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JUDITH M. PERSICHILLI, RN, BSN, MA Commissioner

November 19, 2021

**TO:** Points of Dispensing participating in the COVID-19 Vaccination Program

**FROM:** Judith M. Persichilli, R.N., B.S.N., M.A. Commissioner

SUBJECT: Expansion of Pfizer, Moderna, and Janssen COVID-19 Vaccine Booster Doses to All Adults

Effective Friday, November 19, 2021, Points of Dispensing (PODs) can begin administration of booster doses of COVID-19 vaccines to all adults aged 18 years and older. Persons may receive any of the Food and Drug Administration (FDA) authorized or approved COVID-19 vaccines (Pfizer-BioNTech, Moderna, and Johnson & Johnson/Janssen) as a booster.

Everyone aged 18 years and older who completed their initial vaccination series with Pfizer-BioNTech or with Moderna at least six months ago are eligible for a booster. Everyone aged 18 years and older who received a Johnson & Johnson/Janssen vaccine continue to be recommended to receive a booster at least two months after their primary dose.

Today, the FDA authorized the use of the Moderna and Pfizer-BioNTech vaccines for a single booster dose for all persons aged 18 years and older after completion of primary vaccination with any FDA-authorized or approved COVID-19 vaccine. In accordance with the Advisory Committee on Immunization Practices (ACIP) and the Centers for Disease Control and Prevention (CDC), all persons aged 18 years and older are eligible to receive a COVID-19 booster doses at least six months after completion of the primary series.

Persons may receive any of the FDA authorized or approved COVID-19 vaccines (Pfizer-BioNTech, Moderna, and Johnson & Johnson/Janssen) as a booster.

This memo serves to alert you to the amended use of the Pfizer, Moderna, and Johnson & Johnson COVID-19 vaccines in New Jersey. **Please share this information with appropriate staff in your program.** 

Thank you for your partnership in this initiative. This memo supplements the other materials circulated by the New Jersey Department of Health (NJDOH) regarding the provision, delivery, and administration of COVID-19 vaccines. <a href="https://www.state.nj.us/health/cd/topics/covid2019\_vaccination.shtml">https://www.state.nj.us/health/cd/topics/covid2019\_vaccination.shtml</a>. Points of Dispensing may contact the New Jersey Department of Health Vaccine Operations Center at <a href="https://www.state.nj.us/health/cd/topics/covid2019\_vaccination.shtml">\text{Vax.Operations@doh.nj.gov}</a> with any further questions.

#### Eligible population

Effective immediately, all adults (aged 18 years and older) are eligible to receive a booster dose:

- All adults who received a primary series with Pfizer-BioNTech and/or Moderna COVID-19 vaccine at least six months ago.
- All adults who received one Johnson & Johnson/Janssen vaccine at least two months ago.

For mRNA recipeints, persons aged 50 years or older and persons aged 18 years or older who reside in long-term care facilities should receive a booster dose. All other persons aged 18 years or older may receive a booster based on their individual risks and benefits.

Meanwhile, the following remain eligible for the primary vaccine series:

- All persons aged 5 through 11 years are eligible for the Pfizer-BioNTech pediatric formulation COVID-19 two-dose series.
- All persons aged 12 through 17 years are eligible for the Pfizer-BioNTech adolescent/adult formulation COVID-19 two-dose series.
- All persons aged 18 and older are eligible for the Pfizer-BioNTech and Moderna two-dose series, and for the Johnson & Johnson/Janssen one-dose vaccine.
- Persons who are moderately to severely immune compromised are eligible for an additional (third) dose of the Pfizer-BioNTech or Moderna vaccines 28 days after their second dose.

On Wednesday, November 17, the CDC also issued Emergency Use Instructions (EUI) to provide information about the use of the COVID-19 vaccine by Pfizer-BioNTech, which is FDA-approved (licensed) for individuals aged 16 years and older. The EUI include the use of the Pfizer-BioNTech vaccine as an additional (third) dose in certain immunocompromised persons aged 12 years and older and as a booster dose in certain adults aged 18 years and older after completion of primary vaccination with certain non-FDA authorized or approved COVID-19 vaccines.

All points of dispensing are advised to make booster doses available to those who self identify as eligible under the above criterion. Providers must accept self-reported eligibility from the vaccine recipient as sufficient and are not permitted to require individuals to present documentation or a note from a medical provider to demonstrate eligibility.

