: Ahmad, Shafqat et al. Gene x Physical Activity Interactions in Obesity: Combined Analysis of 111,421 Individuals of European Ancestry, PLOS GENET 9:

DIRECT: Nica, Alexandra C. et al. Cell-type, allelic, and genetic signatures in the human pancreatic beta cell transcriptome, GENOME RES 23: 1554-1562

DIRECT: Pasquali, Lorenzo et al. Pancreatic islet enhancer clusters enriched in type 2 diabetes risk-associated variants, NAT GENET 46: 136-+

DIRECT: Breier, Michaela et al. Targeted Metabolomics Identifies Reliable and Stable Metabolites in Human Serum and Plasma Samples, PLOS ONE 9:

DIRECT: Pedersen, Helle Krogh et al. Human gut microbes impact host serum metabolome and insulin sensitivity, NATURE 535: 376-+

DIRECT: Franks, Paul W. et al. Exposing the exposures responsible for type 2 diabetes and obesity, SCIENCE 354: 69-73

DIRECT: McCarthy, Mark I. et al. Painting a new picture of personalised medicine for diabetes, DIABETOLOGIA 60: 793-799

DIRECT: Hocher, Berthold et al. Metabolomics for clinical use and research in chronic kidney disease, NAT REV NEPHROL 13: 269-284



DIRECT: Dujic, T. et al. Variants in Pharmacokinetic Transporters and Glycemic Response to Metformin: A MetGen Meta-Analysis, CLIN PHARMACOL THER 101: 763-772

DIRECT: Wood, Andrew R. et al. A Genome-Wide Association Study of IVGTT-Based Measures of First-Phase Insulin Secretion Refines the Underlying Physiology of Type 2 Diabetes Variants, DIABETES 66: 2296-2309

DIRECT: Haid, Mark et al. Long-Term Stability of Human Plasma Metabolites during Storage at -80 degrees C, J PROTEOME RES 17: 203-211

DIRECT: Allin, Kristine H. et al. Aberrant intestinal microbiota in individuals with prediabetes, DIABETOLOGIA 61: 810-820

DIRECT: Pedersen, Helle Krogh et al. A computational framework to integrate high-throughput -omics datasets for the identification of potential mechanistic links, NAT PROTOC 13: 2781-2800

DIRECT: Ji, Yingjie et al. Genome-Wide and Abdominal MRI Data Provide Evidence That a Genetically Determined Favorable Adiposity Phenotype Is Characterized by Lower Ectopic Liver Fat and Lower Risk of Type 2 Diabetes, Heart Disease, and Hypertension, DIABETES 68: 207-219

DIRECT: Pearson, Ewan R. et al. Type 2 diabetes: a multifaceted disease, DIABETOLOGIA 62: 1107-1112

DIRECT: Chung, Wendy K. et al. Precision medicine in diabetes: a Consensus Report from the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETOLOGIA 63: 1671-1693

DIRECT: Chung, Wendy K. et al. Precision Medicine in Diabetes: A Consensus Report From the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETES CARE 43: 1617-1635

DIRECT: Atabaki-Pasdar, Naeimeh et al. Predicting and elucidating the etiology of fatty liver disease: A machine learning modeling and validation study in the IMI DIRECT cohorts, PLOS MED 17:

DIRECT: Suhre, Karsten et al. Genetics meets proteomics: perspectives for large population-based studies, NAT REV GENET 22: 19-37

DIRECT: Aguet, Francois et al. The GTEx Consortium atlas of genetic regulatory effects across human tissues, SCIENCE 369: 1318-1330

DIRECT: Oliva, Meritxell et al. The impact of sex on gene expression across human tissues, SCIENCE 369: 1331-+

DIRECT: Demanelis, Kathryn et al. Determinants of telomere length across human tissues, SCIENCE 369: 1333-+

DIRECT: Ferraro, Nicole M. et al. Transcriptomic signatures across human tissues identify functional rare genetic variation, SCIENCE 369: 1334-+

DIRECT: Pividori, Milton et al. PhenomeXcan: Mapping the genome to the phenome through the transcriptome, SCI ADV 6:

DIRECT: Vinuela, Ana et al. Genetic variant effects on gene expression in human pancreatic islets and their implications for T2D, NAT COMMUN 11:

DIRECT: Bar, Noam et al. A reference map of potential determinants for the human serum metabolome, NATURE 588: 135-140

DIRECT: Bizzotto, Roberto et al. Processes Underlying Glycemic Deterioration in Type 2 Diabetes: An IMI DIRECT Study, DIABETES CARE 44: 511-518

DO->IT: Kalkman, Shona et al. Responsible data sharing in international health research: a systematic review of principles and norms, BMC MED ETHICS 20:

DRAGON: Hu, Shaoping et al. Weakly Supervised Deep Learning for COVID-19 Infection Detection and Classification From CT Images, IEEE ACCESS 8: 118869-118883

DRAGON: Wang, Chengjia et al. DiCyc: GAN-based deformation invariant cross-domain information fusion for medical image synthesis, INFORM FUSION 67: 147-160

DRAGON: Lv, Jun et al. PIC-GAN: A Parallel Imaging Coupled Generative Adversarial Network for Accelerated Multi-Channel MRI Reconstruction, DIAGNOSTICS 11:

DRAGON: Guiot, Julien et al. Development and Validation of an Automated Radiomic CT Signature for Detecting COVID-19, DIAGNOSTICS 11:

DRAGON: Wu, Yinzhe et al. Fast and Automated Segmentation for the Three-Directional Multi-Slice Cine Myocardial Velocity Mapping, DIAGNOSTICS 11:

DRAGON: Zhang, Weiwei et al. Multi-task learning with Multi-view Weighted Fusion Attention for artery-specific calcification analysis, INFORM FUSION 71: 64-76

DRAGON: Lv, Jun et al. Which GAN? A comparative study of generative adversarial network-based fast MRI reconstruction, PHILOS T R SOC A 379:

DRAGON: Ma, Huijing et al. Can Clinical Symptoms and Laboratory Results Predict CT Abnormality? Initial Findings Using Novel Machine Learning Techniques in Children With COVID-19 Infections, FRONT MED-LAUSANNE 8:

DRAGON: Yang, Guang et al. Unbox the black-box for the medical explainable AI via multi-modal and multi-centre data fusion: A mini-review, two showcases and beyond, INFORM FUSION 77: 29-52

DRIVE: Alvarez-Uria, Gerardo et al. Global forecast of antimicrobial resistance in invasive isolates of Escherichia coli and Klebsiella pneumoniae, INT J INFECT DIS 68: 50-53

DRIVE: Panatto, Donatella et al. Influenza Vaccination in Italian Healthcare Workers (2018-2019 Season): Strengths and Weaknesses. Results of a Cohort Study in Two Large Italian Hospitals, VACCINES-BASEL 8:

DRIVE-AB: Harbarth, S. et al. Antibiotic research and development: business as usual?, J ANTIMICROB CHEMOTH 70: 1604-1607

DRIVE-AB: Teillant, Aude et al. Potential burden of antibiotic resistance on surgery and cancer chemotherapy antibiotic prophylaxis in the USA: a literature review and modelling study, *LANCET INFECT DIS* 15: 1429-1437

DRIVE-AB: Friedman, N. D. et al. The negative impact of antibiotic resistance, *CLIN MICROBIOL INFECC* 22: 416-422

DRIVE-AB: Tacconelli, Evelina et al. STROBE-AMS: recommendations to optimise reporting of epidemiological studies on antimicrobial resistance and informing improvement in antimicrobial stewardship, *BMJ OPEN* 6:

DRIVE-AB: Deak, Dalia et al. Progress in the Fight Against Multidrug-Resistant Bacteria? A Review of US Food and Drug Administration-Approved Antibiotics, 2010-2015, *ANN INTERN MED* 165: 363-+

DRIVE-AB: de Kraker, M. E. A. et al. Good epidemiological practice: a narrative review of appropriate scientific methods to evaluate the impact of antimicrobial stewardship interventions, *CLIN MICROBIOL INFECC* 23: 819-825

DRIVE-AB: Muller, Anouk E. et al. Therapeutic Drug Monitoring of Beta-Lactams and Other Antibiotics in the Intensive Care Unit: Which Agents, Which Patients and Which Infections?, *DRUGS* 78: 439-451

DRIVE-AB: Alvarez-Uria, Gerardo et al. Global forecast of antimicrobial resistance in invasive isolates of *Escherichia coli* and *Klebsiella pneumoniae*, *INT J INFECT DIS* 68: 50-53

DRIVE-AB: Zanichelli, Veronica et al. Variation in antibiotic use among and within different settings: a systematic review, *J ANTIMICROB CHEMOTH* 73: 17-29

DRIVE-AB: Le Marechal, Marion et al. Quality indicators assessing antibiotic use in the outpatient setting: a systematic review followed by an international multidisciplinary consensus procedure, *J ANTIMICROB CHEMOTH* 73: 40-49

DRIVE-AB: Benic, Mirjana Stanic et al. Metrics for quantifying antibiotic use in the hospital setting: results from a systematic review and international multidisciplinary consensus procedure, *J ANTIMICROB CHEMOTH* 73: 50-58

DRIVE-AB: Savic, Miloje et al. A Grant Framework as a Push Incentive to Stimulate Research and Development of New Antibiotics, *J LAW MED ETHICS* 46: 9-24

DRIVE-AB: Baraldi, Enrico et al. Antibiotic Pipeline Coordinators, *J LAW MED ETHICS* 46: 25-31

DRIVE-AB: Okhravi, Christopher et al. Simulating Market Entry Rewards for Antibiotics Development, *J LAW MED ETHICS* 46: 32-42

DRIVE-AB: Temkin, Elizabeth et al. Estimating the number of infections caused by antibiotic-resistant *Escherichia coli* and *Klebsiella pneumoniae* in 2014: a modelling study, *LANCET GLOB HEALTH* 6: E969-E979

DRIVE-AB: Zanichelli, V. et al. Patient-related determinants of antibiotic use: a systematic review, *CLIN MICROBIOL INFECC* 25: 48-53

DRIVE-AB: Huttner, Benedikt et al. How to improve antibiotic awareness campaigns: findings of a WHO global survey, *BMJ GLOB HEALTH* 4:

EBiSC: Zerbino, Daniel R. et al. Ensembl 2018, *NUCLEIC ACIDS RES* 46: D754-D761

EBiSC: Maffioletti, Sara Martina et al. Three-Dimensional Human iPSC-Derived Artificial Skeletal Muscles Model Muscular Dystrophies and Enable Multilineage Tissue Engineering, *CELL REP* 23: 899-908

EBiSC: Cunningham, Fiona et al. Ensembl 2019, *NUCLEIC ACIDS RES* 47: D745-D751

EBiSC: Laugsch, Magdalena et al. Modeling the Pathological Long-Range Regulatory Effects of Human Structural Variation with Patient-Specific hiPSCs, *CELL STEM CELL* 24: 736-+

EBiSC: Hasselmann, Jonathan et al. Development of a Chimeric Model to Study and Manipulate Human Microglia In Vivo, *NEURON* 103: 1016-+

EBODAC: Anywaine, Zacchaeus et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccination Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Uganda and Tanzania, *J INFECT DIS* 220: 46-56

EBODAC: Mutua, Gaudensia et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccine Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Nairobi, Kenya, *J INFECT DIS* 220: 57-67

EbolaMoDRAD: Guedj, Jeremie et al. Antiviral efficacy of favipiravir against Ebola virus: A translational study in cynomolgus macaques, *PLOS MED* 15:

EbolaMoDRAD: Forbes, Kristian M. et al. Bombali Virus in Mops condylurus Bat, Kenya, *EMERG INFECT DIS* 25: 955-957

EbolaMoDRAD: Ciftci, Sibel et al. Digital Rolling Circle Amplification-Based Detection of Ebola and Other Tropical Viruses, *J MOL DIAGN* 22: 272-283

EbolaMoDRAD: Ciftci, Sibel et al. The sweet detection of rolling circle amplification: Glucose-based electrochemical genosensor for the detection of viral nucleic acid, *BIOSENS BIOELECTRON* 151:

EBOMAN: Anywaine, Zacchaeus et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccination Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Uganda and Tanzania, *J INFECT DIS* 220: 46-56

EBOMAN: Mutua, Gaudensia et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccine Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Nairobi, Kenya, *J INFECT DIS* 220: 57-67

EBOVAC: Huttner, Angela et al. Determinants of antibody persistence across doses and continents after single-dose rVSV-ZEBOV vaccination for Ebola virus disease: an observational cohort study, *LANCET INFECT DIS* 18: 738-748

EBOVAC1: Kucharski, Adam J. et al. Effectiveness of Ring Vaccination as Control Strategy for Ebola Virus Disease, *EMERG INFECT DIS* 22: 105-108

EBOVAC1: Milligan, Iain D. et al. Safety and Immunogenicity of Novel Adenovirus Type 26-and Modified Vaccinia Ankara-Vectored Ebola Vaccines A Randomized Clinical Trial, JAMA-J AM MED ASSOC 315: 1610-1623

EBOVAC1: Enria, Luisa et al. Power, fairness and trust: understanding and engaging with vaccine trial participants and communities in the setting up the EBOVAC-Salone vaccine trial in Sierra Leone, BMC PUBLIC HEALTH 16:

EBOVAC1: Funk, Sebastian et al. Comparative Analysis of Dengue and Zika Outbreaks Reveals Differences by Setting and Virus, PLOS NEGLECT TROP D 10:

EBOVAC1: Sissoko, Daouda et al. Persistence and clearance of Ebola virus RNA from seminal fluid of Ebola virus disease survivors: a longitudinal analysis and modelling study, LANCET GLOB HEALTH 5: E80-E88

EBOVAC1: Funk, Sebastian et al. Real-time forecasting of infectious disease dynamics with a stochastic semi-mechanistic model, EPIDEMICS-NETH 22: 56-61

EBOVAC1: Funk, Sebastian et al. Assessing the performance of real-time epidemic forecasts: A case study of Ebola in the Western Area region of Sierra Leone, 2014-15, PLOS COMPUT BIOL 15:

EBOVAC1: Anywaine, Zacchaeus et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccination Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Uganda and Tanzania, J INFECT DIS 220: 46-56

EBOVAC1: Mutua, Gaudensia et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccine Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Nairobi, Kenya, J INFECT DIS 220: 57-67

EBOVAC2: Rechten, Anne et al. Systems Vaccinology Identifies an Early Innate Immune Signature as a Correlate of Antibody Responses to the Ebola Vaccine rVSV-ZEBOV, CELL REP 20: 2251-2261

EBOVAC2: Goodier, Martin R. et al. CMV and natural killer cells: shaping the response to vaccination, EUR J IMMUNOL 48: 50-65

EBOVAC2: Anywaine, Zacchaeus et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccination Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Uganda and Tanzania, J INFECT DIS 220: 46-56

EBOVAC2: Mutua, Gaudensia et al. Safety and Immunogenicity of a 2-Dose Heterologous Vaccine Regimen With Ad26.ZEBOV and MVA-BN-Filo Ebola Vaccines: 12-Month Data From a Phase 1 Randomized Clinical Trial in Nairobi, Kenya, J INFECT DIS 220: 57-67

EBOVAC2: Pollard, Andrew J. et al. Safety and immunogenicity of a two-dose heterologous Ad26.ZEBOV and MVA-BN-Filo Ebola vaccine regimen in adults in Europe (EBOVAC2): a randomised, observer-blind, participant-blind, placebo-controlled, phase 2 trial, LANCET INFECT DIS 21: 493-506

EHDEN: Burn, Edward et al. Deep phenotyping of 34,128 adult patients hospitalised with COVID-19 in an international network study, NAT COMMUN 11:

EHDEN: Lane, Jennifer C. E. et al. Risk of hydroxychloroquine alone and in combination with azithromycin in the treatment of rheumatoid arthritis: a multinational, retrospective study, LANCET RHEUMATOL 2: E698-E711

EHDEN: Markus, Aniek F. et al. The role of explainability in creating trustworthy artificial intelligence for health care: A comprehensive survey of the terminology, design choices, and evaluation strategies, J BIOMED INFORM 113:

EHDEN: Burn, Edward et al. The natural history of symptomatic COVID-19 during the first wave in Catalonia, NAT COMMUN 12:

EHDEN: Li, Xintong et al. Characterising the background incidence rates of adverse events of special interest for covid-19 vaccines in eight countries: multinational network cohort study, BMJ-BRIT MED J 373:

EHDEN: Lane, Jennifer C. E. et al. Risk of depression, suicide and psychosis with hydroxychloroquine treatment for rheumatoid arthritis: a multinational network cohort study, RHEUMATOLOGY 60: 3222-3234

EHDEN: Tan, Eng Hooi et al. COVID-19 in patients with autoimmune diseases: characteristics and outcomes in a multinational network of cohorts across three countries, RHEUMATOLOGY 60: SI37-SI50

EHR4CR: Coorevits, P. et al. Electronic health records: new opportunities for clinical research, J INTERN MED 274: 547-560

EHR4CR: De Moor, Georges et al. Using electronic health records for clinical research: The case of the EHR4CR project, J BIOMED INFORM 53: 162-173

EHR4CR: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), GENOME MED 12:

ELF: Besnard, Jeremy et al. The Joint European Compound Library: boosting precompetitive research, DRUG DISCOV TODAY 20: 181-186

ELF: Picazo, Edwige et al. Small molecule inhibitors of ebola virus infection, DRUG DISCOV TODAY 20: 277-286

ELF: Neochoritis, Constantinos G. et al. Efficient Isocyanide-less Isocyanide-Based Multicomponent Reactions, ORG LETT 17: 2002-2005

ELF: Liao, George P. et al. Versatile Multicomponent Reaction Macrocyclic Synthesis Using alpha-Isocyano-omega-carboxylic Acids, ORG LETT 17: 4980-4983

ELF: Zarganes-Tzitzikas, Tryfon et al. Multicomponent Reactions, Union of MCRs and Beyond, CHEM REC 15: 981-996

ELF: Zarganes-Tzitzikas, Tryfon et al. Modern multicomponent reactions for better drug syntheses, ORG CHEM FRONT 1: 834-U178

ELF: Karawajczyk, Anna et al. Expansion of chemical space for collaborative lead generation and drug discovery: the European Lead Factory Perspective, DRUG DISCOV TODAY 20: 1310-1316

ELF: Garcia-Castro, Miguel et al. Scaffold Diversity Synthesis and Its Application in Probe and Drug Discovery, *ANGEW CHEM INT EDIT* 55: 7586-7605

ELF: Mueller, Gerhard et al. Charting Biologically Relevant Spirocyclic Compound Space, *CHEM-EUR J* 23: 703-710

ELF: Zak, Krzysztof M. et al. Structural Biology of the Immune Checkpoint Receptor PD-1 and Its Ligands PD-L1/PD-L2, *STRUCTURE* 25: 1163-1174

ELF: Konstantinidou, Markella et al. Immune Checkpoint PD-1/PD-L1: Is There Life Beyond Antibodies?, *ANGEW CHEM INT EDIT* 57: 4840-4848

ELF: Shaabani, Shabnam et al. A patent review on PD-1/PD-L1 antagonists: small molecules, peptides, and macrocycles (2015-2018), *EXPERT OPIN THER PAT* 28: 665-678

ELF: Krajnc, Alen et al. Will morphing boron-based inhibitors beat the beta-lactamases?, *CURR OPIN CHEM BIOL* 50: 101-110

ELF: Krajnc, Alen et al. Bicyclic Boronate VNRX-5133 Inhibits Metallo- and Serine-beta-Lactamases, *J MED CHEM* 62: 8544-8556

ELF: Mock, Elliot D. et al. Discovery of a NAPE-PLD inhibitor that modulates emotional behavior in mice, *NAT CHEM BIOL* 16: 667-+

ELF: Domling, Alexander et al. Chemistry and Biology of SARS-CoV-2, *CHEM-US* 6: 1283-1295

ELF: Tselepis, Lucas et al. In vitro efficacy of imipenem-relebactam and cefepime-AAI101 against a global collection of ESBL-positive and carbapenemase-producing Enterobacteriaceae, *INT J ANTIMICROB AG* 56:

ELF: Sutanto, Fandi et al. Covalent inhibitors: a rational approach to drug discovery, *RSC MED CHEM* 11: 876-884

EMIF: Oresic, Matej et al. Prediction of non-alcoholic fatty-liver disease and liver fat content by serum molecular lipids, *DIABETOLOGIA* 56: 2266-2274

EMIF: Vos, Stephanie J. B. et al. Preclinical Alzheimers disease and its outcome: a longitudinal cohort study, *LANCET NEUROL* 12: 957-965

EMIF: Hyysalo, Jenni et al. A population-based study on the prevalence of NASH using scores validated against liver histology, *J HEPATOL* 60: 839-846

EMIF: Payne, Felicity et al. Mutations disrupting the Kennedy phosphatidylcholine pathway in humans with congenital lipodystrophy and fatty liver disease, *P NATL ACAD SCI USA* 111: 8901-8906

EMIF: Payne, Felicity et al. Hypomorphism in human NSMCE2 linked to primordial dwarfism and insulin resistance, *J CLIN INVEST* 124: 4028-4038

EMIF: Sattar, Naveed et al. Type 2 diabetes as a disease of ectopic fat?, *BMC MED* 12:

EMIF: Van der Musselle, Stefan et al. Depression in Mild Cognitive Impairment is associated with Progression to Alzheimers Disease: A Longitudinal Study, *J ALZHEIMERS DIS* 42: 1239-1250

EMIF: Hye, Abdul et al. Plasma proteins predict conversion to dementia from prodromal disease, ALZHEIMERS DEMENT 10: 799-807

EMIF: Swerdlow, Daniel I. et al. HMG-coenzyme A reductase inhibition, type 2 diabetes, and bodyweight: evidence from genetic analysis and randomised trials, LANCET 385: 351-361

EMIF: Zhou, You et al. Circulating triacylglycerol signatures and insulin sensitivity in NAFLD associated with the E167K variant in TM6SF2, J HEPATOL 62: 657-663

EMIF: Struyfs, Hanne et al. Diagnostic Accuracy of Cerebrospinal Fluid Amyloid-beta Isoforms for Early and Differential Dementia Diagnosis, J ALZHEIMERS DIS 45: 813-822

EMIF: Vos, Stephanie J. B. et al. Prevalence and prognosis of Alzheimers disease at the mild cognitive impairment stage, BRAIN 138: 1327-1338

EMIF: Le Bastard, Nathalie et al. Importance and Impact of Preanalytical Variables on Alzheimer Disease Biomarker Concentrations in Cerebrospinal Fluid, CLIN CHEM 61: 734-743

EMIF: Jansen, Willemijn J. et al. Prevalence of Cerebral Amyloid Pathology in Persons Without Dementia A Meta-analysis, JAMA-J AM MED ASSOC 313: 1924-1938

EMIF: Ossenkoppele, Rik et al. Prevalence of Amyloid PET Positivity in Dementia Syndromes A Meta-analysis, JAMA-J AM MED ASSOC 313: 1939-1949

EMIF: Dahlman, I. et al. The fat cell epigenetic signature in post-obese women is characterized by global hypomethylation and differential DNA methylation of adipogenesis genes, INT J OBESITY 39: 910-919

EMIF: Ostergaard, Soren D. et al. Associations between Potentially Modifiable Risk Factors and Alzheimer Disease: A Mendelian Randomization Study, PLOS MED 12:

EMIF: Tang, Eugene Y. H. et al. Current Developments in Dementia Risk Prediction Modelling: An Updated Systematic Review, PLOS ONE 10:

EMIF: Sood, Sanjana et al. A novel multi-tissue RNA diagnostic of healthy ageing relates to cognitive health status, GENOME BIOL 16:

EMIF: Toledo, Jon B. et al. Alzheimers disease cerebrospinal fluid biomarker in cognitively normal subjects, BRAIN 138: 2701-2715

EMIF: Skillback, Tobias et al. Cerebrospinal fluid tau and amyloid-beta(1-42) in patients with dementia, BRAIN 138: 2716-2731

EMIF: Gutierrez-Sacristan, Alba et al. PsyGeNET: a knowledge platform on psychiatric disorders and their genes, BIOINFORMATICS 31: 3075-3077

EMIF: Nead, Kevin T. et al. Evidence of a Causal Association Between Insulinemia and Endometrial Cancer: A Mendelian Randomization Analysis, JNCI-J NATL CANCER I 107:

EMIF: Brookes, Anthony J. et al. Human genotype-phenotype databases: aims, challenges and opportunities, NAT REV GENET 16: 702-715



EMIF: Yki-Jarvinen, Hannele et al. Nutritional Modulation of Non-Alcoholic Fatty Liver Disease and Insulin Resistance, *NUTRIENTS* 7: 9127-9138

EMIF: Slegers, Kristel et al. A 22-single nucleotide polymorphism Alzheimers disease risk score correlates with family history, onset age, and cerebrospinal fluid A beta(42), *ALZHEIMERS DEMENT* 11: 1452-1460

EMIF: De Vos, Ann et al. C-terminal neurogranin is increased in cerebrospinal fluid but unchanged in plasma in Alzheimers disease, *ALZHEIMERS DEMENT* 11: 1461-1469

EMIF: Hellwig, Konstantin et al. Neurogranin and YKL-40: independent markers of synaptic degeneration and neuroinflammation in Alzheimers disease, *ALZHEIMERS RES THER* 7:

EMIF: Sattar, Naveed et al. Type 2 diabetes in migrant south Asians: mechanisms, mitigation, and management, *LANCET DIABETES ENDO* 3: 1004-1016

EMIF: Hellmuth, Christian et al. Tyrosine Is Associated with Insulin Resistance in Longitudinal Metabolomic Profiling of Obese Children, *J DIABETES RES* 2016:

EMIF: Jack, Clifford R., Jr. et al. Suspected non-Alzheimer disease pathophysiology - concept and controversy, *NAT REV NEUROL* 12: 117-124

EMIF: Hyotylainen, Tuulia et al. Genome-scale study reveals reduced metabolic adaptability in patients with non-alcoholic fatty liver disease, *NAT COMMUN* 7:

EMIF: Rowe, Emily R. et al. Conserved Amphipathic Helices Mediate Lipid Droplet Targeting of Perilipins 1-3, *J BIOL CHEM* 291: 6664-6678

EMIF: Luukkonen, Panu K. et al. Hepatic ceramides dissociate steatosis and insulin resistance in patients with non-alcoholic fatty liver disease, *J HEPATOL* 64: 1167-1175

EMIF: Van Cauwenberghe, Caroline et al. The genetic landscape of Alzheimer disease: clinical implications and perspectives, *GENET MED* 18: 421-430

EMIF: Suarez-Calvet, Marc et al. sTREM2 cerebrospinal fluid levels are a potential biomarker for microglia activity in early-stage Alzheimers disease and associate with neuronal injury markers, *EMBO MOL MED* 8: 466-476

EMIF: Cuyvers, Elise et al. Genetic variations underlying Alzheimers disease: evidence from genome-wide association studies and beyond, *LANCET NEUROL* 15: 857-868

EMIF: Yki-Jarvinen, Hannele et al. Diagnosis of non-alcoholic fatty liver disease (NAFLD), *DIABETOLOGIA* 59: 1104-1111

EMIF: Vos, Stephanie J. B. et al. NIA-AA staging of preclinical Alzheimer disease: discordance and concordance of CSF and imaging biomarkers, *NEUROBIOL AGING* 44: 1-8

EMIF: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, *GENOME MED* 8:

EMIF: Loomis, A. Katrina et al. Body Mass Index and Risk of Nonalcoholic Fatty Liver Disease: Two Electronic Health Record Prospective Studies, *J CLIN ENDOCR METAB* 101: 945-952

EMIF: Lee, Sunjae et al. Integrated Network Analysis Reveals an Association between Plasma Mannose Levels and Insulin Resistance, *CELL METAB* 24: 172-184

EMIF: Lallukka, S. et al. Non-alcoholic fatty liver disease and risk of type 2 diabetes, *BEST PRACT RES CL EN* 30: 385-395

EMIF: Queralt-Rosinach, Nuria et al. DisGeNET-RDF: harnessing the innovative power of the Semantic Web to explore the genetic basis of diseases, *BIOINFORMATICS* 32: 2236-2238

EMIF: Pini, Lorenzo et al. Brain atrophy in Alzheimers Disease and aging, *AGEING RES REV* 30: 25-48

EMIF: Zhou, You et al. Noninvasive Detection of Nonalcoholic Steatohepatitis Using Clinical Markers and Circulating Levels of Lipids and Metabolites, *CLIN GASTROENTEROL H* 14: 1463-+

EMIF: Lewczuk, Piotr et al. Cerebrospinal Fluid A beta(42/40) Corresponds Better than A beta(42) to Amyloid PET in Alzheimers Disease, *J ALZHEIMERS DIS* 55: 813-822

EMIF: Lotta, Luca A. et al. Integrative genomic analysis implicates limited peripheral adipose storage capacity in the pathogenesis of human insulin resistance, *NAT GENET* 49: 17-26

EMIF: Lotta, Luca A. et al. Genetic Predisposition to an Impaired Metabolism of the Branched-Chain Amino Acids and Risk of Type 2 Diabetes: A Mendelian Randomisation Analysis, *PLOS MED* 13:

EMIF: Proitsi, Petroula et al. Association of blood lipids with Alzheimers disease: A comprehensive lipidomics analysis, *ALZHEIMERS DEMENT* 13: 140-151

EMIF: Mardinoglu, Adil et al. Personal model-assisted identification of NAD(+) and glutathione metabolism as intervention target in NAFLD, *MOL SYST BIOL* 13:

EMIF: ten Kate, Mara et al. Clinical validity of medial temporal atrophy as a biomarker for Alzheimers disease in the context of a structured 5-phase development framework, *NEUROBIOL AGING* 52: 167-182

EMIF: Snowden, Stuart G. et al. Association between fatty acid metabolism in the brain and Alzheimer disease neuropathology and cognitive performance: A nontargeted metabolomic study, *PLOS MED* 14:

EMIF: Lunnon, Katie et al. Mitochondrial genes are altered in blood early in Alzheimers disease, *NEUROBIOL AGING* 53: 36-47

EMIF: Kuhlmann, Julia et al. CSF A beta(1-42) - an excellent but complicated Alzheimers biomarker - a route to standardisation, *CLIN CHIM ACTA* 467: 27-33

EMIF: Vos, Stephanie J. B. et al. Modifiable Risk Factors for Prevention of Dementia in Midlife, Late Life and the Oldest-Old: Validation of the LIBRA Index, *J ALZHEIMERS DIS* 58: 537-547

EMIF: Luukkonen, Panu K. et al. Impaired hepatic lipid synthesis from polyunsaturated fatty acids in TM6SF2 E167K variant carriers with NAFLD, *J HEPATOL* 67: 128-136

EMIF: Frisoni, Giovanni B. et al. Strategic roadmap for an early diagnosis of Alzheimers disease based on biomarkers, *LANCET NEUROL* 16: 661-676

EMIF: Chiasserini, Davide et al. Differential role of CSF fatty acid binding protein 3, alpha-synuclein, and Alzheimers disease core biomarkers in Lewy body disorders and Alzheimers dementia, ALZHEIMERS RES THER 9:

EMIF: Niemantsverdriet, Ellis et al. Alzheimers disease CSF biomarkers: clinical indications and rational use, ACTA NEUROL BELG 117: 591-602

EMIF: Isokuortti, Elina et al. Use of HOMA-IR to diagnose non-alcoholic fatty liver disease: a population-based and inter-laboratory study, DIABETOLOGIA 60: 1873-1882

EMIF: Alshahrani, Mona et al. Neuro-symbolic representation learning on biological knowledge graphs, BIOINFORMATICS 33: 2723-2730

EMIF: Niemantsverdriet, Ellis et al. The Cerebrospinal Fluid A beta(1-42)/A beta(1-40) Ratio Improves Concordance with Amyloid-PET for Diagnosing Alzheimers Disease in a Clinical Setting, J ALZHEIMERS DIS 60: 561-576

EMIF: Mroczko, Barbara et al. Amyloid beta oligomers (A beta Os) in Alzheimers disease, J NEURAL TRANSM 125: 177-191

EMIF: Perera, Gayan et al. Dementia prevalence and incidence in a federation of European Electronic Health Record databases: The European Medical Informatics Framework resource, ALZHEIMERS DEMENT 14: 130-139

EMIF: Tsimihodimos, Vasilis et al. Hypertension and Diabetes Mellitus Coprediction and Time Trajectories, HYPERTENSION 71: 422-428

EMIF: Lee, Sunjae et al. Network analyses identify liver-specific targets for treating liver diseases, MOL SYST BIOL 13:

EMIF: Mardinoglu, Adil et al. An Integrated Understanding of the Rapid Metabolic Benefits of a Carbohydrate-Restricted Diet on Hepatic Steatosis in Humans, CELL METAB 27: 559-+

EMIF: Giannoula, Alexia et al. Identifying temporal patterns in patient disease trajectories using dynamic time warping: A population-based study, SCI REP-UK 8:

EMIF: Goossens, Joery et al. Diagnostic value of cerebrospinal fluid tau, neurofilament, and progranulin in definite frontotemporal lobar degeneration, ALZHEIMERS RES THER 10:

EMIF: Dennis, John M. et al. Precision Medicine in Type 2 Diabetes: Clinical Markers of Insulin Resistance Are Associated With Altered Short- and Long-term Glycemic Response to DPP-4 Inhibitor Therapy, DIABETES CARE 41: 705-712

EMIF: Singh, Gurparkash et al. Real world big data for clinical research and drug development, DRUG DISCOV TODAY 23: 652-660

EMIF: Wild, Sarah H. et al. Cardiovascular Disease, Cancer, and Mortality Among People With Type 2 Diabetes and Alcoholic or Nonalcoholic Fatty Liver Disease Hospital Admission, DIABETES CARE 41: 341-347

EMIF: Lewczuk, Piotr et al. Cerebrospinal fluid and blood biomarkers for neurodegenerative dementias: An update of the Consensus of the Task Force on Biological Markers in Psychiatry of the World Federation of Societies of Biological Psychiatry, *WORLD J BIOL PSYCHIA* 19: 244-328

EMIF: Iliodromiti, Stamatina et al. The impact of confounding on the associations of different adiposity measures with the incidence of cardiovascular disease: a cohort study of 296 535 adults of white European descent, *EUR HEART J* 39: 1514-+

EMIF: Mardinoglu, Adil et al. Systems biology in hepatology: approaches and applications, *NAT REV GASTRO HEPAT* 15: 365-377

EMIF: Luukkonen, Panu K. et al. Saturated Fat Is More Metabolically Harmful for the Human Liver Than Unsaturated Fat or Simple Sugars, *DIABETES CARE* 41: 1732-1739

EMIF: Lewczuk, Piotr et al. Plasma neurofilament light as a potential biomarker of neurodegeneration in Alzheimers disease, *ALZHEIMERS RES THER* 10:

EMIF: Dennis, John M. et al. Sex and BMI Alter the Benefits and Risks of Sulfonylureas and Thiazolidinediones in Type 2 Diabetes: A Framework for Evaluating Stratification Using Routine Clinical and Individual Trial Data, *DIABETES CARE* 41: 1844-1853

EMIF: Legdeur, N. et al. Age dependency of risk factors for cognitive decline, *BMC GERIATR* 18:

EMIF: ten Kate, Mara et al. MRI predictors of amyloid pathology: results from the EMIF-AD Multimodal Biomarker Discovery study, *ALZHEIMERS RES THER* 10:

EMIF: Hansson, Oskar et al. The impact of preanalytical variables on measuring cerebrospinal fluid biomarkers for Alzheimers disease diagnosis: A review, *ALZHEIMERS DEMENT* 14: 1313-1333

EMIF: Sliz, Eeva et al. Metabolomic Consequences of Genetic Inhibition of PCSK9 Compared With Statin Treatment, *CIRCULATION* 138: 2499-2512

EMIF: Lotta, Luca A. et al. Association of Genetic Variants Related to Gluteofemoral vs Abdominal Fat Distribution With Type 2 Diabetes, Coronary Disease, and Cardiovascular Risk Factors, *JAMA-J AM MED ASSOC* 320: 2553-2563

EMIF: ten Kate, Mara et al. Atrophy subtypes in prodromal Alzheimers disease are associated with cognitive decline, *BRAIN* 141: 3443-3456

EMIF: Oeckl, Patrick et al. Glial Fibrillary Acidic Protein in Serum is Increased in Alzheimers Disease and Correlates with Cognitive Impairment, *J ALZHEIMERS DIS* 67: 481-488

EMIF: Wittemans, Laura B. L. et al. Assessing the causal association of glycine with risk of cardio-metabolic diseases, *NAT COMMUN* 10:

