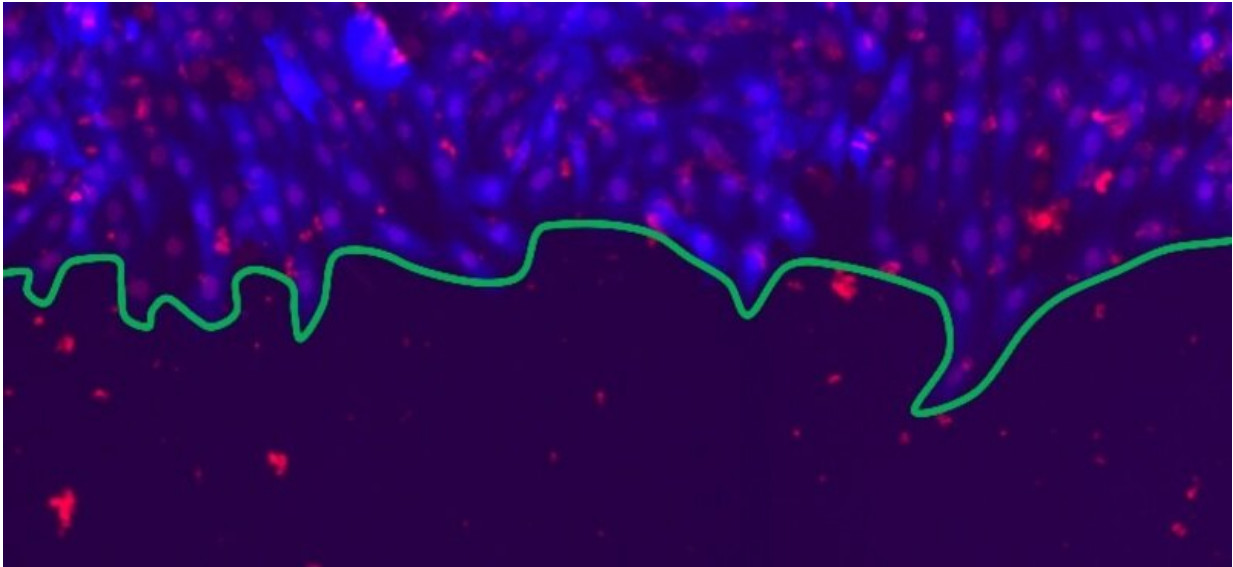


# A physics perspective on wound healing

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Fluorescent microscopy image of a proliferating cell front, whose edge is indicated in green. The rat epithelial cells, with cytoplasm (blue) and nuclei (red), form a 2-dimensional culture which can be used to model wound healing. Credit: © Guillaume Rapin, UNIGE

In material physics understanding how systems interact across the interfaces separating them is of central interest. But can physical models clarify similar concepts in living systems, such as cells? Physicists at the University of Geneva (UNIGE), in collaboration with the University of Zurich (UZH), used the framework of disordered elastic systems to study the process of wound healing—the proliferation of cell fronts which eventually join to close a lesion. Their study identified the scales

of the dominant interactions between cells which determine this process. The results, published in the journal *Scientific Reports*, will allow better analysis of cell front behavior, in terms of both wound healing and tumor development. In the future, this approach may offer personalized diagnostics to classify cancers and better target their treatment, and identify new pharmacological targets for transplantation.

By focusing on macroscopic properties of large datasets, statistical physics makes it possible to extract an overview of system behavior independent of its specific microscopic character. Applied to biological elements, such as the cell fronts bordering a wound, this approach makes it possible to identify the various interactions which play a defining role during tissue growth, differentiation, and healing, but above all to highlight their hierarchy at the different scales observed. Patrycja Paruch, professor in the Department of Quantum Matter Physics at the UNIGE Faculty of Science, explains: "For cancer tumor invasion, or in the event of a wound, cell front proliferation is crucial, but the speed and morphology of the front is highly variable. However, we believe that only a few dominant interactions during this process will define the dynamics and the shape—smooth or rough, for example—of the cell colony edge. Experimental observations across multiple lengthscales to extract general behaviors can allow us to identify these interactions in healthy tissue and diagnose at what level pathological changes can occur, to help combat them. This is where [statistical physics](#) comes in."

## The many scales of wound healing

In this multidisciplinary study, the UNIGE physicists collaborated with the team of Professor Steven Brown from the UZH. Using rat epithelial [cells](#), they established flat colonies (2D) in which the cells grow around a silicone insert, subsequently removed to mimic an open lesion. The cell fronts then proliferate to fill the opening and heal the tissue. "We reproduced five possible scenarios by 'handicapping' the cells in

different ways, in order to see what impact this has on wound healing, i.e. on the speed and roughness of the cell front," explains Guillaume Rapin, a researcher in Patrycja Paruch's team. The idea is to see what happens in normal healthy tissue, or when processes such as cell division and communication between neighboring cells are inhibited, when cell mobility is reduced or when cells are permanently pharmacologically stimulated. "We took some 300 images every four hours for about 80 hours, which allowed us to observe the proliferating cell fronts at very different scales," continues Guillaume Rapin. "By applying high-performance computational techniques, we were able to compare our [experimental observations](#) with the results of numerical simulations," adds Nirvana Caballero, another researcher in Patrycja Paruch's team.

## Zooming out for greater effect

The scientists observed two distinct roughness regimes: at less than 15 micrometers, below the size of a single cell, and between 80 and 200 micrometers, when several cells are involved. "We have analyzed how the roughness exponent evolves over time to reach its natural dynamic equilibrium, depending on the pharmacochemical conditions we have imposed on the cells, and how this roughness increases depending on the scale at which we look," emphasizes Nirvana Caballero. "In a system with a single dominant interaction, we expect to see the same roughness exponent at all scales. Here, we see a changing roughness if we look at the scale of one cell or of 10 cells."

The Geneva and Zurich teams revealed only minor variations in the roughness exponent below 15 micrometers, whatever the conditions imposed on the cell fronts. On the other hand, they found that between 80 and 150 micrometers, the roughness is altered by all pharmacological inhibitors, significantly reducing the [roughness](#) exponent. Moreover, they observed that proliferation speed varied greatly between the different pharmacochemical conditions, slowing when cell division and

motility were inhibited, and accelerating when cells were stimulated. "More surprisingly, the fastest proliferation speed was achieved when gap-junction communication between cells was blocked," says Guillaume Rapin. This observation suggests that such communication may be targeted in future therapies, either to promote healing of burns or wounds, or to slow cancer tumor invasion.

These results show that medium-scale interactions play a crucial role in determining the healthy proliferation of a cell front. "We now know at what scale biologists should look for problematic behavior of cell fronts, which can lead to the development of tumors," says Nirvana Caballero. Now scientists will be able to focus on these key lengthscales to probe tumor cells fronts, and directly compare their pathological interactions with this of healthy cells.

**More information:** Guillaume Rapin et al. Roughness and dynamics of proliferating cell fronts as a probe of cell–cell interactions, *Scientific Reports* (2021). [DOI: 10.1038/s41598-021-86684-3](https://doi.org/10.1038/s41598-021-86684-3)

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