Vaccine Sumit

September 7, 2023: Rockford



Illinois Chapter

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American Academy of Pediatrics

DEDICATED TO THE HEALTH OF ALL CHILDREN®

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SESSION 1: IDPH VFC PROGRAM

Stephanie Atella



- MPH, CHES
- Chief Program Officer
- She/Her/Hers

After this Session

Use the Illinois
Comprehensive
Automated
Immunization Registry
Exchange (I-CARE)
portal to order and
maintain vaccines.

2

Describe inventory reconciliation best practices for I-CARE and VFC I-CARE mandates.

3

Implement effective vaccine storage and handling practices and keep vaccine waste to a minimum.

Overview of the VFC Program

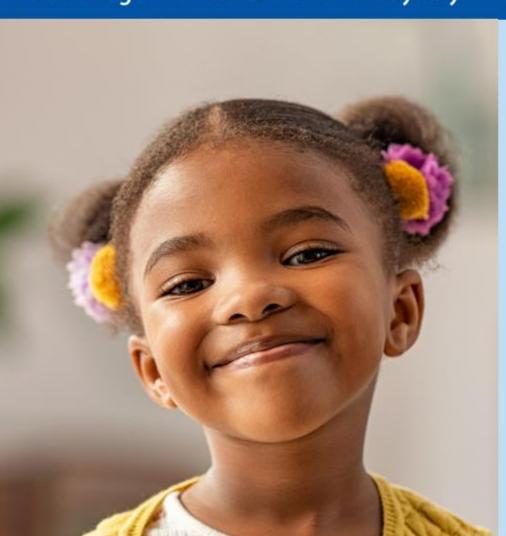
- IL's VFC program is federally-funded by the Centers for Disease Control and Prevention (CDC).
- Provides vaccines at no cost to children who may not otherwise get them due to cost.
 - Children who are eligible to receive VFC vaccines need to be 18 years and younger and one of the following:
 - Uninsured or Underinsured
 - o Medicaid Title 19 or 21 eligible.
 - American Indian or Alaskan Native.
- Children who are underinsured can access VFC vaccines at federally qualified health centers (FQHCs), rural health clinics, and some local health departments that are deputized by FQHCs or RHCs.

Benefits

Vaccines for Children

Protecting America's children every day

The Vaccines for Children (VFC) program helps ensure that all children have a better chance of getting their recommended vaccines. VFC has helped prevent disease and save lives.



CDC estimates that vaccination of children born between 1994 and 2021 will:

prevent **472 million** illnesses



(29.8 million hospitalizations)

help avoid 1,052,000 deaths





save nearly \$2.2

trillion in total
societal costs
(that includes \$479 billion in direct costs)





more than \$5,000 for each American

polated 2021 analysis using methods from Benefits from Immunication during the Vaccines for Children Program Era.—United States, 1994-2021:

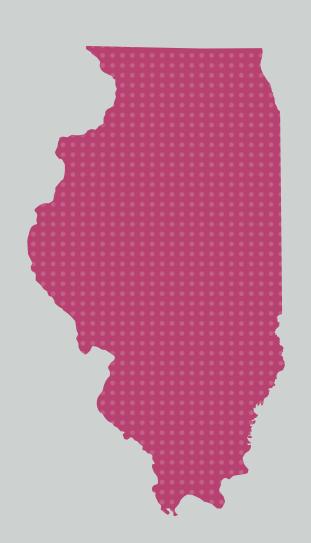


www.cdc.gov/vaccines/vfcprogram/

VFC Provider Requirements

VFC providers must:

- ✓ Be licensed in Illinois to administer vaccines to children aged 18 and younger.
- ✓ Be willing and able to follow all VFC program requirements, policies, and procedures, including participation in site visits and educational opportunities.
- ✓ Have capacity to order, receive, manage, store, and monitor the temperature of public vaccines.
- ✓ Be open at least four consecutive hours for three days a week to receive VFC vaccines.