EMIF: Collij, Lyduine E. et al. Assessing Amyloid Pathology in Cognitively Normal Subjects Using F-18-Flutemetamol PET: Comparing Visual Reads and Quantitative Methods, *J NUCL MED* 60: 541-547

EMIF: Hansson, Oskar et al. Advantages and disadvantages of the use of the CSF Amyloid (A) 42/40 ratio in the diagnosis of Alzheimers Disease, *ALZHEIMERS RES THER* 11:

EMIF: Bos, Isabelle et al. Cerebrospinal fluid biomarkers of neurodegeneration, synaptic integrity, and astroglial activation across the clinical Alzheimers disease spectrum, ALZHEIMERS DEMENT 15: 644-654

EMIF: Vangipurapu, Jagadish et al. Nine Amino Acids Are Associated With Decreased Insulin Secretion and Elevated Glucose Levels in a 7.4-Year Follow-up Study of 5,181 Finnish Men, DIABETES 68: 1353-1358

EMIF: Kim, Min et al. Primary fatty amides in plasma associated with brain amyloid burden, hippocampal volume, and memory in the European Medical Information Framework for Alzheimers Disease biomarker discovery cohort, ALZHEIMERS DEMENT 15: 817-827

EMIF: Bridel, Claire et al. Diagnostic Value of Cerebrospinal Fluid Neurofilament Light Protein in Neurology: A Systematic Review and Meta-analysis, JAMA NEUROL 76: 1035-1048

EMIF: van Maurik, Ingrid S. et al. Biomarker-based prognosis for people with mild cognitive impairment (ABIDE): a modelling study, LANCET NEUROL 18: 1034-1044

EMIF: Alexander, Myriam et al. Non-alcoholic fatty liver disease and risk of incident acute myocardial infarction and stroke: findings from matched cohort study of 18 million European adults, BMJ-BRIT MED J 367:

EMIF: Kjolbaek, Louise et al. Arabinoxylan oligosaccharides and polyunsaturated fatty acid effects on gut microbiota and metabolic markers in overweight individuals with signs of metabolic syndrome: A randomized cross-over trial, CLIN NUTR 39: 67-79

EMIF: Hagenbeek, Fiona A. et al. Heritability estimates for 361 blood metabolites across 40 genome-wide association studies, NAT COMMUN 11:

EMIF: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), GENOME MED 12:

EMIF: van de Kreeke, Jacoba Alida et al. Optical coherence tomography angiography in preclinical Alzheimers disease, BRIT J OPHTHALMOL 104: 157-161

EMIF: Jian, Ching et al. Quantitative PCR provides a simple and accessible method for quantitative microbiota profiling, PLOS ONE 15:

EMIF: Vangipurapu, Jagadish et al. Microbiota-Related Metabolites and the Risk of Type 2 Diabetes, DIABETES CARE 43: 1319-1325

EMIF: Collij, Lyduine E. et al. Multitracer model for staging cortical amyloid deposition using PET imaging, NEUROLOGY 95: E1538-E1553

EMIF: Rodrigues, Filipe B. et al. Mutant huntingtin and neurofilament light have distinct longitudinal dynamics in Huntingtons disease, SCI TRANSL MED 12:

EMIF: Jian, Ching et al. Impact of short-term overfeeding of saturated or unsaturated fat or sugars on the gut microbiota in relation to liver fat in obese and overweight adults, CLIN NUTR 40: 207-216

EMIF: Vojinovic, Dina et al. Association of Circulating Metabolites in Plasma or Serum and Risk of Stroke Meta-analysis From 7 Prospective Cohorts, NEUROLOGY 96: E1110-E1123

EMIF: Lane, Jennifer C. E. et al. Risk of depression, suicide and psychosis with hydroxychloroquine treatment for rheumatoid arthritis: a multinational network cohort study, RHEUMATOLOGY 60: 3222-3234

EMIF: Tan, Eng Hooi et al. COVID-19 in patients with autoimmune diseases: characteristics and outcomes in a multinational network of cohorts across three countries, RHEUMATOLOGY 60: SI37-SI50

EMIF: Stamate, Daniel et al. A metabolite-based machine learning approach to diagnose Alzheimer-type dementia in blood: Results from the European Medical Information Framework for Alzheimer disease biomarker discovery cohort, ALZH DEMENT-TRCI 5: 933-938

ENABLE: Rabanal, Francesc et al. A bioinspired peptide scaffold with high antibiotic activity and low in vivo toxicity, SCI REP-UK 5:

ENABLE: Hughes, Diarmaid et al. Evolutionary consequences of drug resistance: shared principles across diverse targets and organisms, NAT REV GENET 16: 459-471

ENABLE: Kitchen, Philip et al. Beyond water homeostasis: Diverse functional roles of mammalian aquaporins, BBA-GEN SUBJECTS 1850: 2410-2421

ENABLE: Rabanal, Francesc et al. Recent advances and perspectives in the design and development of polymyxins, NAT PROD REP 34: 886-908

ENABLE: Pantel, Lucile et al. Odilorhabdins, Antibacterial Agents that Cause Miscoding by Binding at a New Ribosomal Site, MOL CELL 70: 83+

ENABLE: Dilworth, Marvin V. et al. Microbial expression systems for membrane proteins, METHODS 147: 3-39

ENABLE: Juhas, Mario et al. In vitro activity of apramycin against multidrug-, carbapenem- and aminoglycoside-resistant Enterobacteriaceae and Acinetobacter baumannii, J ANTIMICROB CHEMOTH 74: 944-952

ENABLE: Krajnc, Alen et al. Will morphing boron-based inhibitors beat the beta-lactamases?, CURR OPIN CHEM BIOL 50: 101-110

ENABLE: Moynie, Lucile et al. The complex of ferric-enterobactin with its transporter from Pseudomonas aeruginosa suggests a two-site model, NAT COMMUN 10:

ENABLE: Tooke, Catherine L. et al. Molecular Basis of Class A beta-Lactamase Inhibition by Relebactam, ANTIMICROB AGENTS CH 63:

ENABLE: Krajnc, Alen et al. Bicyclic Boronate VNRX-5133 Inhibits Metallo- and Serine-beta-Lactamases, J MED CHEM 62: 8544-8556

ENABLE: Tselepis, Lucas et al. In vitro efficacy of imipenem-relebactam and cefepime-AAI101 against a global collection of ESBL-positive and carbapenemase-producing Enterobacteriaceae, INT J ANTIMICROB AG 56:

EPAD: Ritchie, Karen et al. Recommended cognitive outcomes in preclinical Alzheimers disease: Consensus statement from the European Prevention of Alzheimers Dementia project, ALZHEIMERS DEMENT 13: 186-195

EPAD: Mortamais, Marion et al. Detecting cognitive changes in preclinical Alzheimers disease: A review of its feasibility, ALZHEIMERS DEMENT 13: 468-492

EPAD: Crous-Bou, Marta et al. Alzheimers disease prevention: from risk factors to early intervention, ALZHEIMERS RES THER 9:

EPAD: Bunnik, Eline M. et al. On the personal utility of Alzheimers disease-related biomarker testing in the research context, J MED ETHICS 44: 830-834

EPAD: Arabi, Hossein et al. Deep learning-guided joint attenuation and scatter correction in multitracer neuroimaging studies, HUM BRAIN MAPP 41: 3667-3679

EPAD: Mutsaerts, Henk J. M. M. et al. EXploreASL: An image processing pipeline for multi-center ASL perfusion MRI studies, NEUROIMAGE 219:

EPAD: Solomon, Alina et al. Multidomain interventions: state-of-the-art and future directions for protocols to implement precision dementia risk reduction. A user manual for Brain Health Services-part 4 of 6, ALZHEIMERS RES THER 13:

EPAD: Ranson, Janice M. et al. Modifiable risk factors for dementia and dementia risk profiling. A user manual for Brain Health Services-part 2 of 6, ALZHEIMERS RES THER 13:

EPAD: Visser, Leonie N. C. et al. Dementia risk communication. A user manual for Brain Health Services-part 3 of 6, ALZHEIMERS RES THER 13:

EPAD: Milne, Richard et al. Societal and equity challenges for Brain Health Services. A user manual for Brain Health Services-part 6 of 6, ALZHEIMERS RES THER 13:

EQIPD: Voelkl, Bernhard et al. Reproducibility of animal research in light of biological variation, NAT REV NEUROSCI 21: 384-393

EQIPD: Loescher, Wolfgang et al. Drug Resistance in Epilepsy: Clinical Impact, Potential Mechanisms, and New Innovative Treatment Options, PHARMACOL REV 72: 606-638

EQIPD: Bernalov, Anton et al. Introduction to the EQIPD quality system, ELIFE 10:

ERA4TB: Kar, Tamalika et al. A candidate multi-epitope vaccine against SARS-CoV-2, SCI REP-UK 10:

eTOX: Bauer-Mehren, Anna et al. DisGeNET: a Cytoscape plugin to visualize, integrate, search and analyze gene-disease networks, BIOINFORMATICS 26: 2924-2926

eTOX: Obiol-Pardo, Cristian et al. A Multiscale Simulation System for the Prediction of Drug-Induced Cardiotoxicity, J CHEM INF MODEL 51: 483-492

eTOX: Klepsch, Freya et al. Exhaustive Sampling of Docking Poses Reveals Binding Hypotheses for Propafenone Type Inhibitors of P-Glycoprotein, PLOS COMPUT BIOL 7:

eTOX: Bauer-Mehren, Anna et al. Gene-Disease Network Analysis Reveals Functional Modules in Mendelian, Complex and Environmental Diseases, PLOS ONE 6:

eTOX: Enoch, S. J. et al. A review of the electrophilic reaction chemistry involved in covalent protein binding relevant to toxicity, CRIT REV TOXICOL 41: 783-802

eTOX: Przybylak, Katarzyna R. et al. In silico models for drug-induced liver injury - current status, EXPERT OPIN DRUG MET 8: 201-217

eTOX: Chiche, Johanna et al. In vivo pH in metabolic-defective Ras-transformed fibroblast tumors: Key role of the monocarboxylate transporter, MCT4, for inducing an alkaline intracellular pH, INT J CANCER 130: 1511-1520

eTOX: Arighi, Cecilia N. et al. Overview of the BioCreative III Workshop, BMC BIOINFORMATICS 12:

eTOX: van Mulligen, Erik M. et al. The EU-ADR corpus: Annotated drugs, diseases, targets, and their relationships, J BIOMED INFORM 45: 879-884

eTOX: Canzar, Stefan et al. Charge Group Partitioning in Biomolecular Simulation, J COMPUT BIOL 20: 188-198

eTOX: Furlong, Laura I. et al. Human diseases through the lens of network biology, TRENDS GENET 29: 150-159

eTOX: Oomen, Agnes G. et al. Concern-driven integrated approaches to nanomaterial testing and assessment - report of the NanoSafety Cluster Working Group 10, NANOTOXICOLOGY 8: 334-348

eTOX: Klepsch, Freya et al. Ligand and Structure-Based Classification Models for Prediction of P-Glycoprotein Inhibitors, J CHEM INF MODEL 54: 218-229

eTOX: Bento, A. Patricia et al. The ChEMBL bioactivity database: an update, NUCLEIC ACIDS RES 42: D1083-D1090

eTOX: Bravo, Alex et al. Extraction of relations between genes and diseases from text and large-scale data analysis: implications for translational research, BMC BIOINFORMATICS 16:

eTOX: Queralt-Rosinach, Nuria et al. DisGeNET-RDF: harnessing the innovative power of the Semantic Web to explore the genetic basis of diseases, BIOINFORMATICS 32: 2236-2238

eTOX: Krallinger, Martin et al. CHEMDNER: The drugs and chemical names extraction challenge, J CHEMINFORMATICS 7:

eTOX: Krallinger, Martin et al. The CHEMDNER corpus of chemicals and drugs and its annotation principles, J CHEMINFORMATICS 7:

eTOX: Mendez, David et al. ChEMBL: towards direct deposition of bioassay data, NUCLEIC ACIDS RES 47: D930-D940

eTRANSAFE: Pawar, Gopal et al. In Silico Toxicology Data Resources to Support Read-Across and (Q)SAR, FRONT PHARMACOL 10:



eTRANSafe: Hiemstra, Steven et al. High-throughput confocal imaging of differentiated 3D liver-like spheroid cellular stress response reporters for identification of drug-induced liver injury liability, ARCH TOXICOL 93: 2895-2911

eTRANSafe: Pinero, Janet et al. The DisGeNET knowledge platform for disease genomics: 2019 update, NUCLEIC ACIDS RES 48: D845-D855

eTRIKS: Agusti, Alvar et al. Personalized Respiratory Medicine: Exploring the Horizon, Addressing the Issues Summary of a BRN-AJRCCM Workshop Held in Barcelona on June 12, 2014, AM J RESP CRIT CARE 191: 391-401

eTRIKS: Shaw, Dominick E. et al. Clinical and inflammatory characteristics of the European U-BIOPRED adult severe asthma cohort, EUR RESPIR J 46: 1308-1321

eTRIKS: Fleming, Louise et al. The burden of severe asthma in childhood and adolescence: results from the paediatric U-BIOPRED cohorts, EUR RESPIR J 46: 1322-1333

eTRIKS: Rocca-Serra, Philippe et al. Data standards can boost metabolomics research, and if there is a will, there is a way, METABOLOMICS 12:

eTRIKS: Debray, Thomas P. A. et al. Get real in individual participant data (IPD) meta-analysis: a review of the methodology, RES SYNTH METHODS 6: 293-309

eTRIKS: Nordon, Clementine et al. The Efficacy-Effectiveness Gap : Historical Background and Current Conceptualization, VALUE HEALTH 19: 75-81

eTRIKS: McQuilton, Peter et al. BioSharing: curated and crowd-sourced metadata standards, databases and data policies in the life sciences, DATABASE-OXFORD :

eTRIKS: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

eTRIKS: Lysenko, Artem et al. Representing and querying disease networks using graph databases, BIODATA MIN 9:

eTRIKS: Efthimiou, Orestis et al. GetReal in network meta-analysis: a review of the methodology, RES SYNTH METHODS 7: 236-263

eTRIKS: Lefauieux, Diane et al. U-BIOPRED clinical adult asthma clusters linked to a subset of sputum omics, J ALLERGY CLIN IMMUN 139: 1797-1807

eTRIKS: Rossios, Christos et al. Sputum transcriptomics reveal upregulation of IL-1 receptor family members in patients with severe asthma, J ALLERGY CLIN IMMUN 141: 560-570

eTRIKS: Gawron, Piotr et al. MINERVA-a platform for visualization and curation of molecular interaction networks, NPJ SYST BIOL APPL 2:

eTRIKS: Mazein, Alexander et al. Systems medicine disease maps: community-driven comprehensive representation of disease mechanisms, NPJ SYST BIOL APPL 4:

eTRIKS: Schofield, James P. R. et al. Stratification of asthma phenotypes by airway proteomic signatures, J ALLERGY CLIN IMMUN 144: 70-82

eTRIKS: Ostaszewski, Marek et al. Community-driven roadmap for integrated disease maps, BRIEF BIOINFORM 20: 659-670

eTRIKS: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), GENOME MED 12:

Eu2P: Dreischulte, Tobias et al. Combined use of nonsteroidal anti-inflammatory drugs with diuretics and/or renin-angiotensin system inhibitors in the community increases the risk of acute kidney injury, KIDNEY INT 88: 396-403

EU-AIMS: Meyer-Lindenberg, Andreas et al. Neural mechanisms of social risk for psychiatric disorders, NAT NEUROSCI 15: 663-668

EU-AIMS: Stein, Jason L. et al. Identification of common variants associated with human hippocampal and intracranial volumes, NAT GENET 44: 552-+

EU-AIMS: Whelan, Robert et al. Adolescent impulsivity phenotypes characterized by distinct brain networks, NAT NEUROSCI 15: 920-U153

EU-AIMS: Kong, Augustine et al. Rate of de novo mutations and the importance of fathers age to disease risk, NATURE 488: 471-475

EU-AIMS: Baudouin, Stephane J. et al. Shared Synaptic Pathophysiology in Syndromic and Nonsyndromic Rodent Models of Autism, SCIENCE 338: 128-132

EU-AIMS: Spooren, Will et al. Synapse dysfunction in autism: a molecular medicine approach to drug discovery in neurodevelopmental disorders, TRENDS PHARMACOL SCI 33: 669-684

EU-AIMS: Budreck, Elaine C. et al. Neuroligin-1 controls synaptic abundance of NMDA-type glutamate receptors through extracellular coupling, P NATL ACAD SCI USA 110: 725-730

EU-AIMS: Persico, Antonio M. et al. Urinary p-cresol in autism spectrum disorder, NEUROTOXICOL TERATOL 36: 82-90

EU-AIMS: Delorme, Richard et al. Progress toward treatments for synaptic defects in autism, NAT MED 19: 685-694

EU-AIMS: El-Kordi, Ahmed et al. Development of an autism severity score for mice using Nlgn4 null mutants as a construct-valid model of heritable monogenic autism, BEHAV BRAIN RES 251: 41-49

EU-AIMS: Persico, Antonio M. et al. Autism genetics, BEHAV BRAIN RES 251: 95-112

EU-AIMS: Siddiqui, Tabrez J. et al. An LRRTM4-HSPG Complex Mediates Excitatory Synapse Development on Dentate Gyrus Granule Cells, NEURON 79: 680-695

EU-AIMS: Lai, Meng-Chuan et al. Biological sex affects the neurobiology of autism, BRAIN 136: 2799-2815

EU-AIMS: Zuko, Amila et al. Contactins in the neurobiology of autism, EUR J PHARMACOL 719: 63-74

EU-AIMS: Ey, Elodie et al. The Autism ProSAP1/Shank2 mouse model displays quantitative and structural abnormalities in ultrasonic vocalisations, *BEHAV BRAIN RES* 256: 677-689

EU-AIMS: Webb, Sara Jane et al. The motivation for very early intervention for infants at high risk for autism spectrum disorders, *INT J SPEECH-LANG PA* 16: 36-42

EU-AIMS: Murray, Macey L. et al. Pharmacological treatments prescribed to people with autism spectrum disorder (ASD) in primary health care, *PSYCHOPHARMACOLOGY* 231: 1011-1021

EU-AIMS: Kleijer, Kristel T. E. et al. Neurobiology of autism gene products: towards pathogenesis and drug targets, *PSYCHOPHARMACOLOGY* 231: 1037-1062

EU-AIMS: Kas, Martien J. et al. Assessing behavioural and cognitive domains of autism spectrum disorders in rodents: current status and future perspectives, *PSYCHOPHARMACOLOGY* 231: 1125-1146

EU-AIMS: Ruggeri, Barbara et al. Biomarkers in autism spectrum disorder: the old and the new, *PSYCHOPHARMACOLOGY* 231: 1201-1216

EU-AIMS: Lai, Meng-Chuan et al. Autism, *LANCET* 383: 896-910

EU-AIMS: Jones, Emily J. H. et al. Developmental pathways to autism: A review of prospective studies of infants at risk, *NEUROSCI BIOBEHAV R* 39: 1-33

EU-AIMS: Piras, I. S. et al. Anti-brain antibodies are associated with more severe cognitive and behavioral profiles in Italian children with Autism Spectrum Disorder, *BRAIN BEHAV IMMUN* 38: 91-99

EU-AIMS: Gabriele, Stefano et al. Blood serotonin levels in autism spectrum disorder: A systematic review and meta-analysis, *EUR NEUROPSYCHOPHARM* 24: 919-929

EU-AIMS: Whelan, Robert et al. Neuropsychosocial profiles of current and future adolescent alcohol misusers, *NATURE* 512: 185-+

EU-AIMS: Gabriele, Stefano et al. Urinary p-cresol is elevated in young French children with autism spectrum disorder: a replication study, *BIOMARKERS* 19: 463-470

EU-AIMS: Baron-Cohen, Simon et al. Attenuation of Typical Sex Differences in 800 Adults with Autism vs. 3,900 Controls, *PLOS ONE* 9:

EU-AIMS: Wilson, C. Ellie et al. The Neuropsychology of Male Adults With High-Functioning Autism or Asperger Syndrome, *AUTISM RES* 7: 568-581

EU-AIMS: Schreiner, Dietmar et al. Targeted Combinatorial Alternative Splicing Generates Brain Region-Specific Repertoires of Neurexins, *NEURON* 84: 386-398

EU-AIMS: Distler, Ute et al. In-depth protein profiling of the postsynaptic density from mouse hippocampus using data-independent acquisition proteomics, *PROTEOMICS* 14: 2607-2613

EU-AIMS: Basil, P. et al. Prenatal maternal immune activation causes epigenetic differences in adolescent mouse brain, *TRANSL PSYCHIAT* 4:

EU-AIMS: Castellanos-Ryan, Natalie et al. Neural and Cognitive Correlates of the Common and Specific Variance Across Externalizing Problems in Young Adolescence, *AM J PSYCHIAT* 171: 1310-1319

EU-AIMS: Orekhova, Elena V. et al. EEG hyper-connectivity in high-risk infants is associated with later autism, *J NEURODEV DISORD* 6:

EU-AIMS: Lai, Meng-Chuan et al. Sex/Gender Differences and Autism: Setting the Scene for Future Research, *J AM ACAD CHILD PSY* 54: 11-24

EU-AIMS: Wass, Sam V. et al. Shorter spontaneous fixation durations in infants with later emerging autism, *SCI REP-UK* 5:

EU-AIMS: Jedlicka, Peter et al. Neuroligin-1 regulates excitatory synaptic transmission, LTP and EPSP-spike coupling in the dentate gyrus in vivo, *BRAIN STRUCT FUNCT* 220: 47-58

EU-AIMS: Man, Kenneth K. C. et al. Exposure to selective serotonin reuptake inhibitors during pregnancy and risk of autism spectrum disorder in children: A systematic review and meta-analysis of observational studies, *NEUROSCI BIOBEHAV R* 49: 82-89

EU-AIMS: Johnson, Mark H. et al. Annual Research Review: Infant development, autism, and ADHD - early pathways to emerging disorders, *J CHILD PSYCHOL PSYC* 56: 228-247

EU-AIMS: Johnson, Mark H. et al. Brain adaptation and alternative developmental trajectories, *DEV PSYCHOPATHOL* 27: 425-442

EU-AIMS: Richiardi, Jonas et al. Correlated gene expression supports synchronous activity in brain networks, *SCIENCE* 348: 1241-1244

EU-AIMS: Gliga, Teodora et al. Enhanced Visual Search in Infancy Predicts Emerging Autism Symptoms, *CURR BIOL* 25: 1727-1730

EU-AIMS: Schmeisser, Michael J. et al. Translational neurobiology in Shank mutant mice - Model systems for neuropsychiatric disorders, *ANN ANAT* 200: 115-117

EU-AIMS: Bourgeron, Thomas et al. From the genetic architecture to synaptic plasticity in autism spectrum disorder, *NAT REV NEUROSCI* 16: 551-563

EU-AIMS: Tost, Heike et al. Environmental influence in the brain, human welfare and mental health, *NAT NEUROSCI* 18: 1421-1431

EU-AIMS: Vulser, Helene et al. Subthreshold Depression and Regional Brain Volumes in Young Community Adolescents, *J AM ACAD CHILD PSY* 54: 832-840

EU-AIMS: French, Leon et al. Early Cannabis Use, Polygenic Risk Score for Schizophrenia, and Brain Maturation in Adolescence, *JAMA PSYCHIAT* 72: 1002-1011

EU-AIMS: Ecker, Christine et al. Neuroimaging in autism spectrum disorder: brain structure and function across the lifespan, *LANCET NEUROL* 14: 1121-1134

EU-AIMS: Lai, Meng-Chuan et al. Identifying the lost generation of adults with autism spectrum conditions, *LANCET PSYCHIAT* 2: 1013-1027

EU-AIMS: Sacco, Roberto et al. Head circumference and brain size in autism spectrum disorder: A systematic review and meta-analysis, *PSYCHIAT RES-NEUROIM* 234: 239-251

EU-AIMS: Ortuno-Sierra, Javier et al. New evidence of factor structure and measurement invariance of the SDQ across five European nations, *EUR CHILD ADOLES PSY* 24: 1523-1534

EU-AIMS: Stringaris, Argyris et al. The Brains Response to Reward Anticipation and Depression in Adolescence: Dimensionality, Specificity, and Longitudinal Predictions in a Community-Based Sample, *AMJ PSYCHIAT* 172: 1215-1223

EU-AIMS: Auyeung, B. et al. Oxytocin increases eye contact during a real-time, naturalistic social interaction in males with and without autism, *TRANSL PSYCHIAT* 5:

EU-AIMS: Floris, Dorothea L. et al. Atypically Rightward Cerebral Asymmetry in Male Adults With Autism Stratifies Individuals With and Without Language Delay, *HUM BRAIN MAPP* 37: 230-253

EU-AIMS: Catani, Marco et al. Frontal networks in adults with autism spectrum disorder, *BRAIN* 139: 616-630

EU-AIMS: Constantino, John N. et al. Diagnosis of autism spectrum disorder: reconciling the syndrome, its diverse origins, and variation in expression, *LANCET NEUROL* 15: 279-291

EU-AIMS: Franke, Barbara et al. Genetic influences on schizophrenia and subcortical brain volumes: large-scale proof of concept, *NAT NEUROSCI* 19: 420-+

EU-AIMS: Salomone, Erica et al. Use of early intervention for young children with autism spectrum disorder across Europe, *AUTISM* 20: 233-249

EU-AIMS: Traunmuller, Lisa et al. Control of neuronal synapse specification by a highly dedicated alternative splicing program, *SCIENCE* 352: 982-986

EU-AIMS: Visser, Janne C. et al. Autism spectrum disorder and attention-deficit/hyperactivity disorder in early childhood: A review of unique and shared characteristics and developmental antecedents, *NEUROSCI BIOBEHAV R* 65: 229-263

EU-AIMS: Elsabbagh, Mayada et al. Autism and the Social Brain: The First-Year Puzzle, *BIOL PSYCHIAT* 80: 94-99

EU-AIMS: Murphy, Clodagh M. et al. Autism spectrum disorder in adults: diagnosis, management, and health services development, *NEUROPSYCH DIS TREAT* 12: 1669-1686

EU-AIMS: Heinrich, Angela et al. Prediction of alcohol drinking in adolescents: Personality-traits, behavior, brain responses, and genetic variations in the context of reward sensitivity, *BIOL PSYCHOL* 118: 79-87

EU-AIMS: Ashwood, K. L. et al. Predicting the diagnosis of autism in adults using the Autism-Spectrum Quotient (AQ) questionnaire, *PSYCHOL MED* 46: 2595-2604

EU-AIMS: Ellie Wilson, C. et al. Does sex influence the diagnostic evaluation of autism spectrum disorder in adults?, *AUTISM* 20: 808-819

EU-AIMS: Ecker, C. et al. Relationship Between Cortical Gyrfication, White Matter Connectivity, and Autism Spectrum Disorder, *CEREB CORTEX* 26: 3297-3309

EU-AIMS: Peter, Sasa et al. Dysfunctional cerebellar Purkinje cells contribute to autism-like behaviour in Shank2-deficient mice, *NAT COMMUN* 7:

EU-AIMS: Braun, Urs et al. Dynamic brain network reconfiguration as a potential schizophrenia genetic risk mechanism modulated by NMDA receptor function, *P NATL ACAD SCI USA* 113: 12568-12573

EU-AIMS: Castellanos-Ryan, Natalie et al. The Structure of Psychopathology in Adolescence and Its Common Personality and Cognitive Correlates, *J ABNORM PSYCHOL* 125: 1039-1052

EU-AIMS: Gevi, Federica et al. Urinary metabolomics of young Italian autistic children supports abnormal tryptophan and purine metabolism, *MOL AUTISM* 7:

EU-AIMS: Evans, David W. et al. Development of Two Dimensional Measures of Restricted and Repetitive Behavior in Parents and Children, *J AM ACAD CHILD PSY* 56: 51-58

EU-AIMS: Naaijen, J. et al. Glutamatergic and GABAergic gene sets in attention-deficit/hyperactivity disorder: association to overlapping traits in ADHD and autism, *TRANSL PSYCHIAT* 7:

EU-AIMS: Thompson, Abigail et al. Impaired Communication Between the Motor and Somatosensory Homunculus Is Associated With Poor Manual Dexterity in Autism Spectrum Disorder, *BIOL PSYCHIAT* 81: 211-219

EU-AIMS: Lilja, Johanna et al. SHANK proteins limit integrin activation by directly interacting with Rap1 and R-Ras, *NAT CELL BIOL* 19: 292-+

EU-AIMS: Ecker, Christine et al. Association Between the Probability of Autism Spectrum Disorder and Normative Sex-Related Phenotypic Diversity in Brain Structure, *JAMA PSYCHIAT* 74: 329-338

EU-AIMS: Shephard, Elizabeth et al. Mid-childhood outcomes of infant siblings at familial high-risk of autism spectrum disorder, *AUTISM RES* 10: 546-557

EU-AIMS: Vicidomini, C. et al. Pharmacological enhancement of mGlu5 receptors rescues behavioral deficits in SHANK3 knock-out mice, *MOL PSYCHIATR* 22: 689-702

EU-AIMS: Sokolova, Elena et al. A Causal and Mediation Analysis of the Comorbidity Between Attention Deficit Hyperactivity Disorder (ADHD) and Autism Spectrum Disorder (ASD), *J AUTISM DEV DISORD* 47: 1595-1604

EU-AIMS: Ulfarsson, M. O. et al. 15q11.2 CNV affects cognitive, structural and functional correlates of dyslexia and dyscalculia, *TRANSL PSYCHIAT* 7:

EU-AIMS: Ajram, L. A. et al. Shifting brain inhibitory balance and connectivity of the prefrontal cortex of adults with autism spectrum disorder, *TRANSL PSYCHIAT* 7:

EU-AIMS: Arora, Manish et al. Fetal and postnatal metal dysregulation in autism, *NAT COMMUN* 8:

EU-AIMS: Loth, Eva et al. The EU-AIMS Longitudinal European Autism Project (LEAP): design and methodologies to identify and validate stratification biomarkers for autism spectrum disorders, MOL AUTISM 8:

EU-AIMS: Charman, Tony et al. The EU-AIMS Longitudinal European Autism Project (LEAP): clinical characterisation, MOL AUTISM 8:

EU-AIMS: Sethna, Vaheshta et al. Mother-infant interactions and regional brain volumes in infancy: an MRI study, BRAIN STRUCT FUNCT 222: 2379-2388

EU-AIMS: Lai, Meng-Chuan et al. Quantifying and exploring camouflaging in men and women with autism, AUTISM 21: 690-702

EU-AIMS: Visser, Janne C. et al. Variation in the Early Trajectories of Autism Symptoms Is Related to the Development of Language, Cognition, and Behavior Problems, J AM ACAD CHILD PSY 56: 659-668

EU-AIMS: Gur, R. E. et al. A neurogenetic model for the study of schizophrenia spectrum disorders: the International 22q11.2 Deletion Syndrome Brain Behavior Consortium, MOL PSYCHIATR 22: 1664-1672

EU-AIMS: Loth, E. et al. Facial expression recognition as a candidate marker for autism spectrum disorder: how frequent and severe are deficits?, MOL AUTISM 9:

EU-AIMS: Kathuria, A. et al. Stem cell-derived neurons from autistic individuals with SHANK3 mutation show morphogenetic abnormalities during early development, MOL PSYCHIATR 23: 735-746

EU-AIMS: Chatham, C. H. et al. Adaptive behavior in autism: Minimal clinically important differences on the Vineland-II, AUTISM RES 11: 270-283

EU-AIMS: OHalloran, Laura et al. Neural circuitry underlying sustained attention in healthy adolescents and in ADHD symptomatology, NEUROIMAGE 169: 395-406

EU-AIMS: Berry-Kravis, Elizabeth M. et al. Drug development for neurodevelopmental disorders: lessons learned from fragile X syndrome, NAT REV DRUG DISCOV 17: 280-298

EU-AIMS: Huguet, Guillaume et al. Measuring and Estimating the Effect Sizes of Copy Number Variants on General Intelligence in Community-Based Samples, JAMA PSYCHIAT 75: 447-457

EU-AIMS: Nystrom, Par et al. Enhanced pupillary light reflex in infancy is associated with autism diagnosis in toddlerhood, NAT COMMUN 9:

EU-AIMS: Horder, Jamie et al. Glutamate and GABA in autism spectrum disorder-a translational magnetic resonance spectroscopy study in man and rodent models, TRANSL PSYCHIAT 8:

EU-AIMS: Bussu, G. et al. Prediction of Autism at 3 Years from Behavioural and Developmental Measures in High-Risk Infants: A Longitudinal Cross-Domain Classifier Analysis, J AUTISM DEV DISORD 48: 2418-2433

EU-AIMS: Tillmann, J. et al. Evaluating Sex and Age Differences in ADI-R and ADOS Scores in a Large European Multi-site Sample of Individuals with Autism Spectrum Disorder, *J AUTISM DEV DISORD* 48: 2490-2505

EU-AIMS: Howells, Henrietta et al. Frontoparietal Tracts Linked to Lateralized Hand Preference and Manual Specialization, *CEREB CORTEX* 28: 2482-2494

EU-AIMS: van Rooij, Daan et al. Cortical and Subcortical Brain Morphometry Differences Between Patients With Autism Spectrum Disorder and Healthy Individuals Across the Lifespan: Results From the ENIGMA ASD Working Group, *AM J PSYCHIAT* 175: 359-369

EU-AIMS: Falck-Ytter, Terje et al. Reduced orienting to audiovisual synchrony in infancy predicts autism diagnosis at 3 years of age, *J CHILD PSYCHOL PSYC* 59: 872-880

EU-AIMS: Bariselli, Sebastiano et al. Role of VTA dopamine neurons and neuroligin 3 in sociability traits related to nonfamiliar conspecific interaction, *NAT COMMUN* 9:

EU-AIMS: Yorke, Isabel et al. The Association Between Emotional and Behavioral Problems in Children with Autism Spectrum Disorder and Psychological Distress in Their Parents: A Systematic Review and Meta-analysis, *J AUTISM DEV DISORD* 48: 3393-3415

EU-AIMS: Wolfers, Thomas et al. Mapping the Heterogeneous Phenotype of Schizophrenia and Bipolar Disorder Using Normative Models, *JAMA PSYCHIAT* 75: 1146-1155

EU-AIMS: Scott, Ricardo et al. Loss of *Cntnap2* Causes Axonal Excitability Deficits, Developmental Delay in Cortical Myelination, and Abnormal Stereotyped Motor Behavior, *CEREB CORTEX* 29: 586-597

EU-AIMS: Cao, Zhipeng et al. Mapping adolescent reward anticipation, receipt, and prediction error during the monetary incentive delay task, *HUM BRAIN MAPP* 40: 262-283

EU-AIMS: Haartsen, Rianne et al. Functional EEG connectivity in infants associates with later restricted and repetitive behaviours in autism, a replication study, *TRANSL PSYCHIAT* 9:

EU-AIMS: Holiga, Stefan et al. Patients with autism spectrum disorders display reproducible functional connectivity alterations, *SCI TRANSL MED* 11:

EU-AIMS: Orr, Catherine et al. Grey Matter Volume Differences Associated with Extremely Low Levels of Cannabis Use in Adolescence, *J NEUROSCI* 39: 1817-1827

EU-AIMS: Silva, Ana, I et al. Reciprocal White Matter Changes Associated With Copy Number Variation at 15q11.2 BP1-BP2: A Diffusion Tensor Imaging Study, *BIOL PSYCHIAT* 85: 563-572

EU-AIMS: Bolte, Sven et al. The contribution of environmental exposure to the etiology of autism spectrum disorder, *CELL MOL LIFE SCI* 76: 1275-1297

EU-AIMS: Tillmann, Julian et al. Investigating the factors underlying adaptive functioning in autism in the EU-AIMS Longitudinal European Autism Project, *AUTISM RES* 12: 645-657

EU-AIMS: Leblond, Claire S. et al. Both rare and common genetic variants contribute to autism in the Faroe Islands, *NPJ GENOM MED* 4:



EU-AIMS: Pretzsch, Charlotte Marie et al. Effects of cannabidiol on brain excitation and inhibition systems, a randomised placebo-controlled single dose trial during magnetic resonance spectroscopy in adults with and without autism spectrum disorder, *NEUROPSYCHOPHARMACOL* 44: 1398-1405

EU-AIMS: Lai, Meng-Chuan et al. Neural self-representation in autistic women and association with compensatory camouflaging, *AUTISM* 23: 1210-1223

EU-AIMS: Camm-Crosbie, Louise et al. People like me dont get support: Autistic adults experiences of support and treatment for mental health difficulties, self-injury and suicidality, *AUTISM* 23: 1431-1441

