The Medical Letter[®]

on Drugs and Therapeutics

Volume 63

December 27, 2021

ISSUE No.

IN THIS ISSUE

In Brief: Booster Doses of mRNA-Based COVID-19 Vaccines for All Adults......p 201

Important Copyright Message

FORWARDING OR COPYING IS A VIOLATION OF U.S. AND INTERNATIONAL COPYRIGHT LAWS

The Medical Letter, Inc. publications are protected by U.S. and international copyright laws. Forwarding, copying or any distribution of this material is prohibited.

Sharing a password with a non-subscriber or otherwise making the contents of this site available to third parties is strictly prohibited.

By accessing and reading the attached content I agree to comply with U.S. and international copyright laws and these terms and conditions of The Medical Letter, Inc.

For further information click: Subscriptions, Site Licenses, Reprints or call customer service at: 800-211-2769 The Medical Letter publications are protected by US and international copyright laws. Forwarding, copying or any other distribution of this material is strictly prohibited. For further information call: 800-211-2769

The Medical Letter[®] on Drugs and Therapeutics

Volume 63 (Issue 1640)

December 27, 2021

Take CME Exams

IN BRIEF

Booster Doses of mRNA-Based COVID-19 Vaccines for All Adults

Original online publication date: November 22, 2021 Updated: December 9, 2021

On November 19, the FDA expanded the Emergency Use Authorizations (EUAs) for the mRNA-based COVID-19 vaccines manufactured by Pfizer/ BioNTech (*Comirnaty*) and Moderna (*Spikevax*) to include administration of a booster dose for all adults \geq 18 years old after primary immunization with either the same COVID-19 vaccine or a different one.¹ Booster doses of these vaccines were previously authorized only for select populations (age \geq 65 years or persons at high risk for severe COVID-19). The EUA for the adenovirus-based vaccine manufactured by Johnson & Johnson was amended in October 2021 to include administration of a booster dose for all adults \geq 18 years old after primary immunization with the Johnson & Johnson vaccine.^{2,3} **Note:** On December 9, the FDA expanded the EUA of the Pfizer/ BioNTech COVID-19 vaccine to include a booster dose for adolescents 16-17 years old; the CDC endorsed use of booster doses in this age group. A full review of such use will be published in a future issue.

The CDC recommends a COVID-19 booster dose for all adults \geq 18 years old.⁴ A single booster dose of either **mRNA-based** COVID-19 vaccine can now be administered \geq 6 months after a primary series of an mRNA-based vaccine or \geq 2 months after a single primary dose of the Johnson & Johnson vaccine in all adults \geq 18 years old.^{5,6} A single booster dose of the **adenovirus-based** COVID-19 vaccine can be administered to any adult \geq 2 months after a single primary dose of the Johnson & Johnson vaccine or \geq 6 months after a primary series of an mRNA-based vaccine (see Table 1).⁷

Additional Content Available Online

COVID-19 Vaccine Dosing Recommendations and Comparison Chart http://medicalletter.org/downloads/1636f_table.pdf

COVID-19 Vaccine Comparison Chart http://medicalletter.org/downloads/1621g_table.pdf

Indication	Pfizer-BioNTech (Comirnaty)	Moderna (Spikevax)	Johnson & Johnson/Janssen
Primary immunization	≥16 yrs: 30 mcg (0.3 mL) IM at 0 and 3 weeks 12-15 yrs: 30 mcg (0.3 mL) IM at 0 and 3 weeks 5-11 yrs: 10 mcg (0.2 mL) IM at 0 and 3 weeks	≥18 yrs: 100 mcg (0.5 mL) IM at 0 and 4 weeks	≥18 yrs: 5x10¹º vp (0.5 mL) IM once
Additional primary dose for immunocompromised persons	≥12 yrs: 30 mcg (0.3 mL) IM ≥4 weeks after second primary dose	≥18 yrs: 100 mcg (0.5 mL) IM ≥4 weeks after second primary dose	Not authorized
Booster dose for all adults³ after a Pfizer-BioNTech or Moderna primary series⁴	30 mcg (0.3 mL) IM ≥6 months after last primary dose	50 mcg (0.25 mL) IM ≥6 months after last primary dose	5x10¹º vp (0.5 mL) IM ≥6 months after last primary dose
Booster dose for all adults after a Johnson & Johnson primary dose	30 mcg (0.3 mL) IM ≥2 months after primary dose	50 mcg (0.25 mL) IM ≥2 months after primary dose	5x10 ¹⁰ vp (0.5 mL) IM ≥2 months after primary dose

vp = viral particles

1. CDC. COVID-19 vaccine booster shots. November 29, 2021. Available at: https://bit.ly/3GvAkUw. Accessed December 9, 2021

 The Pfizer-BioNTech vaccine has received full FDA licensure for use as a 2-dose primary series in patients ≥16 years old. All other recommendations are based on FDA Emergency Use Authorizations (EUAs).

3. Adolescents 16-17 years old who completed a primary series with the Pfizer/BioNTech COVID-19 vaccine can receive a booster dose (30 mcg) of the same vaccine \geq 6 months after the last primary dose.

