



COVID-19 Provider Update

Wednesday, July 6, 2022



Discontinuation of Monthly Calls

Monthly CoVP calls will no longer be conducted. Calls will be scheduled as needed for relevant updates.

Please continue to monitor your communications for up-to-date information.



COVID-19 Updates

- On June 18th, [CDC](#) recommended to expand COVID-19 vaccination to children younger than 5 years of age. This decision now universally recommends COVID-19 vaccination for everyone 6 months and older. Kids under 5 can get either the Pfizer-BioNTech (3 dose) or Moderna (2 dose) primary vaccine series.
- On June 23rd, CDC recommended to [expand the use of Moderna COVID-19 vaccines to include children ages 6 through 17 years](#). A (2 dose) series is recommended for this age group.
- June 28th, [FDA's VRPBAC met to discuss COVID strain selection/future boosters](#).
 - Committee voted 19-2 in favor of including a SARS-CoV-2 omicron component in COVID-19 vaccines that would be used for boosters in the U.S.
 - FDA has advised manufacturers seeking to update their COVID-19 vaccines to develop modified vaccines that add an omicron BA.4/5 spike protein component to the current vaccine composition to create a two component (bivalent) booster vaccine, so that the modified vaccines can potentially be used starting in early to mid-fall 2022 (tentatively in October).
- Novavax Update

Interim Clinical Considerations Update for Pediatric COVID-19 Vaccines



Moderna Pediatric Schedule: People Who Are NOT Moderately or Severely Immunocompromised

Moderna
(6 months–
5 years)

Dose 1
(primary)

4-8 weeks

Dose 2
(primary)



0.25 mL (25 mcg)

Moderna
(6–11 years)

Dose 1
(primary)

4-8 weeks

Dose 2
(primary)



0.50 mL (50 mcg)

Moderna
(12–17 years)

Dose 1
(primary)

4-8 weeks

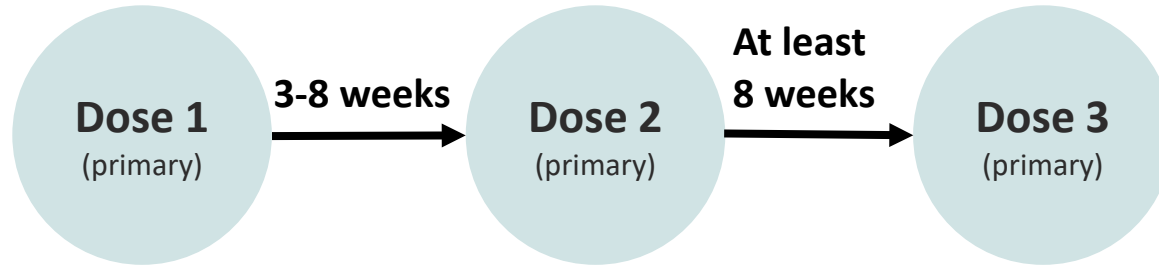
Dose 2
(primary)



0.50 mL (100 mcg)

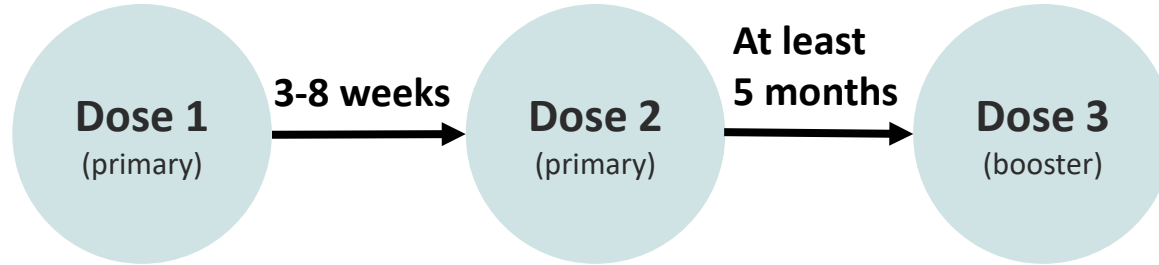
Pfizer-BioNTech Pediatric Schedule: People Who Are NOT Moderately or Severely Immunocompromised

Pfizer-BioNTech
(6 months–
4 years)



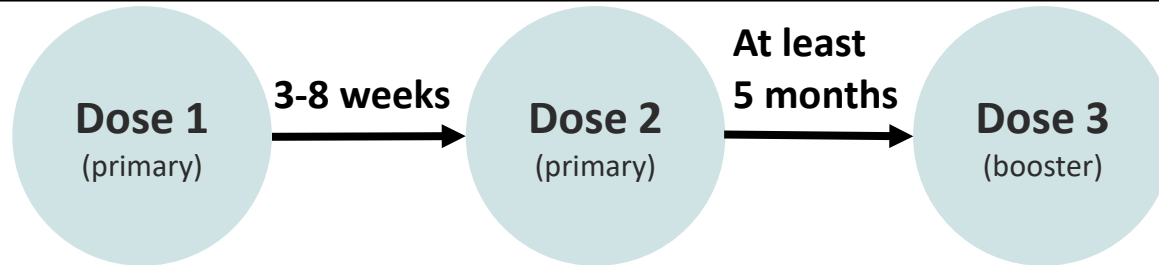
0.2 mL (3 mcg)

Pfizer-BioNTech
(5–11 years)



0.2 mL (10 mcg)

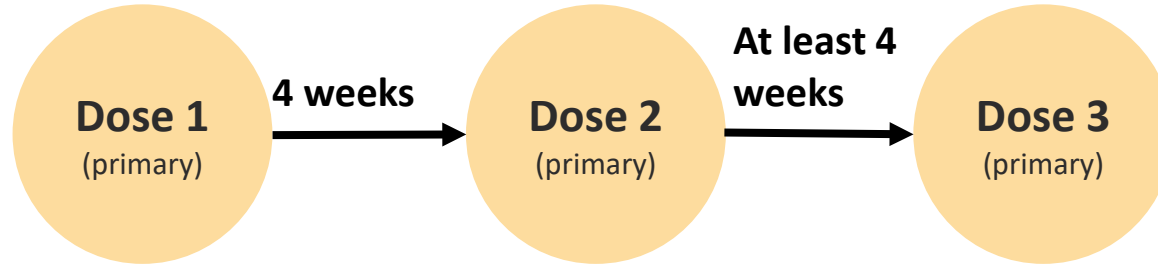
Pfizer-BioNTech
(12–17 years)



0.3 mL (30 mcg)

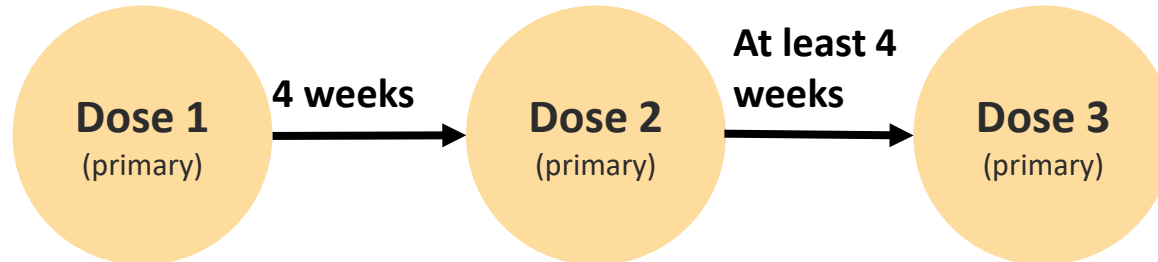
Moderna Pediatric Schedule: People Who ARE Moderately or Severely Immunocompromised

Moderna
(6 months–
5 years)