EU-AIMS: Au-Yeung, Sheena K. et al. Experience of mental health diagnosis and perceived misdiagnosis in autistic, possibly autistic and non-autistic adults, *AUTISM* 23: 1508-1518

EU-AIMS: Jollans, Lee et al. Quantifying performance of machine learning methods for neuroimaging data, *NEUROIMAGE* 199: 351-365

EU-AIMS: Di Lorenzo, Renata et al. Recommendations for motion correction of infant fNIRS data applicable to multiple data sets and acquisition systems, *NEUROIMAGE* 200: 511-527

EU-AIMS: Tost, Heike et al. Neural correlates of individual differences in affective benefit of real-life urban green space exposure, *NAT NEUROSCI* 22: 1389+

EU-AIMS: Warriar, Varun et al. Social and non-social autism symptoms and trait domains are genetically dissociable, *COMMUN BIOL* 2:

EU-AIMS: Wolfers, Thomas et al. From pattern classification to stratification: towards conceptualizing the heterogeneity of Autism Spectrum Disorder, *NEUROSCI BIOBEHAV R* 104: 240-254

EU-AIMS: Evangelou, Evangelos et al. New alcohol-related genes suggest shared genetic mechanisms with neuropsychiatric disorders, *NAT HUM BEHAV* 3: 950-961

EU-AIMS: Gudmundsson, Olafur O. et al. Attention-deficit hyperactivity disorder shares copy number variant risk with schizophrenia and autism spectrum disorder, *TRANSL PSYCHIAT* 9:

EU-AIMS: Postema, Merel C. et al. Altered structural brain asymmetry in autism spectrum disorder in a study of 54 datasets, *NAT COMMUN* 10:

EU-AIMS: Oldehinkel, Marianne et al. Altered Connectivity Between Cerebellum, Visual, and Sensory-Motor Networks in Autism Spectrum Disorder: Results from the EU-AIMS Longitudinal European Autism Project, *BIOL PSYCHIAT-COGN N* 4: 260-270

EU-AIMS: Zabihi, Mariam et al. Dissecting the Heterogeneous Cortical Anatomy of Autism Spectrum Disorder Using Normative Models, *BIOL PSYCHIAT-COGN N* 4: 567-578

EU-AIMS: de Chaumont, Fabrice et al. Real-time analysis of the behaviour of groups of mice via a depth-sensing camera and machine learning, *NAT BIOMED ENG* 3: 930-942

EU-AIMS: Jonsson, B. A. et al. Brain age prediction using deep learning uncovers associated sequence variants, *NAT COMMUN* 10:

EU-AIMS: Bakker-Huvenaars, M. J. et al. Saliva oxytocin, cortisol, and testosterone levels in adolescent boys with autism spectrum disorder, oppositional defiant disorder/conduct disorder and typically developing individuals, *EUR NEUROPSYCHOPHARM* 30: 87-101

EU-AIMS: McDonald, Nicole M. et al. Developmental Trajectories of Infants With Multiplex Family Risk for Autism A Baby Siblings Research Consortium Study, *JAMA NEUROL* 77: 73-81

EU-AIMS: Holz, Nathalie E. et al. Resilience and the brain: a key role for regulatory circuits linked to social stress and support, *MOL PSYCHIATR* 25: 379-396

EU-AIMS: Bossier, Han et al. The empirical replicability of task-based fMRI as a function of sample size, *NEUROIMAGE* 212:

EU-AIMS: Begum Ali, Jannath et al. Early Motor Differences in Infants at Elevated Likelihood of Autism Spectrum Disorder and/or Attention Deficit Hyperactivity Disorder, *J AUTISM DEV DISORD* 50: 4367-4384

EU-AIMS: Yu, Tao et al. Cannabis-Associated Psychotic-like Experiences Are Mediated by Developmental Changes in the Parahippocampal Gyrus, *J AM ACAD CHILD PSY* 59: 642-649

EU-AIMS: Lukito, Steve et al. Comparative meta-analyses of brain structural and functional abnormalities during cognitive control in attention-deficit/hyperactivity disorder and autism spectrum disorder, *PSYCHOL MED* 50: 894-919

EU-AIMS: Hoogman, Martine et al. Consortium neuroscience of attention deficit/hyperactivity disorder and autism spectrum disorder: The ENIGMA adventure, *HUM BRAIN MAPP* 43: 37-55

EU-AIMS: Reichert, Markus et al. Studying the impact of built environments on human mental health in everyday life: methodological developments, state-of-the-art and technological frontiers, *CURR OPIN PSYCHOL* 32: 158-164

EU-AIMS: Maricic, Lea Mascarell et al. The IMAGEN study: a decade of imaging genetics in adolescents, *MOL PSYCHIATR* 25: 2648-2671

EU-AIMS: Ching, Christopher R. K. et al. Mapping Subcortical Brain Alterations in 22q11.2 Deletion Syndrome: Effects of Deletion Size and Convergence With Idiopathic Neuropsychiatric Illness, *AM J PSYCHIAT* 177: 589-600

EU-AIMS: van den Berk-Smeekens, Iris et al. Adherence and acceptability of a robot-assisted Pivotal Response Treatment protocol for children with autism spectrum disorder, *SCI REP-UK* 10:

EU-AIMS: Hornberg, Hanna et al. Rescue of oxytocin response and social behaviour in a mouse model of autism, *NATURE* 584: 252+

EU-AIMS: Oakley, Bethany F. M. et al. How do core autism traits and associated symptoms relate to quality of life? Findings from the Longitudinal European Autism Project, *AUTISM* 25: 389-404

EU-AIMS: Dimitrova, Ralica et al. Heterogeneity in Brain Microstructural Development Following Preterm Birth, *CEREB CORTEX* 30: 4800-4810

EU-AIMS: Chaarani, Bader et al. Neural Correlates of Adolescent Irritability and Its Comorbidity With Psychiatric Disorders, *J AM ACAD CHILD PSY* 59: 1371-1379

EU-AIMS: Piccardi, Elena Serena et al. Behavioural and neural markers of tactile sensory processing in infants at elevated likelihood of autism spectrum disorder and/or attention deficit hyperactivity disorder, *J NEURODEV DISORD* 13:

EU-AIMS: Douard, Elise et al. Effect Sizes of Deletions and Duplications on Autism Risk Across the Genome, *AM J PSYCHIAT* 178: 87-98

EU-AIMS: Gomez, Andrea M. et al. Neurexins: molecular codes for shaping neuronal synapses, *NAT REV NEUROSCI* 22: 137-151

EU-AIMS: Adhya, Dwaipayan et al. Atypical Neurogenesis in Induced Pluripotent Stem Cells From Autistic Individuals, *BIOL PSYCHIAT* 89: 486-496

EU-AIMS: Floris, Dorothea L. et al. Towards robust and replicable sex differences in the intrinsic brain function of autism, *MOL AUTISM* 12:

EU-AIMS: Xie, Chao et al. Reward Versus Nonreward Sensitivity of the Medial Versus Lateral Orbitofrontal Cortex Relates to the Severity of Depressive Symptoms, *BIOL PSYCHIAT-COGN N* 6: 259-269

EU-AIMS: Gui, Anna et al. Attentive brain states in infants with and without later autism, *TRANSL PSYCHIAT* 11:

EU-AIMS: Floris, Dorothea L. et al. Atypical Brain Asymmetry in Autism-A Candidate for Clinically Meaningful Stratification, *BIOL PSYCHIAT-COGN N* 6: 802-812

EU-AIMS: Del Bianco, Teresa et al. Temporal Profiles of Social Attention Are Different Across Development in Autistic and Neurotypical People, *BIOL PSYCHIAT-COGN N* 6: 813-824

EU-AIMS: Eyre, Michael et al. The Developing Human Connectome Project: typical and disrupted perinatal functional connectivity, *BRAIN* 144: 2199-2213

EuBOPEN: Adhikari, Bikash et al. PROTAC-mediated degradation reveals a non-catalytic function of AURORA-A kinase, *NAT CHEM BIOL* 16: 1179-+

EuBOPEN: Deniston, C. K. et al. Structure of LRRK2 in Parkinsons disease and model for microtubule interaction, *NATURE* 588:

EuBOPEN: Youhanna, Sonia et al. The Past, Present and Future of Intestinal In Vitro Cell Systems for Drug Absorption Studies, *J PHARM SCI-US* 110: 50-65

EuBOPEN: Stebbing, Justin et al. JAK inhibition reduces SARS-CoV-2 liver infectivity and modulates inflammatory responses to reduce morbidity and mortality, *SCI ADV* 7:

EuBOPEN: Wright, Nathan David et al. The low-cost Shifter microscope stage transforms the speed and robustness of protein crystal harvesting, *ACTA CRYSTALLOGR D* 77: 62-74

EuBOPEN: Wells, Carrow I. et al. The Kinase Chemogenomic Set (KCGS): An Open Science Resource for Kinase Vulnerability Identification, *INT J MOL SCI* 22:

EUbOPEN: Kemas, Aurino M. et al. Insulin-dependent glucose consumption dynamics in 3D primary human liver cultures measured by a sensitive and specific glucose sensor with nanoliter input volume, *FASEB J* 35:

EUbOPEN: Richters, Andre et al. Modulating Androgen Receptor-Driven Transcription in Prostate Cancer with Selective CDK9 Inhibitors, *CELL CHEM BIOL* 28: 134-+

EUbOPEN: DAmico, Francesca et al. Targeting TRIM Proteins: A Quest towards Drugging an Emerging Protein Class, *CHEMBIOCHEM* 22: 2011-2031

EUbOPEN: Wu, Qin et al. Protein arginine methylation: from enigmatic functions to therapeutic targeting, *NAT REV DRUG DISCOV* 20: 509-530

EUbOPEN: Desta, Zeruesenay et al. PharmVar GeneFocus: CYP2B6, *CLIN PHARMACOL THER* 110: 82-97

EUbOPEN: Perveen, Sumera et al. A High-Throughput RNA Displacement Assay for Screening SARS-CoV-2 nsp10-nsp16 Complex toward Developing Therapeutics for COVID-19, *SLAS DISCOV* 26: 620-627

EUbOPEN: Ishida, Tasuku et al. E3 Ligase Ligands for PROTACs: How They Were Found and How to Discover New Ones, *SLAS DISCOV* 26: 484-502

EUbOPEN: Ni, Xiaomin et al. Structural Insights into Plasticity and Discovery of Remdesivir Metabolite GS-441524 Binding in SARS-CoV-2 Macrodomain, *ACS MED CHEM LETT* 12: 603-609

EUbOPEN: Fossati, Andrea et al. PCprophet: a framework for protein complex prediction and differential analysis using proteomic data, *NAT METHODS* 18: 520-+

EUbOPEN: Williams, Eleanor et al. Saracatinib is an efficacious clinical candidate for fibrodysplasia ossificans progressiva, *JCI INSIGHT* 6:

EUbOPEN: Yazdi, Aliakbar Khalili et al. A High-Throughput Radioactivity-Based Assay for Screening SARS-CoV-2 nsp10-nsp16 Complex, *SLAS DISCOV* 26: 757-765

EUbOPEN: Ingelman-Sundberg, Magnus et al. 3D human liver spheroids for translational pharmacology and toxicology, *BASIC CLIN PHARMACOL* 130: 5-15

EUbOPEN: Ramachandran, Sarath et al. Building ubiquitination machineries: E3 ligase multi-subunit assembly and substrate targeting by PROTACs and molecular glues, *CURR OPIN STRUC BIOL* 67: 110-119

EUbOPEN: Russell, Laura E. et al. Pharmacogenomics in the era of next generation sequencing - from byte to bedside, *DRUG METAB REV* 53: 253-278

EUbOPEN: Berger, Benedict-Tilman et al. Structure-kinetic relationship reveals the mechanism of selectivity of FAK inhibitors over PYK2, *CELL CHEM BIOL* 28: 686-+

EUbOPEN: Devkota, Kanchan et al. Probing the SAM Binding Site of SARS-CoV-2 Nsp14 In Vitro Using SAM Competitive Inhibitors Guides Developing Selective Bisubstrate Inhibitors, *SLAS DISCOV* 26: 1200-1211

EuBOPEN: Attwood, Misty M. et al. Trends in kinase drug discovery: targets, indications and inhibitor design, *NAT REV DRUG DISCOV* 20: 839-861

EuBOPEN: Schmidt, Sven H. et al. Conformation and dynamics of the kinase domain drive subcellular location and activation of LRRK2, *P NATL ACAD SCI USA* 118:

EuBOPEN: Otava, Tomas et al. The Structure-Based Design of SARS-CoV-2 nsp14 Methyltransferase Ligands Yields Nanomolar Inhibitors, *ACS INFECT DIS* 7: 2214-2220

EU-PEARL: Stallard, Nigel et al. Efficient Adaptive Designs for Clinical Trials of Interventions for COVID-19, *STAT BIOPHARM RES* 12: 483-497

EU-PEARL: Lugo-Marin, Jorge et al. COVID-19 pandemic effects in people with Autism Spectrum Disorder and their caregivers: Evaluation of social distancing and lockdown impact on mental health and general status, *RES AUTISM SPECT DIS* 83:

EUROPAIN: Sikandar, Shafaq et al. Visceral pain: the ins and outs, the ups and downs, *CURR OPIN SUPPORT PA* 6: 17-26

EUROPAIN: Aasvang, Eske K. et al. Predictive Risk Factors for Persistent Postherniotomy Pain, *ANESTHESIOLOGY* 112: 957-969

EUROPAIN: Finnerup, Nanna Brix et al. The evidence for pharmacological treatment of neuropathic pain, *PAIN* 150: 573-581

EUROPAIN: Baastrup, Cathrine et al. Spinal-, brainstem- and cerebrally mediated responses at- and below-level of a spinal cord contusion in rats: Evaluation of pain-like behavior, *PAIN* 151: 670-679

EUROPAIN: Wildgaard, K. et al. Consequences of persistent pain after lung cancer surgery: a nationwide questionnaire study, *ACTA ANAESTH SCAND* 55: 60-68

EUROPAIN: Phillips, Tudor J. C. et al. Pharmacological Treatment of Painful HIV-Associated Sensory Neuropathy: A Systematic Review and Meta-Analysis of Randomised Controlled Trials, *PLOS ONE* 5:

EUROPAIN: Marinus, Johan et al. Clinical features and pathophysiology of complex regional pain syndrome, *LANCET NEUROL* 10: 637-648

EUROPAIN: Andersen, Kenneth Geving et al. Persistent Pain After Breast Cancer Treatment: A Critical Review of Risk Factors and Strategies for Prevention, *J PAIN* 12: 725-746

EUROPAIN: Dawes, John M. et al. CXCL5 Mediates UVB Irradiation-Induced Pain, *SCI TRANSL MED* 3:

EUROPAIN: Serra, Jordi et al. Microneurographic identification of spontaneous activity in C-nociceptors in neuropathic pain states in humans and rats, *PAIN* 153: 42-55

EUROPAIN: Haeuser, Winfried et al. The Role of Antidepressants in the Management of Fibromyalgia Syndrome A Systematic Review and Meta-Analysis, *CNS DRUGS* 26: 297-307

EUROPAIN: Finnerup, Nanna Brix et al. Spinal Cord Injury Pain: Mechanisms and Management, *CURR PAIN HEADACHE R* 16: 207-216

EUROPAIN: Petersen, Gitte Laue et al. Placebo manipulations reduce hyperalgesia in neuropathic pain, PAIN 153: 1292-1300

EUROPAIN: Andrews, N. et al. Spontaneous burrowing behaviour in the rat is reduced by peripheral nerve injury or inflammation associated pain, EUR J PAIN 16: 485-495

EUROPAIN: Calvo, Margarita et al. The role of the immune system in the generation of neuropathic pain, LANCET NEUROL 11: 629-642

EUROPAIN: Quick, Kathryn et al. TRPC3 and TRPC6 are essential for normal mechanotransduction in subsets of sensory neurons and cochlear hair cells, OPEN BIOL 2:

EUROPAIN: Baron, Ralf et al. Subgrouping of patients with neuropathic pain according to pain-related sensory abnormalities: a first step to a stratified treatment approach, LANCET NEUROL 11: 999-1005

EUROPAIN: Haroutiunian, Simon et al. The neuropathic component in persistent postsurgical pain: A systematic literature review, PAIN 154: 95-102

EUROPAIN: Derry, Sheena et al. Topical capsaicin (high concentration) for chronic neuropathic pain in adults, COCHRANE DB SYST REV :

EUROPAIN: Huang, Wenlong et al. A clinically relevant rodent model of the HIV antiretroviral drug stavudine induced painful peripheral neuropathy, PAIN 154: 560-575

EUROPAIN: Mejdahl, Mathias Kvist et al. Persistent pain and sensory disturbances after treatment for breast cancer: six year nationwide follow-up study, BMJ-BRIT MED J 346:

EUROPAIN: Eijkelkamp, N. et al. A role for Piezo2 in EPAC1-dependent mechanical allodynia, NAT COMMUN 4:

EUROPAIN: Ellis, A. et al. Neuroinflammation and the generation of neuropathic pain, BRIT J ANAESTH 111: 26-37

EUROPAIN: Sikandar, Shafaq et al. Neural coding of nociceptive stimuli-from rat spinal neurones to human perception, PAIN 154: 1263-1273

EUROPAIN: Denk, Franziska et al. HDAC inhibitors attenuate the development of hypersensitivity in models of neuropathic pain, PAIN 154: 1668-1679

EUROPAIN: Dworkin, Robert H. et al. Interventional management of neuropathic pain: NeuPSIG recommendations, PAIN 154: 2249-2261

EUROPAIN: Gilron, Ian et al. Combination pharmacotherapy for management of chronic pain: from bench to bedside, LANCET NEUROL 12: 1084-1095

EUROPAIN: Rutten, K. et al. Burrowing as a non-reflex behavioural readout for analgesic action in a rat model of sub-chronic knee joint inflammation, EUR J PAIN 18: 204-212

EUROPAIN: Minett, Michael S. et al. Pain without Nociceptors? Nav1.7-Independent Pain Mechanisms, CELL REP 6: 301-312

EUROPAIN: Serra, Jordi et al. Hyperexcitable C nociceptors in fibromyalgia, ANN NEUROL 75: 196-208

EUROPAIN: Haroutounian, Simon et al. Primary afferent input critical for maintaining spontaneous pain in peripheral neuropathy, PAIN 155: 1272-1279

EUROPAIN: Petersen, Gitte Laue et al. The magnitude of placebo effects in pain: A meta-analysis, PAIN 155: 1426-1434

EUROPAIN: Jensen, Troels S. et al. Allodynia and hyperalgesia in neuropathic pain: clinical manifestations and mechanisms, LANCET NEUROL 13: 924-935

EUROPAIN: Gierthmuehlen, Janne et al. Mechanism-based treatment in complex regional pain syndromes, NAT REV NEUROL 10: 518-528

EUROPAIN: Caspani, Ombretta et al. Tramadol reduces anxiety-related and depression-associated behaviors presumably induced by pain in the chronic constriction injury model of neuropathic pain in rats, PHARMACOL BIOCHEM BE 124: 290-296

EUROPAIN: Demant, Dyveke T. et al. The effect of oxcarbazepine in peripheral neuropathic pain depends on pain phenotype: A randomised, double-blind, placebo-controlled phenotype-stratified study, PAIN 155: 2263-2273

EUROPAIN: Petersen, Gitte L. et al. Expectations and positive emotional feelings accompany reductions in ongoing and evoked neuropathic pain following placebo interventions, PAIN 155: 2687-2698

EUROPAIN: Sisignano, Marco et al. Mechanism-based treatment for chemotherapy-induced peripheral neuropathic pain, NAT REV NEUROL 10: 694-707

EUROPAIN: Segerdahl, Andrew R. et al. The dorsal posterior insula subserves a fundamental role in human pain, NAT NEUROSCI 18: 499-+

EUROPAIN: Treede, Rolf-Detlef et al. A classification of chronic pain for ICD-11, PAIN 156: 1003-1007

EUROPAIN: Vase, Lene et al. Predictors of the placebo analgesia response in randomized controlled trials of chronic pain: a meta-analysis of the individual data from nine industrially sponsored trials, PAIN 156: 1795-1802

EUROPAIN: Gierthmuehlen, Janne et al. Who is healthy? Aspects to consider when including healthy volunteers in QST-based studies-a consensus statement by the EUROPAIN and NEUROPPAIN consortia, PAIN 156: 2203-2211

EUROPAIN: Demant, Dyveke T. et al. Pain relief with lidocaine 5% patch in localized peripheral neuropathic pain in relation to pain phenotype: a randomised, double-blind, and placebo-controlled, phenotype panel study, PAIN 156: 2234-2244

EUROPAIN: van Hecke, Oliver et al. Neuropathic pain phenotyping by international consensus (NeuroPPIC) for genetic studies: a NeuPSIG systematic review, Delphi survey, and expert panel recommendations, PAIN 156: 2337-2353

EUROPAIN: Andersen, Kenneth Geving et al. Predictive factors for the development of persistent pain after breast cancer surgery, PAIN 156: 2413-2422

EUROPAIN: Wijayasinghe, Nelun et al. Ultrasound Guided Intercostobrachial Nerve Blockade in Patients with Persistent Pain after Breast Cancer Surgery: A Pilot Study, PAIN PHYSICIAN 19: E309-E317

EUROPAIN: Wildgaard, K. et al. Persistent postsurgical pain after video-assisted thoracic surgery - an observational study, ACTA ANAESTH SCAND 60: 650-658

EUROPAIN: McDonnell, Aoibhinn et al. Inherited erythromelalgia due to mutations in SCN9A: natural history, clinical phenotype and somatosensory profile, BRAIN 139: 1052-1065

EUROPAIN: Ventzel, Lise et al. Chemotherapy-induced pain and neuropathy: a prospective study in patients treated with adjuvant oxaliplatin or docetaxel, PAIN 157: 560-568

EUROPAIN: Vollert, Jan et al. Quantitative sensory testing using DFNS protocol in Europe: an evaluation of heterogeneity across multiple centers in patients with peripheral neuropathic pain and healthy subjects, PAIN 157: 750-758

EUROPAIN: Kosek, Eva et al. Do we need a third mechanistic descriptor for chronic pain states?, PAIN 157: 1382-1386

EUROPAIN: Finnerup, Nanna B. et al. Neuropathic pain: an updated grading system for research and clinical practice, PAIN 157: 1599-1606

EUROPAIN: Wodarski, Rachel et al. Cross-centre replication of suppressed burrowing behaviour as an ethologically relevant pain outcome measure in the rat: a prospective multicentre study, PAIN 157: 2350-2365

EUROPAIN: Colloca, Luana et al. Neuropathic pain, NAT REV DIS PRIMERS 3:

EUROPAIN: Kemp, Harriet I. et al. Use of Corneal Confocal Microscopy to Evaluate Small Nerve Fibers in Patients With Human Immunodeficiency Virus, JAMA OPHTHALMOL 135: 795-799

EUROPAIN: Vollert, Jan et al. Stratifying patients with peripheral neuropathic pain based on sensory profiles: algorithm and sample size recommendations, PAIN 158: 1446-1455

EUROPAIN: Segerdahl, Andrew R. et al. A brain-based pain facilitation mechanism contributes to painful diabetic polyneuropathy, BRAIN 141: 357-364

EUROPAIN: Forstenpointner, Julia et al. Individualized neuropathic pain therapy based on phenotyping: are we there yet?, PAIN 159: 569-575

EUROPAIN: Wanigasekera, V. et al. Disambiguating pharmacological mechanisms from placebo in neuropathic pain using functional neuroimaging, BRIT J ANAESTH 120: 299-307

EUROPAIN: Vollert, Jan et al. Pathophysiological mechanisms of neuropathic pain: comparison of sensory phenotypes in patients and human surrogate pain models, PAIN 159: 1090-1102

EUROPAIN: Finnerup, Nanna B. et al. Neuropathic pain clinical trials: factors associated with decreases in estimated drug efficacy, PAIN 159: 2339-2346



EUROPAIN: Baskozos, Georgios et al. Comprehensive analysis of long noncoding RNA expression in dorsal root ganglion reveals cell-type specificity and dysregulation after nerve injury, PAIN 160: 463-485

EUROPAIN: Treede, Rolf-Detlef et al. The role of quantitative sensory testing in the prediction of chronic pain, PAIN 160: S66-S69

EUROPAIN: Bannister, Kirsty et al. Neuropathic Pain: Mechanism-Based Therapeutics, ANNU REV PHARMACOL 60: 257-274

EUROPAIN: Forstenpointner, Julia et al. No pain, still gain (of function): the relation between sensory profiles and the presence or absence of self-reported pain in a large multicenter cohort of patients with neuropathy, PAIN 162: 718-727

EUROPAIN: Kosek, Eva et al. Chronic nociplastic pain affecting the musculoskeletal system: clinical criteria and grading system, PAIN 162: 2629-2634

FAIRplus: Rodriguez-Espigares, Ismael et al. GPCRmd uncovers the dynamics of the 3D-GPCRome, NAT METHODS 17: 777-+

FAIRplus: Bernal-Llinares, Manuel et al. Identifiers.org: Compact Identifier services in the cloud, BIOINFORMATICS 37: 1781-1782

FLUCOP: Sridhar, Saranya et al. Influenza Vaccination Strategies: Comparing Inactivated and Live Attenuated Influenza Vaccines, VACCINES 3: 373-389

FLUCOP: Pebody, R. et al. Effectiveness of seasonal influenza vaccine for adults and children in preventing laboratory-confirmed influenza in primary care in the United Kingdom: 2015/16 end-of-season results, EUROSURVEILLANCE 21: 41-51

FLUCOP: de Vries, Rory D. et al. Influenza virus-specific antibody dependent cellular cytotoxicity induced by vaccination or natural infection, VACCINE 35: 238-247

FLUCOP: Mohn, Kristin G. I. et al. Boosting of Cross-Reactive and Protection-Associated T Cells in Children After Live Attenuated Influenza Vaccination, J INFECT DIS 215: 1527-1535

FLUCOP: Mohn, Kristin G. -I. et al. Immune responses after live attenuated influenza vaccination, HUM VACC IMMUNOTHER 14: 571-578

FLUCOP: Trieu, Mai-Chi et al. SARS-CoV-2-Specific Neutralizing Antibody Responses in Norwegian Health Care Workers After the First Wave of COVID-19 Pandemic: A Prospective Cohort Study, J INFECT DIS 223: 589-599

FLUCOP: Blomberg, Bjorn et al. Long COVID in a prospective cohort of home-isolated patients, NAT MED 27: 1607-+

GETREAL: Nordon, Clementine et al. The Efficacy-Effectiveness Gap : Historical Background and Current Conceptualization, VALUE HEALTH 19: 75-81

GETREAL: Efthimiou, Orestis et al. GetReal in network meta-analysis: a review of the methodology, RES SYNTH METHODS 7: 236-263

GETREAL: Efthimiou, Orestis et al. Combining randomized and nonrandomized evidence in network meta-analysis, *STAT MED* 36: 1210-1226

GETREAL: Makady, Amr et al. Policies for Use of Real-World Data in Health Technology Assessment (HTA): A Comparative Study of Six HTA Agencies, *VALUE HEALTH* 20: 520-532

GETREAL: Makady, Amr et al. What Is Real-World Data? A Review of Definitions Based on Literature and Stakeholder Interviews, *VALUE HEALTH* 20: 858-865

GETREAL: Zuidgeest, Mira G. P. et al. Series: Pragmatic trials and real world evidence: Paper 1. Introduction, *J CLIN EPIDEMIOL* 88: 7-13

GETREAL: Rengerink, Katrien Oude et al. Series: Pragmatic trials and real world evidence: Paper 3. Patient selection challenges and consequences, *J CLIN EPIDEMIOL* 89: 173-180

GETREAL: Debray, Thomas P. A. et al. An overview of methods for network meta-analysis using individual participant data: when do benefits arise?, *STAT METHODS MED RES* 27: 1351-1364

GETREAL: Van den Berg, Sanne J. P. et al. Cross-species extrapolation of chemical sensitivity, *SCI TOTAL ENVIRON* 753:

HARMONY: Malcikova, J. et al. ERIC recommendations for TP53 mutation analysis in chronic lymphocytic leukemia-update on methodological approaches and results interpretation, *LEUKEMIA* 32: 1070-1080

HARMONY PLUS: Barbui, Tiziano et al. Among classic myeloproliferative neoplasms, essential thrombocythemia is associated with the greatest risk of venous thromboembolism during COVID-19, *BLOOD CANCER J* 11:

HIPPOCRATES: Simon, David et al. Humoral and Cellular Immune Responses to SARS-CoV-2 Infection and Vaccination in Autoimmune Disease Patients With B Cell Depletion, *ARTHRITIS RHEUMATOL* 74: 33-37

HIPPOCRATES: Simon, David et al. Association of Structural Enteseal Lesions With an Increased Risk of Progression From Psoriasis to Psoriatic Arthritis, *ARTHRITIS RHEUMATOL* 74: 253-262

Hypo-RESOLVE: Cherkas, Andriy et al. Glucose as a Major Antioxidant: When, What for and Why It Fails?, *ANTIOXIDANTS-BASEL* 9:

Hypo-RESOLVE: Chatwin, Hannah et al. The impact of hypoglycaemia on quality of life outcomes among adults with type 1 diabetes: A systematic review, *DIABETES RES CLIN PR* 174:

iABC: Schaedle, Thomas et al. Mid-Infrared Waveguides: A Perspective, *APPL SPECTROSC* 70: 1625-1638

iABC: Aliberti, Stefano et al. Research priorities in bronchiectasis: a consensus statement from the EMBARC Clinical Research Collaboration, *EUR RESPIR J* 48: 632-647

iABC: Loebinger, Michael R. et al. Efficacy and safety of TOBI Podhaler in *Pseudomonas aeruginosa*-infected bronchiectasis patients: iBEST study, *EUR RESPIR J* 57:

iABC: Diez-Aguilar, Maria et al. Murepavadin antimicrobial activity against and resistance development in cystic fibrosis *Pseudomonas aeruginosa* isolates, *J ANTIMICROB CHEMOTH* 76: 984-992

iABC: Crichton, Megan L. et al. Validation of the Bronchiectasis Impact Measure (BIM): a novel patient-reported outcome measure, *EUR RESPIR J* 57:

iCONSENSUS: Pinto, Ines F. et al. Multiplexed Microfluidic Cartridge for At-Line Protein Monitoring in Mammalian Cell Culture Processes for Biopharmaceutical Production, *ACS SENSORS* 6: 842-851

IDEA-FAST: Faundez-Zanuy, Marcos et al. Handwriting Biometrics: Applications and Future Trends in e-Security and e-Health, *COGN COMPUT* 12: 940-953

IM2PACT: Goldman, C. et al. Human induced pluripotent stem cells (BIONi010-C) generate tight cell monolayers with blood-brain barrier traits and functional expression of large neutral amino acid transporter 1 (SLC7A5), *EUR J PHARM SCI* 156:

IM2PACT: Mae, Maarja A. et al. Single-Cell Analysis of Blood-Brain Barrier Response to Pericyte Loss, *CIRC RES* 128: E46-E62

IM2PACT: Hudak, Anett et al. Contribution of Syndecans to the Cellular Entry of SARS-CoV-2, *INT J MOL SCI* 22:

IMIDIA: Marciniak, Anja et al. Using pancreas tissue slices for in situ studies of islet of Langerhans and acinar cell biology, *NAT PROTOC* 9: 2809-2822

IMIDIA: Roggli, Elodie et al. Involvement of MicroRNAs in the Cytotoxic Effects Exerted by Proinflammatory Cytokines on Pancreatic beta-Cells, *DIABETES* 59: 978-986

IMIDIA: Gonzalez, Claudio D. et al. The emerging role of autophagy in the pathophysiology of diabetes mellitus, *AUTOPHAGY* 7: 2-11

IMIDIA: Woodfin, Abigail et al. The junctional adhesion molecule JAM-C regulates polarized transendothelial migration of neutrophils in vivo, *NAT IMMUNOL* 12: 761-U145

IMIDIA: Ravassard, Philippe et al. A genetically engineered human pancreatic beta cell line exhibiting glucose-inducible insulin secretion, *J CLIN INVEST* 121: 3589-3597

IMIDIA: Santiago, Marcelo F. et al. Targeting Pannexin1 Improves Seizure Outcome, *PLOS ONE* 6:

IMIDIA: Bosco, Domenico et al. CONNEXINS: KEY MEDIATORS OF ENDOCRINE FUNCTION, *PHYSIOL REV* 91: 1393-1445

IMIDIA: Roggli, Elodie et al. Changes in MicroRNA Expression Contribute to Pancreatic beta-Cell Dysfunction in Prediabetic NOD Mice, *DIABETES* 61: 1742-1751

IMIDIA: Hodson, David J. et al. Lipotoxicity disrupts incretin-regulated human beta cell connectivity, *J CLIN INVEST* 123: 4182-4194

IMIDIA: Huch, Meritxell et al. Unlimited in vitro expansion of adult bi-potent pancreas progenitors through the Lgr5/R-spondin axis, *EMBO J* 32: 2708-2721

IMIDIA: Marselli, Lorella et al. Are we overestimating the loss of beta cells in type 2 diabetes?, *DIABETOLOGIA* 57: 362-365

IMIDIA: Scharfmann, Raphael et al. Development of a conditionally immortalized human pancreatic beta cell line, *J CLIN INVEST* 124: 2087-2098

IMIDIA: Lenzen, Sigurd et al. A Fresh View of Glycolysis and Glucokinase Regulation: History and Current Status, *J BIOL CHEM* 289: 12189-12194

IMIDIA: Hodson, David J. et al. ADCY5 Couples Glucose to Insulin Secretion in Human Islets, *DIABETES* 63: 3009-3021

IMIDIA: Chabosseau, Pauline et al. Mitochondrial and ER-Targeted eCALWY Probes Reveal High Levels of Free Zn<sup>2+</sup>, *ACS CHEM BIOL* 9: 2111-2120

IMIDIA: Broichhagen, Johannes et al. Optical control of insulin release using a photoswitchable sulfonylurea, *NAT COMMUN* 5:

IMIDIA: Rutter, Guy A. et al. SLC30A8 mutations in type 2 diabetes, *DIABETOLOGIA* 58: 31-36

IMIDIA: Mitchell, Ryan K. et al. Selective disruption of Tcf7l2 in the pancreatic beta cell impairs secretory function and lowers beta cell mass, *HUM MOL GENET* 24: 1390-1399

IMIDIA: Rutter, Guy A. et al. Pancreatic beta-cell identity, glucose sensing and the control of insulin secretion, *BIOCHEM J* 466: 203-218

IMIDIA: Johnston, Natalie R. et al. Beta Cell Hubs Dictate Pancreatic Islet Responses to Glucose, *CELL METAB* 24: 389-401

IMIDIA: Chabosseau, Pauline et al. Zinc and diabetes, *ARCH BIOCHEM BIOPHYS* 611: 79-85

IMIDIA: Chakravarthy, Harini et al. Converting Adult Pancreatic Islet alpha Cells into beta Cells by Targeting Both Dnmt1 and Arx, *CELL METAB* 25: 622-634

IMIDIA: Wigger, Leonore et al. Plasma Dihydroceramides Are Diabetes Susceptibility Biomarker Candidates in Mice and Humans, *CELL REP* 18: 2269-2279

IMIDIA: Gerber, Philipp A. et al. The Role of Oxidative Stress and Hypoxia in Pancreatic Beta-Cell Dysfunction in Diabetes Mellitus, *ANTIOXID REDOX SIGN* 26: 501-+

IMIDIA: Cohrs, Christian M. et al. Vessel Network Architecture of Adult Human Islets Promotes Distinct Cell-Cell Interactions In Situ and Is Altered After Transplantation, *ENDOCRINOLOGY* 158: 1373-1385

IMIDIA: Solimena, Michele et al. Systems biology of the IMIDIA biobank from organ donors and pancreatectomised patients defines a novel transcriptomic signature of islets from individuals with type 2 diabetes, *DIABETOLOGIA* 61: 641-657

IMIDIA: Fine, Nicholas H. F. et al. Glucocorticoids Reprogram beta-Cell Signaling to Preserve Insulin Secretion, *DIABETES* 67: 278-290

IMIDIA: Cohrs, Christian M. et al. Dysfunction of Persisting beta Cells Is a Key Feature of Early Type 2 Diabetes Pathogenesis, *CELL REP* 31:

IMI-PainCare: Treede, Rolf-Detlef et al. The role of quantitative sensory testing in the prediction of chronic pain, *PAIN* 160: S66-S69

IMI-PainCare: Quesada, Charles et al. Human surrogate models of central sensitization: A critical review and practical guide, *EUR J PAIN* 25: 1389-1428

