# **EV-MIME**

Use of therapeutic extracellular vesicles as advanced treatment for metastatic triple negative breast cancer and pancreatic cancer.



L'AVENIR EST FAIT D'AUDACE

### **PRESENTATION**

Extracellular Vesicles (EVs) are produced from HEK293T cells that highly express NFAT3 transcription factor that inhibits cancer cell motility and are loaded with a combination of miRNAs inhibiting tumor growth and cell motility. *In vitro* evaluation revealed that these EVs significantly (80%) decrease invasive capacity of triple negative breast (MDA-MB-231, SUM-59PT) and pancreatic (BXPC3, MIA-PACA-2) cancer cell lines. These results were confirmed *in vivo* in a triple negative breast cancer mouse model.

## **APPLICATIONS**

- Adjuvant therapy for triple negative breast cancer or pancreatic cancer as a single agent or in combination with other drugs
- Neoadjuvant therapy for triple negative breast cancer or pancreatic cancer as a single agent or in combination with other drugs

### **COMPETITIVE ADVANTAGES**

Mostly, the potential competitors' approaches are solely at the early stage (proof of concept (PoC) to preclinical stages) and absence of available data did not allows direct comparison with this product.

## INTELLECTUAL PROPERTY

Two patent applications: WO2017167788A1 and WO2022136226A1

### CONTACT

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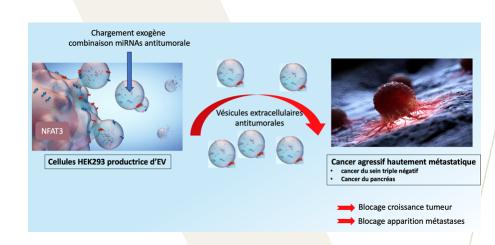
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Ref. project: 637

Extracellular Vesicles - NFAT3 - miRNA

Triple negative breast cancer - Pancreatic cancer



### **DEVELOPMENT PHASE**

- Completed *in vivo* PoC for EVs derived from HEK293T overexpressing NFAT3 in an athymic nude mouse model xenografted with a triple negative breast cancer line (MDA-MB—231) (TLR5)
- Ongoing *in vivo* PoC for EVs loaded with an anti-tumor combination of miRNAs in an athymic nude mouse model xenografted with a triple negative breast cancer line (MDA-MB—231) (TLR4)

### **PUBLICATIONS**

- de Camargo LCB *et al.* Sci Rep 2020 (10), 8964.
- Fougère M. *et al.* Oncogene. 2010 (15), 2292-301.
- Coillard L. *et al.* Front Oncol. 2022 (12), 804868.
   Erratum in: Front Oncol. 2022 (12), 1016189.
- de AKA S. *et al*. Adv Drug Deliv Rev. 2021 (179), 114001.

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