

# New anti-biofilm compounds show promise against drug-resistant bacteria linked to hospital infections

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Researchers at the University of Michigan Life Sciences Institute and School of Public Health have discovered a new class of anti-biofilm compounds derived from marine microorganisms that show promise against a drug-resistant bacterium commonly associated with hospital-acquired infections.

In cell cultures, the new compounds, known as cathitamylicins, are able to stop the *Acinetobacter baumannii* bacteria from gathering together in a sticky cluster known as a biofilm, according to findings scheduled to be published Feb. 16 in *Nature Communications*. Biofilms cling to surfaces and form complex structures that make them far more resistant to drugs than free-floating bacteria.

This property makes biofilms a deadly risk to patients in a hospital setting, where they can cling to medical devices, prosthetic implants and other surfaces, making them resistant to sterilization with traditional antimicrobial agents.

There are currently no drugs that specifically target [biofilm formation](#) on the market or in clinical trials, the researchers noted.

"This is why preventing biofilm formation is such an important research target," said Ashootosh Tripathi, a research fellow in the lab of LSI faculty member David Sherman. Tripathi was co-first author of the study, along with Sung Ryeol Park, a former postdoctoral research fellow in the Sherman lab.

Moreover, *A. baumannii*'s resistance to front-line antibiotics has been a growing concern worldwide and this bacterium can form a biofilm easily, which makes the treatment even more challenging, said co-senior author Chuanwu Xi, associate professor of [environmental health sciences](#) at the U-M

School of Public Health.

Treatment of (multidrug resistant *A. baumannii*) infections is a great challenge for physicians as is control of MDRAB spread, according to a recent article in the International Journal of *Environmental Research and Public Health*.

The new anti-biofilm agents were discovered by conducting high-throughput screening of compounds in the LSI's library of "natural product" extracts—which includes thousands of drug-like substances derived from [marine microorganisms](#) collected by Sherman and his collaborators during marine field collection expeditions in locations across the globe.

These particular compounds were derived from the bacterium *Streptomyces gandocaensis*, which was isolated from marine sediment collected on a 2007 expedition to Punta Mona Island in Costa Rica. The screening was conducted in the LSI's high-throughput lab, the Center for Chemical Genomics.

Once an extract was found that inhibited *A. baumannii*'s ability to form biofilms, the researchers conducted experiments to find its most potent forms; they also created two new more potent analogs.

Anti-biofilm drugs might be applied either directly to objects like medical implants, catheters or dental implants, or through systemic application, like antibiotics. Still in its early stages, the discovery will need further compound optimization and preclinical development before human clinical trials could be considered.

"This new class of biofilm inhibitors provides a foundation toward the development of safe and effective drugs to limit or prevent biofilm formation,"

said Sherman, co-senior author of the study and the Hans W. Vahlteich Professor of Medicinal Chemistry and associate dean for research and graduate education at the U-M College of Pharmacy, professor of chemistry in the College of Literature, Science, and the Arts, and of microbiology and immunology in the Medical School.

"As antibiotic resistance becomes an increasingly important global health concern, marine microorganisms have a great—and largely untapped—potential to provide new classes of antibiotics and anti-biofilm compounds."

Provided by University of Michigan

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