

Ebola: New trial launched in west Africa to evaluate three vaccination strategies

April 6 2017



A study volunteer receives inoculation at Redemption Hospital in Monrovia on the opening day in Liberia of PREVAC, a Phase 2 Ebola vaccine trial in West Africa. Credit: NIAID

The French National Institute of Health and Medical Research (Inserm),

the US National Institutes of Health (NIH) and the London School of Hygiene & Tropical Medicine (LSHTM), in collaboration with health authorities in Guinea and Liberia, are launching a large clinical trial of candidate Ebola vaccines under the aegis of the PREVAC international consortium (Partnership for Research on Ebola VACCination).

This trial seeks to identify vaccination regimens that hold the most promise to protect people from Ebola virus disease in order to prevent or quickly control a future outbreak. More than 5,000 adults and children living in countries at the epicenter of the 2014-16 West Africa Ebola epidemic will be enrolled. An additional site in Sierra Leone is also being planned. The PREVAC trial results from a research partnership that involves Inserm, NIH, LSHTM, and the West African Clinical Research Consortium¹. The pharmaceutical companies Janssen Vaccines & Prevention B.V., part of the Janssen Pharmaceutical Companies of Johnson & Johnson, Bavarian Nordic and Merck Sharp & Dohme, Corp (MSD outside USA and Canada), are supplying the experimental vaccines being tested in the PREVAC trial.

The trial will compare three experimental Ebola [vaccination strategies](#) with placebo regimens. It will be conducted in two stages, the first stage to take place in Guinea and Liberia. In Guinea, the trial is being conducted under the sponsorship of Inserm with the support of NIH and in collaboration with the Guinean authorities, and involves a partnership with the non-governmental organization ALIMA (The Alliance for International Medical Action). The NIH is sponsoring the trial based in Liberia, under its collaboration with the Liberia Ministry of Health in the Partnership for Research on Ebola Virus in Liberia (PREVAIL). Pending confirmation of funding, LSHTM will sponsor the planned site in Sierra Leone working with the University of Sierra Leone's College of Medicine and Allied Health Sciences, which would conduct the study.

In its first stage, the trial will evaluate one of the three strategies, a prime-

boost vaccination combining two different vaccines (one made by Janssen and the other by Bavarian Nordic) compared with a similar placebo regimen. Enrollment into this stage in Guinea and Liberia began on 27th March and 3rd April 2017, respectively. In a second stage, which is expected to start in the second half of 2017, the trial will evaluate all three vaccination strategies, including two additional strategies involving the Merck Sharp & Dohme, Corp vaccine.

On 29 March 2016, the Director-General of the World Health Organization (WHO) announced the end of the Public Health Emergency of International Concern caused by the Ebola outbreak.² At the end of this epidemic, although important advances have been achieved, several questions remain regarding the durability of the immune response of candidate vaccines under development. PREVAC trial will evaluate the rapidity, intensity and duration of the immune responses generated by the various vaccination strategies, and the safety and tolerability of the various vaccines, particularly in children.

According to Yves Levy, Inserm Chairman and CEO; Anthony S. Fauci, Director of the National Institute of Allergy and Infectious Diseases at NIH; and Peter Piot, Director of the LSHTM: "We, Inserm, NIH and LSHTM, have designed and launched this unique international research partnership with our colleagues from the West African countries most affected by Ebola virus disease to answer remaining questions regarding the safety and immunogenicity of candidate Ebola vaccine strategies and thereby enable us to better fight future Ebola outbreaks."

The first stage of the PREVAC study will evaluate a prime-boost strategy that includes the Janssen and Bavarian Nordic experimental vaccines:

- Ad26.ZEBOV administered as a first dose followed 8 weeks later

by MVA-BN-Filo as a booster dose.³ This vaccination strategy will be compared to an identical regimen in terms of dosage and duration, but made up of two placebos.

The second stage of the PREVAC study will evaluate all three strategies: the one used in the first stage, and two different regimens involving the experimental Merck Sharp & Dohme, Corp vaccine:

- rVSV?G-ZEBOV-GP⁴ administered as a first dose followed 8 weeks later by a booster dose of the same vaccine
- rVSV?G-ZEBOV-GP administered as a first dose followed 8 weeks later by an inert placebo Each of these vaccination strategies will be compared to an identical regimen in terms of dosage and duration, but made up of two placebos.