To establish the interval, if a person does not present with a vaccination record or the Docket app, providers can query the New Jersey Immunization Information System (NJIIS).

### Vaccine types

Federal authorizations and recommendations allow for mix and match (heterologous) dosing for booster shots. To offer choice at point of dispensing and while maintaining proper vaccine stewardship, sites are encouraged to carry as many types of COVID-19 vaccines as capable.

Eligible individuals may choose which vaccine they receive as a booster dose. Some people may have a preference for the vaccine type that they originally received and others may prefer to get a different booster. Vaccine recipients may make this individual self-assessment through their consideration of their own risks, may weigh information in the EUA Fact Sheets for Recipients and Caregivers, may consult a healthcare provider, or may seek out other sources of guidance. Points of dispensing must not add barriers to access for anyone who presents at a vaccination site and self-identifies as eligible.

PODs may use the currently available Pfizer-BioNTech and Moderna vaccines for primary, second, and third dose recipients. Currently available Johnson & Johnson/Janssen vaccine may be used for primary and second dose recipients.

- The booster Pfizer-BioNTech COVID-19 vaccine dose is the same product and dosage as the initial two-dose mRNA COVID-19 primary vaccine series.
- The booster Moderna vaccine dose is the same product and half the dosage as the initial two-dose mRNA COVID-19 primary vaccine series.
- The booster Johnson & Johnson/Janssen is the same product and dosage as the initial COVID-19 primary vaccine.

# Administration timeline

If received mRNA (Pfizer-BioNTech or Moderna) primary series, at least six months until booster. If Johnson & Johnson/Janssen primary series, at least two months until booster. If immunocompromised mRNA vaccine recipient, additional dose 28 days after second dose.

Note: "Fully vaccinated" remains defined as at least two weeks after their second dose in a two-dose series, such as the Pfizer-BioNTech or Moderna vaccines, or two weeks after a single-dose vaccine, such as the Johnson & Johnson/Janssen vaccine. This definition applies to all people, including those who receive an additional dose as recommended for moderate to severely immunocompromised people and those who receive a booster dose.

#### Co-administration

COVID-19 and other vaccines (e.g., influenza vaccines, childhood vaccines) may be administered without regard to timing. This includes administration of COVID-19 and other vaccines on the same day, as well as co-administration within 14 days.

## **Scheduling**

During the initial booster rollout, in addition to appointments, walk-ins should still be accommodated. This is especially important to ensure timely access without barriers for those who are unvaccinated, those newly eligible (e.g. just turned 5 years old), and those due for their second dose of an initial two-dose mRNA vaccine series.

Update all appointment schedulers to account for booster dose eligibility. Points of dispensing must continue to be listed on the <u>New Jersey COVID-19 Information Hub</u> and your POD must update NJDOH in a timely manner if any information on the state website is incomplete or out-of-date.

PODs must use recall and reminder methods to message eligibility for boosters to your Pfizer-BioNTech and Moderna vaccine recipients who received their second dose at least six months ago. PODs also must use recall and reminder methods to message eligibility for boosters to your Johnson & Johnson/Janssen vaccine recipients who received their first dose at least two months ago.

## **Site Readiness**

To accommodate as many eligible vaccine recipients as soon as possible, all active sites should prepare to have additional vaccines on hand at all vaccine events, provide night and weekend hours, accommodate walk-ins, update appointment availability reflected on the New Jersey Vaccine Appointment Finder

(<a href="https://covid19.nj.gov/finder">https://covid19.nj.gov/finder</a>), and share information with NJDOH about pop-up vaccination clinics to be included in the COVID-19 Community Calendar (<a href="https://covid19.nj.gov/pages/communitycalendar">https://covid19.nj.gov/pages/communitycalendar</a>). Sites are also encouraged to actively promote the availability of booster doses. If additional vaccine supply is needed at a site, please be in touch with New Jersey Department of Health to arrange for allocation and/or transfer.

To the extent possible while maintaining appropriate vaccine stewardship, all points of dispensing are expected to optimize use of all COVID-19 vaccines. No vaccination opportunity should be missed, so sites should request and carry sufficient inventory to vaccinate at capacity.