IMI-PainCare: Kosek, Eva et al. Chronic nociplastic pain affecting the musculoskeletal system: clinical criteria and grading system, *PAIN* 162: 2629-2634

IMMUCAN: Sompairac, Nicolas et al. Independent Component Analysis for Unraveling the Complexity of Cancer Omics Datasets, *INT J MOL SCI* 20:

Immune-Image: Foray, Claudia et al. Imaging temozolomide-induced changes in the myeloid glioma microenvironment, *THERANOSTICS* 11: 2020-2033

Immune-Image: Barca, Cristina et al. Impact of hydroxytyrosol on stroke: tracking therapy response on neuroinflammation and cerebrovascular parameters using PET-MR imaging and on functional outcomes, *THERANOSTICS* 11: 4030-4049

IMPRiND: Fitzpatrick, Anthony W. P. et al. Cryo-EM structures of tau filaments from Alzheimers disease, *NATURE* 547: 185-+

IMPRiND: Tofaris, George K. et al. A Critical Assessment of Exosomes in the Pathogenesis and Stratification of Parkinsons Disease, *J PARKINSON DIS* 7: 569-576

IMPRiND: McInnes, Joseph et al. Synaptogyrin-3 Mediates Presynaptic Dysfunction Induced by Tau, *NEURON* 97: 823-+

IMPRiND: Peelaerts, W. et al. E-Synuclein strains and seeding in Parkinsons disease, incidental Lewy body disease, dementia with Lewy bodies and multiple system atrophy: similarities and differences, *CELL TISSUE RES* 373: 195-212

IMPRiND: Kundel, Franziska et al. Measurement of Tau Filament Fragmentation Provides Insights into Prion-like Spreading, *ACS CHEM NEUROSCI* 9: 1276-1282

IMPRiND: Wilkinson, Mark D. et al. Comment: A design framework and exemplar metrics for FAIRness, *SCI DATA* 5:

IMPRiND: Falcon, Benjamin et al. Structures of filaments from Picks disease reveal a novel tau protein fold, *NATURE* 561: 137-+

IMPRiND: Jucker, Mathias et al. Propagation and spread of pathogenic protein assemblies in neurodegenerative diseases, *NAT NEUROSCI* 21: 1341-1349

IMPRiND: Falcon, Benjamin et al. Tau filaments from multiple cases of sporadic and inherited Alzheimers disease adopt a common fold, *ACTA NEUROPATHOL* 136: 699-708

IMPRiND: Shrivastava, Amulya Nidhi et al. Clustering of Tau fibrils impairs the synaptic composition of alpha 3-Na+/K+-ATPase and AMPA receptors, *EMBO J* 38:

IMPRiND: Gribaudo, Simona et al. Propagation of alpha-Synuclein Strains within Human Reconstructed Neuronal Network, *STEM CELL REP* 12: 230-244

IMPRiND: Zhang, Wenjuan et al. Heparin-induced tau filaments are polymorphic and differ from those in Alzheimers and Picks diseases, *ELIFE* 8:

IMPRiND: Falcon, Benjamin et al. Novel tau filament fold in chronic traumatic encephalopathy encloses hydrophobic molecules, *NATURE* 568: 420-+

IMPRiND: Vasili, Eftychia et al. Spreading of alpha-Synuclein and Tau: A Systematic Comparison of the Mechanisms Involved, *FRONT MOL NEUROSCI* 12:

IMPRiND: Bieri, Gregor et al. LRRK2 modifies alpha-syn pathology and spread in mouse models and human neurons, *ACTA NEUROPATHOL* 137: 961-980

IMPRiND: Fenyi, Alexis et al. Detection of alpha-synuclein aggregates in gastrointestinal biopsies by protein misfolding cyclic amplification, *NEUROBIOL DIS* 129: 38-43

IMPRiND: Alam, Parvez et al. alpha-synuclein oligomers and fibrils: a spectrum of species, a spectrum of toxicities, *J NEUROCHEM* 150: 522-534

IMPRiND: Mavroedi, Panagiota et al. Endogenous oligodendroglial alpha-synuclein and TPPP/p25 alpha orchestrate alpha-synuclein pathology in experimental multiple system atrophy models, *ACTA NEUROPATHOL* 138: 415-441

IMPRiND: Roesler, Thomas W. et al. Four-repeat tauopathies, *PROG NEUROBIOL* 180:

IMPRiND: Rey, Nolwen L. et al. alpha-Synuclein conformational strains spread, seed and target neuronal cells differentially after injection into the olfactory bulb, *ACTA NEUROPATHOL COM* 7:

IMPRiND: Guerrero-Ferreira, Ricardo et al. Two new polymorphic structures of human full-length alpha-synuclein fibrils solved by cryo-electron microscopy, *ELIFE* 8:

IMPRiND: Shrivastava, Amulya Nidhi et al. Differential Membrane Binding and Seeding of Distinct alpha-Synuclein Fibrillar Polymorphs, *BIOPHYS J* 118: 1301-1320

IMPRiND: Zhang, Wenjuan et al. Novel tau filament fold in corticobasal degeneration, *NATURE* 580: 283-+

IMPRiND: Shrivastava, Amulya Nidhi et al. Cell biology and dynamics of Neuronal Na<sup>+</sup>/K<sup>+</sup>-ATPase in health and diseases, *NEUROPHARMACOLOGY* 169:

IMPRiND: Van der Perren, Anke et al. The structural differences between patient-derived alpha-synuclein strains dictate characteristics of Parkinsons disease, multiple system atrophy and dementia with Lewy bodies, *ACTA NEUROPATHOL* 139: 977-1000

IMPRiND: Teil, Margaux et al. Targeting alpha-Synuclein for PD Therapeutics: A Pursuit on All Fronts, *BIOMOLECULES* 10:

IMPRiND: Nachman, Eliana et al. Disassembly of Tau fibrils by the human Hsp70 disaggregation machinery generates small seeding-competent species, *J BIOL CHEM* 295: 9676-9690

IMPRiND: Courte, Josquin et al. The expression level of alpha-synuclein in different neuronal populations is the primary determinant of its prion-like seeding, *SCI REP-UK* 10:

IMPRiND: Shi, Yang et al. Cryo-EM structures of tau filaments from Alzheimers disease with PET ligand APN-1607, *ACTA NEUROPATHOL* 141: 697-708

IMPRiND: Russ, Kaspar et al. TNF-alpha and alpha-synuclein fibrils differently regulate human astrocyte immune reactivity and impair mitochondrial respiration, *CELL REP* 34:

IMPRiND: Shi, Yang et al. Structure-based classification of tauopathies, *NATURE* 598: 359-+

imSAVAR: Prommersberger, Sabrina et al. CARAMBA: a first-in-human clinical trial with SLAMF7 CAR-T cells prepared by virus-free Sleeping Beauty gene transfer to treat multiple myeloma, *GENE THER* 28: 560-571

imSAVAR: Maulana, Tengku Ibrahim et al. Immunocompetent cancer-on-chip models to assess immuno-oncology therapy, *ADV DRUG DELIVER REV* 173: 281-305

INNODIA: Grieco, Fabio Arturo et al. MicroRNAs miR-23a-3p, miR-23b-3p, and miR-149-5p Regulate the Expression of Proapoptotic BH3-Only Proteins DP5 and PUMA in Human Pancreatic beta-Cells, *DIABETES* 66: 100-112

INNODIA: Schwandt, Anke et al. Longitudinal Trajectories of MetabolicControl From Childhood to Young Adulthood in Type 1 Diabetes From a Large German/Austrian Registry: A Group-Based Modeling Approach, *DIABETES CARE* 40: 309-316

INNODIA: Marroqui, Laura et al. Interferon-alpha mediates human beta cell HLA class I overexpression, endoplasmic reticulum stress and apoptosis, three hallmarks of early human type 1 diabetes, *DIABETOLOGIA* 60: 656-667

INNODIA: Schwandt, Anke et al. Comparison of MDRD, CKD-EPI, and Cockcroft-Gault equation in relation to measured glomerular filtration rate among a large cohort with diabetes, *J DIABETES COMPLICAT* 31: 1376-1383

INNODIA: Charalampopoulos, Dimitrios et al. Exploring Variation in Glycemic Control Across and Within Eight High-Income Countries: A Cross-sectional Analysis of 64,666 Children and Adolescents With Type 1 Diabetes, *DIABETES CARE* 41: 1180-1187

INNODIA: Culina, Slobodan et al. Islet-reactive CD8(+) T cell frequencies in the pancreas, but not in blood, distinguish type 1 diabetic patients from healthy donors, *SCI IMMUNOL* 3:

INNODIA: Yeo, Lorraine et al. Autoreactive T effector memory differentiation mirrors beta cell function in type 1 diabetes, *J CLIN INVEST* 128: 3460-3474

INNODIA: DeSalvo, Daniel J. et al. Continuous glucose monitoring and glycemic control among youth with type 1 diabetes: International comparison from the T1D Exchange and DPV Initiative, *PEDIATR DIABETES* 19: 1271-1275

INNODIA: Colli, Maikel L. et al. PDL1 is expressed in the islets of people with type 1 diabetes and is up-regulated by interferons-alpha and-gamma via IRF1 induction, *EBIOMEDICINE* 36: 367-375

INNODIA: Gonzalez-Duque, Sergio et al. Conventional and Neo-antigenic Peptides Presented by beta Cells Are Targeted by Circulating Naive CD8+T Cells in Type 1 Diabetic and Healthy Donors, CELL METAB 28: 946-+

INNODIA: Balboa, Diego et al. Insulin mutations impair beta-cell development in a patient-derived iPSC model of neonatal diabetes, ELIFE 7:

INNODIA: Atkinson, Mark A. et al. The challenge of modulating beta-cell autoimmunity in type 1 diabetes, LANCET DIABETES ENDO 7: 52-64

INNODIA: Roep, Bart O. et al. Antigen-based immune modulation therapy for type 1 diabetes: the era of precision medicine, LANCET DIABETES ENDO 7: 65-74

INNODIA: Balboa, Diego et al. Concise Review: Human Pluripotent Stem Cells for the Modeling of Pancreatic beta-Cell Pathology, STEM CELLS 37: 33-41

INNODIA: Henriksson, Johan et al. Genome-wide CRISPR Screens in T Helper Cells Reveal Pervasive Crosstalk between Activation and Differentiation, CELL 176: 882-+

INNODIA: Reinehr, Thomas et al. Worse Metabolic Control and Dynamics of Weight Status in Adolescent Girls Point to Eating Disorders in the First Years after Manifestation of Type 1 Diabetes Mellitus: Findings from the Diabetes Patienten Verlaufsdokumentation Registry, J PEDIATR-US 207: 205-212

INNODIA: Danne, Thomas et al. International Consensus on Risk Management of Diabetic Ketoacidosis in Patients With Type 1 Diabetes Treated With Sodium-Glucose Cotransporter (SGLT) Inhibitors, DIABETES CARE 42: 1147-1154

INNODIA: Hermann, J. M. et al. The Transatlantic HbA(1c) gap: differences in glycaemic control across the lifespan between people included in the US T1D Exchange Registry and those included in the German/Austrian DPV registry, DIABETIC MED 37: 848-855

INNODIA: Mirza, Aashiq H. et al. Breast Milk-Derived Extracellular Vesicles Enriched in Exosomes From Mothers With Type 1 Diabetes Contain Aberrant Levels of microRNAs, FRONT IMMUNOL 10:

INNODIA: Ramos-Rodriguez, Mireia et al. The impact of proinflammatory cytokines on the beta-cell regulatory landscape provides insights into the genetics of type 1 diabetes, NAT GENET 51: 1588-+

INNODIA: Boss, Marti et al. PET-Based Human Dosimetry of Ga-68-NODAGA-Exendin-4, a Tracer for beta-Cell Imaging, J NUCL MED 61: 112-116

INNODIA: Nakayasu, Ernesto S. et al. Comprehensive Proteomics Analysis of Stressed Human Islets Identifies GDF15 as a Target for Type 1 Diabetes Intervention, CELL METAB 31: 363-+

INNODIA: Demine, Stephane et al. Pro-inflammatory cytokines induce cell death, inflammatory responses, and endoplasmic reticulum stress in human iPSC-derived beta cells, STEM CELL RES THER 11:

INNODIA: Cohrs, Christian M. et al. Dysfunction of Persisting beta Cells Is a Key Feature of Early Type 2 Diabetes Pathogenesis, CELL REP 31:



INNODIA: Anderzen, Johan et al. International benchmarking in type 1 diabetes: Large difference in childhood HbA1c between eight high-income countries but similar rise during adolescence-A quality registry study, *PEDIATR DIABETES* 21: 621-627

INNODIA: Eizirik, Decio L. et al. Pancreatic beta-cells in type 1 and type 2 diabetes mellitus: different pathways to failure, *NAT REV ENDOCRINOL* 16: 349-362

INNODIA: Colli, Maikel L. et al. An integrated multi-omics approach identifies the landscape of interferon-alpha-mediated responses of human pancreatic beta cells, *NAT COMMUN* 11:

INNODIA: James, Eddie A. et al. T-Cell Epitopes and Neo-epitopes in Type 1 Diabetes: A Comprehensive Update and Reappraisal, *DIABETES* 69: 1311-1335

INNODIA: Fignani, Daniela et al. SARS-CoV-2 Receptor Angiotensin I-Converting Enzyme Type 2 (ACE2) Is Expressed in Human Pancreatic beta-Cells and in the Human Pancreas Microvasculature, *FRONT ENDOCRINOL* 11:

INNODIA: De Franco, Elisa et al. YIPF5 mutations cause neonatal diabetes and microcephaly through endoplasmic reticulum stress, *J CLIN INVEST* 130: 6338-6353

INNODIA: Szymczak, F. et al. Gene expression signatures of target tissues in type 1 diabetes, lupus erythematosus, multiple sclerosis, and rheumatoid arthritis, *SCI ADV* 7:

INNODIA: Sodre, Fernanda M. C. et al. Peptidylarginine Deiminase Inhibition Prevents Diabetes Development in NOD Mice, *DIABETES* 70: 516-528

INNODIA: Grieco, Giuseppina Emanuela et al. The Landscape of microRNAs in beta Cell: Between Phenotype Maintenance and Protection, *INT J MOL SCI* 22:

INNODIA: Mueller, Andreas et al. 3D FIB-SEM reconstruction of microtubule-organelle interaction in whole primary mouse beta cells, *J CELL BIOL* 220:

INNODIA: Alvelos, Maria Ines et al. The RNA-binding profile of the splicing factor SRSF6 in immortalized human pancreatic beta-cells, *LIFE SCI ALLIANCE* 4:

INNODIA HARVEST: Mallone, Roberto et al. Presumption of innocence for beta cells: why are they vulnerable autoimmune targets in type 1 diabetes?, *DIABETOLOGIA* 63: 1999-2006

INNODIA HARVEST: Fignani, Daniela et al. SARS-CoV-2 Receptor Angiotensin I-Converting Enzyme Type 2 (ACE2) Is Expressed in Human Pancreatic beta-Cells and in the Human Pancreas Microvasculature, *FRONT ENDOCRINOL* 11:

INNODIA HARVEST: De Franco, Elisa et al. YIPF5 mutations cause neonatal diabetes and microcephaly through endoplasmic reticulum stress, *J CLIN INVEST* 130: 6338-6353

INNODIA HARVEST: Szymczak, F. et al. Gene expression signatures of target tissues in type 1 diabetes, lupus erythematosus, multiple sclerosis, and rheumatoid arthritis, *SCI ADV* 7:

INNODIA HARVEST: Sodre, Fernanda M. C. et al. Peptidylarginine Deiminase Inhibition Prevents Diabetes Development in NOD Mice, *DIABETES* 70: 516-528

INNODIA HARVEST: Grieco, Giuseppina Emanuela et al. The Landscape of microRNAs in beta Cell: Between Phenotype Maintenance and Protection, INT J MOL SCI 22:

INNODIA HARVEST: Alvelos, Maria Ines et al. The RNA-binding profile of the splicing factor SRSF6 in immortalized human pancreatic beta-cells, LIFE SCI ALLIANCE 4:

iPiE: Queralt-Rosinach, Nuria et al. DisGeNET-RDF: harnessing the innovative power of the Semantic Web to explore the genetic basis of diseases, BIOINFORMATICS 32: 2236-2238

iPiE: Escher, Beate I. et al. General baseline toxicity QSAR for nonpolar, polar and ionisable chemicals and their mixtures in the bioluminescence inhibition assay with *Aliivibrio fischeri*, ENVIRON SCI-PROC IMP 19: 414-428

iPiE: Burns, Emily E. et al. Temporal and spatial variation in pharmaceutical concentrations in an urban river system, WATER RES 137: 72-85

iPiE: Miller, Thomas H. et al. A review of the pharmaceutical exposome in aquatic fauna, ENVIRON POLLUT 239: 129-146

iPiE: Bittner, Lisa et al. Influence of pH on the uptake and toxicity of beta-blockers in embryos of zebrafish, *Danio rerio*, AQUAT TOXICOL 201: 129-137

iPiE: Kluever, Nils et al. QSAR for baseline toxicity and classification of specific modes of action of ionizable organic chemicals in the zebrafish embryo toxicity test, AQUAT TOXICOL 207: 110-119

iPiE: Gunnarsson, Lina et al. Pharmacology beyond the patient - The environmental risks of human drugs, ENVIRON INT 129: 320-332

ITCC-P4: Theruvath, Johanna et al. Locoregionally administered B7-H3-targeted CAR T cells for treatment of atypical teratoid/rhabdoid tumors, NAT MED 26: 712-+

ITCC-P4: Gojo, Johannes et al. Single-Cell RNA-Seq Reveals Cellular Hierarchies and Impaired Developmental Trajectories in Pediatric Ependymoma, CANCER CELL 38: 44-+

ITCC-P4: Loetsch, Daniela et al. Targeting fibroblast growth factor receptors to combat aggressive ependymoma, ACTA NEUROPATHOL 142: 339-360

ITCC-P4: Surdez, Didier et al. STAG2 mutations alter CTCF-anchored loop extrusion, reduce cis-regulatory interactions and EWSR1-FLI1 activity in Ewing sarcoma, CANCER CELL 39: 810-+

K4DD: Hoffmann, C. et al. Ligand Residence Time at G-protein-Coupled Receptors-Why We Should Take Our Time To Study It, MOL PHARMACOL 88: 552-560

K4DD: Hothersall, J. Daniel et al. Can residence time offer a useful strategy to target agonist drugs for sustained GPCR responses?, DRUG DISCOV TODAY 21: 90-96

K4DD: Stank, Antonia et al. Protein Binding Pocket Dynamics, ACCOUNTS CHEM RES 49: 809-815

K4DD: Segala, Elena et al. Controlling the Dissociation of Ligands from the Adenosine A(2A) Receptor through Modulation of Salt Bridge Strength, J MED CHEM 59: 6470-6479

K4DD: de Witte, Wilhelmus E. A. et al. In vivo Target Residence Time and Kinetic Selectivity: The Association Rate Constant as Determinant, *TRENDS PHARMACOL SCI* 37: 831-842

K4DD: Schuetz, Doris A. et al. Kinetics for Drug Discovery: an industry-driven effort to target drug residence time, *DRUG DISCOV TODAY* 22: 896-911

K4DD: Cheng, Robert K. Y. et al. Structures of Human A(1) and A(2A) Adenosine Receptors with Xanthines Reveal Determinants of Selectivity, *STRUCTURE* 25: 1275-+

K4DD: Rucktooa, Prakash et al. Towards high throughput GPCR crystallography: In Meso soaking of Adenosine A(2A) Receptor crystals, *SCI REP-UK* 8:

K4DD: Stoddart, Leigh A. et al. Development of novel fluorescent histamine H-1-receptor antagonists to study ligand-binding kinetics in living cells, *SCI REP-UK* 8:

K4DD: Stoddart, Leigh A. et al. NanoBRET Approaches to Study Ligand Binding to GPCRs and RTKs, *TRENDS PHARMACOL SCI* 39: 136-147

K4DD: Bruce, Neil J. et al. New approaches for computing ligand-receptor binding kinetics, *CURR OPIN STRUC BIOL* 49: 1-10

K4DD: Kokh, Daria B. et al. Estimation of Drug-Target Residence Times by tau-Random Acceleration Molecular Dynamics Simulations, *J CHEM THEORY COMPUT* 14: 3859-3869

K4DD: Schuetz, Doris A. et al. Predicting Residence Time and Drug Unbinding Pathway through Scaled Molecular Dynamics, *J CHEM INF MODEL* 59: 535-549

K4DD: Sykes, David A. et al. Binding kinetics of ligands acting at GPCRs, *MOL CELL ENDOCRINOL* 485: 9-19

K4DD: Berger, Benedict-Tilman et al. Structure-kinetic relationship reveals the mechanism of selectivity of FAK inhibitors over PYK2, *CELL CHEM BIOL* 28: 686-+

KRONO: Rueca, Martina et al. Investigation of Nasal/Oropharyngeal Microbial Community of COVID-19 Patients by 16S rDNA Sequencing, *INT J ENV RES PUB HE* 18:

KRONO: Colavita, Francesca et al. COVID-19 Rapid Antigen Test as Screening Strategy at Points of Entry: Experience in Lazio Region, Central Italy, August-October 2020, *BIOMOLECULES* 11:

KRONO: Amendola, Alessandra et al. Saliva Is a Valid Alternative to Nasopharyngeal Swab in Chemiluminescence-Based Assay for Detection of SARS-CoV-2 Antigen, *J CLIN MED* 10:

LITMUS: Karsdal, Morten A. et al. Assessment of liver fibrosis progression and regression by a serological collagen turnover profile, *AM J PHYSIOL-GASTR L* 316: G25-G31

LITMUS: Stoelzel, Ulrich et al. Clinical Guide and Update on Porphyrias, *GASTROENTEROLOGY* 157: 365-+

LITMUS: Luukkonen, Panu K. et al. Human PNPLA3-I148M variant increases hepatic retention of polyunsaturated fatty acids, *JCI INSIGHT* 4:

LITMUS: Stols-Goncalves, Daniela et al. NAFLD and Atherosclerosis: Two Sides of the Same Dysmetabolic Coin?, *TRENDS ENDOCRIN MET* 30: 891-902

LITMUS: Lazarus, Jeffrey, V et al. A cross-sectional study of the public health response to non-alcoholic fatty liver disease in Europe, *J HEPATOL* 72: 14-24

LITMUS: Aron-Wisnewsky, Judith et al. Gut microbiota and human NAFLD: disentangling microbial signatures from metabolic disorders, *NAT REV GASTRO HEPAT* 17: 279-297

LITMUS: Luukkonen, Panu K. et al. Hydroxysteroid 17-beta dehydrogenase 13 variant increases phospholipids and protects against fibrosis in nonalcoholic fatty liver disease, *JCI INSIGHT* 5:

LITMUS: Luukkonen, Panu K. et al. Effect of a ketogenic diet on hepatic steatosis and hepatic mitochondrial metabolism in nonalcoholic fatty liver disease, *P NATL ACAD SCI USA* 117: 7347-7354

LITMUS: Azzu, Vian et al. Adipose Tissue-Liver Cross Talk in the Control of Whole-Body Metabolism: Implications in Nonalcoholic Fatty Liver Disease, *GASTROENTEROLOGY* 158: 1899-1912

LITMUS: Gehrke, Nadine et al. Metabolic Inflammation-A Role for Hepatic Inflammatory Pathways as Drivers of Comorbidities in Nonalcoholic Fatty Liver Disease?, *GASTROENTEROLOGY* 158: 1929+

LITMUS: Vali, Yasaman et al. Enhanced liver fibrosis test for the non-invasive diagnosis of fibrosis in patients with NAFLD: A systematic review and meta-analysis, *J HEPATOL* 73: 252-262

LITMUS: Hardy, Timothy et al. The European NAFLD Registry: A real-world longitudinal cohort study of nonalcoholic fatty liver disease, *CONTEMP CLIN TRIALS* 98:

LITMUS: Govaere, Olivier et al. Transcriptomic profiling across the nonalcoholic fatty liver disease spectrum reveals gene signatures for steatohepatitis and fibrosis, *SCI TRANSL MED* 12:

LITMUS: Teo, Kevin et al. rs641738C>T near MBOAT7 is associated with liver fat, ALT and fibrosis in NAFLD: A meta-analysis, *J HEPATOL* 74: 20-30

LITMUS: Luukkonen, Panu K. et al. The PNPLA3-I148M Variant Confers an Antiatherogenic Lipid Profile in Insulin-resistant Patients, *J CLIN ENDOCR METAB* 106: E300-E315

LITMUS: Lee, Jenny et al. Prognostic accuracy of FIB-4, NAFLD fibrosis score and APRI for NAFLD-related events: A systematic review, *LIVER INT* 41: 261-270

LITMUS: Pfister, Dominik et al. NASH limits anti-tumour surveillance in immunotherapy-treated HCC, *NATURE* 592: 450-456

LITMUS: Armandi, Angelo et al. Insulin Resistance across the Spectrum of Nonalcoholic Fatty Liver Disease, *METABOLITES* 11:

LITMUS: Eldafashi, Nardeen et al. A PDCD1 Role in the Genetic Predisposition to NAFLD-HCC?, *CANCERS* 13:

LITMUS: Geh, Daniel et al. NAFLD-Associated HCC: Progress and Opportunities, *J HEPATOCELL CARCINO* 8: 223-239

LITMUS: Lazarus, Jeffrey, V et al. European NAFLD Preparedness Index - Is Europe ready to meet the challenge of fatty liver disease?, JHEP REP 3:

LITMUS: Geier, Andreas et al. From the origin of NASH to the future of metabolic fatty liver disease, GUT 70: 1570-1579

LITMUS: Selvaraj, Emmanuel Anandraj et al. Diagnostic accuracy of elastography and magnetic resonance imaging in patients with NAFLD: A systematic review and meta-analysis, J HEPATOL 75: 770-785

LITMUS: Mozes, Ferenc Emil et al. Diagnostic accuracy of non-invasive tests for advanced fibrosis in patients with NAFLD: an individual patient data meta-analysis, GUT 71: 1006-1019

MACUSTAR: Finger, Robert P. et al. MACUSTAR: Development and Clinical Validation of Functional, Structural, and Patient-Reported Endpoints in Intermediate Age-Related Macular Degeneration, OPHTHALMOLOGICA 241: 61-72

MACUSTAR: Terheyden, Jan Henrik et al. Use of Composite End Points in Early and Intermediate Age-Related Macular Degeneration Clinical Trials: State-of-the-Art and Future Directions, OPHTHALMOLOGICA 244: 387-395

MAD-CoV 2: Monteil, Vanessa et al. Human soluble ACE2 improves the effect of remdesivir in SARS-CoV-2 infection, EMBO MOL MED 13:

MAD-CoV 2: Ziv, Omer et al. The Short- and Long-Range RNA-RNA Interactome of SARS-CoV-2, MOL CELL 80: 1067-+

MAD-CoV 2: Saccon, Elisa et al. Cell-type-resolved quantitative proteomics map of interferon response against SARS-CoV-2, ISCIENCE 24:

MAD-CoV 2: Han, Namshik et al. Identification of SARS-CoV-2-induced pathways reveals drug repurposing strategies, SCI ADV 7:

MAD-CoV 2: Hoffmann, David et al. Identification of lectin receptors for conserved SARS-CoV-2 glycosylation sites, EMBO J 40:

MARCAR: Sproul, Duncan et al. Genomic insights into cancer-associated aberrant CpG island hypomethylation, BRIEF FUNCT GENOMICS 12: 174-190

MARCAR: Reddington, James P. et al. Redistribution of H3K27me3 upon DNA hypomethylation results in de-repression of Polycomb target genes, GENOME BIOL 14:

MARCAR: Luisier, Raphaelle et al. Phenobarbital Induces Cell Cycle Transcriptional Responses in Mouse Liver Humanized for Constitutive Androstane and Pregnane X Receptors, TOXICOL SCI 139: 501-511

MARCAR: Nestor, Colm E. et al. Rapid reprogramming of epigenetic and transcriptional profiles in mammalian culture systems, GENOME BIOL 16:

MARCAR: Treindl, Fridolin et al. A bead-based western for high-throughput cellular signal transduction analyses, NAT COMMUN 7:

MARCAR: Meehan, Richard R. et al. DNA methylation as a genomic marker of exposure to chemical and environmental agents, *CURR OPIN CHEM BIOL* 45: 48-56

MIP-DILI: Ivanov, M. et al. Epigenomics and Interindividual Differences in Drug Response, *CLIN PHARMACOL THER* 92: 727-736

MIP-DILI: Wink, Steven et al. Quantitative High Content Imaging of Cellular Adaptive Stress Response Pathways in Toxicity for Chemical Safety Assessment, *CHEM RES TOXICOL* 27: 338-355

MIP-DILI: Fredriksson, Lisa et al. Drug-Induced Endoplasmic Reticulum and Oxidative Stress Responses Independently Sensitize Toward TNF alpha-Mediated Hepatotoxicity, *TOXICOL SCI* 140: 144-159

MIP-DILI: Ivanov, Maxim et al. Epigenetic mechanisms of importance for drug treatment, *TRENDS PHARMACOL SCI* 35: 384-396

MIP-DILI: Kamalian, Laleh et al. The utility of HepG2 cells to identify direct mitochondrial dysfunction in the absence of cell death, *TOXICOL IN VITRO* 29: 732-740

MIP-DILI: Bachour-EI Azzi, Pamela et al. Comparative Localization and Functional Activity of the Main Hepatobiliary Transporters in HepaRG Cells and Primary Human Hepatocytes, *TOXICOL SCI* 145: 157-168

MIP-DILI: Kim, Seung-Hyun et al. Characterization of amoxicillin- and clavulanic acid-specific T cells in patients with amoxicillin-clavulanate-induced liver injury, *HEPATOLOGY* 62: 887-899

MIP-DILI: Sison-Young, Rowena L. C. et al. Comparative Proteomic Characterization of 4 Human Liver-Derived Single Cell Culture Models Reveals Significant Variation in the Capacity for Drug Disposition, Bioactivation, and Detoxication, *TOXICOL SCI* 147: 412-424

MIP-DILI: Bell, Catherine C. et al. Characterization of primary human hepatocyte spheroids as a model system for drug-induced liver injury, liver function and disease, *SCI REP-UK* 6:

MIP-DILI: Sharanek, Ahmad et al. Rho-kinase/myosin light chain kinase pathway plays a key role in the impairment of bile canaliculi dynamics induced by cholestatic drugs, *SCI REP-UK* 6:

MIP-DILI: Sutherland, Jeffrey J. et al. Assessing Concordance of Drug-Induced Transcriptional Response in Rodent Liver and Cultured Hepatocytes, *PLOS COMPUT BIOL* 12:

MIP-DILI: Oorts, Marlies et al. Drug-induced cholestasis risk assessment in sandwich-cultured human hepatocytes, *TOXICOL IN VITRO* 34: 179-186

MIP-DILI: Maiwald, Tim et al. Driving the Model to Its Limit: Profile Likelihood Based Model Reduction, *PLOS ONE* 11:

MIP-DILI: Hendriks, Delilah F. G. et al. Hepatic 3D spheroid models for the detection and study of compounds with cholestatic liability, *SCI REP-UK* 6:

MIP-DILI: Lauschke, Volker M. et al. Massive rearrangements of cellular MicroRNA signatures are key drivers of hepatocyte dedifferentiation, *HEPATOLOGY* 64: 1743-1756

MIP-DILI: Lauschke, Volker M. et al. The Importance of Patient-Specific Factors for Hepatic Drug Response and Toxicity, INT J MOL SCI 17:

MIP-DILI: Lauschke, Volker M. et al. Novel 3D Culture Systems for Studies of Human Liver Function and Assessments of the Hepatotoxicity of Drugs and Drug Candidates, CHEM RES TOXICOL 29: 1936-1955

MIP-DILI: Wink, Steven et al. High-content imaging-based BAC-GFP toxicity pathway reporters to assess chemical adversity liabilities, ARCH TOXICOL 91: 1367-1383

MIP-DILI: Sison-Young, Rowena L. et al. A multicenter assessment of single-cell models aligned to standard measures of cell health for prediction of acute hepatotoxicity, ARCH TOXICOL 91: 1385-1400

MIP-DILI: Goldring, Christopher et al. Stem Cell-Derived Models to Improve Mechanistic Understanding and Prediction of Human Drug-Induced Liver Injury, HEPATOLOGY 65: 710-721

MIP-DILI: Bell, Catherine C. et al. Transcriptional, Functional, and Mechanistic Comparisons of Stem Cell-Derived Hepatocytes, HepaRG Cells, and Three-Dimensional Human Hepatocyte Spheroids as Predictive In Vitro Systems for Drug-Induced Liver Injury, DRUG METAB DISPOS 45: 419-429

MIP-DILI: Vorrink, Sabine U. et al. Endogenous and xenobiotic metabolic stability of primary human hepatocytes in long-term 3D spheroid cultures revealed by a combination of targeted and untargeted metabolomics, FASEB J 31: 2696-2708

MIP-DILI: Proctor, William R. et al. Utility of spherical human liver microtissues for prediction of clinical drug-induced liver injury, ARCH TOXICOL 91: 2849-2863

MIP-DILI: Bell, Catherine C. et al. Comparison of Hepatic 2D Sandwich Cultures and 3D Spheroids for Long-term Toxicity Applications: A Multicenter Study, TOXICOL SCI 162: 655-666

MIP-DILI: Wink, Steven et al. Dynamic imaging of adaptive stress response pathway activation for prediction of drug induced liver injury, ARCH TOXICOL 92: 1797-1814

MIP-DILI: Vorrink, Sabine U. et al. Prediction of Drug-Induced Hepatotoxicity Using Long-Term Stable Primary Hepatic 3D Spheroid Cultures in Chemically Defined Conditions, TOXICOL SCI 163: 655-665

MIP-DILI: Hiemstra, Steven et al. High-throughput confocal imaging of differentiated 3D liver-like spheroid cellular stress response reporters for identification of drug-induced liver injury liability, ARCH TOXICOL 93: 2895-2911

MIP-DILI: Weaver, Richard J. et al. Managing the challenge of drug-induced liver injury: a roadmap for the development and deployment of preclinical predictive models, NAT REV DRUG DISCOV 19: 131-148

MOBILISE-D: Viceconti, Marco et al. Credibility of In Silico Trial Technologies-A Theoretical Framing, IEEE J BIOMED HEALTH 24: 4-13

MOBILISE-D: Flachenecker, Felix et al. Objective sensor-based gait measures reflect motor impairment in multiple sclerosis patients: Reliability and clinical validation of a wearable sensor device, MULT SCLER RELAT DIS 39:

MOBILISE-D: Shema-Shiratzky, Shirley et al. A wearable sensor identifies alterations in community ambulation in multiple sclerosis: contributors to real-world gait quality and physical activity, *J NEUROL* 267: 1912-1921

MOBILISE-D: Angelini, Lorenza et al. Wearable sensors can reliably quantify gait alterations associated with disability in people with progressive multiple sclerosis in a clinical setting, *J NEUROL* 267: 2897-2909

MOBILISE-D: Warmerdam, Elke et al. Long-term unsupervised mobility assessment in movement disorders, *LANCET NEUROL* 19: 462-470

MOBILISE-D: Caruso, Marco et al. Analysis of the Accuracy of Ten Algorithms for Orientation Estimation Using Inertial and Magnetic Sensing under Optimal Conditions: One Size Does Not Fit All, *SENSORS-BASEL* 21:

MOBILISE-D: Mirelman, Anat et al. Detecting Sensitive Mobility Features for Parkinsons Disease Stages Via Machine Learning, *MOVEMENT DISORD* 36: 2144-2155

MOBILISE-D: Del Din, Silvia et al. Body-Worn Sensors for Remote Monitoring of Parkinsons Disease Motor Symptoms: Vision, State of the Art, and Challenges Ahead, *J PARKINSON DIS* 11: S35-S47

MOBILISE-D: Caruso, Marco et al. Orientation Estimation Through Magneto-Inertial Sensor Fusion: A Heuristic Approach for Suboptimal Parameters Tuning, *IEEE SENS J* 21: 3408-3419

MOBILISE-D: Kluge, Felix et al. Consensus based framework for digital mobility monitoring, *PLOS ONE* 16:

MOPEAD: Moreno-Grau, Sonia et al. Genome-wide association analysis of dementia and its clinical endophenotypes reveal novel loci associated with Alzheimers disease and three causality networks: The GR@ACE project, *ALZHEIMERS DEMENT* 15: 1333-1347

MOPEAD: de Rojas, Itziar et al. Common variants in Alzheimers disease and risk stratification by polygenic risk scores, *NAT COMMUN* 12:

