

NANOSTACKS™: novel platforms for Head and Neck Cancer Research



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Background: The FDA Modernization Act 2.0 aims to decrease animal testing significantly and support NAM approaches such as micro-physiological systems (organ-on-a-chip OOAC) to improve preclinical drug discovery. OOAC platforms have recently emerged as a new generation of biomimetic alternatives for initial drug screening. Revivocell platform NANOSTACKS™ is used first time in the development of accurate 3D models of head and neck cancer using human cell lines to understand critical aspects of the tumour microenvironment.

NANOSTACKS™





Key aspects of NANOSTACKS™

Transparent, stackable insertsready-to-use platforms for **3D Cell Cultures**

SBS-standard 24-well plate friendly – **Preclinical Drug Screening**

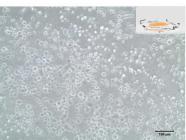
Up to 4 NANOSTACKS™ per well – Organ Modelling

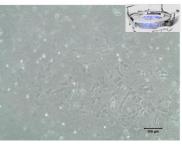
Porous membranes facilitate media circulation – excellent mimic for Microenvironment

Head and neck cancer 3D model

NANOSTACKS™ were used to seed cancer cells from both primary and metastasis sites, and fibroblasts (collected from the same patient) and co-cultures were grown for 72h.







Day 1 (seeding after 2h on NANOSTACKS™

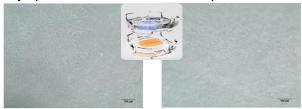
Cancer cells from the primary site of head and neck cancer (P) Cancer Cells from the Primary Site of head and neck cancer (P) Cancer Cells from the Primary Site of from

100 pm

Cancer cells from the Metastasis site from the same patient (M)



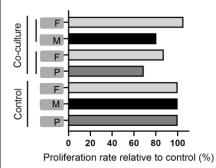
Normal Fibroblasts (F)



Cancer cells from the primary site with fibroblasts (P+F)



Cancer cells from the metastasis site with fibroblasts (M+F)



Proliferation comparison between the controls and co-culture using MTS assay

The proliferation of P and M are lower compared to control when co-culture with fibroblasts, although M cells have slightly higher proliferation (80.44%) compared to P (68.92%)

Fibroblasts co-cultured with P have lower proliferation (87.33%) than the control.

Fibroblasts co-cultured with M have higher proliferation (105.32%) than the control.

The POC study proves that **NANOSTACKS™** can be the best platform to understand the tumour microenvironment and provide essential cues to direct and control tumour progression. Further research plans involving cancer-associated fibroblasts and their implications in cancer malignancy, such as angiogenesis, invasion, and therapy resistance to be studied on **NANOSTACKS™**. Revivocell OOAC platforms will have a tremendous advantage in patient-specific disease modelling in the near future.

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