

Cancer Core Europe

Paving the way for a multi-site virtual European Cancer Institute From New Treatments (POC-studies) to Outcome Research













VALL D'HEBRON Institut d'Oncologia



SUSTAINABILITY

• FP6 EUROCAN+

- inventory of barriers, fragmentation
- Recommendation: create TR Platform
- FP7 EUROCAN TRANSLATION RESEARCH
 PLATFORM
 - 26 partners, 16 wps
 - Recommendation: create Cancer Core Europe

CREATION CANCER CORE EUROPE

- Autofinanced
- Sustainability / Core / Expand later































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Objectives

• Transformative initiative to create sustainable, integrated research-care capabilities

create a multi-site European Cancer Institute

- carry out joint translational and clinical research
- conduct next-generation clinical trials
- develop personalized cancer medicine
- establish standardized academic diagnostic platforms
- create large shared databases
- perform outcomes research















Prerequisites to deal with inherent complexity

- High volume clinical research activity
- Highly developed research infrastructures
- Cutting-edge track records
- Complementarity
 - Basic Research Infrastructure
 - Translational Research Culture
 - Clinical Research Programs
 - Early Clinical Trials Culture
 - Investigator Initiated Trials Culture















Critical Mass

Clinical Activity

- 60,000 new patients/yr
- 250,000 300,000 patients treated/yr
- > 1.2 Million consultations

Common IT system to

- Share Clinical Databases
- Share Research Databases
- Share Research Projects















One Portal

- Create Virtual e-Cancer Institute
 - Consortium transformed into Association
 - Legal Entity
 - One Portal for Clinical Trials
 - Attractive partner for Pharma/Biotech















Common SOPs and Platforms

- Tissue procurement and biobanking SOPs
- Common validated (400 genes) molecular diagnostic platform
- Common immunomonitoring platform
- Functional imaging platform
- Common bioinformatic pipelines
- SAP multi-language unstructured to structured data classifying system















From Early Clinical Trials to Outcomes Research

- Infrastructure
- Culture
- Critical Mass















Cancer Core Europe 6 Working Groups

Data Sharing
 Common Diagnostic Platforms (400 genes / immunomonitoring)
 Functional Imaging
 Preclinical development – Early Clinical trials

 4a Molecular Medicine
 4b Immunotherapy

 Databases- Outcomes Research
 Training













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Task Force #1 Data sharing via common IT platform

- Create platform allowing the exchange of data (genomic, imaging, clinical, treatment outcomes) across centers
- Use common standards to ensure data interoperability
- Implement mandatory metadata directory
- First step will incorporate diseases where using the data for discovery-based research can make an impact















Task Force #2 Molecular Diagnostics Platform

- Common panel of genes available in 6 centers
- Common pipelines for alignment, variant calling, and analysis
- Circulating information on molecularly defined clinical groups to decide on appropriate therapies
- Common controls ensure genomic data is high-quality and uniform















Task Force #3 Functional Imaging

- Incorporate functional imaging into data sharing platform
- Goal 1 Develop predictive modeling
- Goal 2 Identify features of response to treatment with immune checkpoint inhibitors
 - Early response
 - Acquired resistance















Task Force #4 Clinical Trials/Omics

- Evaluate whether matching treatment to molecular abnormalities induces antitumor activity
- Profile patients according to molecular signatures and pathway alterations
- Evaluate tumor heterogeneity, sensitivity, and resistance
- Launch Basket trials
- Follow-up patients using biopsies and cfDNA
- Use innovative statistical designs Cancer Core Europe expertise in biostatistics will be key















Task Force #4b Immunotherapy

- Cancer Core Europe strengths:
 - T cells, T cell repertoire, NK cells, Adoptive Cell Therapies
 - Microenvironment dendritic, vasculature
 - Immunomonitoring platforms
 - **INTEGRATE Genomics**: protein structure and neoantigens, etc.
- Above areas will be incorporated into new clinical trials
- Patient samples to be tested in different institutions according to their expertise after defining Cancer Core Europe immune biomarker panel















Task Force #5 Prospective Fully Anotated Databases: basis for outcome research

- Critical Mass
- Shared / Prospective
- Fully clinically anotated
- Research data anotated















Task Force #6 Training and Mobility

- Molecular Medicine Autumn School
- Mobility grant inter-laboratory
- etc

















1) Funding provided by the 6 members

2) Successful grant applications

- A) EIT Health KIC Innolife
 - » PCM and Biomarkers
- B) Transcan EU projects
 - Heterogeneity in breast and ovarian cancers
 - Triple negative breast cancer
 - Hematological cancers
- C) ANR grant
 - Support coordination inside Cancer Core Europe
- 3) Future aims: IMI2, H2020















Cancer Core Europe Trials

MOLECULAR MEDECIN

- POSEIDON trial
 - (PI3Kinhibitors in Luminal Breast Cancer)
- BASKET OF BASKETS
 - Across tumor types
- From Preclinical into Phase I
 - Trail inhibitor
- IMMUNOTHERAPY
 - SABR-PDL1 trial
 - Stereotaxic Radiotherapy + antiPDL1 metastatic cancers
 - ILSI
 - (CRC Livermets: oxaliplatin (iimunogenic cell death inducer) followed by intralesional checkpoint inhibitors + TLR4inh)















INCLUSIVENESS

- First build core centers, then core structures, then open up for expansion
- Exclusive at first, with goal to become inclusive across Europe
- Precursor for virtual ECI















Objective: Precursor of ECI













