



The problem

Malaria kills one child every two minutes.

Severe malaria causes death in nearly all cases without treatment, especially in children under five years of age.

Though injectable artesunate is far more effective than quinine for treating severe malaria, Unitaid identified that it was not being adopted on a large scale by national health programmes despite a strong recommendation from the World Health Organization (WHO).

The solution

What did Unitaid do?

Through Medicines for Malaria Venture (MMV), Unitaid invested US \$20 million over 2013-2016 to accelerate demand and adoption of quality-approved injectable artesunate in six countries (Cameroon, Ethiopia, Kenya, Malawi, Nigeria and Uganda). MMV partnered with the Clinton Health Access Initiative (CHAI) and the Malaria Consortium (MC) to update treatment guidelines; to quantify, procure and distribute injectable artesunate at affordable prices; and to train healthcare workers to use injectable artesunate in the project countries.

What did the project deliver?

Demand for injectable artesunate increased substantially. Annual procurement increased from very low levels to 10 million vials in 2015, through funding from Unitaid, the Global Fund and the US President's Malaria Initiative (PMI). Donor commitments are expected to exceed 20 million vials per annum in the future. And demand continues to increase. In 2011, only 9 countries in Africa had changed their severe-malaria policy in favour of injectable artesunate. By 2016 this had increased to 30 countries.

What impact will the project have over the long-term?

Based on current projections of demand, injectable artesunate could save an additional 66,000 children's lives each year by 2021 - it is more effective than quinine and potentially cheaper for health systems. Overall, the ISMO project has delivered an excellent return on investment (>\$100 for

Additional impact of the ISMO project - rectal artesunate

every dollar invested).

Where prompt treatment with injectable artesunate isn't available, the WHO recommends the use of rectal artesunate before a child is referred to a healthcare facility for treatment.

Access to rectal artesunate is low due to limited operational guidance, a lack of evidence of its use in real-world settings, and a lack of quality-assured products. The ISMO project supported the acceleration of two rectal artesunate products through WHO prequalification. A complementary investment by Unitaid (through CHAI) is now exploring whether rectal artesunate can be delivered and used safely and effectively in community settings.

The main contributors to Unitaid are: France, United Kingdom, Brazil, Norway, Chile, South Korea, Mauritius, Madagascar, Spain and Bill & Melinda Gates Foundation.

IMPACT DATA: IMPROVING SEVERE MALARIA OUTCOMES



For no extra cost, injectable artesunate can save an additional 2 lives for every 100 children treated

MORE EFFECTIVE



EASIER TO ADMINISTER



FASTER OUTCOMES



PREVIOUSLY

Quinine

Saves around

90

lives out of every 100 children treated.

Painful, toxic and difficult to administer, lasting many hours.

Slower-acting, patients discharged after several days.

NOW Injectable artesunate

92 lives out of every 100 children treated.

Saves around

Treatment can be delivered in only a few minutes, with fewer life-threatening side-effects.

Faster-acting, patients can be discharged after 1-2 days.

EQUIVALENT COST



Whilst the cost of a treatment course of quinine is lower than injectable artesunate, when factoring in additional costs related to severe malaria case management,* injectable artesunate is the same or potentially lower cost than quinine.

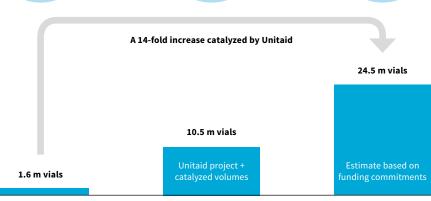
A further 66,000 children's lives saved annually by 2021*



Additional 28,000 children's lives saved

Additional 66,000 children's lives saved

2021 Scale-up by partners and countries



2015

*Demand projections developed for the final evaluation of the ISMO project, based on demand across Africa. Selection of references:

2013

Domdorp et al. (2010), "Artesunate versus quinine in the treatment of severe falciparum malaria in African children (AQUAMAT): an open-label, randomised trial", The Lancet.

Ferrari et al. (2015) "An operational comparative study of quinine and artesunate for the treatment of severe malaria in hospitals and health centres in the Democratic Republic of Congo: the MATIAS study", Malaria Journal.
Lubell et al. (2011), "Cost-effectiveness of parenteral artesunate for treating children with severe malaria in sub-Saharan Africa." WHO Bulletin.

Ntlku et al. (2016), "Feasibility and acceptability of nigectable artesunate for the treatment of severe malaria in the Democratic Republic of Congo.", Malaria Journal.

Sin Kyaw et al. (2014), "Cost of treating inpatient falciparum malaria on the Thai-Myanmar border." Malaria Journal.

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^{*}The total cost including commodity costs, in-patient costs and the cost of health care worker time.