

AI 'scientist' finds that toothpaste ingredient may help fight drug-resistant malaria

18 January 2018



Eve the robot scientist. Credit: Ross King, University of Manchester

An ingredient commonly found in toothpaste could be employed as an anti-malarial drug against strains of malaria parasite that have grown resistant to one of the currently-used drugs. This discovery, led by researchers at the University of Cambridge, was aided by Eve, an artificially-intelligent 'robot scientist'.

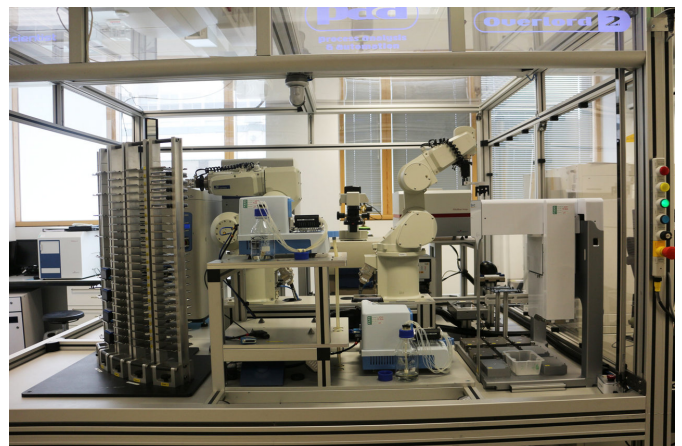
When a mosquito infected with [malaria](#) parasites bites someone, it transfers the parasites into their bloodstream via its saliva. These parasites work their way into the liver, where they mature and reproduce. After a few days, the parasites leave the liver and hijack red blood cells, where they continue to multiply, spreading around the body and causing symptoms, including potentially life-threatening complications.

Malaria kills over half a million people each year, predominantly in Africa and south-east Asia. While a number of medicines are used to treat the disease, malaria parasites are growing increasingly resistant to these drugs, raising the spectre of

untreatable malaria in the future.

Now, in a study published today in the journal *Scientific Reports*, a team of researchers employed the Robot Scientist 'Eve' in a high-throughput screen and discovered that triclosan, an ingredient found in many toothpastes, may help the fight against drug-resistance.

When used in toothpaste, triclosan prevents the build-up of plaque bacteria by inhibiting the action of an enzyme known as enoyl reductase (ENR), which is involved in the production of fatty acids.



Eve the robot scientist. Credit: Ross King, University of Manchester

Scientists have known for some time that triclosan also inhibits the growth in culture of the [malaria parasite](#) *Plasmodium* during the blood-stage, and assumed that this was because it was targeting ENR, which is found in the liver. However, subsequent work showed that improving triclosan's ability to target ENR had no effect on parasite growth in the blood.

Working with 'Eve', the research team discovered

that in fact, triclosan affects parasite growth by specifically inhibiting an entirely different enzyme of the malaria parasite, called DHFR. DHFR is the target of a well-established antimalarial drug, pyrimethamine; however, resistance to the drug among malaria parasites is common, particularly in Africa. The Cambridge team showed that triclosan was able to target and act on this enzyme even in pyrimethamine-resistant [parasites](#).

"Drug-resistant malaria is becoming an increasingly significant threat in Africa and south-east Asia, and our medicine chest of effective treatments is slowly depleting," says Professor Steve Oliver from the Cambridge Systems Biology Centre and the Department of Biochemistry at the University of Cambridge. "The search for new medicines is becoming increasingly urgent."

Because triclosan inhibits both ENR and DHFR, the researchers say it may be possible to target the parasite at both the liver stage and the later blood stage.

Lead author Dr Elizabeth Bilisland, now an assistant professor at the University of Campinas, Brazil, adds: "The discovery by our robot 'colleague' Eve that triclosan is effective against malaria targets offers hope that we may be able to use it to develop a new drug. We know it is a safe compound, and its ability to target two points in the malaria parasite's lifecycle means the parasite will find it difficult to evolve resistance."

Robot scientist Eve was developed by a team of scientists at the Universities of Manchester, Aberystwyth, and Cambridge to automate - and hence speed up - the drug discovery process by automatically developing and testing hypotheses to explain observations, run experiments using laboratory robotics, interpret the results to amend their hypotheses, and then repeat the cycle, automating high-throughput hypothesis-led research.

Professor Ross King from the Manchester Institute of Biotechnology at the University of Manchester, who led the development of Eve, says: "Artificial intelligence and machine learning enables us to create automated scientists that do not just take a

'brute force' approach, but rather take an intelligent approach to science. This could greatly speed up the [drug](#) discovery progress and potentially reap huge rewards."

More information: Elizabeth Bilisland et al, Plasmodium dihydrofolate reductase is a second enzyme target for the antimalarial action of triclosan, *Scientific Reports* (2018). [DOI: 10.1038/s41598-018-19549-x](#)

Provided by University of Cambridge

APA citation: AI 'scientist' finds that toothpaste ingredient may help fight drug-resistant malaria (2018, January 18) retrieved 26 May 2022 from <https://techxplore.com/news/2018-01-ai-scientist-toothpaste-ingredient-drug-resistant.html>

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