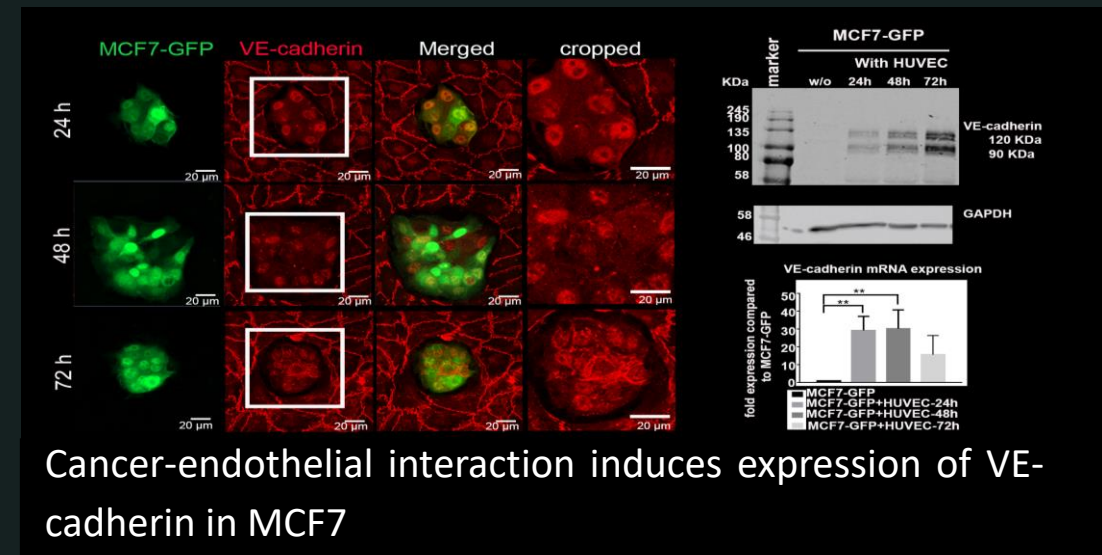
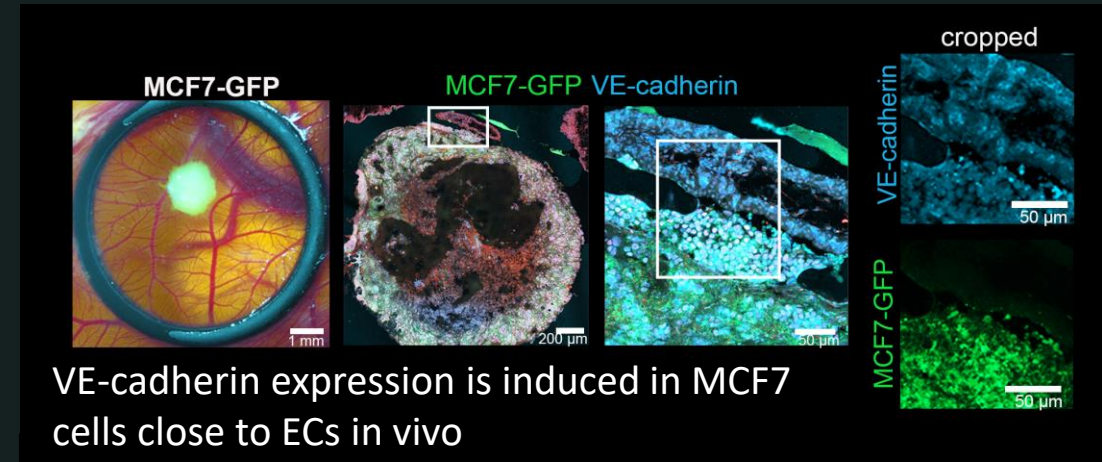
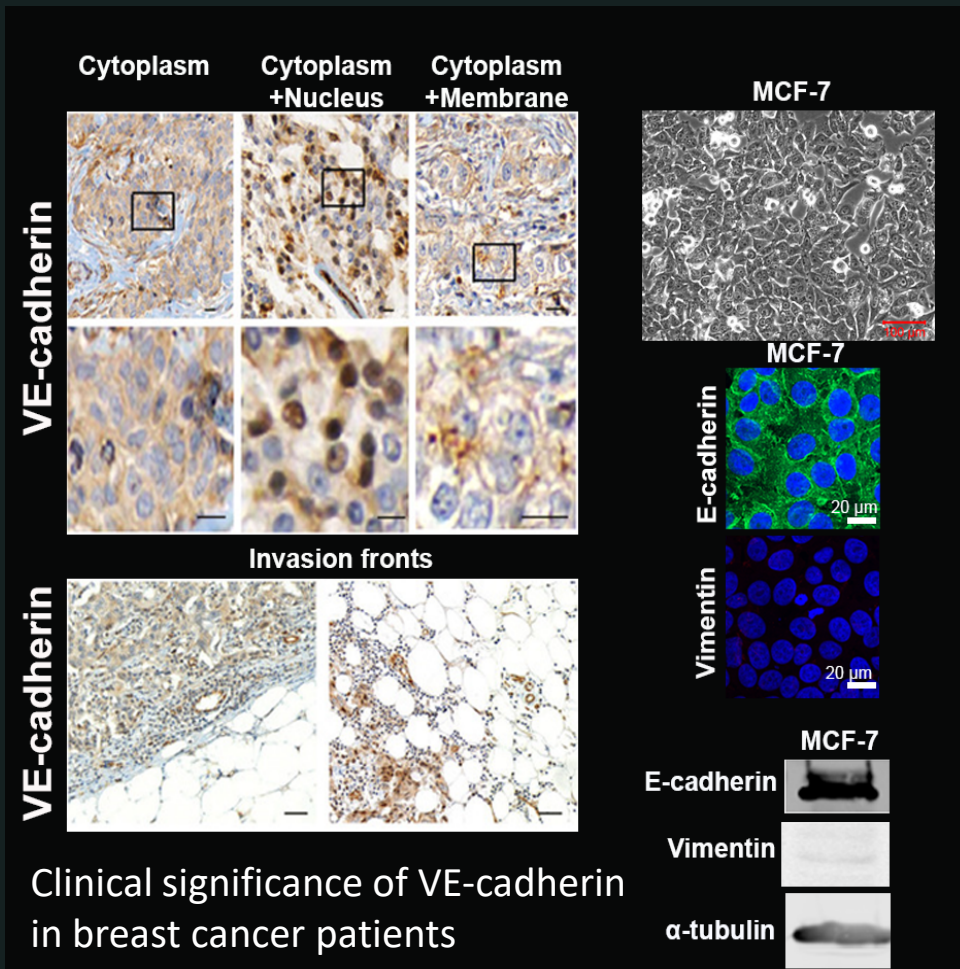