PREVAC in Brief

Currently, there is no approved vaccine to prevent Ebola virus disease, although some vaccines are at late stages of development. To better prepare for Ebola outbreaks and help contain them in future, it is essential to pursue research on vaccination strategies to prevent Ebola virus disease.

PREVAC (Partnership for Research on Ebola VACcination), is a research consortium that brings together the health authorities of three countries in West Africa—Guinea, Liberia and Sierra Leone—and their international partners, the National Institute of Health and Medical Research (Inserm) in France, the National Institute of Allergy and Infectious Diseases of the National Institutes of Health (NIAID/NIH) in the United States, and the London School of Hygiene & Tropical Medicine (LSHTM) in the United Kingdom. Additional partners in the

consortium include the NGO field partner Alliance for International Medical Action (ALIMA). The pharmaceutical companies Janssen Vaccines & Prevention B.V., part of the Janssen Pharmaceutical Companies of Johnson & Johnson, Bavarian Nordic and Merck Sharp & Dohme, Corp (MSD outside USA and Canada), are supplying the experimental vaccines being tested in the PREVAC trial.

The PREVAC trial is a Phase 2 clinical trial being conducted in West Africa. It is a randomised trial that aims to compare three experimental Ebola vaccination strategies with placebos in order to determine whether these strategies are safe and able to trigger a durable immune response that can protect against Ebola virus disease. In the first stage, the trial intends to enroll up to 600 participants 12 years or older. In the second stage, the trial intends to enroll 4,900 participants: 3,500 healthy adults aged 18 years or older, and 1,400 children aged 1-17 years. The study will initially be conducted at two sites in Guinea (Conakry/Landréah and Maferinyah) and one site in Liberia (Monrovia), with an additional site in Sierra Leone awaiting confirmation.

The main objectives of the PREVAC Ebola vaccine trial are as follows:

- To further explore the safety of three Ebola vaccination strategies and their ability to trigger a protective
- To assess these vaccination strategies in the general population and in important groups, such as children, for which there is only limited information available
- To obtain information on the duration of the immune response induced by the vaccines

In Guinea, two vaccination centres have been established, one in Conakry (in an urban area) and the other in Maferinyah (in a rural area). Liberia has one vaccination site in Monrovia.

According to Yazdan Yazdanpanah, PREVAC Principal Investigator, "Our challenge is to identify one or more safe, effective, and durable vaccines in order to prevent or control the next Ebola outbreak, for both adults and children."

"ALIMA's medical teams were at the front line in caring for patients with Ebola virus disease in Guinea. Today we know that this virus may reappear at any time. We therefore want to continue to support the population by pursuing the search for a [vaccine](#) capable of protecting the population from future epidemics," says Solenne Barbe, Programme Manager for ALIMA.

After the enrollment period, the participants will be followed up frequently at regular visits for at least 12 months. Teams of physicians, researchers and anthropologists will work in partnership in the field to ensure that the trial runs smoothly, and to answer questions from study participants and potential volunteers. Study staff will monitor participants' health, record any side effects, and obtain blood samples at follow-up visits after the vaccinations.

An independent Data and Safety Monitoring Board (DSMB) will closely monitor safety and immune response information throughout the trial. Children under the age of 12 years will not be enrolled initially. The DSMB will first examine data in children ages 12 to 17 years to determine if it is safe to expand the trial to children ages five to 11 years. After another review of safety data, the DSMB will consider whether vaccination of children ages one to four years can begin.

There is no risk to participants in this trial of becoming infected with Ebola by the study vaccines. The vaccines being tested are not live Ebola virus. They contain a single gene coding for a single protein from the Ebola virus. This protein cannot cause infection. The principle is the same as for many other existing vaccines.

More information: clinicaltrials.gov/ct2/show/NCT02876328

Provided by NIH/National Institute of Allergy and Infectious Diseases

Citation: Ebola: New trial launched in west Africa to evaluate three vaccination strategies (2017, April 6) retrieved 27 April 2024 from <https://medicalxpress.com/news/2017-04-ebola-trial-west-africa-vaccination.html>

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