

Malaria vaccine that 'could save thousands' to be tested in Africa

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A NEW malaria vaccine will be tested on a large scale in Kenya, Ghana and Malawi, the World Health Organisation has said, with 360,000 children to be vaccinated between 2018 and 2020.

The injectable vaccine RTS,S could provide limited protection against a disease that killed 429,000 people worldwide in 2015, with 92 per cent of victims in Africa and two thirds of them children under five.

"The prospect of a malaria vaccine is great news. Information gathered in the pilot will help us make decisions on

the wider use of this vaccine," said Dr Matshidiso Moeti, the WHO's regional director for Africa, yesterday.

The vaccine should be used alongside other preventative measures such as bed nets, insecticides, repellents and anti-malarial drugs, the WHO said.

"Combined with existing malaria interventions, such a vaccine would have the potential to save tens of thousands of lives in Africa," Dr Moeti added. "This vaccine is a weapon among others, it is one of the tools at our disposal."

The vaccine, also known as Mos-

quirix, has been developed by the British pharmaceutical giant GlaxoSmithKline in partnership with the PATH Malaria Vaccine Initiative, and the three-country pilot will test it on children aged five-to-17 months.

The drug passed previous scientific testing, including a phase three clinical trial between 2009 and 2014, and was approved for the pilot in 2015.

The aim of the trial is to assess the effectiveness of the vaccine as well as the feasibility of its delivery to populations at risk, because four successive doses

must be given on a strict timetable.

The immunisation cycle is not in sync with routine childhood inoculations against diseases such as hepatitis, measles and meningitis, with injections required at five months, six months, seven months and two years.

Symptoms of malaria include fever, muscle pain and headache as well as vomiting and diarrhoea.

While RTS,S does not promise full protection against the mosquito-borne disease, it is the most effective potential vaccine so far developed, reducing

the number of hospitalisations and blood transfusions. Malaria episodes reduced by 40 per cent in tests on 15,000 people in seven countries over five years of clinical trials, and could save hundreds of thousands of lives.

"It's an efficacy rate which is quite low, but given the amount of affected people, the impact will be huge," said Dr Mary Hamel, who is co-ordinating the vaccine's implementation programme.

"There will be other vaccines and they'll be more efficient, but this will have a significant influence." Dr Moeti

emphasised that while the dream is "a vaccine that replaces everything", insecticide-treated bed nets remain "our strongest preventive weapon".

Kenya, Ghana and Malawi were selected for the trial because malaria rates are high and they have a long history of use of bed nets and other interventions.

The large-scale pilot is the latest step in decades of work seeking to eradicate malaria, with the numbers dying falling nearly two thirds since the turn of the century.