# A functional phenomics approach reveals the exchange of material between breast cancer and endothelial cells at single-cell resolution in a 3D co-culture system - Background

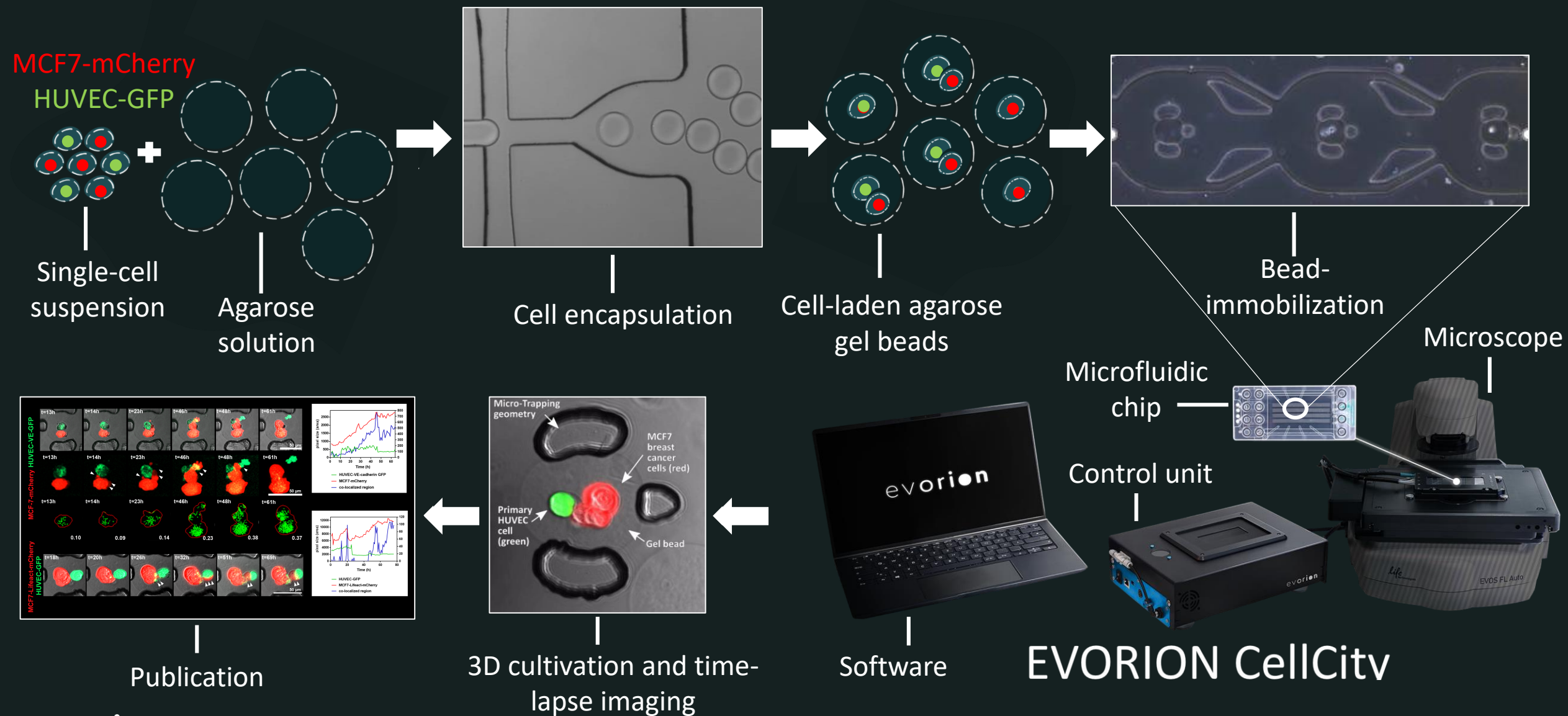
Maryam Rezaei<sup>1</sup>, Franziska S. Bockeloh<sup>1</sup>, Hans K-Brüggeney<sup>1</sup>, Johannes A. Eble<sup>2</sup>, Robert Weingarten<sup>1</sup>

