IHI Call Days | Call 9

EU-Wide Innovative PCM Platform Neoadjuvant Therapy

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- EU-wide innovative PCM Platform
- Nancy Frédérickx



Challenges and Objectives

Topic 2: Boosting innovation through better integration of fragmented health R&I efforts (SO2-SRIA)

Personalized neoadjuvant treatments have the potential to reduce the cancer burden across various type. Yet, its widespread adoption faces significant challenges:

- Complexity of clinical trial organization
- Integration of innovative technologies
- Regulatory hurdles
- Limited sustainable access

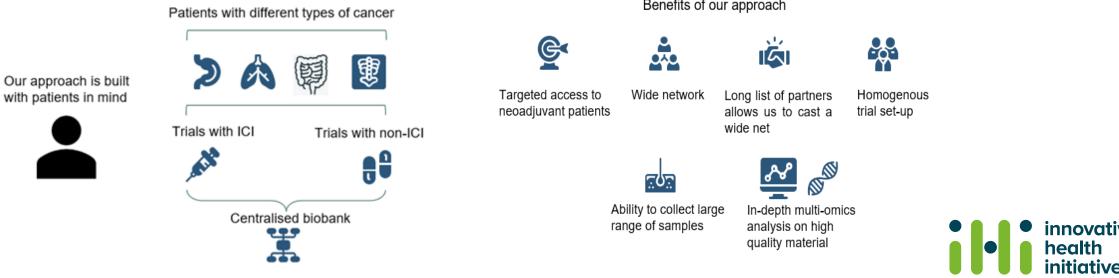
AIM:

To establish an EU-Wide Platform for Neoadjuvant therapy that reshapes clinical trial organization and promote unity and efficiency



Key Elements of the Proposal I

- Create a unified platform for neoadjuvant therapy
 - Support standardized trial setups
 - Integration of cutting-edge technologies
 - Standardise sampling processing method
 - Promote access to skilled personnel



MANIFEST UK Cancer Immunotherapy Response Research Platform

Benefits of our approach



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Key Elements of the Proposal II

- Pave the way to establish a neoadjuvant learning system
 - Accelerate therapy development & promote cancer care
- Foster synergy with EU Initiatives
 - EUnetCCC and upcoming Joint action PCM
- Support and facilitate regulatory alignment





IHI Ideal Instrument

Public-private collaboration within the healthcare system is essential to:

- Share infrastructure,
- Develop robust implementation frameworks,
- Address regulatory challenges

Specific contributions in

- Access to new therapies and clinical trial expertise
- Advanced multi-omics technologies and translational research
- Centralise digital platforms and data integration (AI/ML)
- Insights and guidance for local deployment



Platform Impacts and Outcomes

Impacts

- Streamline Clinical Trials & Speed Development
- Identify Surrogate Outcome Markers to Support Regulatory Framework
- Integrate Advanced Technologies (e.g., Omics)

Outcomes

- Pave the Way for a Neoadjuvant Learning System
- Standardized Sample Collection and Methods
- Collect Robust Evidence for Regulatory Transformation
- Enhance Sustainable Access & Patient Outcome



Expertise and Resources

- We have:
 - Public Health and Clinical Expertise
 - Public-private project management
 - Including communication, dissemination and exploitation
 - Multi-Omics, Molecular Biology, and Data Integration
 - EU oncology research & care Network
 - Strong connection with EU oncology initiative (coordinator of JA PCM)
- We are looking for expertise in:
 - Pharmaceutical Development
 - Advanced Multi-Omics Analysis
 - Data Management and Analytics
 - Al-Driven Cancer Diagnostics
 - Health Informatics
 - Regulatory bodies
 - Patient Advocacy group



This platform will accelerate the field of neoadjuvant therapies, transforming cancer care and improving patient outcomes across Europe





