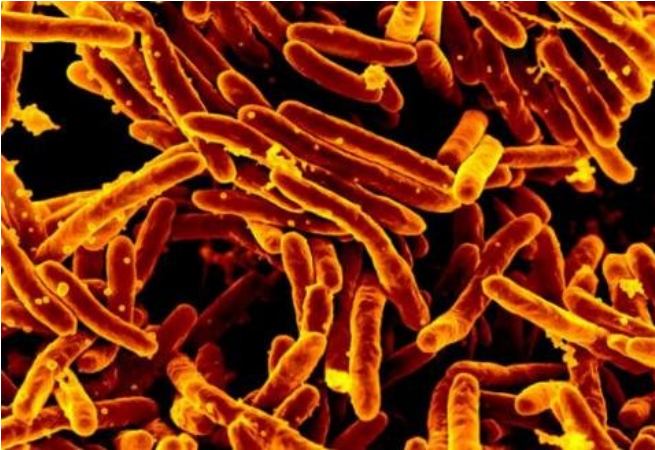


# Drug-resistant tuberculosis: A new study offers new hope

28 March 2019, by Andrew Nunn, I.d. Rusen



Scanning electron micrograph of Mycobacterium tuberculosis bacteria, which cause tuberculosis. Credit: NIAID

Tuberculosis (TB) is the leading cause of death in the world from a single infectious disease, causing more deaths than HIV/AIDS. In 2017, [10 million](#) people developed TB disease globally and an estimated 1.6 million died.

One of the biggest blocks to beating the epidemic is the growing resistance to drugs that have previously cured TB.

The World Health Organisation (WHO) has declared drug-resistant TB a global health crisis. Worldwide in 2017, an estimated [558,000](#) people developed TB that was resistant to the most effective first-line drug – rifampicin (RR-TB). Of these, 82% had multidrug-resistant TB.

But treating drug-resistant TB is still hopelessly inefficient. The 20-month to 24-month regimen used in many countries to treat people is costly and has significant side effects. In addition, the length of the regimen makes it hard for [patients](#) to adhere to, as well as for health systems to sustain.

Globally, the regimen has an average treatment success rate of little more than 50% in real-world treatment settings, although there is considerable variation from country to country.

As a result [researchers](#) around the world have been urgently exploring shorter, more effective, and safer treatments for patients who have [drug-resistant](#) TB.

One of these efforts has begun to bear fruit. Results from Stage 1 of a [clinical trial](#) have recently been published. The news is encouraging. The trial provides evidence that a shorter treatment regimen – of nine to 11 months – is as effective for the treatment of multidrug-resistant TB as the longer, 20-month treatment recommended by the WHO in 2011.

The STREAM trial presents robust evidence for the effectiveness and safety of the nine to 11-month regimen when compared to the much longer 20-month regimen. The results are as good and the fact that the regimen is shorter should make it much more acceptable to patients. It is also likely to result in cost saving to both patients and health services.

## The trial

The STREAM trial (Standardised Treatment Regimen of Anti-TB Drugs for Patients with MDR-TB) is the world's first multi-country randomised phase III clinical trial to test the efficacy, safety and economic impact of shortened MDR-TB treatment regimens. The randomised nature of the trial means that patients were assigned to the long or short regimen in such a way as so as to avoid bias. Treatment allocation is determined by chance not by the choice of a physician.

Phase III [trials](#) are designed to assess the effectiveness and safety of a new intervention in practice.

Stage 1 of the trial was designed to assess whether conditions, and reduces the number of pills patients a nine to 11-month treatment regimen that have to take. demonstrated promising cure rates during a pilot programme in Bangladesh, is as effective as the longer regimen when assessed in other settings under rigorous control trial conditions. Seven sites in Vietnam, Mongolia, South Africa, and Ethiopia participated.

The results from Stage 1 show that the shorter regimen is as good as the 20-month regimen.

Nearly 80% of patients in the trial showed a favourable outcome after two and a half years of follow-up from entry into the trial. The percentage was 79.8% in the 20-month regimen. In the nine to 11-month regimen the percentage was 78.8%.

Results in participants with HIV, although not as good as in those who were HIV-negative, were very similar in the short and long regimen.

The findings on side-effects were also interesting. There were very similar rates of severe side-effects during treatment and follow-up between the two regimens. But there were differences in the types of side-effects. The most common side-effects were cardiac conduction disorders, which increase the risk of serious and potentially fatal arrhythmias, in the nine to 11-month regimen. In the 20-month regimen the most common side effects were metabolic disorders, particularly hypokalemia.

A health economics data analysis is ongoing and will assess potential cost savings to patients and health systems when the nine to 11-month regimen is compared to the 20-month regimen.

## Conclusions

The final results from the trial are encouraging because they show that the nine to 11-month treatment regimen is comparable in efficacy and safety to the 20-month regimen. This supports the use of a shorter regimen for patients with rifampicin-resistant TB.

The nine to 11-month regimen presents substantial advantages. It reduces [treatment](#) times, may improve patient retention under programmatic

However, the required ECG monitoring is an important consideration.

In its latest guidelines issued last year the WHO again highlighted the need to continue to look for regimens that are not only shorter, but are also less toxic for the patient. The STREAM Stage 1 results suggest that some progress is being made.

STREAM Stage 2 is currently evaluating an all-oral regimen that is potentially as effective and more tolerable than the injectable-containing regimens currently used some countries. This, too, would be another major step forward in the battle against MDR-TB.

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