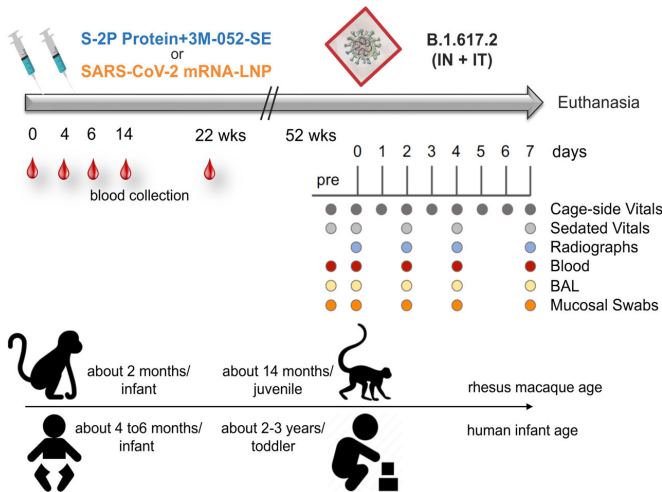


# Experimental COVID-19 vaccine offers long-term protection against severe disease

1 December 2022



**Experimental Design.** Two groups of 2-month old rhesus macaques (RMs) were immunized intramuscularly at weeks (wks) 0 and 4 with stabilized prefusion SARS-CoV-2 S-2P spike (S) protein of the Washington (SARS-CoV-2/human/USA/WA-CDC-02982586-001/2020) strain encoded by mRNA encapsulated in lipid nanoparticles (mRNA-LNP; n = 8) or purified S protein mixed with 3 M-052, a synthetic TLR7/8 agonist, in a squalene emulsion (Protein+3 M-052-SE; n = 8). Immunogenicity data up to week 22 post-immunization have been previously reported (28). RMs were monitored for 1 year post immunization, when vaccinated and an additional group of age-matched unvaccinated control RMs were challenged by the intranasal (IN) and intratracheal (IT) route with heterologous B.1.617.2 (Delta). The clinical monitoring and sample collection schedule until euthanasia on day 7 post-challenge is indicated underneath the time bar. The arrow at the bottom of the figure indicates the approximate age of human infants at the time of the first immunization of RMs and at the time of challenge. The age comparison is based on our data of normal changes in immune parameters in human and RM infants. Credit: *Science Translational Medicine* (2022). DOI: 10.1126/scitranslmed.add6383

In 2021, a group of scientists led by researchers at the University of North Carolina at Chapel Hill,

Weill Cornell Medicine and NewYork-Presbyterian reported that the Moderna mRNA vaccine and a protein-based vaccine candidate containing an adjuvant, a substance that enhances immune responses, elicited durable neutralizing antibody responses to SARS-CoV-2 during infancy in pre-clinical research.

Now a follow-up study by the same group, published in *Science Translational Medicine*, has found that the 2-dose vaccines still provide protection against lung disease in [rhesus macaques](#) one year after their having been vaccinated as infants.

The co-senior authors of the paper are Kristina De Paris, Ph.D., professor of microbiology and immunology at the UNC School of Medicine, Sallie Permar, MD, Ph.D., chair of the Department of Pediatrics at Weill Cornell Medicine, and Koen K.A. Van Rompay, DVM, Ph.D., leader of the Infectious Disease Unit at the California National Primate Research at the University of California, Davis. Co-first authors are Emma C. Milligan at the Children's Research Institute in the UNC School of Medicine and Katherine Olstad at the California National Primate Research Center.

To evaluate SARS-CoV-2 infant vaccination, the researchers immunized two groups of eight infant rhesus macaques at the California National Primate Research Center at 2 months of age and again four weeks later. Each animal received one of two vaccine types: a preclinical version of the Moderna mRNA vaccine or a vaccine combining a protein developed by the Vaccine Research Center of the National Institute of Allergy and Infectious Diseases (NIAID), part of the National Institutes of Health with a potent [adjuvant](#) formulation. Consisting of 3M's molecular adjuvant 3M-052 formulated in a squalene emulsion by the Access to Advanced Health Institute (AAHI), the adjuvant formulation stimulates immune responses by engaging receptors on immune cells.

"Following up on our SARS-CoV-2 infant rhesus macaque study, we gave the animals a high-dose challenge with a SARS-CoV-2 variant one year later to assess durability of vaccine-induced immune responses and their efficacy," Dr. De Paris said. "We found that both vaccines protected against [lung disease](#), despite the fact that the challenge SARS-CoV-2 variants acquired numerous mutations in their spike protein that differed from the vaccine immunogen."

Overall, the adjuvanted protein [vaccine candidate](#) maintained higher levels of [neutralizing antibodies](#) and provided superior protection compared to the mRNA vaccine, Dr. De Paris said. These data imply that these vaccines are safe and highly effective when given to young infant macaques. Furthermore, the results inform the optimization and development of SARS-CoV-2 vaccines in a way that may reduce the need for frequent boosters and protect special populations that don't have fully developed immune systems, such as young children.

"With COVID-19, young infants are one of the most vulnerable pediatric populations. This fall, we are seeing a sharp rise in hospitalizations due to respiratory virus disease in infants as the result of a confluence of SARS-CoV-2, flu, and RSV circulation," said Dr. Permar, who is also the Nancy C. Paduano Professor in Pediatrics at Weill Cornell Medicine and pediatrician-in-chief at NewYork-Presbyterian Komansky Children's Hospital. "We should take every opportunity to provide safe and effective [vaccine](#) immunity to our youngest patients, including considering COVID-19 vaccination earlier than the currently recommended 6 months of age."

"This study emphasizes the need to get human infants immunized against SARS-CoV-2 as much as possible, as the benefits are clear and long-lasting. It also highlights the value of animal models in infectious disease research," Dr. Van Rompay said. "The lessons we learned and the resources and tools that were developed in the current study will be very valuable for future pandemic preparedness, to more effectively combat outbreaks with novel coronaviruses or other respiratory viruses in pediatric populations."

**More information:** Emma C. Milligan et al, Infant rhesus macaques immunized against SARS-CoV-2 are protected against heterologous virus challenge one year later, *Science Translational Medicine* (2022). [DOI: 10.1126/scitranslmed.add6383](https://doi.org/10.1126/scitranslmed.add6383)

Provided by University of North Carolina Health Care

APA citation: Experimental COVID-19 vaccine offers long-term protection against severe disease (2022, December 1) retrieved 7 December 2022 from <https://medicalxpress.com/news/2022-12-experimental-covid-vaccine-long-term-severe.html>

*This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.*