

Answer to antibiotic-resistant infections could already be on the market

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Credit: University of Texas Medical Branch at Galveston

The rise of antibiotic resistant bacterial pathogens is an increasingly global threat to public health. In the United States alone antibiotic resistant bacterial pathogens kill thousands every year.

But non-antibiotic therapeutic drugs already approved for other purposes in people could be effective in fighting the antibiotic-resistant pathogens, according to a new study from researchers at The University of Texas Medical Branch at Galveston.

Antibiotic resistance is increasing due to the over prescription of

[antibiotics](#), said Ashok Chopra, a professor at UTMB and author of the new study in the *Journal of Antimicrobial Agents and Chemotherapy*. But the solution could lie with drugs originally meant for other uses that, until now, no one knew could also help combat bacterial infections.

While antibiotics have been highly effective at treating infectious diseases, [infectious bacteria](#) have adapted to them and antibiotics have become less effective, according to the Centers for disease Control and Prevention. About 2 million people in the United States are infected with [antibiotic resistant bacteria](#) every year and at least 23,000 die, according to the CDC.

"There are no new antibiotics which are being developed and nobody really has given much emphasis to this because everyone feels we have enough antibiotics in the market," Chopra said. "But now the problem is that bugs are becoming resistant to multiple antibiotics. That's why we started thinking about looking at other molecules that could have some effect in killing such antibiotic resistant [bacteria](#)."

By screening a library of 780 Food and Drug Administration approved therapeutics, Chopra, Jourdan Andersson, a graduate student at UTMB, and others on the research team were able to identify as many as 94 drugs that were significantly effective in a cell-culture system when tested against *Yersinia pestis*, the bacteria that cause the plague and which is becoming antibiotic resistant.

After further screening, three drugs, trifluoperazine, an antipsychotic, doxapram, a breathing stimulant, and amoxapine, an anti-depressant, were used in a mouse model and were found to be effective in treating plague. In further experiments, trifluoperazine was successfully used to treat *Salmonella enterica* and *Clostridium difficile* infections, both of which are listed as [drug](#)-resistant bacteria of serious threat by the CDC.

"It is quite possible these drugs are already, unknowingly, treating infections when prescribed for other reasons," Chopra said.

Since these are not antibiotics these drugs are not attacking the bacteria. Instead, they could be dealing with these bacteria in a couple of different ways, Chopra said.

The drugs could somehow be affecting the virulence of these bacteria - although in the case of plague the team found no evidence that the drugs were affecting the destructive strength of the plague-causing bacteria, Chopra said.

Or, the other likely explanation, the drugs are working through the host and could be affecting host proteins or genes so that the bacteria cannot use them to reproduce, Chopra said. There are still more studies needed to answer these and other questions but Chopra said he was hopeful this line of study could lead to a way to combat antibiotic resistant bacteria.

"This area of [antibiotic resistance](#) is a big problem in global terms," Chopra said. "That's why we started thinking of what different ways we can use drugs already available to combat this problem."

Provided by University of Texas Medical Branch at Galveston

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