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**Predicting *in vivo* treatment responses using an *in vitro*-calibrated mathematical model**

**Abstract**

Mathematical models, and their corresponding *in silico* experiments, can be used to simulate both *in vitro* and *in vivo* tumour scenarios. However, the microenvironment in an *in vitro* cell culture is significantly different from the microenvironment in a solid tumour and many details that influence tumour dynamics differ between *in vitro* and *in vivo* settings. These details include cell proliferation, oxygen distribution and drug delivery. It follows that translating quantitative *in vitro* findings to *in vivo* predictions is not straightforward.

In this talk I will present an individual based mathematical cancer model in which one individual corresponds to one cancer cell. This model is governed by a few observable and well-documented principles, or rules. To account for differences between *in vitro* and *in vivo* scenarios, these rules can be appropriately adjusted. By only adjusting the rules (whilst keeping the fundamental framework intact), the mathematical model can first be calibrated by *in vitro* data and thereafter be used to successfully predict treatment responses in mouse xenografts *in vivo*.