#### Reporting

The New Jersey Immunization Information System (NJIIS) has been upgraded to accept booster doses in vaccine recipients' records.

All doses of COVID-19 vaccine administered in New Jersey must be reported to NJIIS. Doses reported to NJIIS will then be available for display in the vaccine recipient's Docket app upon app refresh.

### Federal recommendation

On Friday, November 19, the FDA issued revised EUAs with updated fact sheets for the booster vaccines:

- FDA statement: <a href="https://www.fda.gov/news-events/press-announcements/coronavirus-covid-19-update-fda-expands-eligibility-covid-19-vaccine-boosters">https://www.fda.gov/news-events/press-announcements/coronavirus-covid-19-update-fda-expands-eligibility-covid-19-vaccine-boosters</a>
- For Pfizer-BioNTech vaccine:
  - o Updated EUA: <a href="https://www.fda.gov/media/150386/download">https://www.fda.gov/media/150386/download</a>
  - Updated EUA Fact Sheet for Healthcare Providers:
  - o https://www.fda.gov/media/153713/download
  - Updated EUA Fact Sheet for Recipients and Caregivers: https://www.fda.gov/media/153716/download
  - Translations of the Fact Sheet for Recipients and Caregivers:
     <a href="https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/pfizer-biontech-covid-19-vaccine">https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/pfizer-biontech-covid-19-vaccine</a>
- For Moderna vaccine:
  - o Updated EUA: <a href="https://www.fda.gov/media/144636/download">https://www.fda.gov/media/144636/download</a>
  - Updated EUA Fact Sheet for Healthcare Providers: <a href="https://www.fda.gov/media/144637/download">https://www.fda.gov/media/144637/download</a>
  - Updated EUA Fact Sheet for Recipients and Caregivers: https://www.fda.gov/media/144638/download
  - Translations of the Fact Sheet for Recipients and Caregivers:
     https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/moderna-covid-19-vaccine
- For Johnson & Johnson/Janssen vaccine:
  - Updated EUA: https://www.fda.gov/media/146303/download
  - Updated EUA Fact Sheet for Healthcare Providers: https://www.fda.gov/media/146304/download
  - Updated EUA Fact Sheet for Recipients and Caregivers: https://www.fda.gov/media/146305/download

Translations of the Fact Sheet for Recipients and Caregivers:
 https://www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/janssen-covid-19-vaccine

On Friday, November 19, the CDC adopted recommendations for the booster dose:

CDC statement: https://www.cdc.gov/media/releases/2021/s1119-booster-shots.html

On Wednesday, November 17, the CDC issued Emergency Use Instructions (EUI) for certain people who completed a vaccine primary series with certain COVID-19 vaccines that are not authorized or approved by the U.S. Food and Drug Administration:

- EUI Fact Sheet for Healthcare Providers: <a href="https://www.cdc.gov/vaccines/covid-19/eui/downloads/EUI-HCP.pdf">https://www.cdc.gov/vaccines/covid-19/eui/downloads/EUI-HCP.pdf</a>
- EUI Fact Sheet for Recipients and Caregivers: <a href="https://www.cdc.gov/vaccines/covid-19/eui/downloads/EUI-Caregiver.pdf">https://www.cdc.gov/vaccines/covid-19/eui/downloads/EUI-Caregiver.pdf</a>

### Additional guidance:

Interim Clincal Considerations for Use of COVID-19 Vaccines Currently Approved or Authorized in the United States available at <a href="https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html">https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html</a>

World Health Organization Emergency Use List available at https://extranet.who.int/pgweb/vaccines/vaccinescovid-19-vaccine-eul-issued

### Provider education

Please ensure vaccination providers are trained and well versed in the updated vaccine information, provider fact sheet, and clinical considerations. We expect providers to be prepared to address questions from their patients and their families about the recommended booster dose. Information about COVIDavailable 19 vaccines and training materials is through the CDC website https://www.cdc.gov/vaccines/covid-19/index.html. Please see CDC's Interim Clinical Considerations for COVID-19 Vaccines Currently Authorized https://www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html

## Consumer education

Points of dispensing play a critical role in building trust in vaccination. Please note CDC's dedicated webpage with information on the COVID-19 Vaccine Booster Shot: https://www.cdc.gov/coronavirus/2019-ncov/vaccines/booster-shot.html

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