Marrella et al.: 3D Fluid-Dynamic Ovarian Cancer Model Resembling Systemic Drug Administration for Efficacy Assay

Supplementary Data

Supporting Information



Fig. S1: Schematic representation of the 3D domains used during the simulations (A) MIVO[®] bioreactor set-up in dynamic condition; (B) geometrical configuration used for the simulations in static condition within the well.





Values are reported as mean ± SEM.

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Fig. S3: Kinetics of cisplatin consumption within the hydrogel Comparison between the theoretical model and the experimental data, where R is the reaction term defined according to the Michaelis-Menten kinetics (N = 3 biological replicates).



Fig. S4: Fluorescence images showing immunostaining of Ki67 (green) as index of proliferation and caspase-3 (red) as marker of apoptosis of SKOV-3 cultured within alginate hydrogels without drug and cultured in static and in dynamic conditions with cisplatin 100 μM.

Cells were stained after 2 or 7 days and counter-labeled with DAPI (blue). Untreated controls were cultured in static conditions. Scale bar is 500 μ m. (N = 3 biological replicates; n = 2 technical replicates).







Fig. S6: Cell viability of SKOV-3 cultured in 2D conditions treated with 10 μ M cisplatin assessed by Alamar Blue assay Cell viability was derived as % of live cells normalized to untreated controls. Values are reported as mean ± SD (N = 3 biological replicates; n = 2 technical replicates).