How superbug A. baumannii survives metal stress and resists antibiotics

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The deadly hospital pathogen Acinetobacter baumannii can live for a year on a hospital wall without food and water. Then, when it infects a vulnerable patient, it resists antibiotics as well as the body's built-in infection-fighting response. The World Health Organization (WHO) recognizes it as one of the three top pathogens in critical need of new antibiotic therapies.

Now, an international team, led by Macquarie University researchers Dr. Ram Maharjan and Associate Professor Amy Cain, have discovered how the superbug can survive harsh environments and then rebound, causing deadly infections. They have found a single protein that acts as a master regulator. When the protein is damaged, the bug loses its superpowers allowing it to be controlled, in a lab setting. The research is published in *Nucleic Acids Research*.

"We hope that our paper will encourage researchers worldwide to refocus on developing drugs to fight this superbug, which is spreading through the world's hospitals, and killing already vulnerable people in intensive care units and other high-risk areas," says Associate Professor Cain, the senior author on the paper.

There are six superbugs that scare global health officials. E. coli, Klebsiella pneumoniae and other gram-negative bacteria have common pathways that give them antibiotic resistance. A. baumannii is different. It's particularly tough, and it's one of the most resistant pathogens we encounter. Strangely, we don't know much about how it infects us.

**Breakthrough in a research challenge**

"In the lab we can see this pathogen is very tough. Other researchers have shown that you can desiccate the bug for a year and when they added water, it was still able to infect mice," says Associate Professor Cain.
"The problem had been that A. baumannii is relatively new on the scene, emerging as a problem in hospitals in the 1980s. And it's hard to genetically manipulate with the existing molecular biology toolkit. It usually only infects sick people, but it is very resistant to antibiotics making it incredibly hard to treat and difficult to safely research. So, we don't know much about it. We don't know where it came from, nor how it became so resistant and resilient. Now, thanks to this paper, we know how it deals with stress."

Amy and her colleagues realized about five years ago that they could make a difference by trying to understand the underlying biology of A. baumannii. That led to a major investment by Macquarie University in the research, in biocontainment laboratories for staff safety, and in an ethical animal model using moth caterpillars. The research effort has been strongly supported by the Australian Council and the National Health and Medical Research Council.

During infection our cells fight back by either flooding or starving bacteria of essential metals such as copper and zinc. A. baumannii has strong drug pumps that push antibiotics, metals and other threats out of the cell.

"By studying how this bug deals with infection stresses, we've found an important uncharacterized regulatory protein (DksA). When we disrupt this protein, it leads to changes in about 20 percent of the bug's genome and breaks its pumping system," says Dr. Ram Maharjan, a Macquarie University researcher and first author on the paper.

Not only does this protein control stress response, but it also controls virulence. A. baumannii usually spreads in blood but our disruption also caused it to be completely undetected in the blood of both Galleria mellonella and mice. It also becomes super sticky and harmlessly sticks to organs.

Provided by Macquarie University


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