

Tumor-LN-oC

Tumor and Lymph Node on Chip for cancer studies



This project has received funding from the European Union's Horizon 2020 research and innovation programme under grant agreement No 953234

The Project



Tumor and Lymph Node on Chip for cancer studies



Tumor-LN-oC proposes the development and <a> Molecular analysis validation of **tumor-lymph** node-on-chip platform that will connect primary surgically removed human tumors and lymph node tissue from the same cancer patient.

Project's Goal: Technologies involved:

- ֎ Cancer biology
- Microfluidics
- Mid-IR photothermal spectroscopic sensors
- Realtime imaging sensors
- ML deep learning 8 algorithms
- Laser bioprinting æ



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The consortium



The consortium encompasses key – industrial partners and experts in the relevant interdisciplinary fields and is expected to have substantial impact in EU's economy and healthcare.



Project summary



- Tumor-LN-oC is developing a TRL 5 tumor-lymph node on-chip platform which will mimic the tumor microenvironment and its connection to the lymphatic system.
- Using surgically removed human primary tumors and LN tissue from the same lung cancer patient, the multidisciplinary microfluidic platform will serve as a "biological twin" of the patient.
- This makes Tumor-LN-oC a breakthrough innovation in metastasis diagnosis and drug testing, potentially offering personalized treatment relying on molecular characterization of lymph node metastasizing cells.
- The platform will make it possible to:



monitor the lymph node metastasis process in real time characterise signalling ques facilitating such metastasis,

identify spectral + molecular signatures in metastasizing cells



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Project objectives



The project will significantly advance the fields of **microfluidics, cell biology, cancer biology, physics, and computer programming and software development,** by pursuing the following objectives:

To develop **robust, automated microfluidic chips** optimized for tumor cell and LN culture enabling the study of their crosstalk,



To integrate **Quantum Cascade Laser based mid- IR spectroscopy** for specific chemical signatures,

To **molecularly characterize** both migrating tumor-derived cells attracted to the LN and the soluble signals driving migration,





To demonstrate an **advanced image analysis and signal processing platform** using a micro-optics module and deep learning algorithms to monitor cells,

To **integrate all Tumor-LN-oC technologies in an automated platform** prototype compatible with existing laboratory equipment,



To demonstrate the **Tumor-LN-oC TRL5 platform** and validate it with real patient samples,

To establish regulatory pathways and assure **regulatory compliance** to facilitate exploitation and early market entry.







6 technological modules





Project Concept



- The microfluidic chip has 2 separate chambers for Tumor and LN tissue, connected via microfluidic channels which will facilitate cell migration
- **Microfluidic peripherals** provide the flow mimicking the tumor microenvironment and lymphatic system
- Advanced laser bioprinting technology allows precise placement of tumor cells and organoids in the chip
- The **micro-optics module** makes it possible to visually monitor migrating cells in the channel in real time
- Mid-IR Photothermal spectroscopy facilitates molecular characterisation of migrating cells
- Automatic image analysis and signal processing aid in the interpretation of results



Project impact



- The identification of novel biomarkers characterizing metastatic can be exploited therapeutically and enable the use of existing drugs, or the development of new ones that could inhibit tumor growth and dissemination
- The spectral "fingerprint" of migrating/metastasizing cells can be used for diagnostic purposes in tumor and lymph node biopsies
- The platform will serve as a preclinical setting for parallel testing of drugs for individual cancer lung cancer patients.
- The proposed technologies will provide added value to the EU cancer diagnostics and pharmaceutical industries
- The project will lower the barriers associated with the application of OoC technology in disease diagnosis and therapy.



The Tumor-LN-oC platform will enable the development of novel metastasis-related diagnostics tools and novel therapies in a more targeted and personalized manner.

Expected results







Added value to the EU cancer diagnostics and pharmaceutical industries

Development of novel diagnostic tools and therapeutic approaches

> Verifiable progress in the application of Organ-on-Chip technologies for invitro research

Lower barriers associated with the application of OoC technology in disease diagnosis and therapy

Reduction of the need for animal and clinical testing

Increased awareness and knowledge about medical regulatory policies and requirements, especially by academics and SMEs









Project progress



- Specifications of the Tumor-LN-oC platform completed
- First iteration of the microfluidic chips fabricated
- Tests of microfluidic chip ongoing (co-culturing in static and flow conditions), continuous improvements
- Prototype for low-leak pressure controller developed
- Mechanical design of MIR laser source completed
- Initial setup of micro optics module completed
- Image analysis and signal processing platform established
- Machine learning algorithms
- Experimental Setup of MIR spectroscopy ongoing
- Successful bioprinting process on various membranes
- Regulatory roadmap finalised
- Dissemination strategy and initial commercialisation plan prepared, implementation ongoing



Get in touch!



CONTACTS

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OVERVIEW

Tumor-LN-oC proposes the development and validation of a TRL 5 tumor-lymph node-on-chip platform composed of 3D tissue models and microfluidic chips which will connect surgically removed human primary tumors and lymph node tissue (LN) from the same lung cancer patient. This will serve as a "biological twin" of the patient and will allow Tumor-LN-oC consortiun to study the interaction of primary tumors with LNs for individual patients.

OBJECTIVES

The general objective is to generate a Tumor-LN-oC platform that will allow the monitoring of the LN metastasis process, the characterization of signalling ques facilitating LN metastasis and the identification of spectral and molecular signatures in metastasizing cells. This information could lead to novel diagnostic tools and therapeutic approaches. Moreover, the developed platform will serve as a preclinical setting for parallel testing of drugs for individual cancer lung cancer patients.



Thank you!

Any questions?

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