

Irene Chen and Heather Maynard, UC Los Angeles

What was achieved?:

In this study, polymer chemistry was used to create peptide-mimicking macromolecules with hydrophobic and cationic side chains that can kill bacteria by destabilizing their membranes. The macromolecules were prepared by radical copolymerization of caffeine methacrylate as the hydrophobic monomer and various cationic- or zwitterionic-methacrylate monomers. Copolymers bearing carboxybetaine as the cation exhibited antibacterial activity against both Gram-positive (*S. aureus*) and Gram-negative bacteria (*E. coli*), including methicillin-resistant clinical isolates. The synthesized copolymers also demonstrated good biocompatibility with mouse embryonic fibroblast cells and hemocompatibility with erythrocytes.

Why is it important?:

Antimicrobial resistance is a significant global health concern, and the study's findings offer a novel strategy for combating drug-resistant microbes. The use of caffeine and carboxybetaine as hydrophobic and cationic groups in copolymers could help prepare macromolecules with broad-spectrum antibacterial activity and low cytotoxic effects. These copolymers have potential applications in various fields, including biomedical devices and coatings, and can potentially address the urgent need for effective antimicrobial agents.

How did BioPACIFIC MIP enable this?

BioPACIFIC MIP provided financial support for materials and researchers through the Fellows program. This initiative emerged as a result of the dynamic interactions and collaboration between members of diverse research groups within one of the BioPACIFIC MIP SETs.

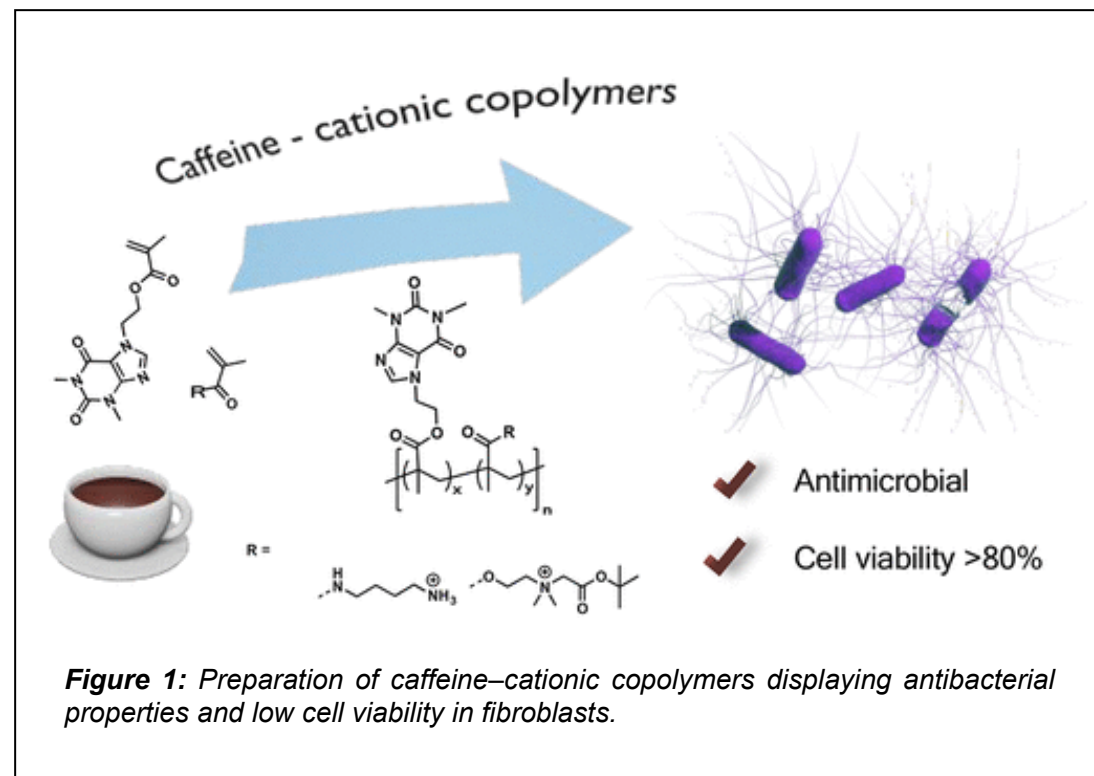


Figure 1: Preparation of caffeine–cationic copolymers displaying antibacterial properties and low cell viability in fibroblasts.

Publication: Pedro Salas-Ambrosio, Shelby Vexler, Rajalakshmi P S, Irene A. Chen, and Heather D. Maynard, ACS Bio Med Chem Au. <https://doi.org/10.1021/acsbiochemau.2c00077>