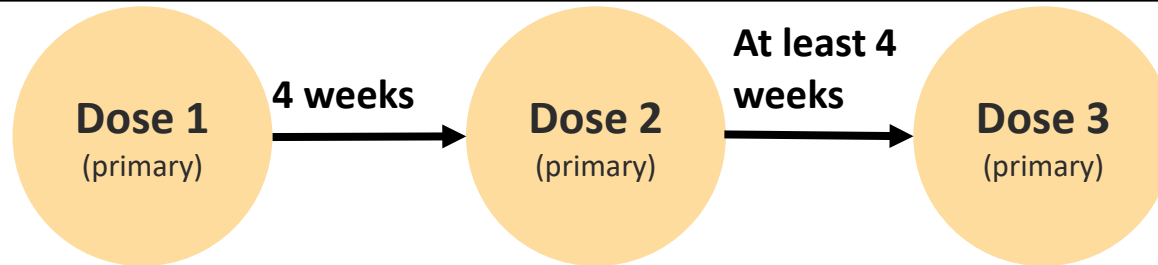
0.25 mL (25 mcg)

Moderna
(6–11 years)



0.50 mL (50 mcg)

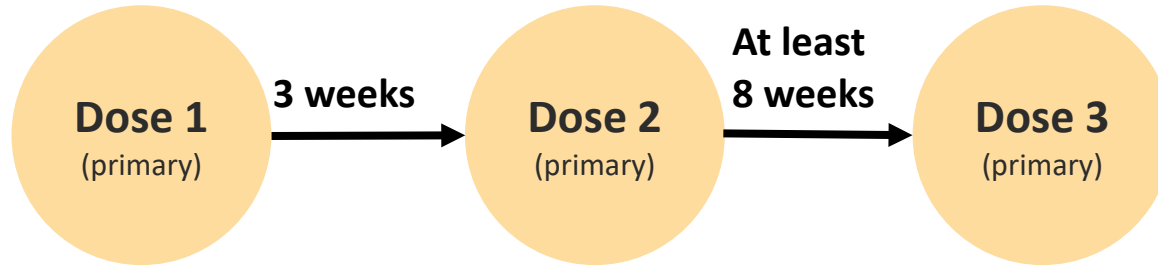
Moderna
(12–17 years)



0.50 mL (100 mcg)

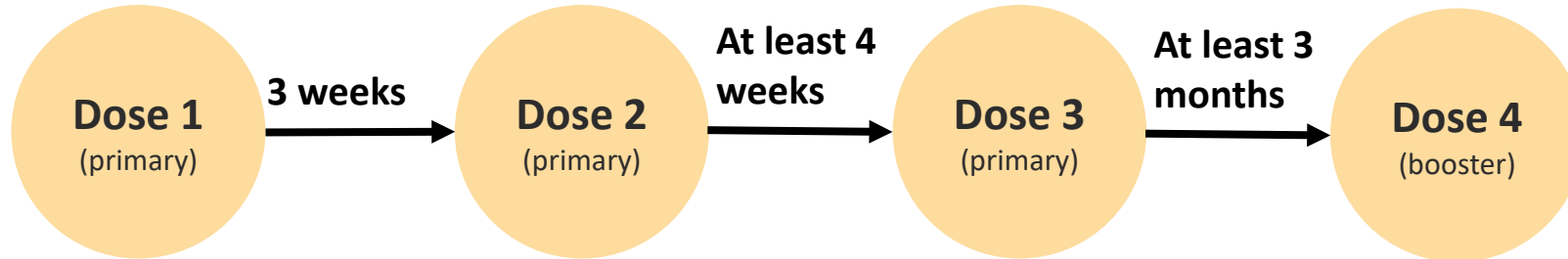
Pfizer-BioNTech Pediatric Schedule: People Who ARE Moderately or Severely Immunocompromised

Pfizer-BioNTech
(6 months–4 years)



0.2 mL (3 mcg)

Pfizer-BioNTech
(5 years–11 years)



0.2 mL (10 mcg)

Pfizer-BioNTech
(12 years–17 years)



0.3 mL (30 mcg)

Schedule Resources

- At-a-glance schedule: <https://www.cdc.gov/vaccines/covid-19/downloads/COVID-19-vacc-schedule-at-a-glance-508.pdf>
- Interim COVID-19 immunization schedule: <https://www.cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-6months-older.pdf>

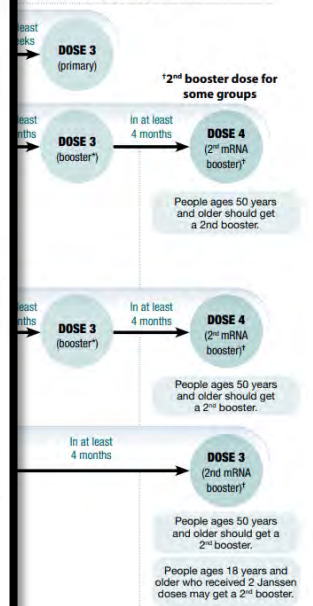
AT-A-GLANCE

COVID-19 Vaccination Schedules

Use the schedules below to determine how many total COVID-19 vaccine doses are recommended based on primary series product, age, and immune status. This schedule does not include clinical details necessary for administering COVID-19 vaccines. For clinical details, see the resources at the end of this document.

COVID-19 Vaccination Schedule for Most People

of COVID-19 vaccine doses



COVID-19 Vaccine

Interim COVID-19 Immunization Schedule for 6 Months of Age and Older

The table below provides guidance for COVID-19 vaccination schedules based on age and medical condition. Scheduling considerations include:

- Administer the appropriate vaccine product based on the recipient's age and the product's age indications.
- COVID-19 vaccines may be administered on the same day as other vaccines.
- Doses administered at any time after the intervals outlined below are valid.

Detailed information can be found in CDC's Interim Clinical Considerations for Use of COVID-19 Vaccines Currently Approved or Authorized in the United States, see: www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html

Table 1. Immunization Schedule for Children 6 Months through 17 Years of Age

| Type | Product* | Recipient Age | For Most People | | Those who ARE Moderately or Severely Immunocompromised | |
|--|--|--------------------------|---------------------------------|--------------------------------------|--|--------------------------------------|
| | | | Doses | Interval Between Doses ^{1†} | Doses | Interval Between Doses ^{1†} |
| mRNA vaccine | Moderna (Blue vial cap with magenta-bordered label) | 6 months through 5 years | Total doses: 2 doses | | Total doses: 3 doses | |
| | | | Dose 1 to 2 | At least 4–8 weeks ¹ | Dose 1 to 2 | At least 4 weeks |
| | Pfizer-BioNTech (Aluminum vial cap with maroon-bordered label) | 6 months through 4 years | Total number: 3 doses | | Total number: 3 doses | |
| | | | Dose 1 to 2 | At least 3–8 weeks ¹ | Dose 1 to 2 | At least 3 weeks |
| Pfizer-BioNTech (Orange vial cap with orange-bordered label) | 5 through 11 years | Total number: 3 doses | | Total number: 4 doses | | |
| | | Dose 1 to 2 | At least 3–8 weeks ¹ | Dose 1 to 2 | At least 3 weeks | |
| Pfizer-BioNTech (Purple vial cap with purple-bordered label or gray vial cap with gray-bordered label) | 12 years through 17 years | Total number: 3 doses | | Total number: 5 doses | | |
| | | Dose 1 to 2 | At least 3–8 weeks ¹ | Dose 1 to 2 | At least 3 weeks | |

* Complete the primary series with same product. If the vaccine product previously administered cannot be determined or is no longer available, any age-appropriate mRNA COVID-19 vaccine product may be administered at least 28 days after the first dose. Any COVID-19 vaccine product (age appropriate) may be administered for a booster dose. † Does not need to be the same product used for the primary series.
¹ Persons with a recent SARS-CoV-2 infection may consider delaying a primary series or booster dose by 3 months from symptom onset or positive test (if infection was asymptomatic).
[†] Some studies in adolescents and adults have shown the small risk of myocarditis associated with mRNA COVID-19 vaccines might be reduced and peak antibody responses and vaccine effectiveness may be increased with an interval longer than 4 weeks. An 8-week interval may be optimal for people who are not moderately or severely immunocompromised and ages 3–6 years, especially for males ages 12–19 years.