I-CARE

- The I-CARE Registry is an electronic web-based immunization data registry operated by the Illinois Department of Public Health (IDPH) as authorized by the Immunization Data Registry Act, 410 ILCS 527.
- All VFC providers must be enrolled in I-CARE.
 - Enrollment and vaccine management is completed in I-CARE.
- Must be able to provide individual patient immunization records on how each VFC vaccine was administered.
- Patient immunization records can be entered manually or electronically through the provider's electronic medical record.

Submitting and Exchanging Data

- Share Electronic Health Record (EHR) with I-CARE using HL7 data exchange.
 - Contact EHR vendor to determine if your system is HL7 compatible.
 - May need to acquire an additional interface for your EHR to send and exchange immunization data.

For more information on HL7 please visit IDPH's <u>I-CARE site</u>. If you have questions, please contact dph.icare@illinois.gov.

Provider Profile

- Providers must submit a Provider Population Profile at initial program enrollment and at least annually or when order patterns indicate a change.
- All VFC programs must determine individual provider populations served and associated vaccine need by fund type.
- This ensures publicly purchased vaccines are distributed in amounts representing the provider population served and to adjust as populations change.

Record Keeping

VFC providers must comply with:

- Distributing the most current vaccine information statements (VISs) for all vaccines included in National Childhood Vaccine Injury Act (NCVIA).
 - o Immunize.org: Vaccine Information Statements (available in 47 languages)
- Reporting adverse reactions to VAERS.

Table 1. G	uidance for l	Jse of Va	ccine Inform	ation Statements

(Source: AAP Committee on Medical Liability. Medical Liability for Pediatricians, 6th Edition. 2004)

Distribution	Documentation in the Patient's Medical Record	
Must be provided each time a National Vaccine Injury Compensation Program (VICP)-covered vaccine is administered*	Vaccine manufacturer, lot number, and date of administration*	
Given to parent, legal guardian, or patient (non-minor) to keep*	Name and business address of the physician administering the vaccine*	
Must be the current version†	Vaccine Information Statement version date and date it is provided†	
Can provide (not substitute) other written or audio- visual materials as necessary‡	Site (eg, deltoid area), route of administration (eg, intramuscular), and expiration date of the vaccine [‡]	

^{*}Required under the National Childhood Vaccine Injury Act.

Required under Centers for Disease Control and Prevention instructions implementing the National Childhood Vaccine Injury Act.

^{*}Recommended by the American Academy of Pediatrics.

Record Keeping

The National Childhood Vaccine Injury Act (NCVIA) and/or CDC requires physicians to document the:

- Name of vaccine administered
- Date of vaccine administration
- Vaccine manufacturer
- Vaccine lot number
- Name, title, and business address of the healthcare professional who administered the vaccine
- Date the VIS was provided to the parent/guardian and VIS version date

The AAP recommends also recording the:

- Site and route of administration
- Vaccine expiration date
- Statement indicating that the VIS was provided and discussed with the parent
- Any vaccine under CDC contract requires a VIS.

The <u>CDC</u> requires that patient VFC eligibility screening must take place with each immunization visit.

Maintain records for a minimum of three years or longer, if required by state law (even in the case of provider retirement or provider location closure).

Required Documentation

- Providers must complete CDC's Provider Agreement.
- The medical director in a group practice must be authorized to administer pediatric vaccines under state law.
- The provider signing the Provider Agreement on behalf of a multi-provider practice must have authority to sign on behalf of the entity.
- All licensed providers in an enrolled practice must be listed with professional license numbers and individual NPI numbers (VFC Enrollment Form).

Recertification of Annual Enrollment

- Provider agreement forms (signed by medical director or equivalent in a group practice*).
- The practitioner will be held accountable for compliance by the entire organization and its VFC providers with the responsible conditions outlined in the Provider Enrollment Agreement.

Recertification of Annual Enrollment

All VFC providers must recertify their enrollment annually to continue participating in the VFC program.

Annual enrollment is submitted in I-CARE.

Additionally, providers should:

- Review and Agree to the VFC Eligibility and the VFC Loss and Replacement Policies.
- Review, sign, and upload the VFC Provider Agreement.