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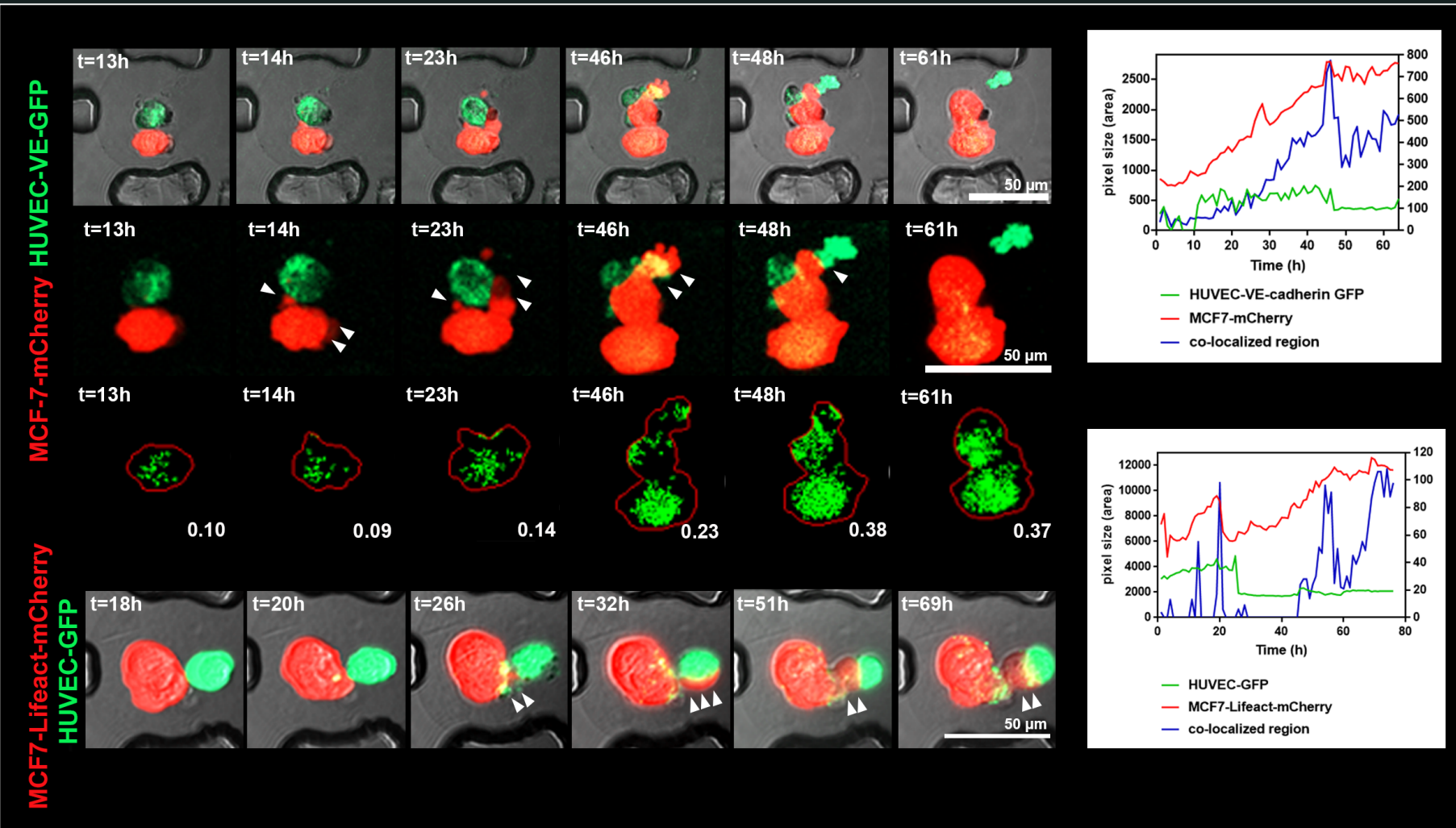