06/17/2022



Pfizer-BioNTech COVID-19 Vaccine Products



Product for ages
6 mons – 4 years



Product for ages
5–11 years

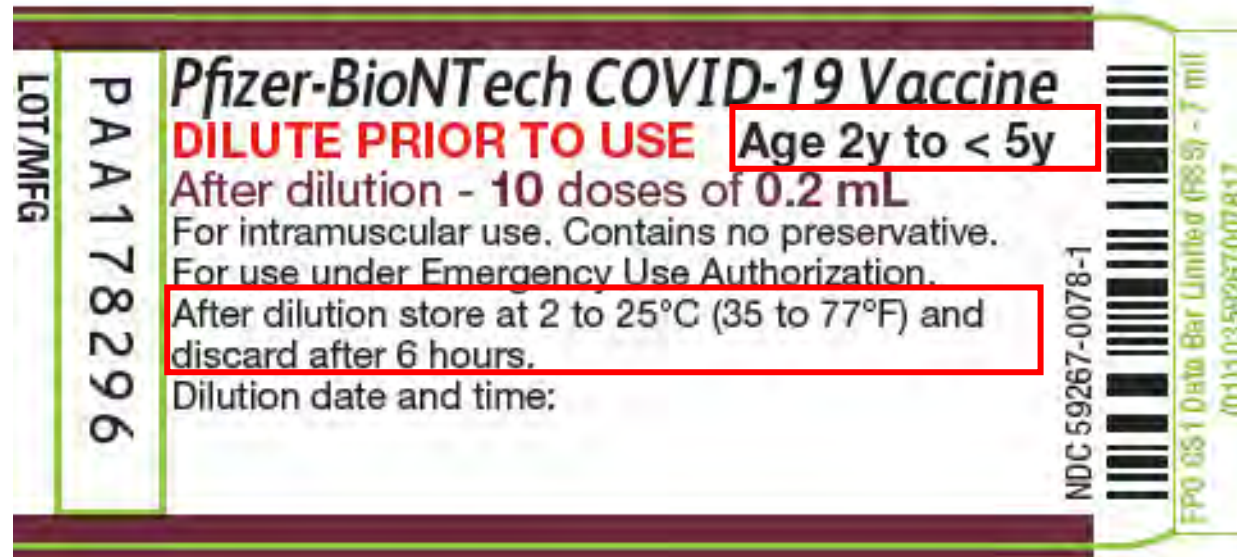


Product for ages
12 years and older

| | Product for ages 6 mons – 4 years | Product for ages 5–11 years | Product for ages 12 years and older |
|----------------------------------|--------------------------------------|--------------------------------|--|
| Authorized for ages | 6 months–4 years | 5–11 years | 12 years and older |
| Vial cap color | Maroon | Orange | Gray |
| Dose (mRNA concentration) | 3 mcg | 10 mcg | 30 mcg |
| Injection volume | 0.2 mL | 0.2 mL | 0.3 mL |
| Dilution required | Yes—2.2 mL | Yes—1.3 mL | No |
| Doses per vial | 10 (after dilution) | 10 (after dilution) | 6 |

Pfizer-BioNTech COVID-19 Vaccine Product for Ages 6 Months–4 Years

Vaccine may be discarded **12 hours** after dilution rather than **6 hours**.

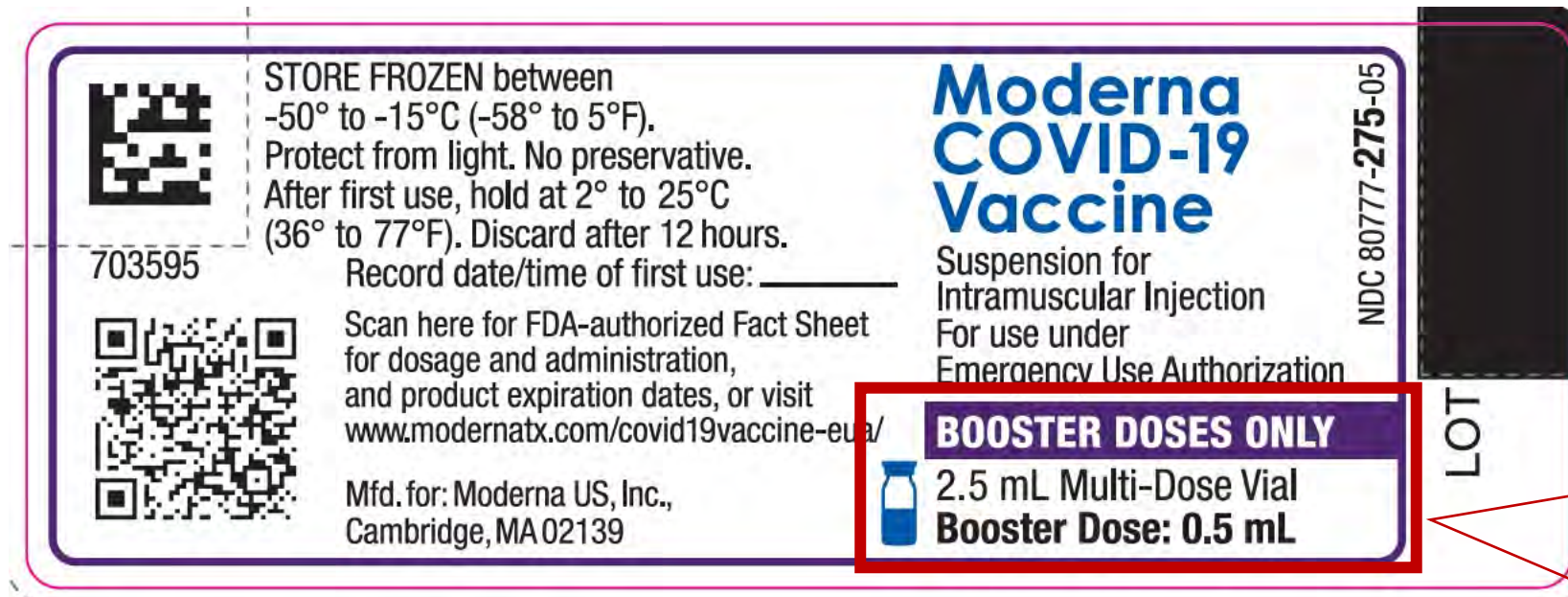


Vial label states Age 2y to <5y but can be used in children ages 6 months–4 years.

Moderna COVID-19 Vaccine Products

| Authorized Age group |  6 months–5 years (primary series) |  <ul style="list-style-type: none"> • 6–11 years (primary series) • 18 years and older (booster doses) |  <ul style="list-style-type: none"> • 12 years and older (primary series) • 18 years and older (booster doses) |
|---------------------------|---|---|---|
| Vial cap color | Dark blue | Dark blue | Red |
| Label border color | Magenta | Purple | Light blue |
| Dose (mRNA concentration) | 25 mcg | 50 mcg | 100 mcg (primary, age 12+); 50mcg (booster, age 18+) |
| Injection volume | 0.25 mL | 0.5 mL | 0.5 mL (primary, age 12+); 0.25mL (booster, age 18+) |
| Dilution required | No | No | No |
| Doses per vial | 10 | 5 | Maximum of 11 |

Moderna COVID-19 Vaccine Product for Ages 6–11 Years



The image shows a rectangular product label for Moderna COVID-19 Vaccine. The label is primarily white with blue and purple text. On the left side, there is a QR code and the number 703595. The top left corner features a barcode. The main text on the label includes storage instructions, a warning to protect from light, and information about the vaccine's use under Emergency Use Authorization. A prominent purple box in the lower right section of the label contains the text "BOOSTER DOSES ONLY" and "2.5 mL Multi-Dose Vial Booster Dose: 0.5 mL". To the right of this box, the word "LOT" is printed vertically. The NDC number 80777-275-05 is also visible on the right side of the label.