Provider Unenrollment

- Either the Provider or the Illinois VFC program may decide to terminate the provider agreement at any time.
- Providers who wish to terminate the provider agreement must:
 - Complete unenrollment form.
 - Stop using VFC vaccines as of the withdrawal date.
 - Return any unused VFC vaccines back within 30 days.
- Examples of why IDPH may terminate the provider agreement include:
 - Provider has not ordered vaccine in the past 12 months.
 - A provider is on the List of Excluded Individual and Entities (LEIE) list maintained by Office of the Inspector General.
 - Failure to comply with requirements.

Vaccine Staff and Training - Vaccine Coordinators

Identify: a primary vaccine coordinator and at least one backup vaccine coordinator for each facility.

The primary and backup vaccine coordinators:

- Responsible for ordering, receiving, rotating, and monitoring vaccines.
- Responsible for ensuring all vaccines are stored and handled correctly.
- Must be fully trained on routine and emergency SOPs for vaccine ordering, storage, handling, transport, and inventory management.

More information about coordinator responsibilities can be found in the <u>Vaccine</u> for <u>Children Program Manual for Illinois VFC Providers</u>.

Notify IDPH when there is a change in vaccine coordinators or medical director.

Education Requirements

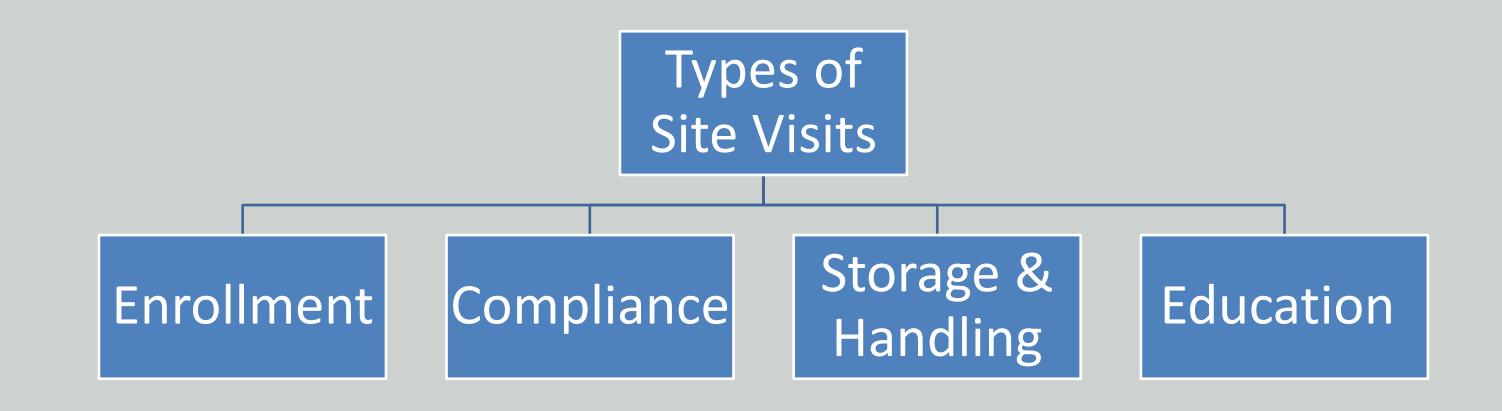
- Annual vaccine storage and handling training.
 - Documentation of training must be retained and submitted with annual enrollment (reviewed during site visits).
- Education is available through:
 - Some VFC site visits.
 - This summit ©
 - CDC online training with both of the following modules:
 - You Call The Shots Module 10 Storage and Handling
 - You Call the Shots <u>Module 16 Vaccines for Children Program</u>

Staff Training

- All staff members who:
 - Receive vaccine deliveries
 - Handle or administer vaccines
 - should be trained in vaccinerelated practices and storage and handling SOPs.
- Training must be documented on the vaccine management plan.



