



# EMA starts rolling review of COVID-19 Vaccine (Vero Cell) Inactivated

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EMA's human medicines committee (CHMP) has started a rolling review of COVID-19 Vaccine (Vero Cell) Inactivated, developed by Sinovac Life Sciences Co., Ltd. The EU applicant for this medicine is Life'On S.r.l.

The CHMP's decision to start the rolling review is based on preliminary results from laboratory studies (non-clinical data) and clinical studies. These studies suggest that the vaccine triggers the production of antibodies that target SARS-CoV-2, the virus that causes COVID-19, and may help protect against the disease.

EMA will evaluate data as they become available to decide if the benefits outweigh the risks. The rolling review will continue until enough evidence is available for a formal marketing authorisation application.

EMA will assess the compliance of COVID-19 Vaccine (Vero Cell) Inactivated with the usual EU standards for effectiveness, safety and quality. While EMA cannot predict the overall timelines, it should take less time than normal to evaluate an eventual application because of the work done during the rolling review.

EMA will communicate further when the marketing authorisation application for the vaccine has been submitted.

## How is the vaccine expected to work?

COVID-19 Vaccine (Vero Cell) Inactivated is expected to prepare the body to defend itself against infection with SARS-CoV-2. The vaccine contains SARS-CoV-2 that has been inactivated (killed) and cannot cause the disease. COVID-19 Vaccine (Vero Cell) Inactivated also contains an 'adjuvant', a substance that helps strengthen the immune response to the vaccine.

When a person is given the vaccine, their immune system identifies the inactivated virus as foreign and makes antibodies against it. If, later, the vaccinated person comes into contact with SARS-CoV-2, the immune system will recognise the virus and be ready to defend the body against it.

## What is a rolling review?

A rolling review is a regulatory tool that EMA uses to speed up the assessment of a promising medicine during a public health emergency. Normally, all data on a medicine or vaccine's effectiveness, safety and quality and all required documents must be ready at the start of the evaluation in a formal application for marketing authorisation. In the case of a rolling review, EMA's human medicines committee (CHMP) reviews data as they become available from ongoing studies. Once the CHMP decides that sufficient data are available, the company can submit a formal application. By reviewing the data as they become available, the CHMP can come to an opinion on the medicine's authorisation sooner.

During the rolling review, and throughout the pandemic, EMA and its scientific committees are supported by the COVID-19 EMA pandemic task force (COVID-ETF). This group brings together experts from across the European medicines regulatory network to advise on the development, authorisation and safety monitoring of medicines and vaccines for COVID-19 and facilitate quick and coordinated regulatory action.

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