

European infrastructure for translational medicine

Introduction

TRANSCAN-3 ERA-NET JTC 2021 International Networking Event

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Anne-Charlotte Fauvel EATRIS Head of European Affairs



Life Sciences Research Infrastructures







Our mission

European infrastructure for translational medicine



To accelerate the translation of research discoveries into patient benefit.

We support academia, industry, patients and policy makers.



Who we are

European infrastructure for translational medicine

Facilities, resources and services to support cutting edge research



eatrıs

What do we offer

European infrastructure for translational medicine



Five Scientific Platforms

eatris



Support for funding applications



EATRIS offers a range of services to help researchers strengthen the translational potential of their research proposals:



Support for funding applications



EATRIS can participate in research funding proposals as a full partner providing various centralised services:



How to make use of EATRIS for your TRANSCAN proposal?





FORMING A CONSORTIUM Submit your consortium-building request using <u>the link</u> or visit eatris.eu/services/support-for-funding-applications/

List of matches with contact information will be provided to you within less than 5 business days.



JOINING AS SUBCONTRACTOR Please check eligibility criteria of your funder.

Contact Anne-Charlotte Fauvel to discuss the needs of your proposal: <u>annecharlottefauvel@eatris.eu</u>

With thanks to our members

eatris



* The contribution in Norway is shared between University of Oslo (UiO), University of Bergen (UiB), Norwegian University of Science and Technology (NTNU), the Arctic University of Norway (UiT) and the four Regional Health Authorities in Southeastern, Western, Central and Northern Norway



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Stay in contact

in EATRIS@EatrisEricwww.eatris.eu

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- Translational Trand
- Translational Trends



Detailed overview of EATRIS' members resources for TRANSCAN-3 JTC 2021



- Multiomics technologies (Genomics: NGS, deep genome sequencing, Mass spectrometry for multi-omic analysis), Expertise in Biomarkers (Discovery, validation, assays), Samples/data cohorts
- Systems level characterization of immune cells in human tissues (multi-parametric flow cytometry, mass cytometry [CyTOF], Helios) and Immune assays (in vitro functional immune assays, high-throughput multiplex immunoassay)
- Identification and isolation of tissue immunological subsets for deep profiling (RNAseq, Spectral Flow, single cell RNA-seq), SNP Array, TCR- sequencing (immunochip, immunoseq) of immune cell subsets
- Large/Medium-scale analysis of the immune proteins (Mass spectrometry; Multiplex immuno-assays)
- Access to 3D culture systems; patient-derived organoids; patient-derived xenografts, Spheroids and Multifluidic Devices for Immune surveillance in TME
- In vivo Imaging Technologies to Monitor the Immune System Mass Cytometry Imaging (MCI), PET-CT, PET-MRI, US modalities for studies of the immune system response
- Epigenetics of immune cells to study genome-wide epigenetic changes including DNA methylation, histone modifications and non-coding RNAs expression.
- Multiomics (epigenomic, transcriptomic, proteomic, metabolomics, study of the microbiome and virome, etc.) to assess functional characteristics of TME-tumour cell interplay within the primary tumour and/or metastases (e.g the underlying signaling, the transcriptional landscape, the cell-cell communication, the network regulation of immune cells, etc.), to identify candidate TME targets and to assess the activity of pathway-targeting agents.
- Artificial Intelligence expertise to develop predictive models based on integrating -omics data and network approaches.
- · Access to Tumour samples collected from retrospective and/or prospective cohorts of patients.
- Expertise in Radiomics, cell-free circulating tumour DNA, miRNA signatures.
- Expertise in cell therapy and genome editing products in targeted cells and tissues (e.g., base editing, prime editing, talens, zinc-finger nucleases, CRISPR).
- Expertise in novel RNA-based therapeutics targeting Cancer.
- Expertise in Regulatory Expertise for Advanced Therapies
- Expertise in upscaling and product development of advanced therapies
- Drug discovery, medicinal chemistry and development expertise (safety, toxicity, pharmacokinetics) all the way up to GLP safety testing and Phase 1 trial support.
- Identification of new drug targets and mechanism of action studies
- · Lead optimisation of targeted small molecule inhibitors of the immune system
- Development of specific molecular tracers to support target and biomarker validation and target engagement in tumour (micro) environment
- Development of in vivo (clinical grade) imaging probes derived from peptides and antibodies involed in immune-oncology (e.g. checkpoint inhibitors)
- Tracking of immune cells
- Chemosensitivity screening using existing drug collections applied to patient samples combined with data sharing/FAIRification.