4. After either a 2- or 3-dose primary series.

The Medical Letter

- 1. FDA News Release. Coronavirus (COVID-19) update: FDA expands eligibility for COVID-19 vaccine boosters. November 19, 2021. Available at: https://bit.ly/3FQS6S7. Accessed December 9, 2021
- 2. Booster doses of COVID-19 vaccines. Med Lett Drugs Ther 2021: 63:186.
- 3. FDA News Release. Coronavirus (COVID-19) update: FDA takes additional actions on the use of a booster dose for COVID-19 vaccines. October 20, 2021. Available at: https:// bit.ly/3vJJPLy. Accessed December 9, 2021.
- 4. CDC News Release. CDC expands eligibility for COVID-19 booster shots to all adults. November 19, 2021. Available at: https://bit.ly/3cFk1aJ. Accessed December 9, 2021.
- 5. FDA. Fact sheet for health care providers administering vaccine (vaccination providers). Emergency Use Authorization (EUA) of the Pfizer-BioNTech COVID-19 vaccine to prevent coronavirus disease 2019 (COVID-19). November 19, 2021. Available at: https://bit.ly/37fX1NG. Accessed December 9, 2021
- 6. FDA. Fact sheet for healthcare providers administering vaccine (vaccination providers). Emergency Use Authorization (EUA) of the Moderna COVID-19 vaccine to prevent coronavirus disease 2019 (COVID-19). November 19, 2021. Available at: https://bit.ly/3nosylA. Accessed December 9, 2021.
- 7. FDA. Fact sheet for healthcare providers administering vaccine (vaccination providers). Emergency Use Authorization (EUA) of the Janssen COVID-19 vaccine to prevent coronavirus disease 2019 (COVID-19). November 19, 2021. Available at: https://bit.ly/3e6KEaD. Accessed December 9, 2021.

PRESIDENT: Mark Abramowicz, M.D.; VICE PRESIDENT AND EXECUTIVE EDITOR: Gianna Zuccotti, M.D., M.P.H., F.A.C.P., Harvard Medical School VICE PRESIDENT AND EDITOR IN CHIEF: Jean-Marie Pflomm, Pharm.D.; ASSOCIATE EDITORS: Susan M. Daron, Pharm.D., Amy Faucard, MLS, Corinne Z. Morrison, Pharm.D., Michael P. Viscusi, Pharm.D. CONSULTING EDITORS: Joanna Esterow, PA-C, Mordechai Sacks, DMSc, PA-C, Brinda M. Shah, Pharm.D., F. Peter Swanson, M.D.

CONTRIBUTING EDITORS: Carl W. Bazil, M.D., Ph.D., Columbia University College of Physicians and Surgeons; Ericka L. Crouse, Pharm.D., B.C.P.P., C.G.P., F.A.S.H.P., F.A.S.C.P., Virginia Commonwealth University; Vanessa K. Dalton, M.D., M.P.H., University of Michigan Medical School; Eric J. Epstein, M.D., Albert Einstein College of Medicine; David N. Juurlink, BPhm, M.D., Ph.D., Sunnybrook Health Sciences Centre; Richard B. Kim, M.D., University of Western Ontario; Sandip K. Mukherjee, M.D., F.A.C.C., Yale School of Medicine; Dan M. Roden, M.D., Vanderbilt University School of Medicine; Esperance A.K. Schaefer, M.D., M.P.H., Harvard Medical School; Neal H. Steigbigel, M.D., New York University School of Medicine; Arthur M. F. Yee, M.D., Ph.D., F.A.C.R., Weill Medical College of Cornell University

MANAGING EDITOR AND DIRECTOR OF CONTENT OPERATIONS: Susie Wong; EDITORIAL ASSISTANT: Karrie Ferrara

FULFILLMENT AND SYSTEMS MANAGER: Cristine Romatowski; EXECUTIVE DIRECTOR OF SALES: Elaine Reaney-Tomaselli

EXECUTIVE DIRECTOR OF MARKETING AND COMMUNICATIONS: Joanne F. Valentino; INTERIM PUBLISHER: Jean-Marie Pflomm, Pharm.D.

Founded in 1959 by Arthur Kallet and Harold Aaron, M.D.

Copyright and Disclaimer. The Medical Letter, Inc. is an independent nonprofit organization that provides healthcare professionals with unbiased drug prescribing recommendations. The editorial process used for its publications relies on a review of published and unpublished literature, with an emphasis on controlled clinical trials, and on the opinions of its consultants. The Medical Letter, Inc. does not sell advertising or receive any commercial support. No part of the material may be reproduced or transmitted by any process in whole or in part without prior permission in writing. The editors do not warrant that all the material in this publication is accurate and complete in every respect. The editors shall not be held responsible for any damage resulting from any error, inaccuracy, or omission.

Subscription Services

Customer Service: Call: 800-211-2769 or 914-235-0500 Fax: 914-632-1733 Subscriptions (US): 1 year - \$159; 2 years - \$298; 3 years - \$398. \$65 per year Permissions: Address The Medical Letter, Inc. 145 Huguenot St. Ste. 312 To reproduce any portion of this issue, please e-mail your request to: New Rochelle, NY 10801-7537 E-mail: custserv@medicalletter.org www.medicalletter.org for students, interns, residents, and fellows in the US and Canada. permissions@medicalletter.org Reprints - \$45 per issue or article Copyright 2021. ISSN 1523-2859

Site License Inquiries: E-mail: SubQuote@medicalletter.org Call: 800-211-2769 Special rates available for bulk subscriptions.

Get	Connected:	y	In
		_	