



Mechanisms of cancer invasion and progression: insights from cellular automaton models

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Tumour invasion and progression may be viewed as collective phenomena emerging from the interplay of biological cells with their environment. Cell-based mathematical models in which cells are regarded as separate discrete entities can be used to decipher the rules of interaction. Here, we focus on the dynamics of glioma and breast cancer.

We introduce lattice-gas cellular automaton models [1, 2] to analyse the role of phenotypic plasticity in cancer invasion, define spatial and non-spatial Moran processes to shed light on the size of the tumour originating niche, and adopt Markov chain models to investigate the origin of genetic heterogeneity in glioblastoma [3, 4, 5].

References

- [1] A. Deutsch, S. Dormann, *Cellular Automaton Modeling of Biological Pattern Formation: Characterization, Examples, and Analysis*, Birkhäuser, Basel, 2005.
- [2] A. Deutsch, J. M. Nava-Sedeño, S. Syga, H. Hatzikirou, BIO-LGCA: a cellular automaton modelling class for analysing collective cell migration, *PLOS Computational Biology*, 17(6):e1009066, 2021.
- [3] T. Buder, A. Deutsch, B. Klink, A. Voss-Böhme, Patterns of tumor progression predict small and tissue-specific tumor-originating niches, *Frontiers in Oncology*, 8:668, 2018.
- [4] A. Dirkse, A. Golebiewska, T. Buder, . . . , A. Deutsch, A. Voss-Böhme, S. P. Niclou, Stem cell-associated heterogeneity in Glioblastoma results from intrinsic tumor plasticity shaped by the microenvironment, *Nature Communications*, 10(1):1787, 2019.
- [5] O. Ilina, P. G. Gritsenko, S. Syga, . . . , J. A. Käs, A. Deutsch, P. Friedl, Cell-cell adhesion and 3D matrix confinement determine jamming transitions in breast cancer invasion, *Nature Cell Biology*, 22:1103-1115, 2020.