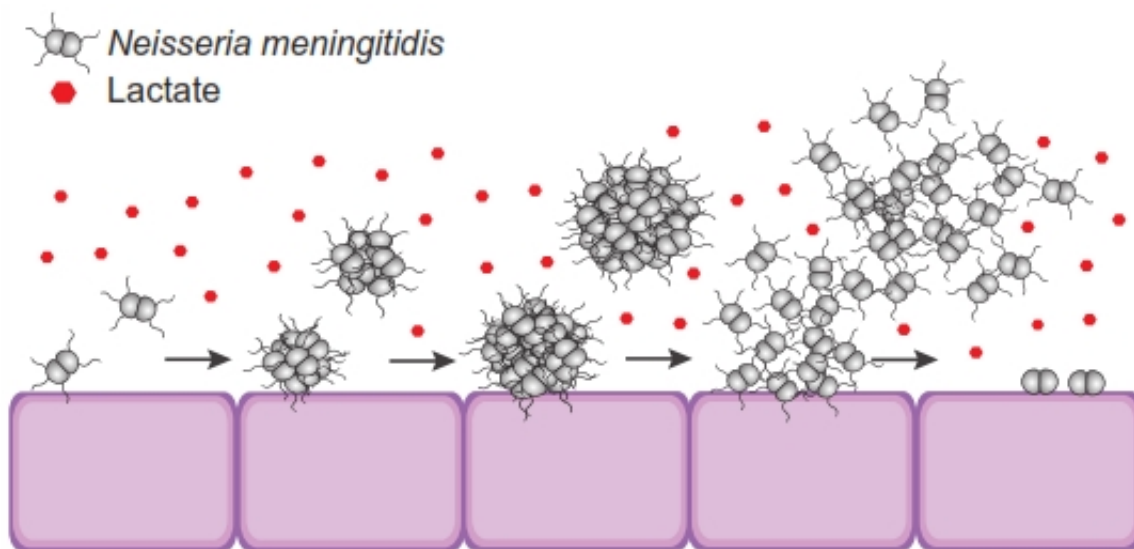


# Lactate from human cells may trigger key step in invasion by meningitis-causing bacteria

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Environmental lactate triggers microcolony dispersal. Credit: Sigurlásdóttir S, et al. (2017)

Lactate produced in the upper throat might trigger meningitis-causing bacterial cells to detach from tiny colonies and spread within the body, according to a new study published in *PLOS Pathogens*.

The research, performed by Sara Sigurlásdóttir and colleagues at Stockholm University in Sweden, focused on the bacterium *Neisseria meningitidis*, which lives in the [upper respiratory tract](#) of 10 to 35

percent of humans. Normally, *N. meningitidis* is harmless, but it can sometimes invade the body and cause meningitis or blood infection.

The initiation of harmful *N. meningitidis* infection is known to follow two steps: First, the bacteria gather in microscopic colonies in the upper throat. Then these microcolonies disperse rapidly, allowing the bacteria to invade the mucosal lining of the throat and enter the blood. However, this dispersal process is poorly understood.

Sigurlásdóttir and her team directed by professor Ann-Beth Jonsson hypothesized that *N. meningitidis* microcolony dispersal is triggered by a substance produced in [human cells](#). To investigate, they used live-cell imaging to show that *N. meningitidis* microcolonies do not need to be in direct contact with human cells in order to disperse; being grown in a cocktail of substances derived from human cells was enough to cause dispersal.

Further experiments narrowed down the possibilities of which specific molecule triggers dispersal. Lactate, a byproduct of glucose fermentation in human cells, emerged as the top candidate. The researchers confirmed that lactate can initiate microcolony dispersal and showed that it is not metabolized by *N. meningitidis*, suggesting that it serves solely as a signaling molecule.

Based on these findings, the scientists propose that *N. meningitidis* microcolonies disperse in response to high concentrations of lactate in their environment. Further research into lactate's role in *N. meningitidis* infection could improve understanding of how harmless *N. meningitidis* bacteria become invasive.

"It will be intriguing to identify the mechanistic details how [lactate](#) serves as signaling molecule. This may not only help to improve understanding of how [harmless bacteria](#) become invasive, but also reveal

novel ways to prevent and treat meningococcal disease." says Ann-Beth Jonsson, professor at Stockholm University.

**More information:** Sigurlásdóttir S, Engman J, Eriksson OS, Saroj SD, Zguna N, Lloris-Garcerá P, et al. (2017) Host cell-derived lactate functions as an effector molecule in *Neisseria meningitidis* microcolony dispersal. *PLoS Pathog* 13(4): e1006251. [DOI: 10.1371/journal.ppat.1006251](https://doi.org/10.1371/journal.ppat.1006251)

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