703595

STORE FROZEN between -50° to -15°C (-58° to 5°F). Protect from light. No preservative. After first use, hold at 2° to 25°C (36° to 77°F). Discard after 12 hours. Record date/time of first use: _____

Scan here for FDA-authorized Fact Sheet for dosage and administration, and product expiration dates, or visit www.modernatx.com/covid19vaccine-eua/

Mfd. for: Moderna US, Inc., Cambridge, MA 02139

Moderna COVID-19 Vaccine

Suspension for Intramuscular Injection
For use under Emergency Use Authorization

NDC 80777-275-05

BOOSTER DOSES ONLY

2.5 mL Multi-Dose Vial
Booster Dose: 0.5 mL

LOT

Labeled for “**BOOSTER DOSES ONLY**” but is authorized for:

- Primary doses in children ages 6–11 years
- Booster doses in adults ages 18 years and older

Product Resources

- Storage, handling, preparation, and administration job aids by product: <https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html>
- Pfizer-BioNTech label infographic: <https://www.cdc.gov/vaccines/covid-19/info-by-product/pfizer/downloads/infant-label-info.pdf>
- Moderna label infographic: <https://www.cdc.gov/vaccines/covid-19/info-by-product/moderna/downloads/Moderna-children-updated-label-iinfo-508.pdf>
- At-a-glance job aids
 - Pfizer-BioNTech: <https://www.cdc.gov/vaccines/covid-19/info-by-product/pfizer/downloads/vaccine-at-a-glance.pdf>
 - Moderna: <https://www.cdc.gov/vaccines/covid-19/info-by-product/moderna/downloads/vaccine-at-a-glance.pdf>

Pfizer-BioNTech COVID-19 Vaccine Products At-A-Glance
 Maroon cap: 6 Months through 4 Years of Age
 Orange cap: 5 through 11 years of age
 Gray or Purple cap: 12 years of age and older

The tables below summarize basic storage, preparation, scheduling, administration, and dosage for Pfizer-BioNTech COVID-19 Vaccine. Use this table in conjunction with materials below:

| | | | |
|--|---|--|---|
| EUA Fact Sheet www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/comrnaty-and-pfizer-biontech-covid-19-vaccine | CDC Interim Clinical Considerations www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html | CDC Clinical Materials www.cdc.gov/vaccines/covid-19/info-by-product/pfizer | Interim COVID-19 Vaccination Schedule www.cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-5yrs-older.pdf |
|--|---|--|---|

Moderna COVID-19 Vaccine At-A-Glance
 Ages 6 months and older

The tables below summarize basic storage, preparation, scheduling, administration, and dosage for Moderna COVID-19 Vaccine.

| | | | |
|--|---|--|---|
| EUA Fact Sheet www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/spillvax-and-moderna-covid-19-vaccine | CDC Interim Clinical Considerations www.cdc.gov/vaccines/covid-19/clinical-considerations/covid-19-vaccines-us.html | CDC Clinical Materials www.cdc.gov/vaccines/covid-19/info-by-product/moderna | Interim COVID-19 Vaccination Schedule https://www.cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-6months-older.pdf |
|--|---|--|---|

Storage and Handling Basics
NOTE: Find additional guidance on storing the vaccine properly at:
 • CDC's COVID-19 Clinical materials for Moderna COVID-19 Vaccine: www.cdc.gov/vaccines/covid-19/info-by-product/moderna/storage.html
 • CDC's Storage and Handling Toolkit: www.cdc.gov/vaccines/hcp/admin/storage/toolkit/storage-handling-toolkit.pdf

| Vial cap color | Blue vial cap with magenta bordered label | Blue vial cap with purple-bordered label | Red vial cap with blue bordered label |
|---|--|--|---|
| Ages | 6 months through 5 years | 6 years through 11 years | 12 years and older |
| Supplied in multidose vial | 10 doses per vial | 5 doses per vial | Per vial: <ul style="list-style-type: none"> 10 primary doses (0.5 mL) or 20 booster doses (0.25 mL) 20 doses: Combination of primary series and booster doses Discard vial and any remaining vaccine after 20 punctures. |
| Storage Temperature: Before Puncture | Between: -50°C and -15°C (-58°F and 5°F) until the expiration date 2°C and 8°C (36°F and 46°F) for up to 30 days 8°C and 25°C (46° and 77°F) for a total of 24 hours. Discard vial and unused vaccine after 24 hours. | | |
| Thawing Frozen Vaccine | Between: 2°C and 8°C (36°F and 46°F) for 2 hours. Let each vial stand at room temperature for 15 minutes before administering. Do NOT refreeze thawed vaccine. OR 15°C and 25°C (46°F and 77°F) for 45 minutes. | | |
| Storage Temperature: After Puncture | Between: 2°C and 25°C (36°F and 77°F) for up to 12 hours. Discard vial and any unused vaccine after 12 hours. | | |

| Cap | Purple Cap |
|------------------------|--|
| | 12 years and older |
| Vial | 6 doses per vial Requires diluent |
| Temperature | Between: -90°C and -60°C (-130°F and -76°F) until the expiration date ¹ -25°C to -15°C (-13°F to 5°F) for up to 2 weeks 2°C and 8°C (36°F and 46°F) for up to 10 weeks |
| Number of Vials | in temperature and number of vials. |
| Discard | Between: 2°C and 25°C (36°F and 77°F) for up to 6 hours. Discard vial and any unused vaccine after 6 hours. |



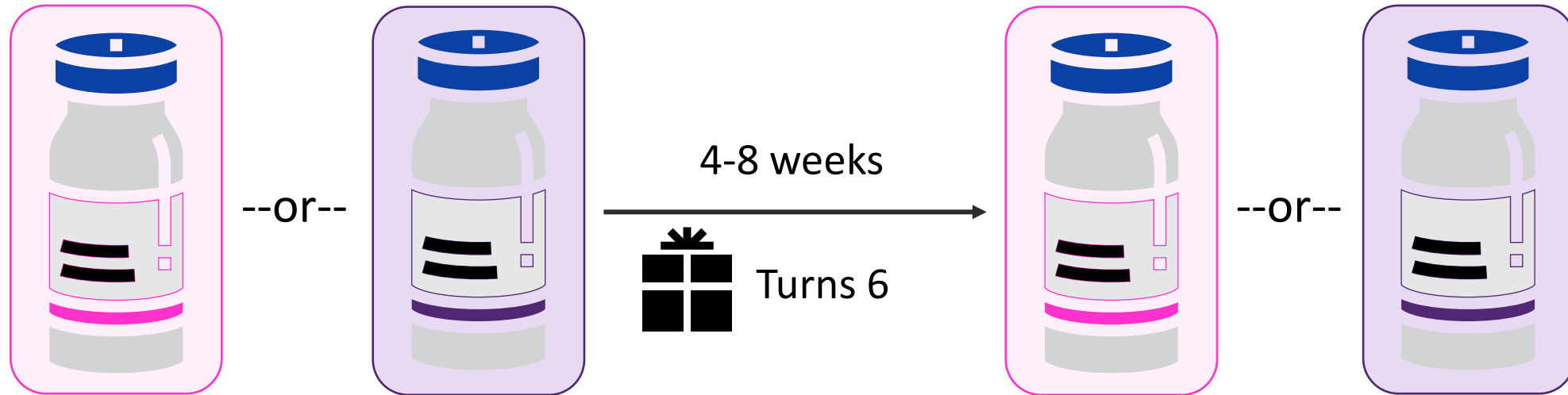
Vaccine Dosage: CDC's Recommendation

- **CDC'S recommendation:** Children should receive the age-appropriate vaccine product and follow the schedule based on their age on the day of vaccination, regardless of their size or weight.
- If a person moves from a younger age group to an older age group during the primary series or between the primary series and receipt of the booster dose(s), they should receive the vaccine dosage for the older age group for all subsequent doses.
- FDA authorization allows for dosing options for certain age transitions. If these options occur, they are not considered errors and the doses “count”.



FDA allowance for Moderna age 5 to 6 years

Children who will turn from age 5 years to 6 years between doses in the primary series to receive, for any primary dose: (1) the Moderna COVID-19 Vaccine product authorized for children ages 6 months–5 years or (2) the Moderna COVID-19 Vaccine product authorized for children ages 6–11 years.



