3D Lung Tumor Spheroids for Oncoimmunological Assays

Commonly used acronym: Lung tumor spheroids


Contact person

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Organisation

Name of the organisation Vrije Universiteit Brussel (VUB)
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Country Belgium
Geographical Area Brussels Region

Partners and collaborations

Institut Curie

SCOPE OF THE METHOD

<table>
<thead>
<tr>
<th>The Method relates to</th>
<th>Animal health, Human health</th>
</tr>
</thead>
<tbody>
<tr>
<td>The Method is situated in</td>
<td>Basic Research, Translational - Applied Research</td>
</tr>
<tr>
<td>Type of method</td>
<td>In vitro - Ex vivo</td>
</tr>
<tr>
<td>Specify the type of cells/tissues/organs</td>
<td>murine lung cancer and fibroblast cell lines or human lung cancer and fibroblast cell lines</td>
</tr>
</tbody>
</table>

DESCRIPTION
Method keywords

lung cancer spheroids
mammary cells
tumour
mice
macrophage polarization
human cell-based model
fluorescence microscope
3D in vitro model
lentiviral reprogramming
immune cell migration
immunology
Immunotherapy
antigen specific T cell killing
cancer-associated fibroblasts
histocompatible

Scientific area keywords

Immunology
Lung cancer
T cell immunity
macrophage polarization
spheroids
immune checkpoint inhibitors
non-small cell lung cancer
tumor stroma
cancer-associated fibroblasts
immunotherapy

Method description

Lung cancer thrives in a complex multicellular tumor microenvironment (TME) that impacts tumor growth, metastasis, response, and resistance to therapy. While orthotopic murine lung cancer models can partly recapitulate this complexity, they do not resonate with high-throughput immunotherapeutic drug screening assays. To
address the current need for relevant and easy-to-use lung tumor models, a protocol is established to generate and evaluate fully histocompatible murine and human lung tumor spheroids, generated by co-culturing lung fibroblasts with tumor cells in ultra-low adherence 96-well plates. A spheroid generation protocol with the murine KrasG12D-p53ko (KP) and Lewis Lung Carcinoma (LLC) cell lines is delivered next to the human lung H1650 adenocarcinoma line. In addition, their application potential to study tumor-stroma organization, T-cell motility, and infiltration as well as distinct macrophage subsets’ behavior using confocal microscopy is described. Finally, a 3D target-specific T-cell killing assay that allows spatiotemporal assessment of different tumor to T-cell ratios and immune checkpoint blockade regimens using flow cytometry and live cell imaging is described. This 3D lung tumor spheroid platform can serve as a blueprint for other solid cancer types to comply with the need for straightforward murine and human oncoimmunology assays.

**Lab equipment**

For generation:
- Laminar air flow;
- Ultra-low adherence (ULA) or cell-repellent U-shaped 96-well plates;
- Biosafety level 2 facility for lentiviral vector production and handling of patient-derived T cells;
- Irradiator for stimulation of T cell feeder cells.

For evaluation:
- Flow cytometry;
- Microscopy (confocal, incucyte technology, ...);
- NGS methods, ...

**Method status**

Published in peer reviewed journal

**PROS, CONS & FUTURE POTENTIAL**

**Advantages**
- Straightforward murine and human multicellular lung tumor spheroid platform that recapitulates the characteristic tumor islet in stroma architecture found in human non-small lung cancer biopsies;
- Platform suitable for macrophage and T-cell infiltration studies;
- Platform suitable for target tumor cell-specific killing evaluation in framework of fundamental and translational immunotherapy research;
- In 96-well format for evaluation on conventional imaging instruments;
- Possibility to perform high-throughput spatiotemporal gene, cell, and/or drug screening;
- Can serve as a blueprint for other solid cancer types to comply with the need for straightforward murine and human oncoimmunology assays.

Challenges

- Generation or access to histocompatible fluorescently-labelled tumor cell lines and tissue-specific fibroblasts;
- Access to histocompatible target specific T cell clones and/or myeloid cells;
- Not every lung tumor cell line is suitable for generation of lung tumor spheroids with relevant tumor-stroma islet architecture.

Future & Other applications

Our 3D lung tumor spheroids can serve as a blueprint for other solid cancer types to comply with the need for straightforward murine and human oncoimmunology assays. Furthermore, we are currently expanding our expertise by optimizing the generation of lung cancer patient-derived organotypic cultures.

REFERENCES, ASSOCIATED DOCUMENTS AND OTHER INFORMATION

Associated documents


Links

LinkedIn profile Cleo Goyvaerts