ND4BB: Abu Kwaik, Yousef et al. Microbial quest for food in vivo: Nutritional virulence as an emerging paradigm, *CELL MICROBIOL* 15: 882-890

ND4BB: Kostyanev, T. et al. The Innovative Medicines Initiatives New Drugs for Bad Bugs programme: European public-private partnerships for the development of new strategies to tackle antibiotic resistance, *J ANTIMICROB CHEMOTH* 71: 290-295

NECESSITY: Hammenfors, Daniel S. et al. Juvenile Sjogrens Syndrome: Clinical Characteristics With Focus on Salivary Gland Ultrasonography, *ARTHRIT CARE RES* 72: 78-87

NECESSITY: Riviere, Elodie et al. Salivary gland epithelial cells from patients with Sjogrens syndrome induce B-lymphocyte survival and activation, *ANN RHEUM DIS* 79: 1468-1477

NECESSITY: Riviere, Elodie et al. Interleukin-7/Interferon Axis Drives T Cell and Salivary Gland Epithelial Cell Interactions in Sjogrens Syndrome, *ARTHRITIS RHEUMATOL* 73: 631-640

NEWMEDS: Meyer-Lindenberg, Andreas et al. From maps to mechanisms through neuroimaging of schizophrenia, *NATURE* 468: 194-202



NEWMEDS: Ingason, Andres et al. Maternally Derived Microduplications at 15q11-q13: Implication of Imprinted Genes in Psychotic Illness, *AM J PSYCHIAT* 168: 408-417

NEWMEDS: Keeler, J. F. et al. Translating cognition from animals to humans, *BIOCHEM PHARMACOL* 81: 1356-1366

NEWMEDS: Meyer-Lindenberg, Andreas et al. Oxytocin and vasopressin in the human brain: social neuropeptides for translational medicine, *NAT REV NEUROSCI* 12: 524-538

NEWMEDS: Smith, Janice W. et al. A comparison of the effects of ketamine and phencyclidine with other antagonists of the NMDA receptor in rodent assays of attention and working memory, *PSYCHOPHARMACOLOGY* 217: 255-269

NEWMEDS: Jacquemont, Sebastien et al. Mirror extreme BMI phenotypes associated with gene dosage at the chromosome 16p11.2 locus, *NATURE* 478: 97-U111

NEWMEDS: Braun, Urs et al. Test-retest reliability of resting-state connectivity network characteristics using fMRI and graph theoretical measures, *NEUROIMAGE* 59: 1404-1412

NEWMEDS: Kirov, G. et al. De novo CNV analysis implicates specific abnormalities of postsynaptic signalling complexes in the pathogenesis of schizophrenia, *MOL PSYCHIATR* 17: 142-153

NEWMEDS: Bussey, T. J. et al. New translational assays for preclinical modelling of cognition in schizophrenia: The touchscreen testing method for mice and rats, *NEUROPHARMACOLOGY* 62: 1191-1203

NEWMEDS: Gilmour, Gary et al. NMDA receptors, cognition and schizophrenia - Testing the validity of the NMDA receptor hypofunction hypothesis, *NEUROPHARMACOLOGY* 62: 1401-1412

NEWMEDS: Gastambide, Francois et al. Selective Remediation of Reversal Learning Deficits in the Neurodevelopmental MAM Model of Schizophrenia by a Novel mGlu5 Positive Allosteric Modulator, *NEUROPSYCHOPHARMACOL* 37: 1057-1066

NEWMEDS: Plichta, Michael M. et al. Test-retest reliability of evoked BOLD signals from a cognitive-emotive fMRI test battery, *NEUROIMAGE* 60: 1746-1758

NEWMEDS: Lyon, L. et al. Spontaneous object recognition and its relevance to schizophrenia: a review of findings from pharmacological, genetic, lesion and developmental rodent models, *PSYCHOPHARMACOLOGY* 220: 647-672

NEWMEDS: Zink, Caroline F. et al. Human neuroimaging of oxytocin and vasopressin in social cognition, *HORM BEHAV* 61: 400-409

NEWMEDS: Uher, R. et al. Depression symptom dimensions as predictors of antidepressant treatment outcome: replicable evidence for interest-activity symptoms, *PSYCHOL MED* 42: 967-980

NEWMEDS: Meyer-Lindenberg, Andreas et al. Neural mechanisms of social risk for psychiatric disorders, *NAT NEUROSCI* 15: 663-668

NEWMEDS: Bortolozzi, A. et al. Selective siRNA-mediated suppression of 5-HT<sub>1A</sub> autoreceptors evokes strong anti-depressant-like effects, *MOL PSYCHIATR* 17: 612-623

NEWMEDS: Llado-Pelfort, Laia et al. 5-HT1A Receptor Agonists Enhance Pyramidal Cell Firing in Prefrontal Cortex Through a Preferential Action on GABA Interneurons, CEREB CORTEX 22: 1487-1497

NEWMEDS: Tansey, Katherine E. et al. Genetic Predictors of Response to Serotonergic and Noradrenergic Antidepressants in Major Depressive Disorder: A Genome-Wide Analysis of Individual-Level Data and a Meta-Analysis, PLOS MED 9:

NEWMEDS: Kapur, S. et al. Why has it taken so long for biological psychiatry to develop clinical tests and what to do about it?, MOL PSYCHIATR 17: 1174-1179

NEWMEDS: Uher, Rudolf et al. SELF-REPORT AND CLINICIAN-RATED MEASURES OF DEPRESSION SEVERITY: CAN ONE REPLACE THE OTHER?, DEPRESS ANXIETY 29: 1043-1049

NEWMEDS: Artigas, Francesc et al. Serotonin receptors involved in antidepressant effects, PHARMACOL THERAPEUT 137: 119-131

NEWMEDS: Doyle, O. M. et al. Quantifying the Attenuation of the Ketamine Pharmacological Magnetic Resonance Imaging Response in Humans: A Validation Using Antipsychotic and Glutamatergic Agents, J PHARMACOL EXP THER 345: 151-160

NEWMEDS: Sullivan, Patrick F. et al. A mega-analysis of genome-wide association studies for major depressive disorder, MOL PSYCHIATR 18: 497-511

NEWMEDS: Tansey, Katherine E. et al. Contribution of Common Genetic Variants to Antidepressant Response, BIOL PSYCHIAT 73: 679-682

NEWMEDS: Anacker, Christoph et al. Role for the kinase SGK1 in stress, depression, and glucocorticoid effects on hippocampal neurogenesis, P NATL ACAD SCI USA 110: 8708-8713

NEWMEDS: Nord, Magdalena et al. Effect of a single dose of escitalopram on serotonin concentration in the non-human and human primate brain, INT J NEUROPSYCHOPH 16: 1577-1586

NEWMEDS: Godsil, Bill P. et al. The hippocampal-prefrontal pathway: The weak link in psychiatric disorders?, EUR NEUROPSYCHOPHARM 23: 1165-1181

NEWMEDS: Horner, Alexa E. et al. The touchscreen operant platform for testing learning and memory in rats and mice, NAT PROTOC 8: 1961-1984

NEWMEDS: Mar, Adam C. et al. The touchscreen operant platform for assessing executive function in rats and mice, NAT PROTOC 8: 1985-2005

NEWMEDS: Oomen, Charlotte A. et al. The touchscreen operant platform for testing working memory and pattern separation in rats and mice, NAT PROTOC 8: 2006-2021

NEWMEDS: Cao, Hengyi et al. Test-retest reliability of fMRI-based graph theoretical properties during working memory, emotion processing, and resting state, NEUROIMAGE 84: 888-900

NEWMEDS: Stefansson, Hreinn et al. CNVs conferring risk of autism or schizophrenia affect cognition in controls, NATURE 505: 361-+

NEWMEDS: Lustig, C. et al. CNTRICS final animal model task selection: Control of attention, NEUROSCI BIOBEHAV R 37: 2099-2110

NEWMEDS: Fejgin, Kim et al. A Mouse Model that Recapitulates Cardinal Features of the 15q13.3 Microdeletion Syndrome Including Schizophrenia- and Epilepsy-Related Alterations, BIOL PSYCHIAT 76: 128-137

NEWMEDS: Plichta, Michael M. et al. Amygdala habituation: A reliable fMRI phenotype, NEUROIMAGE 103: 383-390

NEWMEDS: Artigas, Francesc et al. Developments in the field of antidepressants, where do we go now?, EUR NEUROPSYCHOPHARM 25: 657-670

NEWMEDS: Power, Robert A. et al. Polygenic risk scores for schizophrenia and bipolar disorder predict creativity, NAT NEUROSCI 18: 953+

NEWMEDS: Braun, Urs et al. Dynamic reconfiguration of frontal brain networks during executive cognition in humans, P NATL ACAD SCI USA 112: 11678-11683

NEWMEDS: Kim, Chi Hun et al. The continuous performance test (rCPT) for mice: a novel operant touchscreen test of attentional function, PSYCHOPHARMACOLOGY 232: 3947-3966

NEWMEDS: Grimm, Oliver et al. Acute ketamine challenge increases resting state prefrontal-hippocampal connectivity in both humans and rats, PSYCHOPHARMACOLOGY 232: 4231-4241

NEWMEDS: Paloyelis, Yannis et al. A Spatiotemporal Profile of In Vivo Cerebral Blood Flow Changes Following Intranasal Oxytocin in Humans, BIOL PSYCHIAT 79: 693-705

NEWMEDS: Iniesta, Raquel et al. Combining clinical variables to optimize prediction of antidepressant treatment outcomes, J PSYCHIATR RES 78: 94-102

NEWMEDS: Iniesta, R. et al. Machine learning, statistical learning and the future of biological research in psychiatry, PSYCHOL MED 46: 2455-2465

NEWMEDS: Lo, Min-Tzu et al. Genome-wide analyses for personality traits identify six genomic loci and show correlations with psychiatric disorders, NAT GENET 49: 152-156

NEWMEDS: Ulfarsson, M. O. et al. 15q11.2 CNV affects cognitive, structural and functional correlates of dyslexia and dyscalculia, TRANSL PSYCHIAT 7:

NEWMEDS: Direk, Nese et al. An Analysis of Two Genome-wide Association Meta-analyses Identifies a New Locus for Broad Depression Phenotype, BIOL PSYCHIAT 82: 322-329

NEWMEDS: Silva, Ana, I et al. Reciprocal White Matter Changes Associated With Copy Number Variation at 15q11.2 BP1-BP2: A Diffusion Tensor Imaging Study, BIOL PSYCHIAT 85: 563-572

NEWMEDS: Tost, Heike et al. Neural correlates of individual differences in affective benefit of real-life urban green space exposure, NAT NEUROSCI 22: 1389+

NEWMEDS: Gudmundsson, Olafur O. et al. Attention-deficit hyperactivity disorder shares copy number variant risk with schizophrenia and autism spectrum disorder, TRANSL PSYCHIAT 9:

NEWMEDS: Jonsson, B. A. et al. Brain age prediction using deep learning uncovers associated sequence variants, *NAT COMMUN* 10:

NEWMEDS: Grandjean, Joanes et al. Common functional networks in the mouse brain revealed by multi-centre resting-state fMRI analysis, *NEUROIMAGE* 205:

NEWMEDS: Holz, Nathalie E. et al. Resilience and the brain: a key role for regulatory circuits linked to social stress and support, *MOL PSYCHIATR* 25: 379-396

NEWMEDS: Reichert, Markus et al. Studying the impact of built environments on human mental health in everyday life: methodological developments, state-of-the-art and technological frontiers, *CURR OPIN PSYCHOL* 32: 158-164

NEWMEDS: Coleman, Jonathan R., I et al. The Genetics of the Mood Disorder Spectrum: Genome-wide Association Analyses of More Than 185,000 Cases and 439,000 Controls, *BIOL PSYCHIAT* 88: 169-184

Onco Track: Hildebrandt, Niko et al. Biofunctional Quantum Dots: Controlled Conjugation for Multiplexed Biosensors, *ACS NANO* 5: 5286-5290

Onco Track: Bettermann, Kira et al. SUMOylation in carcinogenesis, *CANCER LETT* 316: 113-125

Onco Track: Algar, W. Russ et al. Quantum Dots as Simultaneous Acceptors and Donors in Time-Gated Forster Resonance Energy Transfer Relays: Characterization and Biosensing, *J AM CHEM SOC* 134: 1876-1891

Onco Track: Taiwo, Oluwatosin et al. Methylome analysis using MeDIP-seq with low DNA concentrations, *NAT PROTOC* 7: 617-636

Onco Track: Jin, Zongwen et al. Semiconductor quantum dots for in vitro diagnostics and cellular imaging, *TRENDS BIOTECHNOL* 30: 394-403

Onco Track: Hoetzer, Benjamin et al. Fluorescence in Nanobiotechnology: Sophisticated Fluorophores for Novel Applications, *SMALL* 8: 2297-2326

Onco Track: Ke, Rongqin et al. In situ sequencing for RNA analysis in preserved tissue and cells, *NAT METHODS* 10: 857-+

Onco Track: Wegner, K. David et al. Quantum-Dot-Based Forster Resonance Energy Transfer Immunoassay for Sensitive Clinical Diagnostics of Low-Volume Serum Samples, *ACS NANO* 7: 7411-7419

Onco Track: Morris, Tiffany J. et al. ChAMP: 450k Chip Analysis Methylation Pipeline, *BIOINFORMATICS* 30: 428-430

Onco Track: Wegner, K. David et al. Nanobodies and Nanocrystals: Highly Sensitive Quantum Dot-Based Homogeneous FRET Immunoassay for Serum-Based EGFR Detection, *SMALL* 10: 734-740

Onco Track: Geissler, Daniel et al. Lanthanides and Quantum Dots as Forster Resonance Energy Transfer Agents for Diagnostics and Cellular Imaging, *INORG CHEM* 53: 1824-1838

Onco Track: Lechner, Matthias et al. Identification and functional validation of HPV-mediated hypermethylation in head and neck squamous cell carcinoma, *GENOME MED* 5:

Onco Track: Feber, Andrew et al. Using high-density DNA methylation arrays to profile copy number alterations, *GENOME BIOL* 15:

Onco Track: Butcher, Lee M. et al. Probe Lasso: A novel method to rope in differentially methylated regions with 450K DNA methylation data, *METHODS* 72: 21-28

Onco Track: Wegner, K. David et al. Quantum dots: bright and versatile in vitro and in vivo fluorescence imaging biosensors, *CHEM SOC REV* 44: 4792-4834

Onco Track: Jin, Zongwen et al. A Rapid, Amplification-Free, and Sensitive Diagnostic Assay for Single-Step Multiplexed Fluorescence Detection of MicroRNA, *ANGEW CHEM INT EDIT* 54: 10024-10029

Onco Track: Qiu, Xue et al. Rapid and Multiplexed MicroRNA Diagnostic Assay Using Quantum Dot-Based Forster Resonance Energy Transfer, *ACS NANO* 9: 8449-8457

Onco Track: Kargl, J. et al. GPR55 promotes migration and adhesion of colon cancer cells indicating a role in metastasis, *BRIT J PHARMACOL* 173: 142-154

Onco Track: Boehnke, Karsten et al. Assay Establishment and Validation of a High-Throughput Screening Platform for Three-Dimensional Patient-Derived Colon Cancer Organoid Cultures, *J BIOMOL SCREEN* 21: 931-941

Onco Track: Schuette, Moritz et al. Molecular dissection of colorectal cancer in pre-clinical models identifies biomarkers predicting sensitivity to EGFR inhibitors, *NAT COMMUN* 8:

Onco Track: Taniguchi, Koji et al. YAP-IL-6ST autoregulatory loop activated on APC loss controls colonic tumorigenesis, *P NATL ACAD SCI USA* 114: 1643-1648

Onco Track: Qiu, Xue et al. Multiplexed Nucleic Acid Hybridization Assays Using Single-FRET-Pair Distance-Tuning, *SMALL* 13:

Onco Track: Golob-Schwarzl, Nicole et al. New liver cancer biomarkers: PI3K/AKT/mTOR pathway members and eukaryotic translation initiation factors, *EUR J CANCER* 83: 56-70

Onco Track: Regan, Joseph L. et al. Non-Canonical Hedgehog Signaling Is a Positive Regulator of the WNT Pathway and Is Required for the Survival of Colon Cancer Stem Cells, *CELL REP* 21: 2813-2828

Onco Track: Schumacher, Dirk et al. Heterogeneous pathway activation and drug response modelled in colorectal-tumor-derived 3D cultures, *PLOS GENET* 15:

Open PHACTS: Williams, Antony J. et al. Towards a gold standard: regarding quality in public domain chemistry databases and approaches to improving the situation, *DRUG DISCOV TODAY* 17: 685-701

Open PHACTS: Williams, Antony J. et al. Open PHACTS: semantic interoperability for drug discovery, *DRUG DISCOV TODAY* 17: 1188-1198

Open PHACTS: Furlong, Laura I. et al. Human diseases through the lens of network biology, *TRENDS GENET* 29: 150-159

Open PHACTS: Bento, A. Patricia et al. The ChEMBL bioactivity database: an update, *NUCLEIC ACIDS RES* 42: D1083-D1090

Open PHACTS: Jupp, Simon et al. The EBI RDF platform: linked open data for the life sciences, *BIOINFORMATICS* 30: 1338-1339

Open PHACTS: Dumontier, Michel et al. The SemanticScience Integrated Ontology (SIO) for biomedical research and knowledge discovery, *J BIOMED SEMANT* 5:

Open PHACTS: Lizio, Marina et al. Gateways to the FANTOM5 promoter level mammalian expression atlas, *GENOME BIOL* 16:

Open PHACTS: Pinero, Janet et al. DisGeNET: a discovery platform for the dynamical exploration of human diseases and their genes, *DATABASE-OXFORD* :

Open PHACTS: Kringelum, Jens et al. ChemProt-3.0: a global chemical biology diseases mapping, *DATABASE-OXFORD* :

Open PHACTS: Kutmon, Martina et al. WikiPathways: capturing the full diversity of pathway knowledge, *NUCLEIC ACIDS RES* 44: D488-D494

Open PHACTS: Moreau, Luc et al. The rationale of PROV, *J WEB SEMANT* 35: 235-257

Open PHACTS: Queralt-Rosinach, Nuria et al. DisGeNET-RDF: harnessing the innovative power of the Semantic Web to explore the genetic basis of diseases, *BIOINFORMATICS* 32: 2236-2238

Open PHACTS: Pinero, Janet et al. DisGeNET: a comprehensive platform integrating information on human disease-associated genes and variants, *NUCLEIC ACIDS RES* 45: D833-D839

Open PHACTS: Gaulton, Anna et al. The ChEMBL database in 2017, *NUCLEIC ACIDS RES* 45: D945-D954

Open PHACTS: Alshahrani, Mona et al. Neuro-symbolic representation learning on biological knowledge graphs, *BIOINFORMATICS* 33: 2723-2730

Open PHACTS: Slenter, Denise N. et al. WikiPathways: a multifaceted pathway database bridging metabolomics to other omics research, *NUCLEIC ACIDS RES* 46: D661-D667

ORBITO: Koziolok, Mirko et al. Intra-gastric Volume Changes after Intake of a High-Caloric, High-Fat Standard Breakfast in Healthy Human Subjects Investigated by MRI, *MOL PHARMACEUT* 11: 1632-1639

ORBITO: Sjogren, Erik et al. In vivo methods for drug absorption - Comparative physiologies, model selection, correlations with in vitro methods (IVIVC), and applications for formulation/API/excipient characterization including food effects, *EUR J PHARM SCI* 57: 99-151

ORBITO: Bergstrom, Christel A. S. et al. Early pharmaceutical profiling to predict oral drug absorption: Current status and unmet needs, *EUR J PHARM SCI* 57: 173-199

ORBITO: Kostewicz, Edmund S. et al. PBPK models for the prediction of in vivo performance of oral dosage forms, EUR J PHARM SCI 57: 300-321

ORBITO: Augustijns, Patrick et al. A review of drug solubility in human intestinal fluids: Implications for the prediction of oral absorption, EUR J PHARM SCI 57: 322-332

ORBITO: Kostewicz, Edmund S. et al. In vitro models for the prediction of in vivo performance of oral dosage forms, EUR J PHARM SCI 57: 342-366

ORBITO: Hens, Bart et al. Gastrointestinal transfer: In vivo evaluation and implementation in in vitro and in silico predictive tools, EUR J PHARM SCI 63: 233-242

ORBITO: Khadra, Ibrahim et al. Statistical investigation of simulated intestinal fluid composition on the equilibrium solubility of biopharmaceutics classification system class II drugs, EUR J PHARM SCI 67: 65-75

ORBITO: Harwood, M. D. et al. Application of an LC-MS/MS method for the simultaneous quantification of human intestinal transporter proteins absolute abundance using a QconCAT technique, J PHARMACEUT BIOMED 110: 27-33

ORBITO: Markopoulos, Constantinos et al. In-vitro simulation of luminal conditions for evaluation of performance of oral drug products: Choosing the appropriate test media, EUR J PHARM BIOPHARM 93: 173-182

ORBITO: Hens, Bart et al. Gastrointestinal behavior of nano- and micro-sized fenofibrate: In vivo evaluation in man and in vitro simulation by assessment of the permeation potential, EUR J PHARM SCI 77: 40-47

ORBITO: Dahlgren, David et al. Direct In Vivo Human Intestinal Permeability (P<sub>eff</sub>) Determined with Different Clinical Perfusion and Intubation Methods, J PHARM SCI-US 104: 2702-2726

ORBITO: Koziolok, Mirko et al. Investigation of pH and Temperature Profiles in the GI Tract of Fasted Human Subjects Using the Intellicap((R)) System, J PHARM SCI-US 104: 2855-2863

ORBITO: Koziolok, M. et al. Intra-gastric pH and pressure profiles after intake of the high-caloric, high-fat meal as used for food effect studies, J CONTROL RELEASE 220: 71-78

ORBITO: Kourentas, Alexandros et al. An in vitro biorelevant gastrointestinal transfer (BioGIT) system for forecasting concentrations in the fasted upper small intestine: Design, implementation, and evaluation, EUR J PHARM SCI 82: 106-114

ORBITO: Verwei, Miriam et al. Evaluation of two dynamic in vitro models simulating fasted and fed state conditions in the upper gastrointestinal tract (TIM-1 and tiny-TIM) for investigating the bioaccessibility of pharmaceutical compounds from oral dosage forms, INT J PHARMACEUT 498: 178-186

ORBITO: Koziolok, Mirko et al. Navigating the human gastrointestinal tract for oral drug delivery: Uncharted waters and new frontiers, ADV DRUG DELIVER REV 101: 75-88

ORBITO: Dahlgren, David et al. Regional Intestinal Permeability of Three Model Drugs in Human, MOL PHARMACEUT 13: 3013-3021

ORBITO: Schneider, Felix et al. Resolving the physiological conditions in bioavailability and bioequivalence studies: Comparison of fasted and fed state, EUR J PHARM BIOPHARM 108: 214-219

ORBITO: Palmelund, Henrik et al. Studying the Propensity of Compounds to Supersaturate: A Practical and Broadly Applicable Approach, J PHARM SCI-US 105: 3021-3029

ORBITO: Van den Abeele, Jens et al. The dynamic gastric environment and its impact on drug and formulation behaviour, EUR J PHARM SCI 96: 207-231

ORBITO: Margolskee, Alison et al. IMI - Oral biopharmaceutics tools project - Evaluation of bottom-up PBPK prediction success part 2: An introduction to the simulation exercise and overview of results, EUR J PHARM SCI 96: 610-625

ORBITO: Ruff, Aaron et al. Prediction of Ketoconazole absorption using an updated in vitro transfer model coupled to physiologically based pharmacokinetic modelling, EUR J PHARM SCI 100: 42-55

ORBITO: Hens, Bart et al. Exploring gastrointestinal variables affecting drug and formulation behavior: Methodologies, challenges and opportunities, INT J PHARMACEUT 519: 79-97

ORBITO: Lennernas, H. et al. In Vivo Predictive Dissolution (IPD) and Biopharmaceutical Modeling and Simulation: Future Use of Modern Approaches and Methodologies in a Regulatory Context, MOL PHARMACEUT 14: 1307-1314

ORBITO: Nguyen, M. A. et al. A survey on IVIVC/IVIVR development in the pharmaceutical industry - Past experience and current perspectives, EUR J PHARM SCI 102: 1-13

ORBITO: Andreas, Cord J. et al. Mechanistic investigation of the negative food effect of modified release zolpidem, EUR J PHARM SCI 102: 284-298

ORBITO: Mann, James et al. Validation of Dissolution Testing with Biorelevant Media: An OrBiTo Study, MOL PHARMACEUT 14: 4192-4201

ORBITO: Pathak, Shriram M. et al. Model-Based Analysis of Biopharmaceutic Experiments To Improve Mechanistic Oral Absorption Modeling: An Integrated in Vitro in Vivo Extrapolation Perspective Using Ketoconazole as a Model Drug, MOL PHARMACEUT 14: 4305-4320

ORBITO: Hens, Bart et al. In Silico Modeling Approach for the Evaluation of Gastrointestinal Dissolution, Supersaturation, and Precipitation of Posaconazole, MOL PHARMACEUT 14: 4321-4333

ORBITO: Grimm, Michael et al. Gastric Emptying and Small Bowel Water Content after Administration of Grapefruit Juice Compared to Water and Isocaloric Solutions of Glucose and Fructose: A Four-Way Crossover MRI Pilot Study in Healthy Subjects, MOL PHARMACEUT 15: 548-559

ORBITO: Berben, Philippe et al. Drug permeability profiling using cell-free permeation tools: Overview and applications, EUR J PHARM SCI 119: 219-233

ORBITO: Grimm, Michael et al. Interindividual and intraindividual variability of fasted state gastric fluid volume and gastric emptying of water, EUR J PHARM BIOPHARM 127: 309-317



ORBITO: Butler, James et al. In vitro models for the prediction of in vivo performance of oral dosage forms: Recent progress from partnership through the IMI OrBiTo collaboration, EUR J PHARM BIOPHARM 136: 70-83

ORBITO: Hossain, Shakhawath et al. Molecular simulation as a computational pharmaceuticals tool to predict drug solubility, solubilization processes and partitioning, EUR J PHARM BIOPHARM 137: 46-55

ORBITO: Couto, Narciso et al. Quantitative Proteomics of Clinically Relevant Drug-Metabolizing Enzymes and Drug Transporters and Their Intercorrelations in the Human Small Intestine, DRUG METAB DISPOS 48: 245-254

PARADIGM: Vat, Lidewij Eva et al. Evaluating the return on patient engagement initiatives in medicines research and development: A literature review, HEALTH EXPECT 23: 5-18

PARADIGM: Vat, Lidewij Eva et al. Evaluation of patient engagement in medicine development: A multi-stakeholder framework with metrics, HEALTH EXPECT 24: 491-506

PD-MitoQUANT: Alam, Parvez et al. alpha-synuclein oligomers and fibrils: a spectrum of species, a spectrum of toxicities, J NEUROCHEM 150: 522-534

PD-MitoQUANT: Rey, Nolwen L. et al. alpha-Synuclein conformational strains spread, seed and target neuronal cells differentially after injection into the olfactory bulb, ACTA NEUROPATHOL COM 7:

PD-MitoQUANT: Shrivastava, Amulya Nidhi et al. Differential Membrane Binding and Seeding of Distinct alpha-Synuclein Fibrillar Polymorphs, BIOPHYS J 118: 1301-1320

PD-MitoQUANT: Courte, Josquin et al. The expression level of alpha-synuclein in different neuronal populations is the primary determinant of its prion-like seeding, SCI REP-UK 10:

PD-MitoQUANT: Dominguez-Meijide, Antonio et al. Doxycycline inhibits alpha -synuclein-associated pathologies in vitro and in vivo, NEUROBIOL DIS 151:

PD-MitoQUANT: Scheiblich, Hannah et al. Microglia jointly degrade fibrillar alpha-synuclein cargo by distribution through tunneling nanotubes, CELL 184: 5089+

PERISCOPE: Wilk, Mieszko M. et al. Immunization with whole cell but not acellular pertussis vaccines primes CD4 T-RM cells that sustain protective immunity against nasal colonization with Bordetella pertussis, EMERG MICROBES INFEC 8: 169-185

PERISCOPE: Dubois, Violaine et al. Suppression of mucosal Th17 memory responses by acellular pertussis vaccines enhances nasal Bordetella pertussis carriage, NPJ VACCINES 6:

PERISCOPE: Versteegen, Pauline et al. Responses to an acellular pertussis booster vaccination in children, adolescents, and young and older adults: A collaborative study in Finland, the Netherlands, and the United Kingdom, EBIOMEDICINE 65:

PERISCOPE: Khatri, Indu et al. Population matched (pm) germline allelic variants of immunoglobulin (IG) loci: Relevance in infectious diseases and vaccination studies in human populations, GENES IMMUN 22: 172-186

PERISCOPE: Evers, Felix et al. Composition and stage dynamics of mitochondrial complexes in *Plasmodium falciparum*, NAT COMMUN 12:

PHAGO: Schlepckow, Kai et al. An Alzheimer-associated TREM2 variant occurs at the ADAM cleavage site and affects shedding and phagocytic function, EMBO MOL MED 9: 1356-1365

PHAGO: Thornton, Peter et al. TREM2 shedding by cleavage at the H157-S158 bond is accelerated for the Alzheimers disease-associated H157Y variant, EMBO MOL MED 9: 1366-1378

PHAGO: Garcia-Reitboeck, Pablo et al. Human Induced Pluripotent Stem Cell-Derived Microglia-Like Cells Harboring TREM2 Missense Mutations Show Specific Deficits in Phagocytosis, CELL REP 24: 2300-2311

PHAGO: Xiang, Xianyuan et al. The Trem2 R47H Alzheimers risk variant impairs splicing and reduces Trem2 mRNA and protein in mice but not in humans, MOL NEURODEGENER 13:

PHAGO: Parhizkar, Samira et al. Loss of TREM2 function increases amyloid seeding but reduces plaque-associated ApoE, NAT NEUROSCI 22: 191-+

PHAGO: Linnartz-Gerlach, Bettina et al. TREM2 triggers microglial density and age-related neuronal loss, GLIA 67: 539-550

PHAGO: Houtman, Judith et al. Beclin1-driven autophagy modulates the inflammatory response of microglia via NLRP3, EMBO J 38:

PHAGO: Boza-Serrano, Antonio et al. Galectin-3, a novel endogenous TREM2 ligand, detrimentally regulates inflammatory response in Alzheimers disease, ACTA NEUROPATHOL 138: 251-273

PHAGO: Brown, Guy C. et al. The endotoxin hypothesis of neurodegeneration, J NEUROINFLAMM 16:

PHAGO: Miles, Luke A. et al. Small Molecule Binding to Alzheimer Risk Factor CD33 Promotes A beta Phagocytosis, ISCIENCE 19: 110-+

PHAGO: Piers, Thomas M. et al. A locked immunometabolic switch underlies TREM2 R47H loss of function in human iPSC-derived microglia, FASEB J 34: 2436-2450

PHAGO: Schlepckow, Kai et al. Enhancing protective microglial activities with a dual function TREM2 antibody to the stalk region, EMBO MOL MED 12:

PHAGO: Puigdellivol, Mar et al. Sialylation and Galectin-3 in Microglia-Mediated Neuroinflammation and Neurodegeneration, FRONT CELL NEUROSCI 14:

PHAGO: Meinhardt, Jenny et al. Olfactory transmucosal SARS-CoV-2 invasion as a port of central nervous system entry in individuals with COVID-19, NAT NEUROSCI 24: 168-175

PHAGO: Lewcock, Joseph W. et al. Emerging Microglia Biology Defines Novel Therapeutic Approaches for Alzheimers Disease, NEURON 108: 801-821

PHAGO: Liu, Wenfei et al. Trem2 promotes anti-inflammatory responses in microglia and is suppressed under pro-inflammatory conditions, HUM MOL GENET 29: 3224-3248

PHAGO: Wissfeld, Jannis et al. Deletion of Alzheimers disease-associated CD33 results in an inflammatory human microglia phenotype, *GLIA* 69: 1393-1412

PHAGO: Butler, Claire A. et al. Microglial phagocytosis of neurons in neurodegeneration, and its regulation, *J NEUROCHEM* 158: 621-639

PHAGO: Ann Butler, Claire et al. CD33M inhibits microglial phagocytosis, migration and proliferation, but the Alzheimers disease-protective variant CD33m stimulates phagocytosis and proliferation, and inhibits adhesion, *J NEUROCHEM* 158: 297-310

PHAGO: Vilalta, Anna et al. Wild-type sTREM2 blocks A beta aggregation and neurotoxicity, but the Alzheimers R47H mutant increases A beta aggregation, *J BIOL CHEM* 296:

Pharma-Cog: Frisoni, Giovanni B. et al. The clinical use of structural MRI in Alzheimer disease, *NAT REV NEUROL* 6: 67-77

Pharma-Cog: Drago, Valeria et al. Disease Tracking Markers for Alzheimers Disease at the Prodromal (MCI) Stage, *J ALZHEIMERS DIS* 26: 159-199

Pharma-Cog: Languille, S. et al. The grey mouse lemur: A non-human primate model for ageing studies, *AGEING RES REV* 11: 150-162

Pharma-Cog: Carrillo, Maria C. et al. Worldwide Alzheimers Disease Neuroimaging Initiative, *ALZHEIMERS DEMENT* 8: 337-342

Pharma-Cog: Babiloni, Claudio et al. Resting state cortical electroencephalographic rhythms are related to gray matter volume in subjects with mild cognitive impairment and Alzheimers disease, *HUM BRAIN MAPP* 34: 1427-1446

Pharma-Cog: Jovicich, Jorge et al. Brain morphometry reproducibility in multi-center 3 T MRI studies: A comparison of cross-sectional and longitudinal segmentations, *NEUROIMAGE* 83: 472-484

Pharma-Cog: Jovicich, Jorge et al. Multisite longitudinal reliability of tract-based spatial statistics in diffusion tensor imaging of healthy elderly subjects, *NEUROIMAGE* 101: 390-403

Pharma-Cog: Jovicich, Jorge et al. Longitudinal reproducibility of default-mode network connectivity in healthy elderly participants: A multicentric resting-state fMRI study, *NEUROIMAGE* 124: 442-454

Pharma-Cog: Galluzzi, S. et al. Clinical and biomarker profiling of prodromal Alzheimers disease in workpackage 5 of the Innovative Medicines Initiative PharmaCog project: a European ADNI study, *J INTERN MED* 279: 576-591

Pharma-Cog: Pini, Lorenzo et al. Brain atrophy in Alzheimers Disease and aging, *AGEING RES REV* 30: 25-48

Pharma-Cog: Albi, Angela et al. Free water elimination improves test-retest reproducibility of diffusion tensor imaging indices in the brain: A longitudinal multisite study of healthy elderly subjects, *HUM BRAIN MAPP* 38: 12-26

Pharma-Cog: Nathan, Pradeep J. et al. Association between CSF biomarkers, hippocampal volume and cognitive function in patients with amnesic mild cognitive impairment (MCI), *NEUROBIOL AGING* 53: 1-10

Pharma-Cog: Lim, Chai K. et al. Involvement of the kynurenine pathway in the pathogenesis of Parkinsons disease, *PROG NEUROBIOL* 155: 76-95

PIONEER: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), *GENOME MED* 12:

PRECISESADS: Alvarez-Errico, Damiana et al. Epigenetic control of myeloid cell differentiation, identity and function, *NAT REV IMMUNOL* 15: 7-17

PRECISESADS: Teruel, Maria et al. The genetic basis of systemic lupus erythematosus: What are the risk factors and what have we learned, *J AUTOIMMUN* 74: 161-175

PRECISESADS: Barturen, Guillermo et al. Moving towards a molecular taxonomy of autoimmune rheumatic diseases, *NAT REV RHEUMATOL* 14: 75-93

PRECISESADS: Toro-Dominguez, Daniel et al. Stratification of Systemic Lupus Erythematosus Patients Into Three Groups of Disease Activity Progression According to Longitudinal Gene Expression, *ARTHRITIS RHEUMATOL* 70: 2025-2035

PRECISESADS: Acosta-Herrera, Marialbert et al. Genome-wide meta-analysis reveals shared new loci in systemic seropositive rheumatic diseases, *ANN RHEUM DIS* 78: 311-319

PRECISESADS: Beretta, Lorenzo et al. Genome-wide whole blood transcriptome profiling in a large European cohort of systemic sclerosis patients, *ANN RHEUM DIS* 79: 1218-1226

PRECISESADS: Bossini-Castillo, Lara et al. Genomic Risk Score impact on susceptibility to systemic sclerosis, *ANN RHEUM DIS* 80: 118-127