Dose 1 (Age 5):

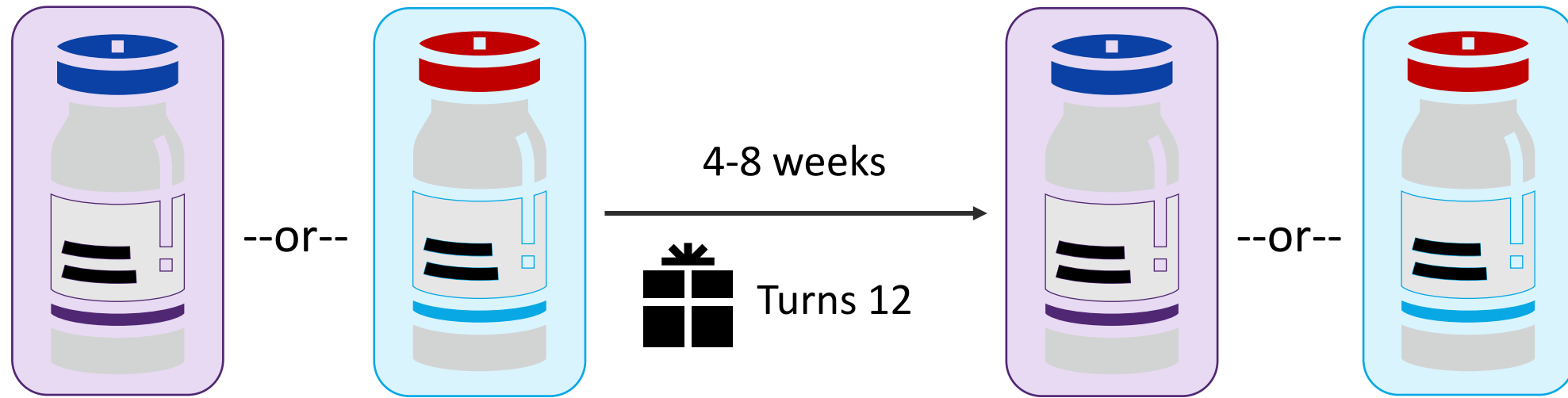
0.25 mL (25 mcg) of the product for ages
6 months–5 years or
0.50 mL (50 mcg) of the product for
ages 6–11 years

Dose 2 (Age 6):

0.25 mL (25 mcg) of the product for ages
6 months–5 years or
0.50 mL (50 mcg) of the product for ages
6–11 years

FDA allowance for Moderna age 11 to 12 years

Children who will turn from age 11 years to 12 years between doses in the primary series to receive, for any primary dose: (1) the Moderna COVID-19 Vaccine product authorized for children ages 6–11 years or (2) the Moderna COVID-19 Vaccine product authorized for people ages 12 years and older.



Dose 1 (Age 11):

0.50 mL (50 mcg) of the product for ages
6–11 years

0.50 mL (100mcg) of the product for ages
12 years and older

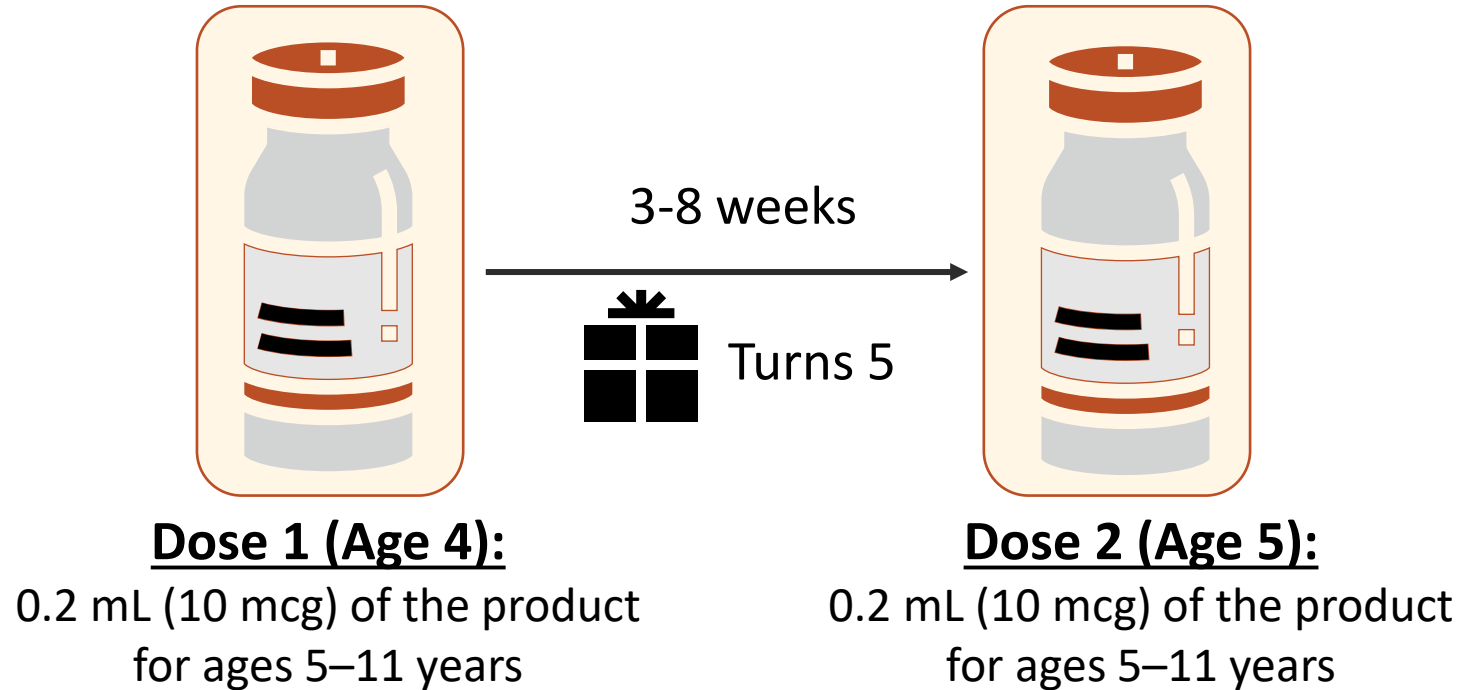
Dose 2 (Age 12):

0.50 mL (50 mcg) of the product for
ages 6–11 years

0.50 mL (100mcg) of the product for
ages 12 years and older

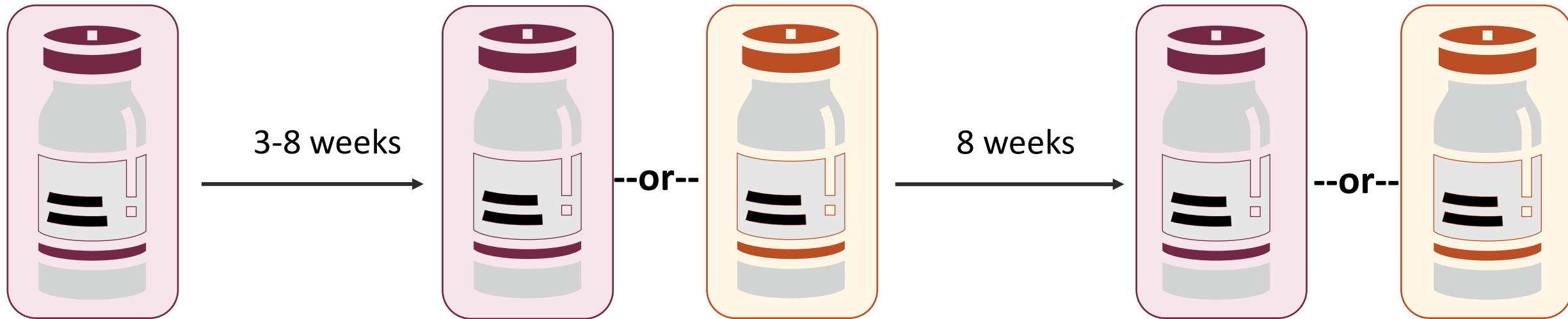
FDA allowance for Pfizer-BioNTech age 4 to 5 years

Scenario 1: A 2-dose primary series using the product for people ages 5–11 years (orange cap)



FDA allowance for Pfizer-BioNTech age 4 to 5 years

Scenario 2: A 3-dose primary series initiated with the product for ages 6 months–4 years. Dose 2 and 3 may be with: the product for ages 6 months–4 years or the product for ages 5–11 years.



