Antibiotics with novel mechanism of action discovered
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Many life-threatening bacteria are becoming increasingly resistant to existing antibiotics. Swiss researchers co-headed by the University of Zurich have now discovered a new class of antibiotics with a unique spectrum of activity and mechanism of action—a major step in the fight against antimicrobial resistance. By disrupting outer membrane synthesis, the antibiotics effectively kill Gram-negative bacteria.

The rapid emergence of antimicrobial resistance is a matter of global concern. According to the World Health Organization (WHO), particularly Gram-

negative bacteria like Acinetobacter baumannii, Pseudomonas aeruginosa and Enterobacteriaceae that are resistant to the carbapenem and cephalosporin antibiotics, pose a growing threat to human health. These pathogens can cause severe and often life-threatening infections. The last new class of antibiotics to reach the market against these microorganisms, the fluoroquinolones, dates back to the 1960s. Novel antibiotics with new mechanisms of action against Gram-negative bacteria are urgently needed, especially because resistance against the last resort antibiotic colistin is on the rise globally.

Swiss research teams headed by the University of Zurich (UZH) and Polyphor AG now report the discovery and characterization of a new family of synthetic antibiotics that possess broad-spectrum anti-Gram-negative antimicrobial activity. "The new antibiotics interact with essential outer membrane proteins in Gram-negative bacteria," says John Robinson from the UZH Department of Chemistry, who co-headed the study. "According to our results, the antibiotics bind to complex fat-like substances called lipopolysaccharides and to BamA, an essential protein of the outer membrane of Gram-negative bacteria," Robinson adds.

Disruption of outer membrane synthesis via unexploited mechanism of action

BamA is the main component of the so-called β-barrel folding complex (BAM), which is essential for outer membrane synthesis. After targeting this essential outer membrane protein, the antibiotics destroy the integrity of the bacterial membranes and the cells burst. The outer membrane of Gram-negative bacteria has the important function to protect the cells from toxic environmental factors, such as antibiotics. It is also responsible for the uptake and export of nutrients and signaling.
molecules. "Despite its critical importance, so far no clinical antibiotics target these key proteins required for outer membrane biogenesis," Robinson says.

**Lead molecule in preclinical studies**

The research was carried out in close collaboration with Polyphor AG, a former UZH start-up company that was founded in 1996. The clinical stage biopharmaceutical company based in Allschwil now plans to progress one compound into human clinical trials. "POL7306, a first lead molecule of the novel antibiotics class, is now in preclinical development," says Daniel Obrecht, chief scientific officer at Polyphor and co-head of the work.

This study is published in *Nature*.

**More information:** Chimeric peptidomimetic antibiotics against Gram-negative bacteria, *Nature*. DOI: 10.1038/s41586-019-1665-6

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