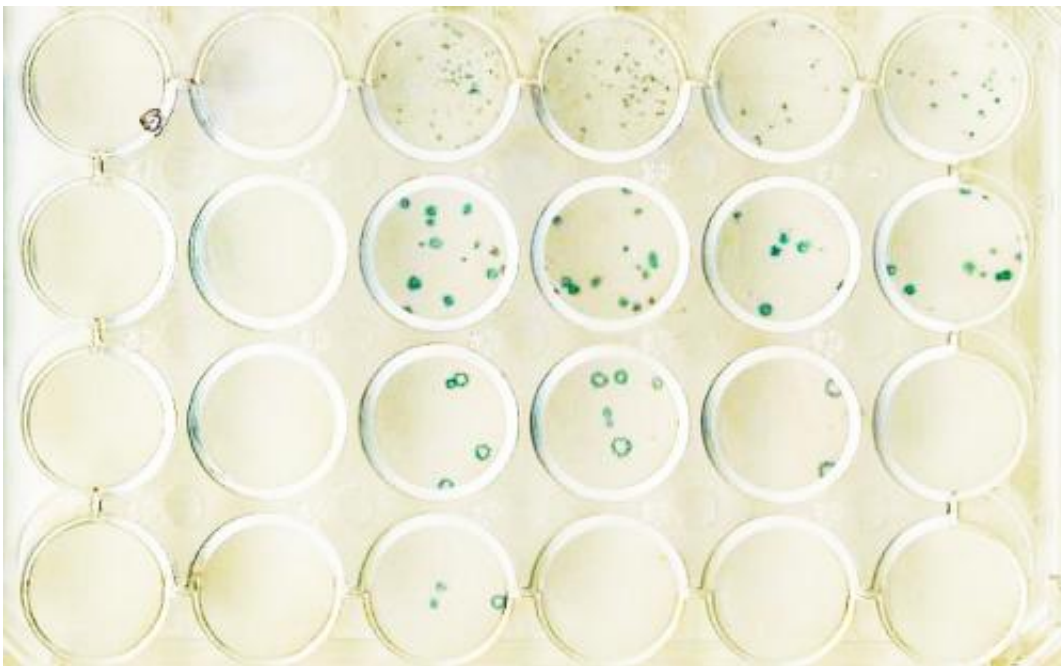


New strategy to disarm the dengue virus brings new hope for a universal dengue vaccine

August 13 2013



A new strategy that cripples the ability of the dengue virus to escape the host immune system has been discovered by A*STAR's Singapore Immunology Network (SIgN). This breakthrough strategy opens a door of hope to what may become the world's first universal dengue vaccine candidate that can give full protection from all four serotypes of the dreadful virus. This research done in collaboration with Singapore's

Novartis Institute of Tropical Diseases (NITD) and Beijing Institute of Microbiology and Epidemiology is published in the *Plos Pathogens* journal, and is also supported by Singapore STOP Dengue Translational and Clinical Research (TCR) Programme grant.

Early studies have shown that a sufficiently weakened [virus](#) that is still strong enough to generate protective immune response offers the best hope for an effective vaccine. However, over the years of vaccine development, scientists have learnt that the path to finding a virus of appropriate strength is fraught with challenges. This hurdle is compounded by the complexity of the [dengue](#) virus. Even though there are only four different [serotypes](#), the fairly high rates of mutation means the virus evolve constantly, and this contributes to the great diversity of the dengue viruses circulating globally. Furthermore, in some cases, the immune response developed following infection by one of the four dengue viruses appears to increase the risk of severe dengue when the same individual is infected with any of the remaining three viruses. With nearly [half the world](#)'s population at risk of dengue infection and an estimated 400 million people getting infected each year, the need for a safe and long-lasting vaccine has never been greater.

The new strategy uncovered in this study overcomes the prevailing challenges of vaccine development by tackling the virus' ability to 'hide' from the host immune system. Dengue virus requires the enzyme called MTase (also known as 2'-O-[methyltransferase](#)) to chemically modify its [genetic material](#) to escape detection. In this study, the researchers discovered that by introducing a genetic mutation to deactivate the MTase enzyme of the virus, initial cells infected by the weakened MTase mutant virus is immediately recognised as foreign. As a result, the desired outcome of a strong protective immune response is triggered yet at the same time the mutant virus hardly has a chance to spread in the host.

Animal models immunised with the weakened MTase mutant virus were fully protected from a challenge with the normal dengue virus. The researchers went on to demonstrate that the MTase mutant dengue virus cannot infect *Aedes* mosquitoes. This means that the mutated virus is unable to replicate in the mosquito, and will not be able to spread through mosquitoes into our natural environment. Taken together, the results confirmed that MTase mutant dengue virus is potentially a safe vaccine approach for developing a universal dengue vaccine that protects from all four serotypes.

The team leader, Dr Katja Fink from SIGN said, "There is still no clinically approved vaccine or specific treatment available for dengue, so we are very encouraged by the positive results with this novel vaccine strategy. Our next step will be to work on a vaccine formulation that will confer full protection from all four serotypes with a single injection. If this proves to be safe in humans, it can be a major breakthrough for the dengue vaccine field."

Associate Professor Leo Yee Sin, Clinical Director of Communicable Diseases Centre and Institute of Infectious Disease and Epidemiology at Tan Tock Seng Hospital who heads the Singapore STOP Dengue Translational and Clinical Research (TCR) Programme said, "We are into the seventh decade of dengue [vaccine development](#), this indeed is an exciting breakthrough that brings us a step closer to an effective vaccine."

Acting Executive Director of SIGN, Associate Professor Laurent Rénia said, "Dengue is a major public health problem in many of the tropical countries. We are very delighted that our collaborative efforts with colleagues in Singapore and China have made a promising step towards a cost-effective and safe [dengue vaccine](#) to combat the growing threat of dengue worldwide."

Mice immunised with the weakened MTase mutant virus or with a placebo were challenged with a normal [dengue virus](#). A plaque assay was used to test how much virus (visualized by the blue spots) was present in the blood of the mice. The placebo-treated mice were not protected and had virus in the blood, whereas the mice immunised with weakened MTase mutant virus were protected and had no virus in the blood.

More information: Zust, R. et al. Rational design of a live attenuated dengue vaccine: 2'-O-methyltransferase mutants are highly attenuated and immunogenic in mice and macaques, *Plos Pathogens*.
[www.plospathogens.org/article/ ... journal.ppat.1003521](http://www.plospathogens.org/article/...journal.ppat.1003521)

Provided by Agency for Science, Technology and Research (A*STAR), Singapore

Citation: New strategy to disarm the dengue virus brings new hope for a universal dengue vaccine (2013, August 13) retrieved 7 May 2024 from <https://medicalxpress.com/news/2013-08-strategy-dengue-virus-universal-vaccine.html>

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