Dose 1 (Age 4):

0.2 mL (3 mcg) of the product for ages 6 months–4 years

Dose 2 (Age 4 or 5):

0.2 mL (3 mcg) of the product for ages 6 months–4 years, or
0.2 mL (10 mcg) of the product for ages 5–11 years

Dose 3 (Age 5):

0.2 mL (3 mcg) of the product for ages 6 months–4 years, or
0.2 mL (10 mcg) of the product for ages 5–11 years


Aging up Resources

- Moderna:

<https://www.cdc.gov/vaccines/covid-19/downloads/Moderna-Child-Age-Transition-508.pdf>
- Pfizer-BioNTech:

<https://www.cdc.gov/vaccines/covid-19/downloads/Pfizer-Child-Age-Transition-508.pdf>


Pfizer-BioNTech COVID-19 Vaccine
for Children who Transition from a Younger to Older Age Group



CDC recommends vaccine recipients receive the recommended age-appropriate vaccine product and dosage **based on their age on the day of vaccination.**

- If a person moves from a younger age group to an older age group during the primary series or between the primary series and receipt of the booster dose(s), they should receive the vaccine product and dosage for the older age group for all subsequent doses.
- FDA emergency use authorization (EUA) allows for different dosing for certain age transitions, which are not considered vaccine administration errors and do not need to be reported to the Vaccine Adverse Event Reporting System (VAERS).

Moderna COVID-19 Vaccine
for Children who Transition from a Younger to Older Age Group



CDC recommends vaccine recipients receive the recommended age-appropriate vaccine product and dosage **based on their age on the day of vaccination.**

- If a person moves from a younger age group to an older age group during the primary series, they should receive the vaccine product and dosage for the older age group for all subsequent doses.
- FDA emergency use authorization (EUA) allows for different dosing for certain age transitions, which are not considered vaccine administration errors and do not need to be reported to the Vaccine Adverse Event Reporting System (VAERS).

Children who turn from age 5 to age 6 years

Recommended: Children who started a primary series and turned from age 5 to age 6 years before completion of the series should receive:

Dose 1 (Age 5): 0.25 mL (25 mcg) of the product authorized for children ages 6 months–5 years (dark blue cap/magenta label border)

4-8 weeks

Dose 2 (Age 6): 0.50 mL (50 mcg) of the product authorized for children ages 6–11 years (dark blue cap/purple label border)

Child turns 6

Acceptable: If the following dosing occurs, it is NOT considered an error and the primary series is considered complete. **Either dose may be:**

- 0.25 mL (25 mcg) of the product authorized for children ages 6 months–5 years (dark blue cap/magenta label border), or
- 0.50 mL (50 mcg) of the product authorized for children ages 6–11 years (dark blue cap/purple label border)

Dose 1: Age 5 OR **Dose 2: Age 6**

4-8 weeks

06/23/2022 02201570-8 1

turned from age 4 to age 5 years between dose 1 and dose 2

8 weeks

Dose 3 (Age 5): 0.20 mL (10 mcg) of the product authorized for children ages 5–11 years (orange cap and label border)

turned from age 4 to age 5 years between dose 2 and dose 3

8 weeks

Dose 3 (Age 5): 0.20 mL (10 mcg) of the product authorized for children ages 5–11 years (orange cap and label border)

Child turns 5

ed an error and the primary series is considered complete. vaccine product authorized for children ages 5–11 years

Dose 2: Age 4 or 5

ks

1



Interchangeability

- COVID-19 vaccines are not interchangeable.
- The same mRNA vaccine product should be used for all doses of the primary series.
- In exceptional situations in which the mRNA vaccine product administered for a previous dose(s) of the primary series cannot be determined or is not available, either age-appropriate available mRNA COVID-19 vaccine product may be administered at a minimum interval of 28 days between doses to complete the mRNA COVID-19 primary vaccination series.



Mixed Series For Children Ages 6 months–4 Years

- Children ages 6 months–4 years who receive different mRNA products for the first 2 doses of an mRNA COVID-19 vaccine series should receive a third dose of either mRNA vaccine 8 weeks after the second dose to complete the 3-dose primary series.



Mixed Series For Children Ages 6 months–4 Years

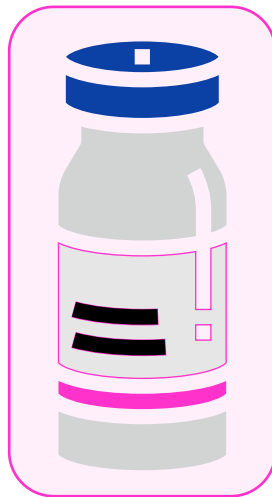
■ Scenario 1:



Dose 1:

0.20 mL (3mcg) of the Pfizer-BioNTech product for ages 6 months through 4 years

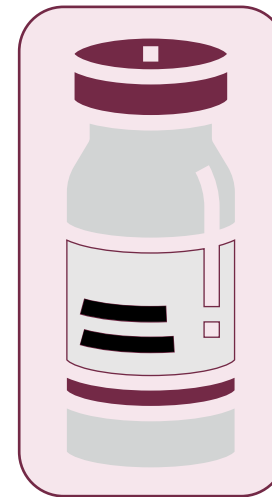
4-8 weeks



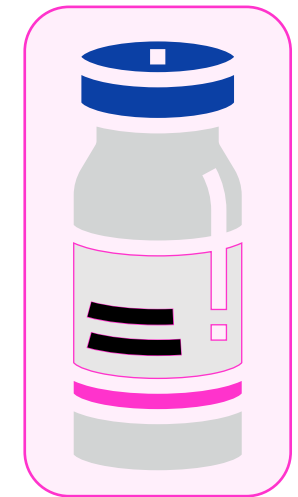
Dose 2:

0.25 mL (25mcg) of the Moderna product for ages 6 months through 5 years

8 weeks



--or--



Dose 3:

0.20 mL (3mcg) of the Pfizer-BioNTech product for ages 6 months through 4 years

--or--

0.25 mL (25mcg) of the Moderna product for ages 6 months through 5 years

Mixed Series For Children Ages 6 months–4 Years

- **Scenario 1:**



Dose 1:

0.25 mL (25mcg) of the Moderna product for ages 6 months through 5 years

Dose 2:

0.20 mL (3mcg) of the Pfizer-BioNTech product for ages 6 months through 4 years

Dose 3:

0.20 mL (3mcg) of the Pfizer-BioNTech product for ages 6 months through 4 years
--or--
0.25 mL (25mcg) of the Moderna product for ages 6 months through 5 years

Preventing Vaccine Administration Errors

- Clinical guidance for errors: <https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#appendix-c>
- Handout: <https://www.cdc.gov/vaccines/hcp/admin/downloads/vaccine-administration-preventing-errors.pdf>

YOU CALL THE SHOTS

Vaccine Administration: Preventing Vaccine Administration Errors

A vaccine administration error is any preventable event that may cause or lead to inappropriate medication use or patient harm.¹ Vaccine administration errors can have many consequences, including inadequate immunological protection, possible injury to the patient, cost, inconvenience, and reduced confidence in the health care delivery system. Take preventive actions to avoid vaccine administration errors and establish an environment that values reporting and investigating errors as part of risk management and quality improvement.