PRECISESADS: Toro-Dominguez, Daniel et al. A survey of gene expression meta-analysis: methods and applications, *BRIEF BIOINFORM* 22: 1694-1705

PRECISESADS: Simon, Quentin et al. A Proinflammatory Cytokine Network Profile in Th1/Type 1 Effector B Cells Delineates a Common Group of Patients in Four Systemic Autoimmune Diseases, *ARTHRITIS RHEUMATOL* 73: 1550-1561

Predict: Tanos, Tamara et al. Progesterone/RANKL Is a Major Regulatory Axis in the Human Breast, *SCI TRANSL MED* 5:

Predict: Nieminen, Anni I. et al. Myc-induced AMPK-phospho p53 pathway activates Bak to sensitize mitochondrial apoptosis, *P NATL ACAD SCI USA* 110: E1839-E1848

Predict: de Jong, Marion et al. Imaging preclinical tumour models: improving translational power, *NAT REV CANCER* 14: 481-493

Predict: Hickman, John A. et al. Three-dimensional models of cancer for pharmacology and cancer cell biology: Capturing tumor complexity in vitro/ex vivo, *BIOTECHNOL J* 9: 1115-1128

Predict: Metsalu, Tauno et al. ClustVis: a web tool for visualizing clustering of multivariate data using Principal Component Analysis and heatmap, *NUCLEIC ACIDS RES* 43: W566-W570

Predict: Davies, Emma et al. Capturing complex tumour biology in vitro: histological and molecular characterisation of precision cut slices, *SCI REP-UK* 5:

Predict: Gualda, Emilio J. et al. SPIM-fluid: open source light-sheet based platform for high-throughput imaging, BIOMED OPT EXPRESS 6: 4447-4456

Predict: Estrada, Marta F. et al. Modelling the tumour microenvironment in long-term microencapsulated 3D co-cultures recapitulates phenotypic features of disease progression, BIOMATERIALS 78: 50-61

Predict: Santo, Vitor E. et al. Adaptable stirred-tank culture strategies for large scale production of multicellular spheroid-based tumor cell models, J BIOTECHNOL 221: 118-129

Predict: Sflomos, George et al. A Preclinical Model for ER alpha-Positive Breast Cancer Points to the Epithelial Microenvironment as Determinant of Luminal Phenotype and Hormone Response, CANCER CELL 29: 407-422

Predict: Stock, Kristin et al. Capturing tumor complexity in vitro: Comparative analysis of 2D and 3D tumor models for drug discovery, SCI REP-UK 6:

Predict: Dobrolecki, Lacey E. et al. Patient-derived xenograft (PDX) models in basic and translational breast cancer research, CANCER METAST REV 35: 547-573

Predict: Santo, Vitor E. et al. Drug screening in 3D in vitro tumor models: overcoming current pitfalls of efficacy read-outs, BIOTECHNOL J 12:

Predict: de Witte, Samantha F. H. et al. Cytokine treatment optimises the immunotherapeutic effects of umbilical cord-derived MSC for treatment of inflammatory liver disease, STEM CELL RES THER 8:

Predict: Blom, Sami et al. Systems pathology by multiplexed immunohistochemistry and whole-slide digital image analysis, SCI REP-UK 7:

Predict: Rebelo, Sofia P. et al. 3D-3-culture: A tool to unveil macrophage plasticity in the tumour microenvironment, BIOMATERIALS 163: 185-197

Predict: Arandkar, Sharathchandra et al. Altered p53 functionality in cancer-associated fibroblasts contributes to their cancer-supporting features, P NATL ACAD SCI USA 115: 6410-6415

PreDiCT-TB: Sisniega, A. et al. Monte Carlo study of the effects of system geometry and antiscatter grids on cone-beam CT scatter distributions, MED PHYS 40:

PreDiCT-TB: Svensson, Elin M. et al. Model-Based Estimates of the Effects of Efavirenz on Bedaquiline Pharmacokinetics and Suggested Dose Adjustments for Patients Coinfected with HIV and Tuberculosis, ANTIMICROB AGENTS CH 57: 2780-2787

PreDiCT-TB: Zumla, Alimuddin I. et al. New antituberculosis drugs, regimens, and adjunct therapies: needs, advances, and future prospects, LANCET INFECT DIS 14: 327-340

PreDiCT-TB: Manina, Giulia et al. Stress and Host Immunity Amplify Mycobacterium tuberculosis Phenotypic Heterogeneity and Induce Nongrowing Metabolically Active Forms, CELL HOST MICROBE 17: 32-46

PreDiCT-TB: Svensson, Elin M. et al. Rifampicin and rifapentine significantly reduce concentrations of bedaquiline, a new anti-TB drug, J ANTIMICROB CHEMOTH 70: 1106-1114

PreDiCT-TB: Ates, Louis S. et al. Essential Role of the ESX-5 Secretion System in Outer Membrane Permeability of Pathogenic Mycobacteria, PLOS GENET 11:

PreDiCT-TB: Hu, Yanmin et al. High-dose rifampicin kills persisters, shortens treatment duration, and reduces relapse rate in vitro and in vivo, FRONT MICROBIOL 6:

PreDiCT-TB: Kaufmann, Stefan H. E. et al. Molecular Determinants in Phagocyte-Bacteria Interactions, IMMUNITY 44: 476-491

PreDiCT-TB: Lipworth, S. et al. Defining dormancy in mycobacterial disease, TUBERCULOSIS 99: 131-142

PreDiCT-TB: Boritsch, Eva C. et al. Key experimental evidence of chromosomal DNA transfer among selected tuberculosis-causing mycobacteria, P NATL ACAD SCI USA 113: 9876-9881

PreDiCT-TB: Boritsch, Eva C. et al. pks5-recombination-mediated surface remodelling in Mycobacterium tuberculosis emergence, NAT MICROBIOL 1:

PreDiCT-TB: Kaufmann, Stefan H. E. et al. Host-directed therapies for bacterial and viral infections, NAT REV DRUG DISCOV 17: 35-56

PreDiCT-TB: Svensson, Robin J. et al. A Population Pharmacokinetic Model Incorporating Saturable Pharmacokinetics and Autoinduction for High Rifampicin Doses, CLIN PHARMACOL THER 103: 674-683

PreDiCT-TB: Svensson, Robin J. et al. Greater Early Bactericidal Activity at Higher Rifampicin Doses Revealed by Modeling and Clinical Trial Simulations, J INFECT DIS 218: 991-999

PreDiCT-TB: Baranowski, Catherine et al. Maturing Mycobacterium smegmatis peptidoglycan requires non-canonical crosslinks to maintain shape, ELIFE 7:

PreDiCT-TB: Pei, Gang et al. Cellular stress promotes NOD1/2-dependent inflammation via the endogenous metabolite sphingosine-1-phosphate, EMBO J 40:

PREFER: van Overbeeke, Eline et al. Factors and situations influencing the value of patient preference studies along the medical product lifecycle: a literature review, DRUG DISCOV TODAY 24: 57-68

PREFER: Russo, Selena et al. Understanding Patients Preferences: A Systematic Review of Psychological Instruments Used in Patients Preference and Decision Studies, VALUE HEALTH 22: 491-501

PREFER: Soekhai, Vikas et al. Methods for exploring and eliciting patient preferences in the medical product lifecycle: a literature review, DRUG DISCOV TODAY 24: 1324-1331

PREFER: Janssens, Rosanne et al. Patient Preferences in the Medical Product Life Cycle: What do Stakeholders Think? Semi-Structured Qualitative Interviews in Europe and the USA, PATIENT 12: 513-526

PREFER: van Overbeeke, Eline et al. Design, Conduct, and Use of Patient Preference Studies in the Medical Product Life Cycle: A Multi-Method Study, FRONT PHARMACOL 10:

PREFER: Durosini, Ilaria et al. Patient Preferences for Lung Cancer Treatment: A Qualitative Study Protocol Among Advanced Lung Cancer Patients, FRONT PUBLIC HEALTH 9:

PRISM: Saris, I. M. J. et al. Social functioning in patients with depressive and anxiety disorders, ACTA PSYCHIAT SCAND 136: 352-361

PRISM: Bralten, J. et al. Autism spectrum disorders and autistic traits share genetics and biology, MOL PSYCHIATR 23: 1205-1212

PRISM: Arango, Celso et al. Preventive strategies for mental health, LANCET PSYCHIAT 5: 591-604

PRISM: Galderisi, Silvana et al. Negative symptoms of schizophrenia: new developments and unanswered research questions, LANCET PSYCHIAT 5: 664-677

PRISM: Downs, Johnny et al. Negative Symptoms in Early-Onset Psychosis and Their Association With Antipsychotic Treatment Failure, SCHIZOPHRENIA BULL 45: 69-79

PRISM: Kas, Martien J. et al. A quantitative approach to neuropsychiatry: The why and the how, NEUROSCI BIOBEHAV R 97: 3-9

PRISM: Porcelli, Stefano et al. Social brain, social dysfunction and social withdrawal, NEUROSCI BIOBEHAV R 97: 10-33

PRISM: Winsky-Sommerer, Raphaelle et al. Disturbances of sleep quality, timing and structure and their relationship with other neuropsychiatric symptoms in Alzheimers disease and schizophrenia: Insights from studies in patient populations and animal models, NEUROSCI BIOBEHAV R 97: 112-137

PRISM: Hornix, Betty E. et al. Multisensory cortical processing and dysfunction across the neuropsychiatric spectrum, NEUROSCI BIOBEHAV R 97: 138-151

PRISM: Fraguas, David et al. Oxidative Stress and Inflammation in First-Episode Psychosis: A Systematic Review and Meta-analysis, SCHIZOPHRENIA BULL 45: 742-751

PRISM: Fraguas, David et al. Dietary Interventions for Autism Spectrum Disorder: A Meta-analysis, PEDIATRICS 144:

PRISM: van Heukelum, Sabrina et al. Where is Cingulate Cortex? A Cross-Species View, TRENDS NEUROSCI 43: 285-299

PRISM: de Pablo, Gonzalo Salazar et al. Impact of coronavirus syndromes on physical and mental health of health care workers: Systematic review and meta-analysis, J AFFECT DISORDERS 275: 48-57

PRISM: Moreno, Carmen et al. How mental health care should change as a consequence of the COVID-19 pandemic, LANCET PSYCHIAT 7: 813-824

PRISM: Davies, Robert W. et al. Using common genetic variation to examine phenotypic expression and risk prediction in 22q11.2 deletion syndrome, NAT MED 26:

PRISM: Fraguas, David et al. Assessment of School Anti-Bullying Interventions A Meta-analysis of Randomized Clinical Trials, JAMA PEDIATR 175: 44-55

PRISM: Fusar-Poli, Paolo et al. Preventive psychiatry: a blueprint for improving the mental health of young people, *WORLD PSYCHIATRY* 20: 200-221

PRISM: van Heukelum, Sabrina et al. A central role for anterior cingulate cortex in the control of pathological aggression, *CURR BIOL* 31: 2321-+

PRISM: Persico, Antonio M. et al. The pediatric psychopharmacology of autism spectrum disorder: A systematic review - Part I: The past and the present, *PROG NEURO-PSYCHOPH* 110:

PROACTIVE: van Remoortel, Hans et al. Validity of Six Activity Monitors in Chronic Obstructive Pulmonary Disease: A Comparison with Indirect Calorimetry, *PLOS ONE* 7:

PROACTIVE: van Remoortel, Hans et al. Validity of activity monitors in health and chronic disease: a systematic review, *INT J BEHAV NUTR PHY* 9:

PROACTIVE: Rabinovich, Roberto A. et al. Validity of physical activity monitors during daily life in patients with COPD, *EUR RESPIR J* 42: 1205-1215

PROACTIVE: Gimeno-Santos, Elena et al. Determinants and outcomes of physical activity in patients with COPD: a systematic review, *THORAX* 69: 731-739

PROACTIVE: Demeyer, Heleen et al. Standardizing the Analysis of Physical Activity in Patients With COPD Following a Pulmonary Rehabilitation Program, *CHEST* 146: 318-327

PROACTIVE: Gimeno-Santos, Elena et al. The PROactive instruments to measure physical activity in patients with chronic obstructive pulmonary disease, *EUR RESPIR J* 46: 988-1000

PROACTIVE: Demeyer, Heleen et al. The Minimal Important Difference in Physical Activity in Patients with COPD, *PLOS ONE* 11:

PROACTIVE: Demeyer, H. et al. Physical activity is increased by a 12-week semiautomated telecoaching programme in patients with COPD: a multicentre randomised controlled trial, *THORAX* 72: 415-423

PROTECT: van Staa, T. P. et al. Glucose-lowering agents and the patterns of risk for cancer: a study with the General Practice Research Database and secondary care data, *DIABETOLOGIA* 55: 654-665

PROTECT: Lalmohamed, Arief et al. Risk of fracture after bariatric surgery in the United Kingdom: population based, retrospective cohort study, *BRIT MED J* 345:

PROTECT: Ryan, Patrick B. et al. Defining a Reference Set to Support Methodological Research in Drug Safety, *DRUG SAFETY* 36: S33-S47

PROTECT: Abbing-Karahagopian, V. et al. Antidepressant prescribing in five European countries: application of common definitions to assess the prevalence, clinical observations, and methodological implications, *EUR J CLIN PHARMACOL* 70: 849-857

PROTECT: Mt-Isa, Shahrul et al. Balancing benefit and risk of medicines: a systematic review and classification of available methodologies, *PHARMACOEPIDEM DR S* 23: 667-678



PROTECT: Ali, M. Sanni et al. Reporting of covariate selection and balance assessment in propensity score analysis is suboptimal: a systematic review, *J CLIN EPIDEMIOL* 68: 122-131

PROTECT: Candore, Gianmario et al. Comparison of Statistical Signal Detection Methods Within and Across Spontaneous Reporting Databases, *DRUG SAFETY* 38: 577-587

PROTECT: Wisniewski, Antoni F. Z. et al. Good Signal Detection Practices: Evidence from IMI PROTECT, *DRUG SAFETY* 39: 469-490

Quic-Concept: Lambin, Philippe et al. Radiomics: Extracting more information from medical images using advanced feature analysis, *EUR J CANCER* 48: 441-446

Quic-Concept: Asselin, Marie-Claude et al. Quantifying heterogeneity in human tumours using MRI and PET, *EUR J CANCER* 48: 447-455

Quic-Concept: van der Heide, Uulke A. et al. Functional MRI for radiotherapy dose painting, *MAGN RESON IMAGING* 30: 1216-1223

Quic-Concept: Lambin, Philippe et al. Predicting outcomes in radiation oncology-multifactorial decision support systems, *NAT REV CLIN ONCOL* 10: 27-40

Quic-Concept: Velazquez, Emmanuel Rios et al. A semiautomatic CT-based ensemble segmentation of lung tumors: Comparison with oncologists delineations and with the surgical specimen, *RADIOTHER ONCOL* 105: 167-173

Quic-Concept: Challapalli, Amarnath et al. F-18-ICMT-11, a Caspase-3-Specific PET Tracer for Apoptosis: Biodistribution and Radiation Dosimetry, *J NUCL MED* 54: 1551-1556

Quic-Concept: Leijenaar, Ralph T. H. et al. Stability of FDG-PET Radiomics features: An integrated analysis of test-retest and inter-observer variability, *ACTA ONCOL* 52: 1391-1397

Quic-Concept: Lambin, Philippe et al. Rapid Learning health care in oncology - An approach towards decision support systems enabling customised radiotherapy, *RADIOTHER ONCOL* 109: 159-164

Quic-Concept: Velazquez, Emmanuel Rios et al. Volumetric CT-based segmentation of NSCLC using 3D-Slicer, *SCI REP-UK* 3:

Quic-Concept: Aerts, Hugo J. W. L. et al. Decoding tumour phenotype by noninvasive imaging using a quantitative radiomics approach, *NAT COMMUN* 5:

Quic-Concept: Parmar, Chintan et al. Robust Radiomics Feature Quantification Using Semiautomatic Volumetric Segmentation, *PLOS ONE* 9:

Quic-Concept: Frings, Virginie et al. Repeatability of Metabolically Active Tumor Volume Measurements with FDG PET/CT in Advanced Gastrointestinal Malignancies: A Multicenter Study, *RADIOLOGY* 273: 539-548

Quic-Concept: Coroller, Thibaud P. et al. CT-based radiomic signature predicts distant metastasis in lung adenocarcinoma, *RADIOTHER ONCOL* 114: 345-350

Quic-Concept: Peeters, Sarah G. J. A. et al. TH-302 in Combination with Radiotherapy Enhances the Therapeutic Outcome and Is Associated with Pretreatment [F-18]HX4 Hypoxia PET Imaging, CLIN CANCER RES 21: 2984-2992

Quic-Concept: Leijenaar, Ralph T. H. et al. The effect of SUV discretization in quantitative FDG-PET Radiomics: the need for standardized methodology in tumor texture analysis, SCI REP-UK 5:

Quic-Concept: Panth, Kranthi Marella et al. Is there a causal relationship between genetic changes and radiomics-based image features? An in vivo preclinical experiment with doxycycline inducible GADD34 tumor cells, RADIOTHER ONCOL 116: 462-466

Quic-Concept: Leijenaar, Ralph T. H. et al. External validation of a prognostic CT-based radiomic signature in oropharyngeal squamous cell carcinoma, ACTA ONCOL 54: 1423-1429

Quic-Concept: Bollineni, V. R. et al. A systematic review on [F-18]FLT-PET uptake as a measure of treatment response in cancer patients, EUR J CANCER 55: 81-97

Quic-Concept: Scrivener, Madeleine et al. Radiomics applied to lung cancer: a review, TRANSL CANCER RES 5: 398-409

Quic-Concept: Huizinga, W. et al. PCA-based groupwise image registration for quantitative MRI, MED IMAGE ANAL 29: 65-78

Quic-Concept: Lambin, Philippe et al. Decision support systems for personalized and participative radiation oncology, ADV DRUG DELIVER REV 109: 131-153

Quic-Concept: OConnor, James P. B. et al. Imaging biomarker roadmap for cancer studies, NAT REV CLIN ONCOL 14: 169-186

Quic-Concept: van Timmeren, Janna E. et al. Survival prediction of non-small cell lung cancer patients using radiomics analyses of cone-beam CT images, RADIOTHER ONCOL 123: 363-369

Quic-Concept: Grossmann, Patrick et al. Defining the biological basis of radiomic phenotypes in lung cancer, ELIFE 6:

Quic-Concept: Larue, Ruben T. H. M. et al. 4DCT imaging to assess radiomics feature stability: An investigation for thoracic cancers, RADIOTHER ONCOL 125: 147-153

Quic-Concept: Lambin, Philippe et al. Radiomics: the bridge between medical imaging and personalized medicine, NAT REV CLIN ONCOL 14: 749-762

Quic-Concept: deSouza, N. M. et al. Implementing diffusion-weighted MRI for body imaging in prospective multicentre trials: current considerations and future perspectives, EUR RADIOL 28: 1118-1131

Quic-Concept: Carvalho, Sara et al. F-18-fluorodeoxyglucose positron-emission tomography (FDG-PET)-Radiomics of metastatic lymph nodes and primary tumor in non-small cell lung cancer (NSCLC) - A prospective externally validated study, PLOS ONE 13:

Quic-Concept: Leijenaar, Ralph T. H. et al. Development and validation of a radiomic signature to predict HPV (p16) status from standard CT imaging: a multicenter study, BRIT J RADIOL 91:

Quic-Concept: Sanduleanu, Sebastian et al. Tracking tumor biology with radiomics: A systematic review utilizing a radiomics quality score, *RADIOTHER ONCOL* 127: 349-360

Quic-Concept: Deist, Timo M. et al. Machine learning algorithms for outcome prediction in (chemo)radiotherapy: An empirical comparison of classifiers, *MED PHYS* 45: 3449-3459

Quic-Concept: Peerlings, Jurgen et al. Stability of radiomics features in apparent diffusion coefficient maps from a multi-centre test-retest trial, *SCI REP-UK* 9:

Quic-Concept: van Timmeren, Janna E. et al. Challenges and caveats of a multi-center retrospective radiomics study: an example of early treatment response assessment for NSCLC patients using FDG-PET/CT radiomics, *PLOS ONE* 14:

Quic-Concept: Zwanenburg, Alex et al. The Image Biomarker Standardization Initiative: Standardized Quantitative Radiomics for High-Throughput Image-based Phenotyping, *RADIOLOGY* 295: 328-338

RADAR-AD: Stavropoulos, Thanos G. et al. IoT Wearable Sensors and Devices in Elderly Care: A Literature Review, *SENSORS-BASEL* 20:

RADAR-AD: Muurling, Marijn et al. Remote monitoring technologies in Alzheimers disease: design of the RADAR-AD study, *ALZHEIMERS RES THER* 13:

RADAR-CNS: Simblett, Sara et al. Barriers to and Facilitators of Engagement With Remote Measurement Technology for Managing Health: Systematic Review and Content Analysis of Findings, *J MED INTERNET RES* 20:

RADAR-CNS: Bruno, Elisa et al. Wearable technology in epilepsy: The views of patients, caregivers, and healthcare professionals, *EPILEPSY BEHAV* 85: 141-149

RADAR-CNS: Cummins, Nicholas et al. Speech analysis for health: Current state-of-the-art and the increasing impact of deep learning, *METHODS* 151: 41-54

RADAR-CNS: Rintala, Aki et al. Response Compliance and Predictors Thereof in Studies Using the Experience Sampling Method, *PSYCHOL ASSESSMENT* 31: 226-235

RADAR-CNS: Simblett, Sara et al. Barriers to and Facilitators of Engagement With mHealth Technology for Remote Measurement and Management of Depression: Qualitative Analysis, *JMIR MHEALTH UHEALTH* 7:

RADAR-CNS: Matcham, F. et al. Remote assessment of disease and relapse in major depressive disorder (RADAR-MDD): a multi-centre prospective cohort study protocol, *BMC PSYCHIATRY* 19:

RADAR-CNS: Ranjan, Yatharth et al. RADAR-Base: Open Source Mobile Health Platform for Collecting, Monitoring, and Analyzing Data Using Sensors, Wearables, and Mobile Devices, *JMIR MHEALTH UHEALTH* 7:

RADAR-CNS: Difrancesco, Sonia et al. Sleep, circadian rhythm, and physical activity patterns in depressive and anxiety disorders: A 2-week ambulatory assessment study, *DEPRESS ANXIETY* 36: 975-986

RADAR-CNS: Zhang, Zixing et al. Snore-GANs: Improving Automatic Snore Sound Classification With Synthesized Data, *IEEE J BIOMED HEALTH* 24: 300-310

RADAR-CNS: Simblett, Sara Katherine et al. Patients experience of wearing multimodal sensor devices intended to detect epileptic seizures: A qualitative analysis, *EPILEPSY BEHAV* 102:

RADAR-CNS: Bruno, Elisa et al. Seizure detection at home: Do devices on the market match the needs of people living with epilepsy and their caregivers?, *EPILEPSIA* 61: S11-S24

RADAR-CNS: Dalla Costa, Gloria et al. Real-time assessment of COVID-19 prevalence among multiple sclerosis patients: a multicenter European study, *NEUROL SCI* 41: 1647-1650

RADAR-CNS: Viana, Pedro F. et al. 230 days of ultra long-term subcutaneous EEG: seizure cycle analysis and comparison to patient diary, *ANN CLIN TRANSL NEUR* 8: 288-293

RADAR-CNS: Sun, Shaoxiong et al. Using Smartphones and Wearable Devices to Monitor Behavioral Changes During COVID-19, *J MED INTERNET RES* 22:

RADAR-CNS: Pegg, Emily J. et al. Interictal electroencephalographic functional network topology in drug-resistant and well-controlled idiopathic generalized epilepsy, *EPILEPSIA* 62: 492-503

RADAR-CNS: Difrancesco, Sonia et al. The day -to -day bidirectional longitudinal association between objective and self-reported sleep and affect: An ambulatory assessment study, *J AFFECT DISORDERS* 283: 165-171

RADAR-CNS: Qian, Kun et al. Can Machine Learning Assist Locating the Excitation of Snore Sound? A Review, *IEEE J BIOMED HEALTH* 25: 1233-1246

RADAR-CNS: Pitharouli, Maria C. et al. Elevated C-Reactive Protein in Patients With Depression, Independent of Genetic, Health, and Psychosocial Factors: Results From the UK Biobank, *AM J PSYCHIAT* 178: 522-529

RADAR-CNS: Viana, Pedro F. et al. Signal quality and power spectrum analysis of remote ultra long-term subcutaneous EEG, *EPILEPSIA* 62: 1820-1828

RADAR-CNS: Bruno, Elisa et al. Wearable devices for seizure detection: Practical experiences and recommendations from the Wearables for Epilepsy And Research (WEAR) International Study Group, *EPILEPSIA* 62: 2307-2321

RAPP-ID: Schechner, Vered et al. Epidemiological Interpretation of Studies Examining the Effect of Antibiotic Usage on Resistance, *CLIN MICROBIOL REV* 26: 289-307

RAPP-ID: Leirs, Karen et al. Bioassay Development for Ultrasensitive Detection of Influenza A Nucleoprotein Using Digital ELISA, *ANAL CHEM* 88: 8450-8458

RAPP-ID: Faridi, Muhammad Asim et al. Elasto-inertial microfluidics for bacteria separation from whole blood for sepsis diagnostics, *J NANOBIOELECTRON* 15:

RAPP-ID: Guha, Arnab et al. Direct detection of small molecules using a nano-molecular imprinted polymer receptor and a quartz crystal resonator driven at a fixed frequency and amplitude, *BIOSENS BIOELECTRON* 158:

RESCEU: Lin, Gu-Lung et al. Epidemiology and Immune Pathogenesis of Viral Sepsis, *FRONT IMMUNOL* 9:

RESCEU: Li, You et al. Global patterns in monthly activity of influenza virus, respiratory syncytial virus, parainfluenza virus, and metapneumovirus: a systematic analysis, LANCET GLOB HEALTH 7: E1031-E1045

RESCEU: Barr, Rachael et al. Respiratory syncytial virus: diagnosis, prevention and management, THER ADV INFECT DIS 6:

RESCEU: Wang, Xin et al. Global burden of respiratory infections associated with seasonal influenza in children under 5 years in 2018: a systematic review and modelling study, LANCET GLOB HEALTH 8: E497-E510

RESCEU: Lowensteyn, Yvette N. et al. Respiratory Syncytial Virus-related Death in Children With Down Syndrome The RSV GOLD Study, PEDIATR INFECT DIS J 39: 665-670

RESCEU: Habibi, Maximillian S. et al. Neutrophilic inflammation in the respiratory mucosa predisposes to RSV infection, SCIENCE 370: 188+

RESCEU: Tabor, David E. et al. Global Molecular Epidemiology of Respiratory Syncytial Virus from the 2017-2018 INFORM-RSV Study, J CLIN MICROBIOL 59:

RESCEU: Shi, Ting et al. The Etiological Role of Common Respiratory Viruses in Acute Respiratory Infections in Older Adults: A Systematic Review and Meta-analysis, J INFECT DIS 222: S563-S569

RESCEU: Shi, Ting et al. Global and Regional Burden of Hospital Admissions for Pneumonia in Older Adults: A Systematic Review and Meta-Analysis, J INFECT DIS 222: S570-S576

RESCEU: Shi, Ting et al. Global Disease Burden Estimates of Respiratory Syncytial Virus-Associated Acute Respiratory Infection in Older Adults in 2015: A Systematic Review and Meta-Analysis, J INFECT DIS 222: S577-S583

RESCEU: Shi, Ting et al. Association Between Respiratory Syncytial Virus-Associated Acute Lower Respiratory Infection in Early Life and Recurrent Wheeze and Asthma in Later Childhood, J INFECT DIS 222: S628-S633

RESCEU: Korsten, Koos et al. Burden of respiratory syncytial virus infection in community-dwelling older adults in Europe (RESCEU): an international prospective cohort study, EUR RESPIR J 57:

RESCEU: Linssen, Rosalie S. et al. Burden of respiratory syncytial virus bronchiolitis on the Dutch pediatric intensive care units, EUR J PEDIATR 180: 3141-3149

RESCEU: Thomas, Emilia et al. Burden of Respiratory Syncytial Virus Infection During the First Year of Life, J INFECT DIS 223: 811-817

RespiriNTM: Kilinc, Gul et al. Host-directed therapy to combat mycobacterial infections\*, IMMUNOL REV 301: 62-83

RespiriTb: Kilinc, Gul et al. Host-directed therapy to combat mycobacterial infections\*, IMMUNOL REV 301: 62-83

RHAPSODY: Franks, Paul W. et al. Exposing the exposures responsible for type 2 diabetes and obesity, SCIENCE 354: 69-73

RHAPSODY: McCarthy, Mark I. et al. Painting a new picture of personalised medicine for diabetes, DIABETOLOGIA 60: 793-799

RHAPSODY: Schmid, Vera et al. Safety of intranasal human insulin: A review, DIABETES OBES METAB 20: 1563-1577

RHAPSODY: Diedisheim, Marc et al. Modeling human pancreatic beta cell dedifferentiation, MOL METAB 10: 74-86

RHAPSODY: Falcon, Benjamin et al. Structures of filaments from Picks disease reveal a novel tau protein fold, NATURE 561: 137-+

RHAPSODY: Falcon, Benjamin et al. Tau filaments from multiple cases of sporadic and inherited Alzheimers disease adopt a common fold, ACTA NEUROPATHOL 136: 699-708

RHAPSODY: Prasad, R. B. et al. Precision medicine in type 2 diabetes, J INTERN MED 285: 40-48

RHAPSODY: Salem, Victoria et al. Leader beta-cells coordinate Ca<sup>2+</sup> dynamics across pancreatic islets in vivo, NAT METAB 1: 615-629

RHAPSODY: Akalestou, Elina et al. Glucocorticoid Metabolism in Obesity and Following Weight Loss, FRONT ENDOCRINOL 11:

RHAPSODY: Cohrs, Christian M. et al. Dysfunction of Persisting beta Cells Is a Key Feature of Early Type 2 Diabetes Pathogenesis, CELL REP 31:

RHAPSODY: Xavier, Gabriela Da Silva et al. Metabolic and Functional Heterogeneity in Pancreatic beta Cells, J MOL BIOL 432: 1395-1406

RHAPSODY: Lytrivi, Maria et al. Recent Insights Into Mechanisms of beta-Cell Lipo- and Glucolipototoxicity in Type 2 Diabetes, J MOL BIOL 432: 1514-1534

RHAPSODY: Georgiadou, Eleni et al. The pore-forming subunit MCU of the mitochondrial Ca<sup>2+</sup> uniporter is required for normal glucose-stimulated insulin secretion in vitro and in vivo in mice, DIABETOLOGIA 63: 1368-1381

RHAPSODY: Eizirik, Decio L. et al. Pancreatic beta-cells in type 1 and type 2 diabetes mellitus: different pathways to failure, NAT REV ENDOCRINOL 16: 349-362

RHAPSODY: Lucey, Maria et al. Disconnect between signalling potency and in vivo efficacy of pharmacokinetically optimised biased glucagon-like peptide-1 receptor agonists, MOL METAB 37:

RHAPSODY: Chung, Wendy K. et al. Precision medicine in diabetes:a Consensus Report from the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETOLOGIA 63: 1671-1693

RHAPSODY: Chung, Wendy K. et al. Precision Medicine in Diabetes: A Consensus Report From the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETES CARE 43: 1617-1635

RHAPSODY: Campbell, Scott A. et al. Human islets contain a subpopulation of glucagon-like peptide-1 secreting alpha cells that is increased in type 2 diabetes, MOL METAB 39:

RHAPSODY: Muniangi-Muhitu, Hermine et al. Covid-19 and Diabetes: A Complex Bidirectional Relationship, FRONT ENDOCRINOL 11:

RHAPSODY: Fang, Zijian et al. The Influence of Peptide Context on Signaling and Trafficking of Glucagon-like Peptide-1 Receptor Biased Agonists, ACS PHARMACOL TRANSL 3: 345-360

RHAPSODY: Mueller, Andreas et al. 3D FIB-SEM reconstruction of microtubule-organelle interaction in whole primary mouse beta cells, J CELL BIOL 220:

RHAPSODY: Jones, Ben et al. Genetic and biased agonist-mediated reductions in beta-arrestin recruitment prolong cAMP signaling at glucagon family receptors, J BIOL CHEM 296:

RHAPSODY: Nasteska, Daniela et al. PDX1(LOW) MAFA(LOW) beta-cells contribute to islet function and insulin release, NAT COMMUN 12:

ROADMAP: Ponjoan, Anna et al. Epidemiology of dementia: prevalence and incidence estimates using validated electronic health records from primary care, CLIN EPIDEMIOL 11: 217-228

RTCure: Rauber, Simon et al. Resolution of inflammation by interleukin-9-producing type 2 innate lymphoid cells, NAT MED 23: 938-+

RTCure: Engdahl, Cecilia et al. Estrogen induces St6gal1 expression and increases IgG sialylation in mice and patients with rheumatoid arthritis: a potential explanation for the increased risk of rheumatoid arthritis in postmenopausal women, ARTHRITIS RES THER 20:

RTCure: Lloyd, Katy A. et al. Differential ACPA Binding to Nuclear Antigens Reveals a PAD-Independent Pathway and a Distinct Subset of Acetylation Cross-Reactive Autoantibodies in Rheumatoid Arthritis, FRONT IMMUNOL 9:

RTCure: Perucha, Esperanza et al. The cholesterol biosynthesis pathway regulates IL-10 expression in human Th1 cells, NAT COMMUN 10:

RTCure: Wohlfahrt, Thomas et al. PU.1 controls fibroblast polarization and tissue fibrosis, NATURE 566: 344-+

RTCure: Chemin, Karine et al. Effector Functions of CD4+T Cells at the Site of Local Autoimmune Inflammation-Lessons From Rheumatoid Arthritis, FRONT IMMUNOL 10:

RTCure: Wehr, P. et al. Dendritic cells, T cells and their interaction in rheumatoid arthritis, CLIN EXP IMMUNOL 196: 12-27

RTCure: Eriksson, Kaja et al. Periodontal Health and Oral Microbiota in Patients with Rheumatoid Arthritis, J CLIN MED 8:

RTCure: Mosanya, Chijioke H. et al. Tolerising cellular therapies: what is their promise for autoimmune disease?, ANN RHEUM DIS 78: 297-310

RTCure: Kampstra, Arieke Suzanna Berendina et al. Different classes of anti-modified protein antibodies are induced on exposure to antigens expressing only one type of modification, ANN RHEUM DIS 78: 908-916

RTCure: Kampylafka, Eleni et al. Disease interception with interleukin-17 inhibition in high-risk psoriasis patients with subclinical joint inflammation-data from the prospective IVEPSA study, *ARTHRITIS RES THER* 21:

RTCure: Daniel, Christoph et al. Extracellular DNA traps in inflammation, injury and healing, *NAT REV NEPHROL* 15: 559-575

RTCure: Schulz, Axel R. et al. Stabilizing Antibody Cocktails for Mass Cytometry, *CYTOM PART A* 95A: 910-916

RTCure: Culemann, Stephan et al. Locally renewing resident synovial macrophages provide a protective barrier for the joint, *NATURE* 572: 670-+

RTCure: Hafkenscheid, Lise et al. N-Linked Glycans in the Variable Domain of IgG Anti-Citrullinated Protein Antibodies Predict the Development of Rheumatoid Arthritis, *ARTHRITIS RHEUMATOL* 71: 1626-1633

RTCure: Cossarizza, Andrea et al. Guidelines for the use of flow cytometry and cell sorting in immunological studies (second edition), *EUR J IMMUNOL* 49: 1457-1973

RTCure: Sun, Meng et al. Anticitrullinated protein antibodies facilitate migration of synovial tissue-derived fibroblasts, *ANN RHEUM DIS* 78: 1621-1631

RTCure: Grueeneboom, Anika et al. A network of trans-cortical capillaries as mainstay for blood circulation in long bones, *NAT METAB* 1: 236-250

RTCure: Nemeth, Tamas et al. Neutrophils as emerging therapeutic targets, *NAT REV DRUG DISCOV* 19: 253-275

RTCure: Steffen, Ulrike et al. IgA subclasses have different effector functions associated with distinct glycosylation profiles, *NAT COMMUN* 11:

RTCure: Adam, Susanne et al. JAK inhibition increases bone mass in steady-state conditions and ameliorates pathological bone loss by stimulating osteoblast function, *SCI TRANSL MED* 12:

RTCure: Sokolova, Maria, V et al. A set of serum markers detecting systemic inflammation in psoriatic skin, enthesal, and joint disease in the absence of C-reactive protein and its link to clinical disease manifestations, *ARTHRITIS RES THER* 22:

RTCure: Kissel, T. et al. Antibodies and B cells recognising citrullinated proteins display a broad cross-reactivity towards other post-translational modifications, *ANN RHEUM DIS* 79: 472-480

RTCure: Klareskog, L. et al. The importance of differences, On environment and its interactions with genes and immunity in the causation of rheumatoid arthritis, *J INTERN MED* 287: 514-533

RTCure: Schett, Georg et al. COVID-19 revisiting inflammatory pathways of arthritis, *NAT REV RHEUMATOL* 16: 465-470

RTCure: Reed, Evan et al. Presence of autoantibodies in seronegative rheumatoid arthritis associates with classical risk factors and high disease activity, *ARTHRITIS RES THER* 22:



RTCure: Simon, David et al. Patients with immune-mediated inflammatory diseases receiving cytokine inhibitors have low prevalence of SARS-CoV-2 seroconversion, NAT COMMUN 11:

RTCure: Tajik, Narges et al. Targeting zonulin and intestinal epithelial barrier function to prevent onset of arthritis, NAT COMMUN 11:

RTCure: Sahlstroem, Peter et al. Different Hierarchies of Anti-Modified Protein Autoantibody Reactivities in Rheumatoid Arthritis, ARTHRITIS RHEUMATOL 72: 1643-1657

RTCure: Kristyanto, Hendy et al. Persistently activated, proliferative memory autoreactive B cells promote inflammation in rheumatoid arthritis, SCI TRANSL MED 12:

RTCure: Schett, Georg et al. Why remission is not enough: underlying disease mechanisms in RA that prevent cure, NAT REV RHEUMATOL 17: 135-144

RTCure: Maschmeyer, Patrick et al. Antigen-driven PD-1(+)/TOX(+)/BHLHE40(+) and PD-1(+)/TOX(+)/EOMES(+) T lymphocytes regulate juvenile idiopathic arthritis in situ, EUR J IMMUNOL 51: 915-929

RTCure: Knitza, Johannes et al. Mobile Health Usage, Preferences, Barriers, and eHealth Literacy in Rheumatology: Patient Survey Study, JMIR MHEALTH UHEALTH 8:

RTCure: Vodencarevic, Asmir et al. Advanced machine learning for predicting individual risk of flares in rheumatoid arthritis patients tapering biologic drugs, ARTHRITIS RES THER 23:

RTCure: Kharlamova, Nastya et al. False Positive Results in SARS-CoV-2 Serological Tests for Samples From Patients With Chronic Inflammatory Diseases, FRONT IMMUNOL 12:

RTCure: Friscic, Jasna et al. The complement system drives local inflammatory tissue priming by metabolic reprogramming of synovial fibroblasts, IMMUNITY 54: 1002+

RTCure: Simon, David et al. SARS-CoV-2 vaccination responses in untreated, conventionally treated and anticytokine-treated patients with immune-mediated inflammatory diseases, ANN RHEUM DIS 80: 1312-1316

RTCure: Haberman, Rebecca H. et al. Methotrexate hampers immunogenicity to BNT162b2 mRNA COVID-19 vaccine in immune-mediated inflammatory disease, ANN RHEUM DIS 80: 1339-1344

RTCure: Simon, David et al. Humoral and Cellular Immune Responses to SARS-CoV-2 Infection and Vaccination in Autoimmune Disease Patients With B Cell Depletion, ARTHRITIS RHEUMATOL 74: 33-37

SafeSciMET: Heslop, James A. et al. Concise Review: Workshop Review: Understanding and Assessing the Risks of Stem Cell-Based Therapies, STEM CELL TRANSL MED 4: 389-400

SAFE-T: Robles-Diaz, Mercedes et al. Use of Hys Law and a New Composite Algorithm to Predict Acute Liver Failure in Patients With Drug-Induced Liver Injury, GASTROENTEROLOGY 147: 109-U204

SAFE-T: Church, Rachel J. et al. Candidate biomarkers for the diagnosis and prognosis of drug-induced liver injury: An international collaborative effort, HEPATOLOGY 69: 760-773

SAFE-T: Regnier, Paul et al. Targeting JAK/STAT pathway in Takayasu arteritis, ANN RHEUM DIS 79: 951-959

SOPHIA: Chung, Wendy K. et al. Precision medicine in diabetes: a Consensus Report from the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETOLOGIA 63: 1671-1693

SOPHIA: Chung, Wendy K. et al. Precision Medicine in Diabetes: A Consensus Report From the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD), DIABETES CARE 43: 1617-1635

SOPHIA: Stefan, Norbert et al. Global pandemics interconnected - obesity, impaired metabolic health and COVID-19, NAT REV ENDOCRINOL 17: 135-149

SPRINTT: Landi, Francesco et al. Sarcopenia as the Biological Substrate of Physical Frailty, CLIN LIVER DIS 19: 367-+

SPRINTT: Calvani, Riccardo et al. Biomarkers for physical frailty and sarcopenia: state of the science and future developments, J CACHEXIA SARCOPEMI 6: 278-286

SPRINTT: von Haehling, Stephan et al. The wasting continuum in heart failure: from sarcopenia to cachexia, P NUTR SOC 74: 367-377

SPRINTT: Landi, Francesco et al. Anorexia of Aging: Risk Factors, Consequences, and Potential Treatments, NUTRIENTS 8:

SPRINTT: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

SPRINTT: Landi, Francesco et al. Protein Intake and Muscle Health in Old Age: From Biological Plausibility to Clinical Evidence, NUTRIENTS 8:

SPRINTT: Landi, Francesco et al. Impact of physical function impairment and multimorbidity on mortality among community-living older persons with sarcopenia: results from the iSIRENTE prospective cohort study, BMJ OPEN 6:

SPRINTT: Collamati, Agnese et al. Sarcopenia in heart failure: mechanisms and therapeutic strategies, J GERIATR CARDIOL 13: 615-624

SPRINTT: Marzetti, Emanuele et al. Altered mitochondrial quality control signaling in muscle of old gastric cancer patients with cachexia, EXP GERONTOL 87: 92-99

SPRINTT: Sirven, Nicolas et al. The cost of frailty in France, EUR J HEALTH ECON 18: 243-253

SPRINTT: Marzetti, Emanuele et al. Sarcopenia: an overview, AGING CLIN EXP RES 29: 11-17

SPRINTT: Tosato, Matteo et al. Measurement of muscle mass in sarcopenia: from imaging to biochemical markers, AGING CLIN EXP RES 29: 19-27

SPRINTT: Calvani, Riccardo et al. Biomarkers for physical frailty and sarcopenia, AGING CLIN EXP RES 29: 29-34

SPRINTT: Marzetti, Emanuele et al. Physical activity and exercise as countermeasures to physical frailty and sarcopenia, *AGING CLIN EXP RES* 29: 35-42

SPRINTT: Cruz-Jentoft, Alfonso J. et al. Nutrition, frailty, and sarcopenia, *AGING CLIN EXP RES* 29: 43-48

SPRINTT: Cesari, Matteo et al. Rationale for a preliminary operational definition of physical frailty and sarcopenia in the SPRINTT trial, *AGING CLIN EXP RES* 29: 81-88

SPRINTT: Landi, Francesco et al. The Sarcopenia and Physical Frailty IN older people: multi-component Treatment strategies (SPRINTT) randomized controlled trial: design and methods, *AGING CLIN EXP RES* 29: 89-100

SPRINTT: Landi, Francesco et al. Age-Related Variations of Muscle Mass, Strength, and Physical Performance in Community-Dwellers: Results From the Milan EXPO Survey, *J AM MED DIR ASSOC* 18:

SPRINTT: Martone, Anna Maria et al. Exercise and Protein Intake: A Synergistic Approach against Sarcopenia, *BIOMED RES INT* 2017:

SPRINTT: Landi, F. et al. The association between sarcopenia and functional outcomes among older patients with hip fracture undergoing in-hospital rehabilitation, *OSTEOPOROSIS INT* 28: 1569-1576

SPRINTT: von Haehling, Stephan et al. Muscle wasting and cachexia in heart failure: mechanisms and therapies, *NAT REV CARDIOL* 14: 323-341

SPRINTT: Picca, Anna et al. Fueling Inflamm-Aging through Mitochondrial Dysfunction: Mechanisms and Molecular Targets, *INT J MOL SCI* 18:

SPRINTT: Picca, Anna et al. Circulating Mitochondrial DNA at the Crossroads of Mitochondrial Dysfunction and Inflammation During Aging and Muscle Wasting Disorders, *REJUV RES* 21: 350-359

SPRINTT: Picca, Anna et al. Gut Dysbiosis and Muscle Aging: Searching for Novel Targets against Sarcopenia, *MEDIAT INFLAMM* 2018:

SPRINTT: Calvani, Riccardo et al. Of Microbes and Minds: A Narrative Review on the Second Brain Aging, *FRONT MED-LAUSANNE* 5:

SPRINTT: Landi, Francesco et al. Sarcopenia: An Overview on Current Definitions, Diagnosis and Treatment, *CURR PROTEIN PEPT SC* 19: 633-638

SPRINTT: Picca, Anna et al. Mitochondrial quality control mechanisms as molecular targets in cardiac ageing, *NAT REV CARDIOL* 15: 543-554

SPRINTT: Marzetti, Emanuele et al. The Sarcopenia and Physical Frailty IN older people: multi-component Treatment strategies (SPRINTT) randomized controlled trial: Case finding, screening and characteristics of eligible participants, *EXP GERONTOL* 113: 48-57

SPRINTT: Calvani, Riccardo et al. A Distinct Pattern of Circulating Amino Acids Characterizes Older Persons with Physical Frailty and Sarcopenia: Results from the BIOSPHERE Study, *NUTRIENTS* 10:

SPRINTT: Picca, Anna et al. Mitochondrial Dysfunction and Aging: Insights from the Analysis of Extracellular Vesicles, INT J MOL SCI 20:

SPRINTT: Marzetti, Emanuele et al. Inflammatory signatures in older persons with physical frailty and sarcopenia: The frailty cytokinome at its core, EXP GERONTOL 122: 129-138

SPRINTT: Picca, Anna et al. Mitochondrial-Derived Vesicles as Candidate Biomarkers in Parkinsons Disease: Rationale, Design and Methods of the EXosomes in PARKinson Disease (EXPAND) Study, INT J MOL SCI 20:

SPRINTT: Picca, Anna et al. Gut Microbial, Inflammatory and Metabolic Signatures in Older People with Physical Frailty and Sarcopenia: Results from the BIOSPHERE Study, NUTRIENTS 12:

SPRINTT: Curcio, Francesco et al. Sarcopenia and Heart Failure, NUTRIENTS 12:

SPRINTT: Picca, Anna et al. Mitochondrial Signatures in Circulating Extracellular Vesicles of Older Adults with Parkinsons Disease: Results from the EXosomes in PARKinsons Disease (EXPAND) Study, J CLIN MED 9:

SPRINTT: Coelho-Junior, Helio J. et al. Protein-Related Dietary Parameters and Frailty Status in Older Community-Dwellers across Different Frailty Instruments, NUTRIENTS 12:

SPRINTT: Picca, Anna et al. Generation and Release of Mitochondrial-Derived Vesicles in Health, Aging and Disease, J CLIN MED 9:

SPRINTT: Angulo, Javier et al. Physical activity and exercise: Strategies to manage frailty, REDOX BIOL 35:

SPRINTT: von Haehling, Stephan et al. Muscle wasting as an independent predictor of survival in patients with chronic heart failure, J CACHEXIA SARCOPENI 11: 1242-1249

SPRINTT: Picca, Anna et al. Mitochondrial Dysfunction, Oxidative Stress, and Neuroinflammation: Intertwined Roads to Neurodegeneration, ANTIOXIDANTS-BASEL 9:

SPRINTT: Billot, Maxime et al. Preserving Mobility in Older Adults with Physical Frailty and Sarcopenia: Opportunities, Challenges, and Recommendations for Physical Activity Interventions, CLIN INTERV AGING 15: 1675-1690

SPRINTT: Coelho-Junior, Helio J. et al. Protein Intake and Frailty: A Matter of Quantity, Quality, and Timing, NUTRIENTS 12:

SPRINTT: Jyvakorpi, S. K. et al. The sarcopenia and physical frailty in older people: multi-component treatment strategies (SPRINTT) project: description and feasibility of a nutrition intervention in community-dwelling older Europeans, EUR GERIATR MED 12: 303-312

SPRINTT: Coelho, Helio J., Jr. et al. Age- and Gender-Related Changes in Physical Function in Community-Dwelling Brazilian Adults Aged 50 to 102 Years, J GERIATR PHYS THER 44: E123-E131

STEMBANCC: Badger, J. L. et al. Parkinsons disease in a dish - Using stem cells as a molecular tool, NEUROPHARMACOLOGY 76: 88-96

STEMBANCC: Kempf, Henning et al. Controlling Expansion and Cardiomyogenic Differentiation of Human Pluripotent Stem Cells in Scalable Suspension Culture, STEM CELL REP 3: 1132-1146

STEMBANCC: Kaye, Jane et al. Dynamic consent: a patient interface for twenty-first century research networks, EUR J HUM GENET 23: 141-146

STEMBANCC: Patsch, Christoph et al. Generation of vascular endothelial and smooth muscle cells from human pluripotent stem cells, NAT CELL BIOL 17: 994-U294

STEMBANCC: Kempf, Henning et al. Cardiac differentiation of human pluripotent stem cells in scalable suspension culture, NAT PROTOC 10: 1345-1361

STEMBANCC: Heywood, Wendy E. et al. Identification of novel CSF biomarkers for neurodegeneration and their validation by a high-throughput multiplexed targeted proteomic assay, MOL NEURODEGENER 10:

STEMBANCC: Kempf, Henning et al. Large-scale production of human pluripotent stem cell derived cardiomyocytes, ADV DRUG DELIVER REV 96: 18-30

STEMBANCC: Viereck, Janika et al. Long noncoding RNA Chast promotes cardiac remodeling, SCI TRANSL MED 8:

STEMBANCC: Handel, Adam E. et al. Assessing similarity to primary tissue and cortical layer identity in induced pluripotent stem cell-derived cortical neurons through single-cell transcriptomics, HUM MOL GENET 25: 989-1000

STEMBANCC: Fernandes, Hugo J. R. et al. ER Stress and Autophagic Perturbations Lead to Elevated Extracellular alpha-Synuclein in GBA-N370S LEParkinsons iPSC-Derived Dopamine Neurons, STEM CELL REP 6: 342-356

STEMBANCC: Cao, Lishuang et al. Pharmacological reversal of a pain phenotype in iPSC-derived sensory neurons and patients with inherited erythromelalgia, SCI TRANSL MED 8:

STEMBANCC: Kropp, Christina et al. Impact of Feeding Strategies on the Scalable Expansion of Human Pluripotent Stem Cells in Single-Use Stirred Tank Bioreactors, STEM CELL TRANSL MED 5: 1289-1301

STEMBANCC: Kuijlaars, Jacobine et al. Sustained synchronized neuronal network activity in a human astrocyte co-culture system, SCI REP-UK 6:

STEMBANCC: Sandor, Cynthia et al. Transcriptomic profiling of purified patient-derived dopamine neurons identifies convergent perturbations and therapeutics for Parkinsons disease, HUM MOL GENET 26: 552-566

STEMBANCC: Clark, Alex J. et al. Co-cultures with stem cell-derived human sensory neurons reveal regulators of peripheral myelination, BRAIN 140: 898-913

STEMBANCC: Hocher, Berthold et al. Metabolomics for clinical use and research in chronic kidney disease, NAT REV NEPHROL 13: 269-284

STEMBANCC: Haenseler, Walther et al. A Highly Efficient Human Pluripotent Stem Cell Microglia Model Displays a Neuronal-Co-culture-Specific Expression Profile and Inflammatory Response, STEM CELL REP 8: 1727-1742

STEMBANCC: Paillusson, Sebastien et al. alpha-Synuclein binds to the ER-mitochondria tethering protein VAPB to disrupt Ca<sup>2+</sup> homeostasis and mitochondrial ATP production, ACTA NEUROPATHOL 134: 129-149

STEMBANCC: Kropp, Christina et al. Progress and challenges in large-scale expansion of human pluripotent stem cells, PROCESS BIOCHEM 59: 244-254

STEMBANCC: Heman-Ackah, Sabrina M. et al. Alpha-synuclein induces the unfolded protein response in Parkinsons disease SNCA triplication iPSC-derived neurons, HUM MOL GENET 26: 4441-4450

STEMBANCC: Kathuria, A. et al. Stem cell-derived neurons from autistic individuals with SHANK3 mutation show morphogenetic abnormalities during early development, MOL PSYCHIATR 23: 735-746

STEMBANCC: Brownjohn, Philip W. et al. Functional Studies of Missense TREM2 Mutations in Human Stem Cell-Derived Microglia, STEM CELL REP 10: 1294-1307

STEMBANCC: Koch, Lothar et al. Laser bioprinting of human induced pluripotent stem cells-the effect of printing and biomaterials on cell survival, pluripotency, and differentiation, BIOFABRICATION 10:

STEMBANCC: Olmer, Ruth et al. Differentiation of Human Pluripotent Stem Cells into Functional Endothelial Cells in Scalable Suspension Culture, STEM CELL REP 10: 1657-1672

STEMBANCC: Ludtmann, Marthe H. R. et al. alpha-synuclein oligomers interact with ATP synthase and open the permeability transition pore in Parkinsons disease, NAT COMMUN 9:

STEMBANCC: Hartlova, Anetta et al. LRRK2 is a negative regulator of Mycobacterium tuberculosis phagosome maturation in macrophages, EMBO J 37:

STEMBANCC: Ramond, Cyrille et al. Understanding human fetal pancreas development using subpopulation sorting, RNA sequencing and single-cell profiling, DEVELOPMENT 145:

STEMBANCC: Volpato, Viola et al. Reproducibility of Molecular Phenotypes after Long-Term Differentiation to Human iPSC-Derived Neurons: A Multi-Site Omics Study, STEM CELL REP 11: 897-911

STEMBANCC: Lee, Heyne et al. LRRK2 in peripheral and central nervous system innate immunity: its link to Parkinsons disease, BIOCHEM SOC T 45: 131-139

STEMBANCC: Lang, Charmaine et al. Single-Cell Sequencing of iPSC-Dopamine Neurons Reconstructs Disease Progression and Identifies HDAC4 as a Regulator of Parkinson Cell Phenotypes, CELL STEM CELL 24: 93+

STEMBANCC: Delsing, Louise et al. Barrier Properties and Transcriptome Expression in Human iPSC-Derived Models of the Blood-Brain Barrier, STEM CELLS 36: 1816-1827

STEMBANCC: Bennett, David L. et al. THE ROLE OF VOLTAGE-GATED SODIUM CHANNELS IN PAIN SIGNALING, *PHYSIOL REV* 99: 1079-1151

STEMBANCC: McDermott, Lucy A. et al. Defining the Functional Role of Na(v)1.7 in Human Nociception, *NEURON* 101: 905-+

STEMBANCC: Baskozos, Georgios et al. Comprehensive analysis of long noncoding RNA expression in dorsal root ganglion reveals cell-type specificity and dysregulation after nerve injury, *PAIN* 160: 463-485

STEMBANCC: Little, Daniel et al. Using stem cell-derived neurons in drug screening for neurological diseases, *NEUROBIOL AGING* 78: 130-141

STEMBANCC: Connor-Robson, Natalie et al. An integrated transcriptomics and proteomics analysis reveals functional endocytic dysregulation caused by mutations in LRRK2, *NEUROBIOL DIS* 127: 512-526

STEMBANCC: Booth, Heather D. E. et al. RNA sequencing reveals MMP2 and TGFB1 downregulation in LRRK2 G2019S Parkinsons iPSC-derived astrocytes, *NEUROBIOL DIS* 129: 56-66

STEMBANCC: Zambon, Federico et al. Cellular alpha-synuclein pathology is associated with bioenergetic dysfunction in Parkinsons iPSC-derived dopamine neurons, *HUM MOL GENET* 28: 2001-2013

STEMBANCC: Halloin, Caroline et al. Continuous WNT Control Enables Advanced hPSC Cardiac Processing and Prognostic Surface Marker Identification in Chemically Defined Suspension Culture, *STEM CELL REP* 13: 366-379

STEMBANCC: Pettingill, Philippa et al. A causal role for TRESK loss of function in migraine mechanisms, *BRAIN* 142: 3852-3867

STEMBANCC: Lee, Heyne et al. LRRK2 Is Recruited to Phagosomes and Co-recruits RAB8 and RAB10 in Human Pluripotent Stem Cell-Derived Macrophages, *STEM CELL REP* 14: 940-955

STEMBANCC: Adhya, Dwaipayan et al. Atypical Neurogenesis in Induced Pluripotent Stem Cells From Autistic Individuals, *BIOL PSYCHIAT* 89: 486-496

STEMBANCC: Chandrasekaran, Vidya et al. Generation and characterization of iPSC-derived renal proximal tubule-like cells with extended stability, *SCI REP-UK* 11:

STOPFOP: Williams, Eleanor et al. Saracatinib is an efficacious clinical candidate for fibrodysplasia ossificans progressiva, *JCI INSIGHT* 6:

SUMMIT: Boekholdt, S. Matthijs et al. Association of LDL Cholesterol, Non-HDL Cholesterol, and Apolipoprotein B Levels With Risk of Cardiovascular Events Among Patients Treated With Statins A Meta-analysis, *JAMA-J AM MED ASSOC* 307: 1302-1309

SUMMIT: Rocca, B. et al. The recovery of platelet cyclooxygenase activity explains interindividual variability in responsiveness to low-dose aspirin in patients with and without diabetes, *J THROMB HAEMOST* 10: 1220-1230

SUMMIT: Boni, Enrico et al. A Reconfigurable and Programmable FPGA-Based System for Nonstandard Ultrasound Methods, IEEE T ULTRASON FERR 59: 1378-1385

SUMMIT: Sandholm, Niina et al. New Susceptibility Loci Associated with Kidney Disease in Type 1 Diabetes, PLOS GENET 8:

SUMMIT: Fall, Tove et al. The Role of Adiposity in Cardiometabolic Traits: A Mendelian Randomization Analysis, PLOS MED 10:

SUMMIT: Zhou, Kaixin et al. Heritability of variation in glycaemic response to metformin: a genome-wide complex trait analysis, LANCET DIABETES ENDO 2: 481-487

SUMMIT: Postmus, Iris et al. Pharmacogenetic meta-analysis of genome-wide association studies of LDL cholesterol response to statins, NAT COMMUN 5:

SUMMIT: Meng, W. et al. A genome-wide association study suggests an association of Chr8p21.3 (GFRA2) with diabetic neuropathic pain, EUR J PAIN 19: 392-399

SUMMIT: Zaccardi, Francesco et al. Platelet mean volume, distribution width, and count in type 2 diabetes, impaired fasting glucose, and metabolic syndrome: a meta-analysis, DIABETES-METAB RES 31: 402-410

SUMMIT: Goncalves, Isabel et al. Elevated Plasma Levels of MMP-12 Are Associated With Atherosclerotic Burden and Symptomatic Cardiovascular Disease in Subjects With Type 2 Diabetes, ARTERIOSCL THROM VAS 35: 1723-1731

SUMMIT: Patrono, Carlo et al. The Multifaceted Clinical Readouts of Platelet Inhibition by Low-Dose Aspirin, J AM COLL CARDIOL 66: 74-85

SUMMIT: Looker, Helen C. et al. Biomarkers of rapid chronic kidney disease progression in type 2 diabetes, KIDNEY INT 88: 888-896

SUMMIT: De Marinis, Yang et al. Epigenetic regulation of the thioredoxin-interacting protein (TXNIP) gene by hyperglycemia in kidney, KIDNEY INT 89: 342-353

SUMMIT: Edsfeldt, Andreas et al. Sphingolipids Contribute to Human Atherosclerotic Plaque Inflammation, ARTERIOSCL THROM VAS 36: 1132-+

SUMMIT: Sandholm, Niina et al. The Genetic Landscape of Renal Complications in Type 1 Diabetes, J AM SOC NEPHROL 28: 557-574

SUMMIT: Fadista, Joao et al. LoFtool: a gene intolerance score based on loss-of-function variants in 60 706 individuals, BIOINFORMATICS 33: 471-474

SUMMIT: Justice, Anne E. et al. Genome-wide meta-analysis of 241,258 adults accounting for smoking behaviour identifies novel loci for obesity traits, NAT COMMUN 8:

SUMMIT: Wain, Louise V. et al. Novel Blood Pressure Locus and Gene Discovery Using Genome-Wide Association Study and Expression Data Sets From Blood and the Kidney, HYPERTENSION 70: E4-+



SUMMIT: van Zuydam, Natalie R. et al. A Genome-Wide Association Study of Diabetic Kidney Disease in Subjects With Type 2 Diabetes, *DIABETES* 67: 1414-1427

SUMMIT: Colombo, Marco et al. Serum kidney injury molecule 1 and (2)-microglobulin perform as well as larger biomarker panels for prediction of rapid decline in renal function in type 2 diabetes, *DIABETOLOGIA* 62: 156-168

SUMMIT: Salem, Rany M. et al. Genome-Wide Association Study of Diabetic Kidney Disease Highlights Biology Involved in Glomerular Basement Membrane Collagen, *J AM SOC NEPHROL* 30: 2000-2016

SUMMIT: Hebert, Harry L. et al. Cohort profile: DOLORisk Dundee: a longitudinal study of chronic neuropathic pain, *BMJ OPEN* 11:

T2EVOLVE: Prommersberger, Sabrina et al. CARAMBA: a first-in-human clinical trial with SLAMF7 CAR-T cells prepared by virus-free Sleeping Beauty gene transfer to treat multiple myeloma, *GENE THER* 28: 560-571

TransBioLine: Attard, Joseph A. et al. Ex situ Normothermic Split Liver Machine Perfusion: Protocol for Robust Comparative Controls in Liver Function Assessment Suitable for Evaluation of Novel Therapeutic Interventions in the Pre-clinical Setting, *FRONT SURG* 8:

TransBioLine: Bozward, Amber G. et al. Natural Killer Cells and Regulatory T Cells Cross Talk in Hepatocellular Carcinoma: Exploring Therapeutic Options for the Next Decade, *FRONT IMMUNOL* 12:

TRANSLOCATION: Ruggerone, Paolo et al. RND Efflux Pumps: Structural Information Translated into Function and Inhibition Mechanisms, *CURR TOP MED CHEM* 13: 3079-3100

TRANSLOCATION: Mislin, Gaetan L. A. et al. Siderophore-dependent iron uptake systems as gates for antibiotic Trojan horse strategies against *Pseudomonas aeruginosa*, *METALLOMICS* 6: 408-420

TRANSLOCATION: Eicher, Thomas et al. Coupling of remote alternating-access transport mechanisms for protons and substrates in the multidrug efflux pump AcrB, *ELIFE* 3:

TRANSLOCATION: Gutschmann, Thomas et al. Protein reconstitution into freestanding planar lipid membranes for electrophysiological characterization, *NAT PROTOC* 10: 188-198

TRANSLOCATION: Davin-Regli, Anne et al. Enterobacter aerogenes and Enterobacter cloacae, versatile bacterial pathogens confronting antibiotic treatment, *FRONT MICROBIOL* 6:

TRANSLOCATION: Anes, Joao et al. The ins and outs of RND efflux pumps in *Escherichia coli*, *FRONT MICROBIOL* 6:

TRANSLOCATION: Gasser, Veronique et al. Cellular organization of siderophore biosynthesis in *Pseudomonas aeruginosa*: Evidence for siderosomes, *J INORG BIOCHEM* 148: 27-34

TRANSLOCATION: Dreier, Juerg et al. Interaction of antibacterial compounds with RND efflux pumps in *Pseudomonas aeruginosa*, *FRONT MICROBIOL* 6:

TRANSLOCATION: Du, Dijun et al. Structure, mechanism and cooperation of bacterial multidrug transporters, *CURR OPIN STRUC BIOL* 33: 76-91

TRANSLOCATION: Daury, Laetitia et al. Tripartite assembly of RND multidrug efflux pumps, NAT COMMUN 7:

TRANSLOCATION: Sjuts, Hanno et al. Molecular basis for inhibition of AcrB multidrug efflux pump by novel and powerful pyranopyridine derivatives, P NATL ACAD SCI USA 113: 3509-3514

TRANSLOCATION: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

TRANSLOCATION: Arunmanee, Wanatchaporn et al. Gram-negative trimeric porins have specific LPS binding sites that are essential for porin biogenesis, P NATL ACAD SCI USA 113: E5034-E5043

TRANSLOCATION: Glenwright, Amy J. et al. Structural basis for nutrient acquisition by dominant members of the human gut microbiota, NATURE 541: 407-+

TRANSLOCATION: Ghai, Ishan et al. General Method to Determine the Flux of Charged Molecules through Nanopores Applied to beta-Lactamase Inhibitors and OmpF, J PHYS CHEM LETT 8: 1295-1301

TRANSLOCATION: Masi, Muriel et al. Mechanisms of envelope permeability and antibiotic influx and efflux in Gram-negative bacteria, NAT MICROBIOL 2:

TRANSLOCATION: Moynie, Lucile et al. Structure and Function of the PiuA and PirA Siderophore-Drug Receptors from *Pseudomonas aeruginosa* and *Acinetobacter baumannii*, ANTIMICROB AGENTS CH 61:

TRANSLOCATION: Abellon-Ruiz, Javier et al. Structural basis for maintenance of bacterial outer membrane lipid asymmetry, NAT MICROBIOL 2: 1616-1623

TRANSLOCATION: Vergalli, Julia et al. Spectrofluorimetric quantification of antibiotic drug concentration in bacterial cells for the characterization of translocation across bacterial membranes, NAT PROTOC 13: 1348-1361

TRANSLOCATION: Du, Dijun et al. Multidrug efflux pumps: structure, function and regulation, NAT REV MICROBIOL 16: 523-539

TRANSLOCATION: Acosta-Gutierrez, Silvia et al. Getting Drugs into Gram-Negative Bacteria: Rational Rules for Permeation through General Porins, ACS INFECT DIS 4: 1487-1498

TRANSLOCATION: Prochnow, Hans et al. Subcellular Quantification of Uptake in Gram-Negative Bacteria, ANAL CHEM 91: 1863-1872

TRANSLOCATION: Moynie, Lucile et al. The complex of ferric-enterobactin with its transporter from *Pseudomonas aeruginosa* suggests a two-site model, NAT COMMUN 10:

TRANSLOCATION: Davin-Regli, Anne et al. Enterobacter spp.: Update on Taxonomy, Clinical Aspect, and Emerging Antimicrobial Resistance, CLIN MICROBIOL REV 32:

TRANSLOCATION: Cama, Jehangir et al. Breaching the Barrier: Quantifying Antibiotic Permeability across Gram-negative Bacterial Membranes, J MOL BIOL 431: 3531-3546

TRANSLOCATION: Tam, Heng-Keat et al. Binding and Transport of Carboxylated Drugs by the Multidrug Transporter AcrB, J MOL BIOL 432: 861-877

TRANSLOCATION: Perraud, Quentin et al. Phenotypic Adaption of Pseudomonas aeruginosa by Hacking Siderophores Produced by Other Microorganisms, MOL CELL PROTEOMICS 19: 589-607

TRANSLOCATION: Bafna, Jayesh Arun et al. Kanamycin Uptake into Escherichia coli Is Facilitated by OmpF and OmpC Porin Channels Located in the Outer Membrane, ACS INFECT DIS 6: 1855-1865

TRANSLOCATION: Prajapati, Jigneshkumar Dahyabhai et al. How to Enter a Bacterium: Bacterial Porins and the Permeation of Antibiotics, CHEM REV 121: 5158-5192

TransQST: Passini, Elisa et al. Human In Silico Drug Trials Demonstrate Higher Accuracy than Animal Models in Predicting Clinical Pro-Arrhythmic Cardiotoxicity, FRONT PHYSIOL 8:

TransQST: del-Toro, N. et al. Capturing variation impact on molecular interactions in the IMEx Consortium mutations data set, NAT COMMUN 10:

TransQST: Albrecht, Wiebke et al. Prediction of human drug-induced liver injury (DILI) in relation to oral doses and blood concentrations, ARCH TOXICOL 93: 1609-1637

TransQST: Liu, Anika et al. From expression footprints to causal pathways: contextualizing large signaling networks with CARNIVAL, NPJ SYST BIOL APPL 5:

TransQST: Buhl, Eva M. et al. Dysregulated mesenchymal PDGFR-beta drives kidney fibrosis, EMBO MOL MED 12:

TransQST: Tomek, Jakub et al. Development, calibration, and validation of a novel human ventricular myocyte model in health, disease, and drug block, ELIFE 8:

TransQST: Liu, Jun et al. Integration of epidemiologic, pharmacologic, genetic and gut microbiome data in a drug-metabolite atlas, NAT MED 26: 110-+

TransQST: Malik-Sheriff, Rahuman S. et al. BioModels-15 years of sharing computational models in life science, NUCLEIC ACIDS RES 48: D407-D415

TransQST: Pinero, Janet et al. The DisGeNET knowledge platform for disease genomics: 2019 update, NUCLEIC ACIDS RES 48: D845-D855

TransQST: Cirillo, Davide et al. Sex and gender differences and biases in artificial intelligence for biomedicine and healthcare, NPJ DIGIT MED 3:

TransQST: Gjerga, Enio et al. Converting networks to predictive logic models from perturbation signalling data with CellNOpt, BIOINFORMATICS 36: 4523-4524

TransQST: Margara, Francesca et al. In-silico human electro-mechanical ventricular modelling and simulation for drug-induced pro-arrhythmia and inotropic risk assessment, PROG BIOPHYS MOL BIO 159: 58-74

TransQST: Tuerei, Denes et al. Integrated intra- and intercellular signaling knowledge for multicellular omics analysis, MOL SYST BIOL 17:

TRISTAN: Heskamp, Sandra et al. Zr-89-Immuno-Positron Emission Tomography in Oncology: State-of-the-Art Zr-89 Radiochemistry, *BIOCONJUGATE CHEM* 28: 2211-2223

TRISTAN: Skeoch, Sarah et al. Drug-Induced Interstitial Lung Disease: A Systematic Review, *J CLIN MED* 7:

TRISTAN: de Vries, Elisabeth G. E. et al. Integrating molecular nuclear imaging in clinical research to improve anticancer therapy, *NAT REV CLIN ONCOL* 16: 241-255

TRISTAN: Weatherley, Nicholas D. et al. Experimental and quantitative imaging techniques in interstitial lung disease, *THORAX* 74: 611-619

TRISTAN: Raave, Rene et al. Direct comparison of the in vitro and in vivo stability of DFO, DFO\* and DFOcyclo\* for Zr-89-immunoPET, *EUR J NUCL MED MOL I* 46: 1966-1977

U-BIOPRED: Bousquet, Jean et al. Systems medicine and integrated care to combat chronic noncommunicable diseases, *GENOME MED* 3:

U-BIOPRED: Auffray, Charles et al. An Integrative Systems Biology Approach to Understanding Pulmonary Diseases, *CHEST* 137: 1410-1416

U-BIOPRED: Bel, Elisabeth H. et al. Diagnosis and definition of severe refractory asthma: an international consensus statement from the Innovative Medicine Initiative (IMI), *THORAX* 66: 910-917

U-BIOPRED: Harris, Jennifer R. et al. Toward a roadmap in global biobanking for health, *EUR J HUM GENET* 20: 1105-1111

U-BIOPRED: Carraro, S. et al. Asthma severity in childhood and metabolomic profiling of breath condensate, *ALLERGY* 68: 110-117

U-BIOPRED: Montuschi, Paolo et al. The Electronic Nose in Respiratory Medicine, *RESPIRATION* 85: 72-84

U-BIOPRED: Wolkenhauer, Olaf et al. The road from systems biology to systems medicine, *PEDIATR RES* 73: 502-507

U-BIOPRED: Wheelock, Craig E. et al. Application of omics technologies to biomarker discovery in inflammatory lung diseases, *EUR RESPIR J* 42: 802-825

U-BIOPRED: Lambrecht, Bart N. et al. Allergens and the airway epithelium response: Gateway to allergic sensitization, *J ALLERGY CLIN IMMUN* 134: 499-507

U-BIOPRED: Schuijs, Martijn J. et al. ALLERGY Farm dust and endotoxin protect against allergy through A20 induction in lung epithelial cells, *SCIENCE* 349: 1106-1110

U-BIOPRED: Durham, Andrew L. et al. Targeted anti-inflammatory therapeutics in asthma and chronic obstructive lung disease, *TRANSL RES* 167: 192-203