# A functional phenomics approach reveals the exchange of material between breast cancer and endothelial cells at single-cell resolution in a 3D co-culture system - Workflow



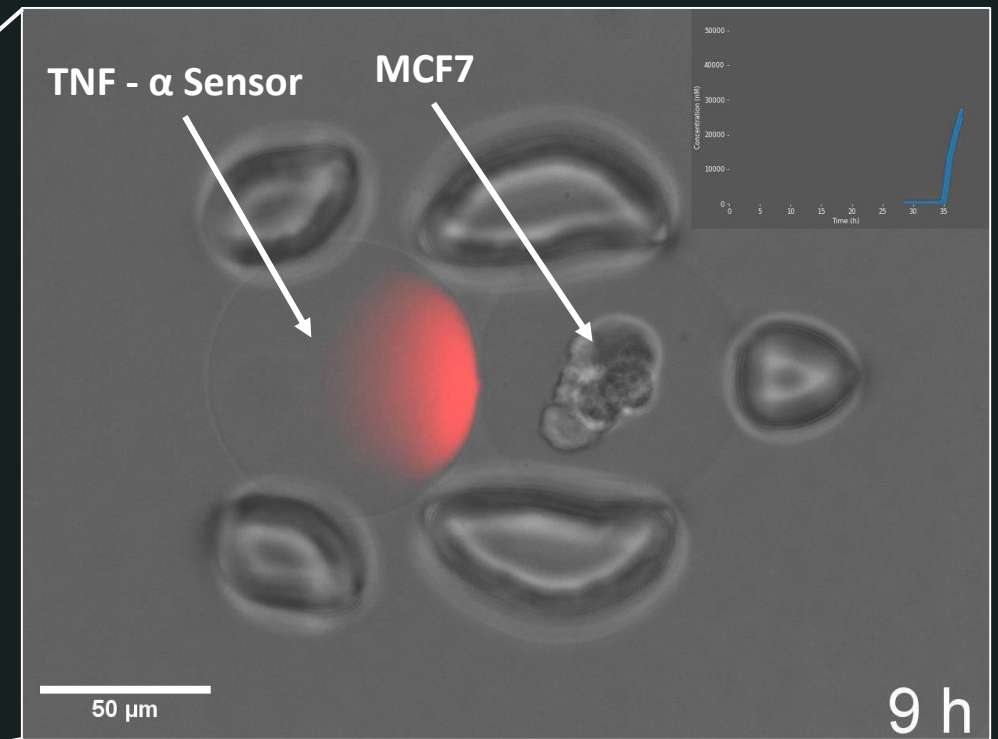
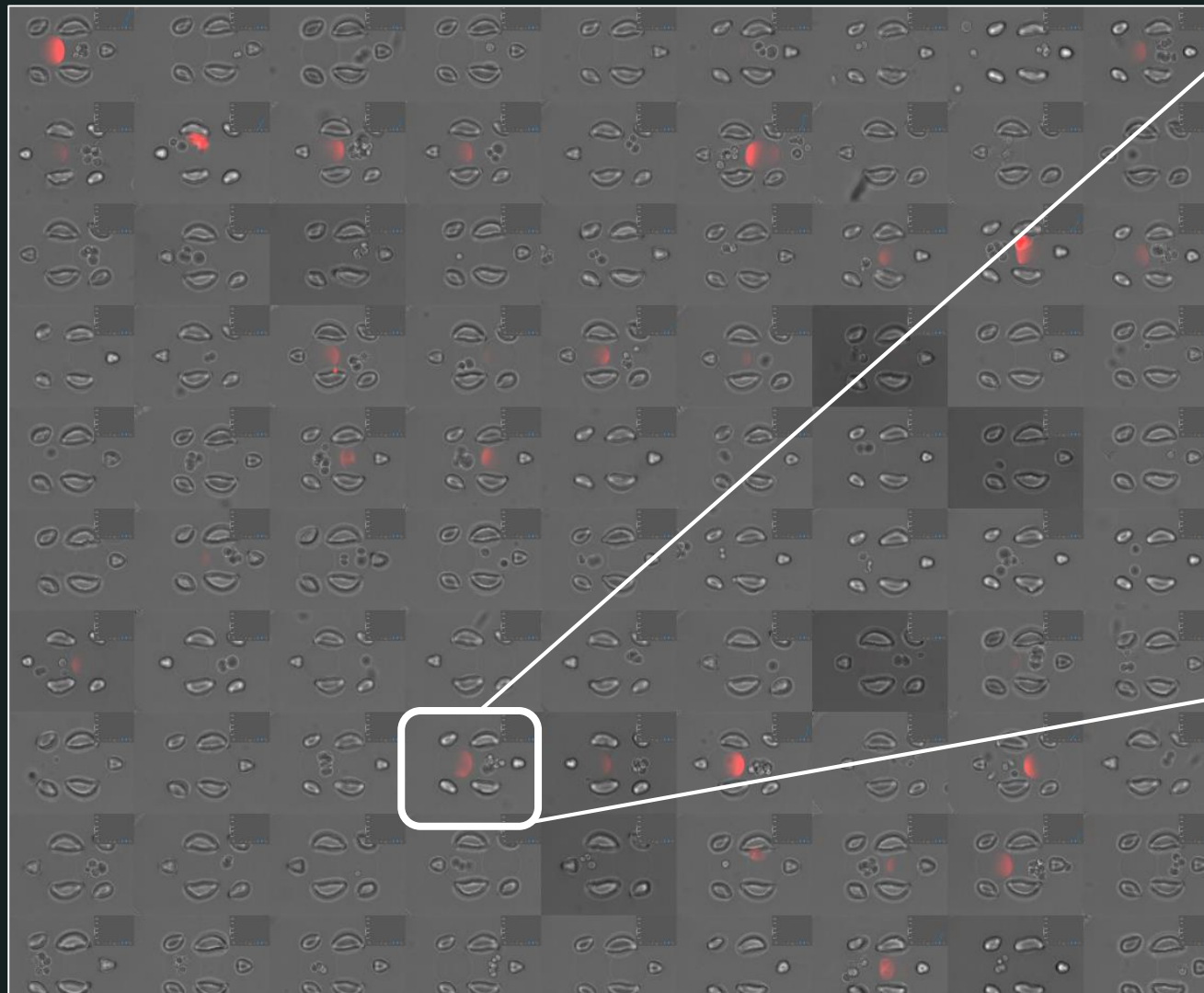
# A functional phenomics approach reveals the exchange of material between breast cancer and endothelial cells at single-cell resolution in a 3D co-culture system - Results

Live cell analysis of cell-cell interaction at single-cell resolution in 3D environment



- Fluorescent areas of mCherry (MCF7 cells), GFP (HUVEC-expressing VE-cadherin-GFP or cytoplasmic GFP) and of both fluorophores in colocalization were quantified time-dependently.
- MCF7 cancer cells form lamellipodia after 32h of co-culturing for phagocytic clearance
- Area of colocalized mCherry and GFP fluorescence increased over time due to the increasing number of HUVEC-derived, GFP-containing vesicles inside MCF7 cells

# A functional phenomics approach reveals the exchange of material between breast cancer and endothelial cells at single-cell resolution in a 3D co-culture system - Outlook



- Phagocytic clearance of dying cells (efferocytosis) is a mechanism by which MCF7 cells could take up extracellular vesicles from dying HUVECs
- EVORION Cytokine Sensors could support the hypothesis that TNF- $\alpha$  secretion by MCF7 cells could induce a targeted necroptosis and phagocytic clearance of HUVECs