Vaccine administration errors may be due to causes such as:

- Insufficient staff training
- Distraction
- Changes in recommendations
- Lack of standardized protocols
- Patient misidentification
- Using nonstandard or error-prone abbreviations
- Easily misidentified products (e.g. DTaP, DT, Tdap, Td)

If an error occurs, determine how it occurred and take the appropriate actions to put strategies in place to prevent it from happening in the future. The following table outlines common vaccine administration errors and possible preventive actions you can take to avoid errors.

| Error(s) | Possible Preventive Actions |
|--|--|
| Wrong vaccine, route, site, or dosage (amount); or improperly prepared. | <ul style="list-style-type: none"> Circle important information on the packaging to emphasize the difference between the vaccines. Include the brand name with the vaccine abbreviation whenever possible (e.g., PCV13 [Pevnar13]) in orders, medical screens, etc. Separate vaccines into bins or other containers according to type and formulation. Use color-coded identification labels on vaccine storage containers. Store look-alike vaccines in different areas of the storage unit (e.g., pediatric and adult formulations of the same vaccine on different shelves in the unit). Do not list vaccines with look-alike names sequentially on computer screens, order forms, or medical records, if possible. Consider using "name alert" or "look-alike" stickers on packaging and areas where these vaccines are stored. Consider purchasing products with look-alike packaging from different manufacturers, if possible. Establish "Do NOT Disturb" or no-interruption areas or times when vaccines are being prepared or administered. Prepare vaccine for one patient at a time. Once prepared, label the syringe with vaccine name. Do not administer vaccines prepared by someone else. Triple-check work before administering a vaccine and ask another staff member to check. Keep reference materials on recommended sites, routes, and needle lengths for each vaccine used in your facility in the medication preparation area. Clearly identify diluents if the manufacturer's label could mislead staff into believing the diluent is the vaccine itself. Integrate vaccine administration training into orientation and other appropriate education requirements. Provide education when new products are added to inventory or recommendations are updated. Use standing orders, if appropriate. |

1. National Coordinating Council for Medication Error Reporting and Prevention, <https://www.nccmerp.org/about-medication-errors>



VAERS website at <https://vaers.hhs.gov/reportevent.html>
 * At this time, COVID-19 vaccination has additional VAERS reporting requirements, including required reporting of vaccine administration errors. Please see <https://vaers.hhs.gov/fao.html> for more information.

Interim Clinical Considerations

- Interim Clinical Considerations for Use of COVID-19 Vaccines Currently Approved or Authorized in the United States:
<https://www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html>
- FAQs for the Interim Clinical Considerations:
<https://www.cdc.gov/vaccines/covid-19/clinical-considerations/faq.html>

The screenshot shows the CDC website page for COVID-19 Vaccination Interim Clinical Considerations. The page has a dark green header with the text "Vaccines & Immunizations". Below the header, there is a breadcrumb trail: "CDC > COVID-19 Vaccination > Interim Clinical Considerations". The main content area is titled "Interim Clinical Considerations for Use of COVID-19 Vaccines Currently Approved or Authorized in the United States". A summary of recent changes (last updated June 24, 2022) is provided, including a bullet point: "New guidance for use of Moderna COVID-19 Vaccine in children and adolescents ages 6-17 years". There is a "Reference Materials" section with several links, including "Summary Document for Interim Clinical Considerations", "Interim COVID-19 Immunization Schedule", "At-A-Glance COVID-19 Vaccination Schedule (NEW 6/24/2022)", "Moderna COVID-19 Vaccine for Children who Transition from a Younger to Older Age Group (NEW 6/24/2022)", and "Pfizer-BioNTech for Children who Transition from a Younger to Older Age Group (NEW 6/24/2022)". A "Get Email Updates" button is also present. On the left side, there is a navigation menu with categories like "Product Info by U.S. Vaccine", "Interim Clinical Considerations", "Clinical Care", "Provider Requirements and Support", "Training and Education", "Vaccine Recipient Education", "Health Departments", "Planning & Partnerships", and "Vaccine Effectiveness Research".



Clinical Resources

- US COVID-19 Vaccine Product Information:

<https://www.cdc.gov/vaccines/covid-19/info-by-product/index.html>

The screenshot displays the CDC Vaccines & Immunizations website for COVID-19. The page title is "U.S. COVID-19 Vaccine Product Information". A navigation menu on the left lists various resources: Product Info by U.S. Vaccine (expanded to show Pfizer-BioNTech Vaccines, Moderna Vaccine, Janssen/J&J Vaccine, EUA, EUI, FAQs for Healthcare Professionals, Interim Clinical Considerations, Clinical Care, Provider Requirements and Support, and Training and Education). The main content area features a search bar, a language selector for "Español", and a descriptive paragraph: "Find a suite of information and materials that are needed for each specific COVID-19 vaccine that cover administration, storage and handling, safety, and reporting." Below this are three filter buttons: "Pfizer-BioNTech", "Moderna", and "Janssen/J&J". Two featured content boxes are visible: "Interim COVID-19 Immunization Schedule for Ages 5+" with a calendar icon and "Prevaccination Screening Form COVID-19 Prevaccination Guidelines" with a checklist icon. The latter includes a link to download a checklist in multiple languages, listing options such as Arabic, Dari, English, French (Canada), Haitian Creole, Korean, Pashto, Portuguese (Portugal), Simplified Chinese, Spanish, Ukrainian, and Vietnamese.





CT WiZ/VAMS Update

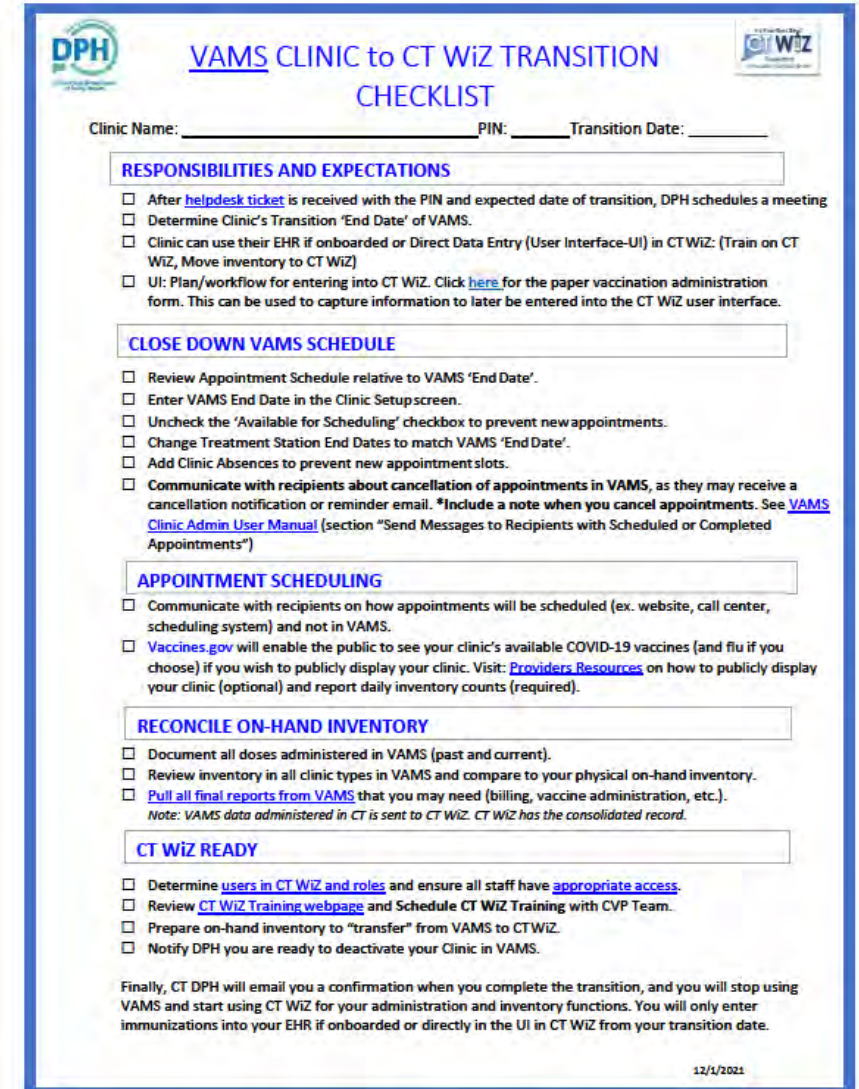
CT WiZ Updates



The Recommender in CT WiZ

New Immunization Program Webpage *Coming Soon!*

VAMS to CT WiZ Transition

- Effective 7/1/2022, the CT WiZ law was amended mandating all vaccinating providers to report electronically to CT WiZ.
- DPH will help VAMS clinics transition to CT WiZ. Submit a [helpdesk ticket](#) with your PIN to start the process.
- DPH will schedule a call to review the [transition process](#) with your clinic.



