

Title: Fighting bacteria using innovative eradication techniques

Synopsis:

In our lab we are committed to fight bacteria that are serious health threats. Most of these bacteria shelter in biofilms, i.e. communities of bacteria inserted in a matrix of biomacromolecules they produce.

Chasing bacteria in these shelters is a demanding endeavor as innovative antibiotics are needed: able to diffuse in the crowded biomolecular medium and able to kill bacteria, even if they are in a dormant state.

We are seeking for enthusiastic and focused students with determination to help to develop a new generation of antibiotics.

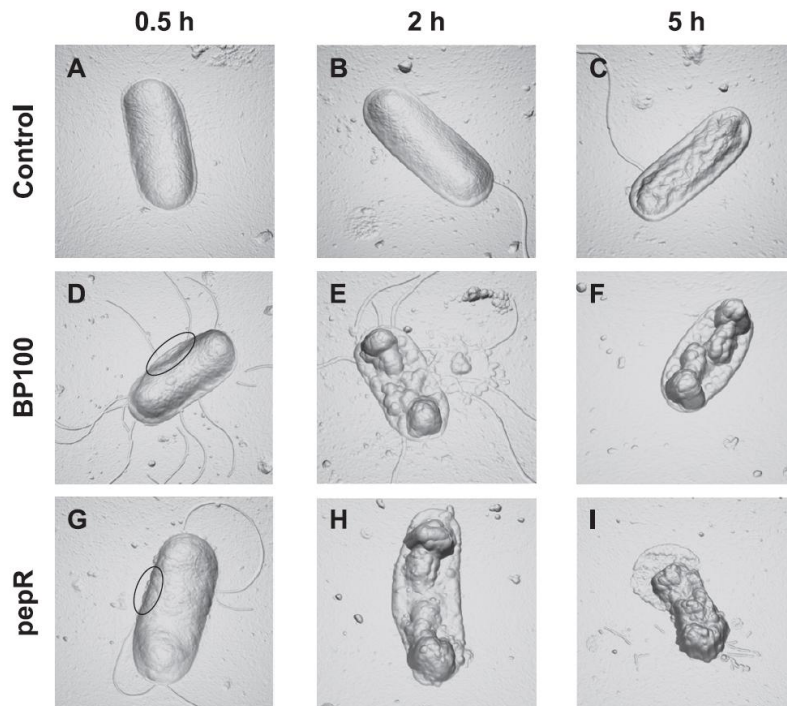


FIGURE 3. Time dependence of AMP effects on *E. coli* imaged by AFM. A–I, three-dimensional orthogonal projection images (derived from the height data) of untreated *E. coli* cells (top row), and *E. coli* cells treated with 3 μM BP100 (middle row), and 5 μM pepR (bottom row). Images were acquired following the treatment of the bacterial cells for 0.5 h (first column), 2 h (second column), and 5 h (third column). Total scanning area for each image: $4 \times 4 \mu\text{m}^2$. See the text for a description of the highlighted areas.

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Bibliography:

<https://www.sciencedirect.com/science/article/pii/S0021925819890841>

<https://academic.oup.com/jac/article/74/9/2617/5498604?login=true>