VFC Site-Visits



VFC Enrollment Visits

- All providers (newly enrolling or re-enrolling after an absence) must have an enrollment site visit **before** being approved to order VFC vaccines.
- The purpose of this visit is to:
 - Educate providers about VFC program requirements.
 - Educate providers on proper vaccine storage and handling.
 - Certify providers have the appropriate resources to implement requirements.
- Providers should be prepared for follow-up visits during the first year.

VFC Compliance Visits

- VFC providers agree to program site visits that will determine compliance with requirements.
- A new provider must be enrolled and active in the VFC program for 3-6 months before a receiving a compliance site visit.
- Compliance visit includes review of and ensuring compliance with:
 - o Provider Profile.
 - Vaccine ordering and inventory management.
 - Policies and procedures and vaccine management plan.
 - Vaccine storage and handling equipment, procedures, and documentation.
 - VFC screening requirements and billing practices.
 - All ACIP vaccines are available to VFC-eligible patients.
 - VFC-related document retention.

VFC Storage & Handling Visits

- Storage and Handling visits may be announced (scheduled) or unannounced.
- IDPH is required to complete unannounced storage and handling site visits for a percentage of providers each year.
- Compliance visit includes review of and ensuring compliance with:
 - Vaccine inventory management.
 - Vaccine storage and handling equipment and monitoring.
 - Vaccine storage and handling procedures and Vaccine management plan.
 - Appropriate storage and handling related documentation.

Vaccine Replacement

 VFC providers agree to replace vaccines purchased with state and federal funds that are deemed non-viable due to provider negligence on a dosefor-dose basis with privately purchased vaccines.



Fraud and Abuse

By enrolling in the VFC program, you agree to comply with all program requirements

Examples of fraud and abuse:

- Providing VFC vaccines to non-VFC eligible children.
- Billing a patient or third party for a VFC vaccine.
- Denying VFC eligible children a VFC vaccine due to inability to pay an administration fee.
- Failing to screen for and document eligibility at each visit.
- o Failing to properly maintain VFC records and requirements.
- Failing to properly store and handle VFC vaccines, etc.
- The Department will investigate to determine intentional or unintentional fraud/misuse of lintentional, further investigation is necessary.

Patient Eligibility Screening and Documentation*

- Providers must screen, document, and verify VFC eligibility with every vaccine visit before administering vaccines.
- Use the <u>MEDI system</u> or equivalent system (with HFS 270/271 electronic transaction data).
- The Patient Eligibility Screening Form provides a means of recording responses to VFC eligibility questions.
 - The provider, parent, or guardian may complete the VFC eligibility portion of the form.
 - Verification of parent/guardian responses is not required.

Memorandum of Understanding (MOU) with FQHC or Rural Health Center

- Underinsured VFC-eligible children can only receive VFC vaccine from a Federally Qualified Health Center (FQHC) or Rural Health Clinic (RHC).
- A local health department (LHD) can be deputized by an FQHC or RHC to allow the LHD to vaccinate underinsured VFC-eligible children.
- A Memorandum of Understanding (MOU) form is available through IDPH. It must be signed by three parties: the LHD, the FQHC or RHC, and IDPH.
- If a provider is not a FQHC, RHC or a deputized local health department, they should refer underinsured VFC-eligible children to a qualified provider location.