 **VAMS CLINIC to CT WiZ TRANSITION CHECKLIST** 

Clinic Name: _____ PIN: _____ Transition Date: _____

RESPONSIBILITIES AND EXPECTATIONS

- After [helpdesk ticket](#) is received with the PIN and expected date of transition, DPH schedules a meeting
- Determine Clinic's Transition 'End Date' of VAMS.
- Clinic can use their EHR if onboarded or Direct Data Entry (User Interface-UI) in CTWiZ: (Train on CT WiZ, Move inventory to CT WiZ)
- UI: Plan/workflow for entering into CT WiZ. Click [here](#) for the paper vaccination administration form. This can be used to capture information to later be entered into the CT WiZ user interface.

CLOSE DOWN VAMS SCHEDULE

- Review Appointment Schedule relative to VAMS 'End Date'.
- Enter VAMS End Date in the Clinic Setupscreen.
- Uncheck the 'Available for Scheduling' checkbox to prevent new appointments.
- Change Treatment Station End Dates to match VAMS 'End Date'.
- Add Clinic Absences to prevent new appointment slots.
- Communicate with recipients about cancellation of appointments in VAMS, as they may receive a cancellation notification or reminder email. *Include a note when you cancel appointments. See [VAMS Clinic Admin User Manual](#) (section "Send Messages to Recipients with Scheduled or Completed Appointments")

APPOINTMENT SCHEDULING

- Communicate with recipients on how appointments will be scheduled (ex. website, call center, scheduling system) and not in VAMS.
- [Vaccines.gov](#) will enable the public to see your clinic's available COVID-19 vaccines (and flu if you choose) if you wish to publicly display your clinic. Visit: [Providers Resources](#) on how to publicly display your clinic (optional) and report daily inventory counts (required).

RECONCILE ON-HAND INVENTORY

- Document all doses administered in VAMS (past and current).
- Review inventory in all clinic types in VAMS and compare to your physical on-hand inventory.
- [Pull all final reports from VAMS](#) that you may need (billing, vaccine administration, etc.).
Note: VAMS data administered in CT is sent to CT WiZ. CT WiZ has the consolidated record.

CT WiZ READY

- Determine [users in CT WiZ and roles](#) and ensure all staff have [appropriate access](#).
- Review [CT WiZ Training webpage](#) and Schedule CT WiZ Training with CVP Team.
- Prepare on-hand inventory to "transfer" from VAMS to CT WiZ.
- Notify DPH you are ready to deactivate your Clinic in VAMS.

Finally, CT DPH will email you a confirmation when you complete the transition, and you will stop using VAMS and start using CT WiZ for your administration and inventory functions. You will only enter immunizations into your EHR if onboarded or directly in the UI in CT WiZ from your transition date.

12/1/2021

COVID-19 Vaccine Updates

- CT WiZ has been updated with the new young pediatric COVID-19 vaccines:

- [Vaccines Supplied By CT CVP 62422.pdf](#)

Please ensure you enter the correct COVID-19 vaccines.



STATE OF CONNECTICUT
DEPARTMENT OF PUBLIC HEALTH
IMMUNIZATION PROGRAM

Vaccines supplied by the Connecticut Vaccine Program as of June 24, 2022

| Vaccine | Brand Name and Packaging | NDC | Manufacturer | CVX | CT WiZ Drop Down Selection |
|------------------------------|---|---------------|---------------------|-----|-----------------------------|
| COVID Tris-Suc (PFR 6mo-4y) | PFR COVID Tris-sucrose (10 x 10(0.2mL/dose) MDV) | 59267-0078-04 | Pfizer, Inc. | 219 | COVID Tris-Suc (PFR 6mo-4y) |
| COVID Tris-Suc (PFR 12+) | PFR COVID Tris-sucrose (10 x 6 (0.3mL/dose) MDV) | 59267-1025-04 | Pfizer, Inc. | 217 | COVID Tris-Suc (PFR 12+) |
| COVID Tris-Suc (PFR 16+) | Pfizer Comirnaty COVID-19 (10 x 2.0mL MDV) | 00069-2025-10 | Pfizer, Inc. | 217 | COVID Tris-Suc (PFR 12+) |
| COVID Tris-Suc (PFR 5-12) | Pfizer COVID-19 (.2mL/dose) MDV) | 59267-1055-04 | Pfizer, Inc. | 218 | COVID Tris-Suc (PFR 5-12) |
| COVID-19 (MOD) 6mo - <6yr | Moderna COVID-19 Ped 6mo - 6yr (10 x 2.5mL Vials) | 80777-0279-99 | Moderna | 228 | COVID-19 (MOD) 6mo - <6yr |
| COVID-19 (MOD) 6-11, Booster | Moderna COVID-19 6-11yr (10 x 2.5mL Vials) | 80777-0275-99 | Moderna | 221 | COVID-19 (MOD) Booster |
| COVID-19 mRNA (MOD) | Moderna COVID-19 (10 x 10 dose 5.0 mL MDV) | 80777-0273-99 | Moderna | 207 | COVID-19 mRNA (MOD) |
| COVID-19 Vector-NR (JSN) | Janssen COVID-19 (10 x 5 dose 5.0 mL MDV) | 59676-0580-15 | Janssen | 212 | COVID-19 Vector-NR (JSN) |
| DTaP | Infanrix (0.5 mL x 10 syr) | 58160-0810-52 | GlaxoSmithKline | 20 | DTaP |
| DTaP (Daptacel) | Daptacel (0.5 mL x 10 vials) | 49281-0286-10 | Sanofi Pasteur | 106 | DTaP (Daptacel) |
| DTaP-HepB-IPV | Pediarix (0.5 mL x 10 syr) | 58160-0811-52 | GlaxoSmithKline | 110 | DTaP-HepB-IPV |
| DTaP-Hib-IPV (Pentac) | PENTACEL (SDV; 5 PACK) | 49281-0511-05 | Sanofi Pasteur | 120 | DTaP-Hib-IPV (Pentac) |
| DTaP-IPV | Kinrix (0.5 mL x 10 syr) | 58160-0812-52 | GlaxoSmithKline | 130 | DTaP-IPV |
| DTaP-IPV | Quadracel 10 pack | 49281-0564-15 | Sanofi Pasteur | 130 | DTaP-IPV |
| DTaP-IPV-Hib-Hep B | Vaxelis (10 x 0.5mL Syringes) | 63361-0243-15 | MSP Vaccine Company | 146 | DTaP-IPV-Hib-Hep B |
| Flu MDCK Quad P-Free Inj | Flucelvax Quad 2021-2022 (10 x 0.5mL Syringes) | 70461-0321-03 | Seqirus | 171 | Flu MDCK Quad P-Free Inj |
| Hep A, adult | Havrix (1 mL x 10 syr) | 58160-0826-52 | GlaxoSmithKline | 52 | Hep A, adult |
| Hep A, ped/adol, 2D | Havrix (0.5 mL x 10 syr) | 58160-0825-52 | GlaxoSmithKline | 83 | Hep A, ped/adol, 2D |
| Hep A, ped/adol, 2D | Vaqta (0.5 mL x 10 syr) | 00006-4095-02 | Merck & Co, Inc. | 83 | Hep A, ped/adol, 2D |
| Hep B, adult | Heplisav-B | 43528-0003-05 | Dynavax | 43 | Hep B, adult |
| Hep B, ped/adol | Enerix B (0.5 mL x 10 syr) | 58160-0820-52 | GlaxoSmithKline | 08 | Hep B, ped/adol |
| Hep B, ped/adol | Recombivax (0.5 mL x 10 vials) | 00006-4981-00 | Merck & Co, Inc. | 08 | Hep B, ped/adol |

CT WiZ Recommender

- When there is a new vaccine recommendation or a change in the vaccine schedule, it requires the **CT WiZ Recommender to be updated in a new CT WiZ release.**
 - If you see a valid dose with an **! (invalid)**, when the Recommender is updated, the **!** will disappear and the dose will display as valid.
- Tip: Print the Full immunization record until the Recommender is updated. (The COVID-19 only record only displays valid doses.)

New Immunization Program Webpage!



Question and Answers

To ask a question, please raise your hand using the hand icon on your screen, type your question in the chat box or if you are on the phone press *6 to unmute yourself.

If you have additional questions after the meeting, please feel free to email them to DPH.Immunizations@ct.gov

You can fill out a help desk ticket by visiting <https://dph-cthelpdesk.ct.gov/Ticket>