U-BIOPRED: James, Anna J. et al. Increased YKL-40 and Chitotriosidase in Asthma and Chronic Obstructive Pulmonary Disease, *AM J RESP CRIT CARE* 193: 131-142

U-BIOPRED: Auffray, Charles et al. Making sense of big data in health research: Towards an EU action plan, GENOME MED 8:

U-BIOPRED: Lysenko, Artem et al. Representing and querying disease networks using graph databases, BIODATA MIN 9:

U-BIOPRED: Loymans, Rik J. B. et al. Identifying patients at risk for severe exacerbations of asthma: development and external validation of a multivariable prediction model, THORAX 71: 838-846

U-BIOPRED: Loza, Matthew J. et al. Validated and longitudinally stable asthma phenotypes based on cluster analysis of the ADEPT study, RESP RES 17:

U-BIOPRED: Kuo, Chih-Hsi Scott et al. A Transcriptome-driven Analysis of Epithelial Brushings and Bronchial Biopsies to Define Asthma Phenotypes in U-BIOPRED, AM J RESP CRIT CARE 195: 443-455

U-BIOPRED: Kuo, Chih-Hsi Scott et al. T-helper cell type 2 (Th2) and non-Th2 molecular phenotypes of asthma using sputum transcriptomics in U-BIOPRED, EUR RESPIR J 49:

U-BIOPRED: Bigler, Jeannette et al. A Severe Asthma Disease Signature from Gene Expression Profiling of Peripheral Blood from U-BIOPRED Cohorts, AM J RESP CRIT CARE 195: 1311-1320

U-BIOPRED: Lefaudeux, Diane et al. U-BIOPRED clinical adult asthma clusters linked to a subset of sputum omics, J ALLERGY CLIN IMMUN 139: 1797-1807

U-BIOPRED: Rossios, Christos et al. Sputum transcriptomics reveal upregulation of IL-1 receptor family members in patients with severe asthma, J ALLERGY CLIN IMMUN 141: 560-570

U-BIOPRED: Hekking, Pieter-Paul et al. Pathway discovery using transcriptomic profiles in adult-onset severe asthma, J ALLERGY CLIN IMMUN 141: 1280-1290

U-BIOPRED: Takahashi, Kentaro et al. Sputum proteomics and airway cell transcripts of current and ex-smokers with severe asthma in U-BIOPRED: an exploratory analysis, EUR RESPIR J 51:

U-BIOPRED: Shrine, Nick et al. Moderate-to-severe asthma in individuals of European ancestry: a genome-wide association study, LANCET RESP MED 7: 20-34

U-BIOPRED: Pavlidis, Stelios et al. T2-high in severe asthma related to blood eosinophil, exhaled nitric oxide and serum periostin, EUR RESPIR J 53:

U-BIOPRED: Jevnikar, Zala et al. Epithelial IL-6 trans-signaling defines a new asthma phenotype with increased airway inflammation, J ALLERGY CLIN IMMUN 143: 577-590

U-BIOPRED: Mazein, Alexander et al. Systems medicine disease maps: community-driven comprehensive representation of disease mechanisms, NPJ SYST BIOL APPL 4:

U-BIOPRED: Ravanetti, Lara et al. IL-33 drives influenza-induced asthma exacerbations by halting innate and adaptive antiviral immunity, J ALLERGY CLIN IMMUN 143: 1355-+

U-BIOPRED: Brinkman, Paul et al. Identification and prospective stability of electronic nose (eNose)-derived inflammatory phenotypes in patients with severe asthma, J ALLERGY CLIN IMMUN 143: 1811-+

U-BIOPRED: Schofield, James P. R. et al. Stratification of asthma phenotypes by airway proteomic signatures, *J ALLERGY CLIN IMMUN* 144: 70-82

U-BIOPRED: Ostaszewski, Marek et al. Community-driven roadmap for integrated disease maps, *BRIEF BIOINFORM* 20: 659-670

U-BIOPRED: Aarestrup, F. M. et al. Towards a European health research and innovation cloud (HRIC), *GENOME MED* 12:

U-BIOPRED: Kolmert, Johan et al. Urinary Leukotriene E-4 and Prostaglandin D-2 Metabolites Increase in Adult and Childhood Severe Asthma Characterized by Type 2 Inflammation A Clinical Observational Study, *AM J RESP CRIT CARE* 203: 37-53

U-BIOPRED: Kermani, Nazanin Zounemat et al. Sputum ACE2, TMPRSS2 and FURIN gene expression in severe neutrophilic asthma, *RESP RES* 22:

U-BIOPRED: Abdel-Aziz, Mahmoud, I et al. Sputum microbiome profiles identify severe asthma phenotypes of relative stability at 12 to 18 months, *J ALLERGY CLIN IMMUN* 147: 123-134

ULTRA-DD: Hammitzsch, Ariane et al. CBP30, a selective CBP/p300 bromodomain inhibitor, suppresses human Th17 responses, *P NATL ACAD SCI USA* 112: 10768-10773

ULTRA-DD: Xu, Chao et al. Structural Basis for the Discriminative Recognition of N-6-Methyladenosine RNA by the Human YT521-B Homology Domain Family of Proteins, *J BIOL CHEM* 290: 24902-24913

ULTRA-DD: Huang, Ling et al. Ductal pancreatic cancer modeling and drug screening using human pluripotent stem cell- and patient-derived tumor organoids, *NAT MED* 21: 1364-1371

ULTRA-DD: Leitner, Alexander et al. Crosslinking and Mass Spectrometry: An Integrated Technology to Understand the Structure and Function of Molecular Machines, *TRENDS BIOCHEM SCI* 41: 20-32

ULTRA-DD: McAllister, Tom E. et al. Recent Progress in Histone Demethylase Inhibitors, *J MED CHEM* 59: 1308-1329

ULTRA-DD: Bavetsias, Vassilios et al. 8-Substituted Pyrido[3,4-d]pyrimidin-4(3H)-one Derivatives As Potent, Cell Permeable, KDM4 (JMJD2) and KDM5 (JARID1) Histone Lysine Demethylase Inhibitors, *J MED CHEM* 59: 1388-1409

ULTRA-DD: Eram, Mohammad S. et al. A Potent, Selective, and Cell-Active Inhibitor of Human Type I Protein Arginine Methyltransferases, *ACS CHEM BIOL* 11: 772-781

ULTRA-DD: Zhang, Wei et al. System-Wide Modulation of HECT E3 Ligases with Selective Ubiquitin Variant Probes, *MOL CELL* 62: 121-136

ULTRA-DD: Faini, Marco et al. The Evolving Contribution of Mass Spectrometry to Integrative Structural Biology, *J AM SOC MASS SPECTR* 27: 966-974

ULTRA-DD: Eggert, Erik et al. Discovery and Characterization of a Highly Potent and Selective Aminopyrazoline-Based in Vivo Probe (BAY-598) for the Protein Lysine Methyltransferase SMYD2, *J MED CHEM* 59: 4578-4600

ULTRA-DD: Kagoya, Yuki et al. BET bromodomain inhibition enhances T cell persistence and function in adoptive immunotherapy models, *J CLIN INVEST* 126: 3479-3494

ULTRA-DD: de Witte, Wilhelmus E. A. et al. In vivo Target Residence Time and Kinetic Selectivity: The Association Rate Constant as Determinant, *TRENDS PHARMACOL SCI* 37: 831-842

ULTRA-DD: Shen, Yudao et al. Discovery of a Potent, Selective, and Cell-Active Dual Inhibitor of Protein Arginine Methyltransferase 4 and Protein Arginine Methyltransferase 6, *J MED CHEM* 59: 9124-9139

ULTRA-DD: Hauri, Simon et al. A High-Density Map for Navigating the Human Polycomb Complexome, *CELL REP* 17: 583-595

ULTRA-DD: Picaud, Sarah et al. Promiscuous targeting of bromodomains by bromosporine identifies BET proteins as master regulators of primary transcription response in leukemia, *SCI ADV* 2:

ULTRA-DD: Vaz, Bruno et al. Metalloprotease SPRTN/DVC1 Orchestrates Replication-Coupled DNA-Protein Crosslink Repair, *MOL CELL* 64: 704-719

ULTRA-DD: Fang, Hai et al. XGR software for enhanced interpretation of genomic summary data, illustrated by application to immunological traits, *GENOME MED* 8:

ULTRA-DD: Moustakim, Moses et al. Chemical probes and inhibitors of bromodomains outside the BET family, *MEDCHEMCOMM* 7: 2246-2264

ULTRA-DD: Veschi, Veronica et al. Epigenetic siRNA and Chemical Screens Identify SETD8 Inhibition as a Therapeutic Strategy for p53 Activation in High-Risk Neuroblastoma, *CANCER CELL* 31: 50-63

ULTRA-DD: Grieben, Mariana et al. Structure of the polycystic kidney disease TRP channel Polycystin-2 (PC2), *NAT STRUCT MOL BIOL* 24: 114-+

ULTRA-DD: Wilkes, Martin et al. Molecular insights into lipid-assisted Ca<sup>2+</sup> regulation of the TRP channel Polycystin-2, *NAT STRUCT MOL BIOL* 24: 123-+

ULTRA-DD: Xiong, Yan et al. Discovery of Potent and Selective Inhibitors for G9a-Like Protein (GLP) Lysine Methyltransferase, *J MED CHEM* 60: 1876-1891

ULTRA-DD: Fujisawa, Takao et al. Functions of bromodomain-containing proteins and their roles in homeostasis and cancer, *NAT REV MOL CELL BIO* 18: 246-262

ULTRA-DD: He, Yupeng et al. The EED protein-protein interaction inhibitor A-395 inactivates the PRC2 complex, *NAT CHEM BIOL* 13: 389-+

ULTRA-DD: Tumber, Anthony et al. Potent and Selective KDM5 Inhibitor Stops Cellular Demethylation of H3K4me<sub>3</sub> at Transcription Start Sites and Proliferation of MM1S Myeloma Cells, *CELL CHEM BIOL* 24: 371-380

ULTRA-DD: Urbanucci, Alfonso et al. Androgen Receptor Deregulation Drives Bromodomain-Mediated Chromatin Alterations in Prostate Cancer, *CELL REP* 19: 2045-2059

ULTRA-DD: Rocklin, Gabriel J. et al. Global analysis of protein folding using massively parallel design, synthesis, and testing, *SCIENCE* 357: 168-174

ULTRA-DD: Drewry, David H. et al. Progress towards a public chemogenomic set for protein kinases and a call for contributions, *PLOS ONE* 12:

ULTRA-DD: Teyra, Joan et al. Comprehensive Analysis of the Human SH3 Domain Family Reveals a Wide Variety of Non-canonical Specificities, *STRUCTURE* 25: 1598-+

ULTRA-DD: de Freitas, Renato Ferreira et al. A systematic analysis of atomic protein-ligand interactions in the PDB, *MEDCHEMCOMM* 8: 1970-1981

ULTRA-DD: Schapira, Matthieu et al. WD40 repeat domain proteins: a novel target class?, *NAT REV DRUG DISCOV* 16: 773-786

ULTRA-DD: Al-Mossawi, M. H. et al. Unique transcriptome signatures and GM-CSF expression in lymphocytes from patients with spondyloarthritis, *NAT COMMUN* 8:

ULTRA-DD: Dahlin, Jayme L. et al. Assay interference and off-target liabilities of reported histone acetyltransferase inhibitors, *NAT COMMUN* 8:

ULTRA-DD: Fernandez-Montalvan, Amaury E. et al. Isoform-Selective ATAD2 Chemical Probe with Novel Chemical Structure and Unusual Mode of Action, *ACS CHEM BIOL* 12: 2730-2736

ULTRA-DD: Xu, Chao et al. DNA Sequence Recognition of Human CXXC Domains and Their Structural Determinants, *STRUCTURE* 26: 85-+

ULTRA-DD: Asquith, Christopher R. M. et al. Identification and Optimization of 4-Anilinoquinolines as Inhibitors of CyclinG Associated Kinase, *CHEMMEDCHEM* 13: 48-66

ULTRA-DD: Clerici, Marcello et al. Structural basis of AAUAAA polyadenylation signal recognition by the human CPSF complex, *NAT STRUCT MOL BIOL* 25: 135-+

ULTRA-DD: Vasta, James D. et al. Quantitative, Wide-Spectrum Kinase Profiling in Live Cells for Assessing the Effect of Cellular ATP on Target Engagement, *CELL CHEM BIOL* 25: 206-+

ULTRA-DD: Kasinath, Vignesh et al. Structures of human PRC2 with its cofactors AEBP2 and JARID2, *SCIENCE* 359: 940-944

ULTRA-DD: Babault, Nicolas et al. Discovery of Bisubstrate Inhibitors of Nicotinamide N-Methyltransferase (NNMT), *J MED CHEM* 61: 1541-1551

ULTRA-DD: Chaikuad, Apirat et al. The Cysteinome of Protein Kinases as a Target in Drug Development, *ANGEW CHEM INT EDIT* 57: 4372-4385

ULTRA-DD: Dong, Cheng et al. Molecular basis of GID4-mediated recognition of degrons for the Pro/N-end rule pathway, *NAT CHEM BIOL* 14: 466-+

ULTRA-DD: Mueller, Susanne et al. Donated chemical probes for open science, *ELIFE* 7:

ULTRA-DD: Harding, Rachel J. et al. Proteostasis in Huntingtons disease: disease mechanisms and therapeutic opportunities, *ACTA PHARMACOL SIN* 39: 754-769



ULTRA-DD: Fulcher, Luke J. et al. The DUF1669 domain of FAM83 family proteins anchor casein kinase 1 isoforms, *SCI SIGNAL* 11:

ULTRA-DD: Kock, Anna et al. Inhibition of Microsomal Prostaglandin E Synthase-1 in Cancer-Associated Fibroblasts Suppresses Neuroblastoma Tumor Growth, *EBIOMEDICINE* 32: 84-92

ULTRA-DD: Favalli, Nicholas et al. A DNA-Encoded Library of Chemical Compounds Based on Common Scaffolding Structures Reveals the Impact of Ligand Geometry on Protein Recognition, *CHEMMEDCHEM* 13: 1303-1307

ULTRA-DD: Hudson, Liam et al. Novel Quinazolinone Inhibitors of ALK2 Flip between Alternate Binding Modes: Structure-Activity Relationship, Structural Characterization, Kinase Profiling, and Cellular Proof of Concept, *J MED CHEM* 61: 7261-7272

ULTRA-DD: Bonday, Zahid Q. et al. LLY-283, a Potent and Selective Inhibitor of Arginine Methyltransferase 5, PRMT5, with Antitumor Activity, *ACS MED CHEM LETT* 9: 612-617

ULTRA-DD: Tu, William B. et al. MYC Interacts with the G9a Histone Methyltransferase to Drive Transcriptional Repression and Tumorigenesis, *CANCER CELL* 34: 579-+

ULTRA-DD: Imrie, Fergus et al. Protein Family-Specific Models Using Deep Neural Networks and Transfer Learning Improve Virtual Screening and Highlight the Need for More Data, *J CHEM INF MODEL* 58: 2319-2330

ULTRA-DD: Kalkat, Manpreet et al. MYC Protein Interactome Profiling Reveals Functionally Distinct Regions that Cooperate to Drive Tumorigenesis, *MOL CELL* 72: 836-+

ULTRA-DD: Scheer, Sebastian et al. A chemical biology toolbox to study protein methyltransferases and epigenetic signaling, *NAT COMMUN* 10:

ULTRA-DD: Christott, Thomas et al. Discovery of a Selective Inhibitor for the YEATS Domains of ENL/AF9, *SLAS DISCOV* 24: 133-141

ULTRA-DD: Lambert, Jean-Philippe et al. Interactome Rewiring Following Pharmacological Targeting of BET Bromodomains, *MOL CELL* 73: 621-+

ULTRA-DD: Schewe, Marcus et al. A pharmacological master key mechanism that unlocks the selectivity filter gate in K<sup>+</sup> channels, *SCIENCE* 363: 875-+

ULTRA-DD: Watts, Ellen et al. Designing Dual Inhibitors of Anaplastic Lymphoma Kinase (ALK) and Bromodomain-4 (BRD4) by Tuning Kinase Selectivity, *J MED CHEM* 62: 2618-2637

ULTRA-DD: Asquith, Christopher R. M. et al. SGC-GAK-1: A Chemical Probe for Cyclin G Associated Kinase (GAK), *J MED CHEM* 62: 2830-2836

ULTRA-DD: Ebrahimi, Ayyub et al. Bromodomain inhibition of the coactivators CBP/EP300 facilitate cellular reprogramming, *NAT CHEM BIOL* 15: 519-+

ULTRA-DD: Wu, Qin et al. A chemical toolbox for the study of bromodomains and epigenetic signaling, *NAT COMMUN* 10:

ULTRA-DD: Asquith, Christopher R. M. et al. Design of a Cyclin G Associated Kinase (GAK)/Epidermal Growth Factor Receptor (EGFR) Inhibitor Set to Interrogate the Relationship of EGFR and GAK in Chordoma, *J MED CHEM* 62: 4772-4778

ULTRA-DD: Carvalho, Diana et al. ALK2 inhibitors display beneficial effects in preclinical models of ACVR1 mutant diffuse intrinsic pontine glioma, *COMMUN BIOL* 2:

ULTRA-DD: Fox, Nicholas G. et al. Structure of the human frataxin-bound iron-sulfur cluster assembly complex provides insight into its activation mechanism, *NAT COMMUN* 10:

ULTRA-DD: Resnick, Efrat et al. Rapid Covalent-Probe Discovery by Electrophile-Fragment Screening, *J AM CHEM SOC* 141: 8951-8968

ULTRA-DD: Celis-Gutierrez, Javier et al. Quantitative Interactomics in Primary T Cells Provides a Rationale for Concomitant PD-1 and BTLA Coinhibitor Blockade in Cancer Immunotherapy, *CELL REP* 27: 3315-+

ULTRA-DD: Fang, Hai et al. A genetics-led approach defines the drug target landscape of 30 immune-related traits, *NAT GENET* 51: 1082-+

ULTRA-DD: Amon, Sabine et al. Sensitive Quantitative Proteomics of Human Hematopoietic Stem and Progenitor Cells by Data-independent Acquisition Mass Spectrometry, *MOL CELL PROTEOMICS* 18: 1454-1467

ULTRA-DD: Fong, Jia Yi et al. Therapeutic Targeting of RNA Splicing Catalysis through Inhibition of Protein Arginine Methylation, *CANCER CELL* 36: 194-+

ULTRA-DD: Bushell, K. Simon R. et al. The structural basis of lipid scrambling and inactivation in the endoplasmic reticulum scramblase TMEM16K, *NAT COMMUN* 10

ULTRA-DD: Alam, Mahmood M. et al. Validation of the protein kinase PfCLK3 as a multistage cross-species malarial drug target, *SCIENCE* 365: 884-+

ULTRA-DD: Shigesu, Nina et al. The association between endometriosis and autoimmune diseases: a systematic review and meta-analysis, *HUM REPROD UPDATE* 25: 486-503

ULTRA-DD: Liu, Lihua et al. UbiHub: a data hub for the explorers of ubiquitination pathways, *BIOINFORMATICS* 35: 2882-2884

ULTRA-DD: Ostrom, Quinn T. et al. Risk factors for childhood and adult primary brain tumors, *NEURO-ONCOLOGY* 21: 1357-1375

ULTRA-DD: Bauer, Matthias R. et al. A structure-guided molecular chaperone approach for restoring the transcriptional activity of the p53 cancer mutant Y220C, *FUTURE MED CHEM* 11: 2491-2504

ULTRA-DD: Schapira, Matthieu et al. Targeted protein degradation: expanding the toolbox, *NAT REV DRUG DISCOV* 18: 949-963

ULTRA-DD: Carter, Adrian J. et al. Target 2035: probing the human proteome, *DRUG DISCOV TODAY* 24: 2111-2115

ULTRA-DD: Ayelen Carabaja, Maria et al. Quinazoline-Based Antivirulence Compounds Selectively Target Salmonella PhoP/PhoQ Signal Transduction System, *ANTIMICROB AGENTS CH* 64

ULTRA-DD: Shukla, Vipul et al. HMCES Functions in the Alternative End-Joining Pathway of the DNA DSB Repair during Class Switch Recombination in B Cells, *MOL CELL* 77: 384-+

ULTRA-DD: Klatt, Felix et al. A precisely positioned MED12 activation helix stimulates CDK8 kinase activity, *P NATL ACAD SCI USA* 117: 2894-2905

ULTRA-DD: Villasenor, Rodrigo et al. ChromID identifies the protein interactome at chromatin marks, *NAT BIOTECHNOL* 38: 728-+

ULTRA-DD: Heusel, Moritz et al. A Global Screen for Assembly State Changes of the Mitotic Proteome by SEC-SWATH-MS, *CELL SYST* 10: 133-+

ULTRA-DD: Fortin, Jerome et al. Mutant ACVR1 Arrests Glial Cell Differentiation to Drive Tumorigenesis in Pediatric Gliomas, *CANCER CELL* 37: 308-+

ULTRA-DD: Gray, Janine L. et al. Targeting the Small GTPase Superfamily through Their Regulatory Proteins, *ANGEW CHEM INT EDIT* 59: 6342-6366

ULTRA-DD: Ward, Jennifer A. et al. Re-Evaluating the Mechanism of Action of alpha,beta-Unsaturated Carbonyl DUB Inhibitors b-AP15 and VLX1570: A Paradigmatic Example of Unspecific Protein Cross-linking with Michael Acceptor Motif-Containing Drugs, *J MED CHEM* 63: 3756-3762

ULTRA-DD: Bergqvist, Filip et al. A review on mPGES-1 inhibitors: From preclinical studies to clinical applications, *PROSTAG OTH LIPID M* 147

ULTRA-DD: Rodstrom, Karin E. J. et al. A lower X-gate in TASK channels traps inhibitors within the vestibule, *NATURE* 582: 443-+

ULTRA-DD: Zhang, Xin et al. Therapeutic targeting of p300/CBP HAT domain for the treatment of NUT midline carcinoma, *ONCOGENE* 39: 4770-4779

ULTRA-DD: Ferla, Matteo P. et al. MichelaNglo: sculpting protein views on web pages without coding, *BIOINFORMATICS* 36: 3268-3270

ULTRA-DD: Michealraj, Kulandaimanuel Antony et al. Metabolic Regulation of the Epigenome Drives Lethal Infantile Ependymoma, *CELL* 181: 1329-+

ULTRA-DD: Sanchez-Duffhues, Gonzalo et al. Bone morphogenetic protein receptors: Structure, function and targeting by selective small molecule kinase inhibitors, *BONE* 138

ULTRA-DD: Szewczyk, Magdalena M. et al. Pharmacological inhibition of PRMT7 links arginine monomethylation to the cellular stress response, *NAT COMMUN* 11

ULTRA-DD: Deblois, Genevieve et al. Epigenetic Switch-Induced Viral Mimicry Evasion in Chemotherapy-Resistant Breast Cancer, *CANCER DISCOV* 10: 1312-1329

ULTRA-DD: Wu, Qin et al. GLUT1 inhibition blocks growth of RB1-positive triple negative breast cancer, *NAT COMMUN* 11

ULTRA-DD: Douangamath, Alice et al. Crystallographic and electrophilic fragment screening of the SARS-CoV-2 main protease, *NAT COMMUN* 11

ULTRA-DD: Stoesser, Nicole et al. Performance characteristics of five immunoassays for SARS-CoV-2: a head-to-head benchmark comparison, *LANCET INFECT DIS* 20: 1390-1400

ULTRA-DD: Kasinath, Vignesh et al. JARID2 and AEBP2 regulate PRC2 in the presence of H2AK119ub1 and other histone modifications, *SCIENCE* 371: 362-+

ULTRA-DD: de Freitas, Renato Ferreira et al. Discovery of Small-Molecule Antagonists of the PWWP Domain of NSD2, *J MED CHEM* 64: 1584-1592

ULTRA-DD: Sachamitr, Patty et al. PRMT5 inhibition disrupts splicing and stemness in glioblastoma, *NAT COMMUN* 12

ULTRA-DD: Eyre, David W. et al. Stringent thresholds in SARS-CoV-2 IgG assays lead to under-detection of mild infections, *BMC INFECT DIS* 21

ULTRA-DD: Schuller, Marion et al. Fragment binding to the Nsp3 macrodomain of SARS-CoV-2 identified through crystallographic screening and computational docking, *SCI ADV* 7

ULTRA-DD: Fossati, Andrea et al. PCprophet: a framework for protein complex prediction and differential analysis using proteomic data, *NAT METHODS* 18: 520-+

ULTRA-DD: Wells, Carrow, I et al. Development of a potent and selective chemical probe for the pleiotropic kinase CK2, *CELL CHEM BIOL* 28: 546-+

ULTRA-DD: Shum, Michael et al. ABCB10 exports mitochondrial biliverdin, driving metabolic maladaptation in obesity, *SCI TRANSL MED* 13

ULTRA-DD: Lumley, Sheila F. et al. The Duration, Dynamics, and Determinants of Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) Antibody Responses in Individual Healthcare Workers, *CLIN INFECT DIS* 73: E699-E709

VAC2VAC: Etna, Marilena P. et al. Human plasmacytoid dendritic cells at the crossroad of type I interferon-regulated B cell differentiation and antiviral response to tick-borne encephalitis virus, *PLOS PATHOG* 17

VALUE-Dx: De Nardo, Pasquale et al. Multi-Criteria Decision Analysis to prioritize hospital admission of patients affected by COVID-19 in low-resource settings with hospital-bed shortage, *INT J INFECT DIS* 98: 494-500

VALUE-Dx: Hellou, Mona Mustafa et al. Nucleic acid amplification tests on respiratory samples for the diagnosis of coronavirus infections: a systematic review and meta-analysis, *CLIN MICROBIOL INFECT* 27: 341-351

VALUE-Dx: Rodriguez-Bano, Jesus et al. Antimicrobial resistance research in a post-pandemic world: Insights on antimicrobial resistance research in the COVID-19 pandemic, *J GLOB ANTIMICROB RE* 25: 5-7

VALUE-Dx: van der Velden, Alike W. et al. Primary care for patients with respiratory tract infection before and early on in the COVID-19 pandemic: an observational study in 16 European countries, *BMJ OPEN* 11

VALUE-Dx: Rodriguez-Bano, Jesus et al. Key considerations on the potential impacts of the COVID-19 pandemic on antimicrobial resistance research and surveillance, *T ROY SOC TROP MED H* 115: 1122-1129

VITAL: Wagner, Angelika et al. Vaccines to Prevent Infectious Diseases in the Older Population: Immunological Challenges and Future Perspectives, *FRONT IMMUNOL* 11

VSV-EBOPLUS: Alter, Galit et al. Antibody glycosylation in inflammation, disease and vaccination, *SEMIN IMMUNOL* 39: 102-110

VSV-EBOPLUS: Mathew, Nimitha R. et al. Single-cell BCR and transcriptome analysis after influenza infection reveals spatiotemporal dynamics of antigen-specific B cells, *CELL REP* 35

VSV-EBOVAC: Huttner, Angela et al. A dose-dependent plasma signature of the safety and immunogenicity of the rVSV-Ebola vaccine in Europe and Africa, *SCI TRANSL MED* 9

WEB-RADR: Sloane, Richard et al. Social media and pharmacovigilance: A review of the opportunities and challenges, *BRIT J CLIN PHARMACO* 80: 910-920

WEB-RADR: Powell, Gregory E. et al. Social Media Listening for Routine Post-Marketing Safety Surveillance, *DRUG SAFETY* 39: 443-454

WEB-RADR: Pierce, Carrie E. et al. Evaluation of Facebook and Twitter Monitoring to Detect Safety Signals for Medical Products: An Analysis of Recent FDA Safety Alerts, *DRUG SAFETY* 40: 317-331

WEB-RADR: Caster, Ola et al. Assessment of the Utility of Social Media for Broad-Ranging Statistical Signal Detection in Pharmacovigilance: Results from the WEB-RADR Project, *DRUG SAFETY* 41: 1355-1369

WEB-RADR: Pierce, Carrie E. et al. Recommendations on the Use of Mobile Applications for the Collection and Communication of Pharmaceutical Product Safety Information: Lessons from IMI WEB-RADR, *DRUG SAFETY* 42: 477-489

ZAPI: Haagmans, Bart L. et al. An orthopoxvirus-based vaccine reduces virus excretion after MERS-CoV infection in dromedary camels, *SCIENCE* 351: 77-81

ZAPI: Ludlow, Martin et al. Neurotropic virus infections as the cause of immediate and delayed neuropathology, *ACTA NEUROPATHOL* 131: 159-184

ZAPI: Widagdo, W. et al. Differential Expression of the Middle East Respiratory Syndrome Coronavirus Receptor in the Upper Respiratory Tracts of Humans and Dromedary Camels, *J VIROL* 90: 4838-4842

ZAPI: Becares, Martina et al. Mutagenesis of Coronavirus nsp14 Reveals Its Potential Role in Modulation of the Innate Immune Response, *J VIROL* 90: 5399-5414

ZAPI: Vergara-Alert, Julia et al. Livestock Susceptibility to Infection with Middle East Respiratory Syndrome Coronavirus, *EMERG INFECT DIS* 23: 232-240

ZAPI: Wernike, Kerstin et al. The N-terminal domain of Schmallenberg virus envelope protein Gc is highly immunogenic and can provide protection from infection, *SCI REP-UK* 7

ZAPI: Morales, Lucia et al. SARS-CoV-Encoded Small RNAs Contribute to Infection-Associated Lung Pathology, *CELL HOST MICROBE* 21: 344-355

ZAPI: Okba, Nisreen M. A. et al. Middle East respiratory syndrome coronavirus vaccines: current status and novel approaches, *CURR OPIN VIROL* 23: 49-58

ZAPI: Munyua, Peninah et al. No Serologic Evidence of Middle East Respiratory Syndrome Coronavirus Infection among Camel Farmers Exposed to Highly Seropositive Camel Herds: A Household Linked Study, Kenya, 2013, *AM J TROP MED HYG* 96: 1318-1324

ZAPI: Li, Wentao et al. Identification of sialic acid-binding function for the Middle East respiratory syndrome coronavirus spike glycoprotein, *P NATL ACAD SCI USA* 114: E8508-E8517

ZAPI: Canton, Javier et al. MERS-CoV 4b protein interferes with the NF-kappa B-dependent innate immune response during infection, *PLOS PATHOG* 14

ZAPI: Widagdo, W. et al. MERS-coronavirus: From discovery to intervention, *ONE HEALTH-AMSTERDAM* 3: 11-16

ZAPI: Rey, Felix A. et al. Common Features of Enveloped Viruses and Implications for Immunogen Design for Next-Generation Vaccines, *CELL* 172: 1319-1334

ZAPI: Castano-Rodriguez, Carlos et al. Role of Severe Acute Respiratory Syndrome Coronavirus Viroporins E, 3a, and 8a in Replication and Pathogenesis, *MBIO* 9

ZAPI: Raj, V. Stalin et al. Chimeric camel/human heavy-chain antibodies protect against MERS-CoV infection, *SCI ADV* 4

ZAPI: Hellert, Jan et al. Orthobunyavirus spike architecture and recognition by neutralizing antibodies, *NAT COMMUN* 10

ZAPI: Widjaja, Ivy et al. Towards a solution to MERS: protective human monoclonal antibodies targeting different domains and functions of the MERS-coronavirus spike glycoprotein, *EMERG MICROBES INFEC* 8: 516-530

ZAPI: Siu, Kam-Leung et al. Severe acute respiratory syndrome coronavirus ORF3a protein activates the NLRP3 inflammasome by promoting TRAF3-dependent ubiquitination of ASC, *FASEB J* 33: 8865-8877

ZAPI: Widagdo, W. et al. Species-Specific Colocalization of Middle East Respiratory Syndrome Coronavirus Attachment and Entry Receptors, *J VIROL* 93

ZAPI: Okba, Nisreen M. A. et al. Sensitive and Specific Detection of Low-Level Antibody Responses in Mild Middle East Respiratory Syndrome Coronavirus Infections, *EMERG INFECT DIS* 25: 1868-1877

ZAPI: Wang, Chunyan et al. A human monoclonal antibody blocking SARS-CoV-2 infection, *NAT COMMUN* 11

ZAPI: Okba, Nisreen M. A. et al. Severe Acute Respiratory Syndrome Coronavirus 2-Specific Antibody Responses in Coronavirus Disease Patients, EMERG INFECT DIS 26: 1478-1488

ZAPI: Gutierrez-Alvarez, Javier et al. Middle East Respiratory Syndrome Coronavirus Gene 5 Modulates Pathogenesis in Mice, J VIROL 95

ZAPI: Wang, Chunyan et al. A conserved immunogenic and vulnerable site on the coronavirus spike protein delineated by cross-reactive monoclonal antibodies, NAT COMMUN 12

Unassigned: Visscher, Peter M. et al. 10 Years of GWAS Discovery: Biology, Function, and Translation, AM J HUM GENET 101: 5-22

Unassigned: Attal, Nadine et al. Diagnosis and assessment of neuropathic pain through questionnaires, LANCET NEUROL 17: 456-466

Unassigned: Ahlqvist, Emma et al. Novel subgroups of adult-onset diabetes and their association with outcomes: a data-driven cluster analysis of six variables, LANCET DIABETES ENDO 6: 361-369

Unassigned: Williams, Tim et al. Recent advances in the utility and use of the General Practice Research Database as an example of a UK Primary Care Data resource, THER ADV DRUG SAF 3: 89-99

Unassigned: Carey, Iain M. et al. Are noise and air pollution related to the incidence of dementia? A cohort study in London, England, BMJ OPEN 8:

Unassigned: Brazel, David M. et al. Exome Chip Meta-analysis Fine Maps Causal Variants and Elucidates the Genetic Architecture of Rare Coding Variants in Smoking and Alcohol Use, BIOL PSYCHIAT 85: 946-955

Unassigned: Heerspink, Hidde J. L. et al. Canagliflozin reduces inflammation and fibrosis biomarkers: a potential mechanism of action for beneficial effects of SGLT2 inhibitors in diabetic kidney disease, DIABETOLOGIA 62: 1154-1166

Unassigned: Gallagher, Arlene M. et al. The accuracy of date of death recording in the Clinical Practice Research Datalink GOLD database in England compared with the Office for National Statistics death registrations, PHARMACOEPIDEM DR S 28: 563-569

Unassigned: Generaal, Ellen et al. Neighbourhood characteristics and prevalence and severity of depression: pooled analysis of eight Dutch cohort studies, BRIT J PSYCHIAT 215: 468-475

Unassigned: Dhar, Raja et al. Bronchiectasis in India: results from the European Multicentre Bronchiectasis Audit and Research Collaboration (EMBARC) and Respiratory Research Network of India Registry, LANCET GLOB HEALTH 7: E1269-E1279

Unassigned: Piot, Peter et al. Immunization: vital progress, unfinished agenda, NATURE 575: 119-129

Unassigned: Cassidy, Sarah A. et al. Measurement Properties of the Suicidal Behaviour Questionnaire-Revised in Autistic Adults, J AUTISM DEV DISORD 50: 3477-3488

Unassigned: Bachert, Claus et al. Staphylococcus aureus and its IgE-inducing enterotoxins in asthma: current knowledge, EUR RESPIR J 55:

Unassigned: Sabiti, Wilber et al. Tuberculosis bacillary load, an early marker of disease severity: the utility of tuberculosis Molecular Bacterial Load Assay, THORAX 75: 606-608

Unassigned: Sheetz, Joshua B. et al. Structural Insights into Pseudokinase Domains of Receptor Tyrosine Kinases, MOL CELL 79: 390+

Unassigned: Ball, Harriet A. et al. Functional cognitive disorder: dementias blind spot, BRAIN 143: 2895-2903

Unassigned: Barreto, Savio George et al. Critical thresholds: key to unlocking the door to the prevention and specific treatments for acute pancreatitis, GUT 70: 194-203

Unassigned: McCrimmon, Rory J. et al. Consequences of recurrent hypoglycaemia on brain function in diabetes, DIABETOLOGIA 64: 971-977

Unassigned: Custers, Jerome et al. Vaccines based on replication incompetent Ad26 viral vectors: Standardized template with key considerations for a risk/benefit assessment, VACCINE 39: 3081-3101

Unassigned: Oliveira, Luis M. A. et al. Alpha-synuclein research: defining strategic moves in the battle against Parkinsons disease, NPJ PARKINSONS DIS 7:

Unassigned: Aman, Jurjan et al. Imatinib in patients with severe COVID-19: a randomised, double-blind, placebo-controlled, clinical trial, LANCET RESP MED 9: 957-968



Prepared by Clarivate on behalf of IHI Programme Office under a public procurement procedure  
document reference: IMI.2018.OP.01.