Important Announcements from IDPH Sent to VFC Providers on 9/6/2023

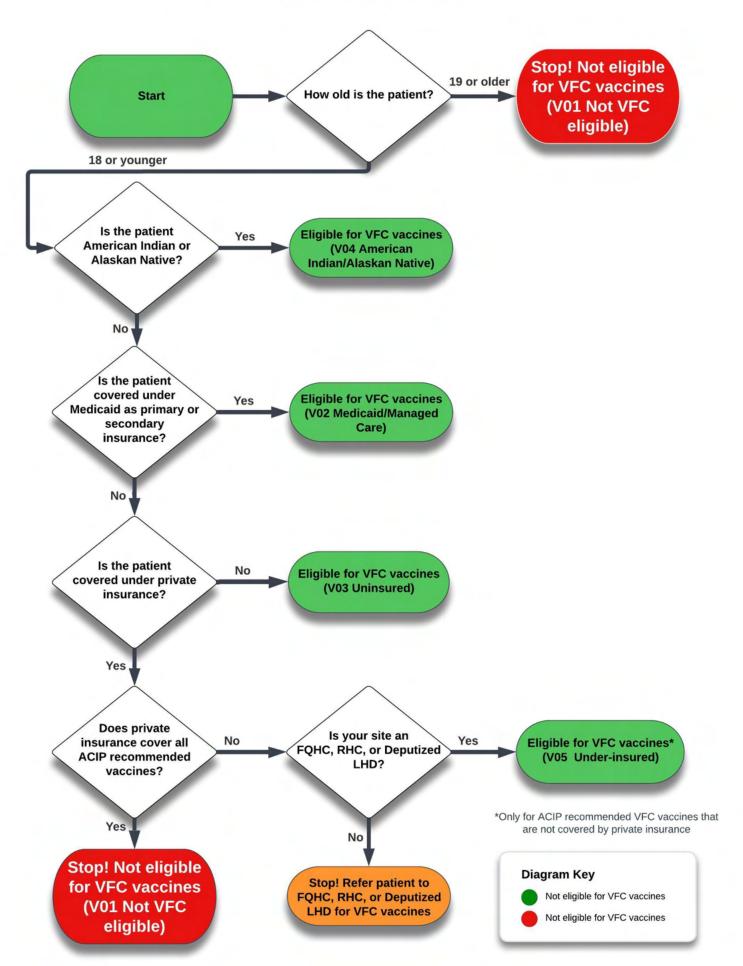
- Because of Medicaid Expansion, all Medicaid-enrolled children 18 years of age or younger are now VFC-eligible regardless of their Title 19, Title 21, or State-Funded Medicaid Status.
- How this impacts providers:
 - Medi should be checked at each encounter to ensure the patient is enrolled in Medicaid, but providers no longer need to check for Title 19, Title 21, or State Funded status.
 - When marking the patient VFC eligibility in your patient's chart and in I-CARE, the VO2 Medicaid/Medicaid Manage Care status should be selected for all Medicaid-enrolled children.

VFC Eligibility Status Code Information and Associated Inventory Deductions

DOGGOTOTIO				
VFC Eligibility Status Code	VFC Code will deduct from this Inventory in I-CARE	Comments		
V00: Eligibility not determined/unknown	Private	Do not use for public vaccine programs.		
V01: Not VFC eligible	Private	Do not use for public vaccine programs.		
V02: Medicaid/Medicaid Managed Care	VFC	Use for all Medicaid-enrolled children through 18 years of age.		
V03: Eligible-Uninsured	VFC			
V04: Eligible-American Indian/Alaska Native	VFC			
V05: Eligible-Federally Qualified Health Center patient (under-insured)	VFC			
V07: Local program eligibility	Private	Not currently used by Illinois or Chicago VFC.		
V22: CHIP	VFC	This status is no longer applicable to Illinois or Chicago VFC patients. It will continue to deduct from VFC while providers transition to this new process, but providers should stop using this code. Use "V02: Medicaid/Medicaid Managed Care" instead.		
V23: 317	317			
V24: Medicare	Private	Not currently used by Illinois or Chicago VFC.		
V25: State program eligibility	Private	Not currently used by Illinois or Chicago VFC.		

ILLINOIS VFC ELIGIBILITY DECISION TREE

Effective 09/06/2023



Eligibility

VFC Eligibility Criteria	Definition
American Indian or Alaska Native (AI/AN)	This population is defined by the Indian Health Care Improvement Act (25 U.S.C. 1603). (AI/AN children are VFC-eligible under any circumstance.)
Medicaid-eligible	Children who are eligible for the Medicaid program. For the purposes of the VFC program, the terms "Medicaid-eligible" and "Medicaid enrolled" are used interchangeably.
Uninsured	Children not covered by any health insurance plan.
Underinsured	 Underinsured means the child has health insurance, but the insurance policy: Does not include any vaccines; Does not include all vaccines recommended by the Advisory Committee on Immunization Practices (ACIP); or Has a fixed dollar limit or cap for vaccines. Underinsured children are only eligible to receive VFC vaccines at a FQHC, RHC, or a deputized provider.

Eligibility - American Indian or Alaska Native (AI/AN)

- Al/AN children are VFC-eligible under any circumstance.
 - Participation is voluntary.
- When an Al/AN child also fits a second VFC eligibility category, the provider should always choose the category that will cost less for the family.
 - Depending on the facility, when an Al/AN parent chooses to have their child vaccinated, the parent may be responsible for the vaccine administration fees if the vaccines are delivered through the VFC program.
 - o If the child has private insurance, it may result in fewer out-of-pocket costs for the child to receive privately purchased vaccinations since the insurance would likely cover the vaccine without cost-sharing.
 - o If the AI/AN child is also Medicaid eligible, Medicaid should be used for the administration fee because it will provide the least out of pocket expense.

Eligibility - Insured Children with Medicaid

- Some children may have a private primary health insurance plan with Medicaid as their secondary insurance.
 - These children are considered VFC-eligible because of their Medicaid enrollment.
 - Their parents are not required to participate in the VFC program.
- There are billing options and the provider should choose the option that is most cost-effective for the family.
 - The parent of a child with Medicaid as secondary insurance should never be billed for a vaccine or an administration fee.

Eligibility - Insured Children with Medicaid

Option 1: The provider can administer VFC vaccines and bill Medicaid for the administration fee.

Considerations:

- Easiest way for a provider to use VFC vaccines and bill Medicaid for the administration fee.
- ➤ No out-of-pocket costs to the parent for the vaccine or the administration fee.

Option 2: The provider can administer private stock vaccines and bill the primary insurance carrier for both the cost of the vaccine and the administration fee.

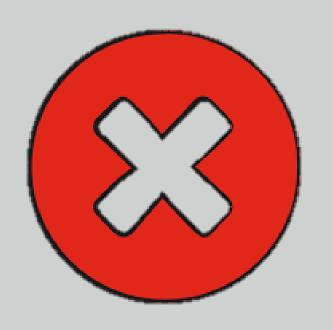
Considerations:

Provider may be reimbursed a higher dollar amount if privately purchased vaccine is administered and both the vaccine and the administration fee are billed to the primary insurer.

Child's Insurance Status	VFC-Eligible?	VFC Eligibility Category
Enrolled in Medicaid	Yes	Medicaid (V02).
Has private health insurance plan with Medicaid as secondary Insurance.	Yes	Medicaid (V02). The provider should choose the option that is most cost-effective for the family.
Has health insurance covering all vaccines but has not yet met plan's deductible or paid for other services received at visit.	No	Insured (V01). This applies even when primary insurer would deny reimbursement for the cost of the vaccine and its administration because the plan's deductible has not been met.
Has health insurance covering all vaccines but has not yet met plan's deductible or paid for other services received at visit and has Medicaid as secondary insurance.	Yes	Medicaid (V02).
Has health insurance covering all vaccines, but the plan has a fixed dollar limit or cap on amount that it will cover.	Yes	Insured (V01) until the fixed dollar limit is met. Underinsured (V05) after the fixed dollar limit is reached.
Has an insurance plan that does not cover all ACIP-recommended vaccines.	Yes	Underinsured (V05). Child can only receive vaccines not covered by the plan.
Has health insurance, but plan does not cover any vaccines.	Yes	Underinsured (V05). With implementation of ACA, this situation should be rare.
Has no health insurance coverage.	Yes	Uninsured (V03).
Has private health insurance that covers all vaccinations and is AI/AN.	Yes	AI/AN (VO4). The provider should choose the eligibility category most cost effective for the family.
Has Medicaid and is Al/AN.	Yes	Medicaid (VO2) or AI/AN (VO4). Providers should use Medicaid for the administration fee (least out-of-pocket expense for family).

No Charge for Vaccines*

- Patient cannot be charged for publicly purchased vaccine.
- