

Malaria under arrest: New drug target prevents deadly transmission

17 December 2019



Mosquito that transmits malaria. Credit: Walter and Eliza Hall Institute of Medical Research

Australian researchers have found a new drug target for stopping the spread of malaria, after successfully blocking the world's deadliest malaria parasite—*Plasmodium falciparum*—from completing the 'transmission stage' of its lifecycle.

Using [small molecule inhibitors](#) developed at the Walter and Eliza Hall Institute, the researchers blocked plasmepsin V, an enzyme essential for the development of gametocytes which are the only form of the parasite that can be transmitted from humans to mosquitoes.

The research, published today in *Cell Reports*, was led by Associate Professor Justin Boddey from the Walter and Eliza Hall Institute and University of Melbourne, in collaboration with Professor Vicky Avery from Griffith University in Queensland.

More than half a million people die from [malaria](#) every year and *Plasmodium falciparum* - the most

lethal of all malaria [parasites](#)—is responsible for 90 percent of infection cases. Due to the parasite's ability to constantly mutate and develop resistance to therapies, new preventions and treatments that act across different stages of the malaria parasite lifecycle—the liver stage, blood stage and transmission stage—are now required.

Arrested development

Associate Professor Boddey said the team had gained new ground towards malaria elimination because blocking the parasite's transmission stage was important for developing preventative therapies that stop the spread of disease.

"It was exciting to find that plasmepsin V plays a role in malaria transmission, and that our inhibitors could target plasmepsin V and block transmission to the mosquito from occurring," Associate Professor Boddey said.

"We showed that an optimal concentration of the inhibitors could kill gametocytes, and that even with a lower dose, the gametocytes made it all the way through their two-week development phase but still couldn't complete the task of transmitting infection to mosquitoes. This shows plasmepsin V is a target for transmission-blocking drugs," he said.

Using the Institute's insectary facilities, the researchers were able to study how gametocytes transmit malaria from human blood to a mosquito. They demonstrated, using gametocyte-specific fluorescent 'tags,' that plasmepsin V was critical for the export of gametocyte proteins—a process essential to gametocyte transmission—before proving their compounds could stop this process in its tracks.



Study lead, Associate Professor Justin Boddey. Credit: Walter and Eliza Hall Institute of Medical Research

Double whammy for disease

The results build on previous Institute studies including in 2014 when researchers discovered plasmepsin V was an effective drug target for killing the malaria parasite in the asexual blood stage of its lifecycle, when malaria symptoms—such as fever, chills, muscle pain and nausea—occur.

Institute chemical biologist Dr. Brad Sleebs, who was involved in both the current and previous plasmepsin V studies, said the enzyme was proving to be an ideal drug target because of its importance for parasite survival at different stages of the malaria lifecycle.

"It's encouraging to observe inhibitors that target plasmepsin V are effective against both the asexual blood and sexual transmission stages of the parasite's lifecycle," he said. "Our research demonstrates that an antimalarial treatment targeting plasmepsin V has potential, not only in treatment of the disease, but also as a preventative population control measure," Dr. Sleebs said.

Associate Professor Boddey said the research had been an example of how basic and translational knowledge was established from each new study building on the last.

"It's been a rewarding journey from identifying the

function of plasmepsin V, to developing inhibitors that block it and kill the malaria parasite, to now validating this enzyme's dual function as an effective blood stage and transmission-blocking drug target."

Sights set on final pillar

The researchers are now turning their attention to the role of plasmepsin V in the remaining pillar of the malaria lifecycle: the liver stage.

Associate Professor Boddey said the aim was to assess plasmepsin V as a multi-stage drug target for treating, as well as preventing, the spread of malaria; and to understand the unique biology occurring during liver infection.

"We are also collaborating with Merck and the Wellcome Trust to develop drugs targeting plasmepsin V in multiple parasite species," he said.

More information: Charlie Jennison et al. Inhibition of Plasmepsin V Activity Blocks *Plasmodium falciparum* Gametocytogenesis and Transmission to Mosquitoes, *Cell Reports* (2019). DOI: 10.1016/j.celrep.2019.11.073

Provided by Walter and Eliza Hall Institute

APA citation: Malaria under arrest: New drug target prevents deadly transmission (2019, December 17)
retrieved 15 April 2021 from
<https://medicalxpress.com/news/2019-12-malaria-drug-deadly-transmission.html>

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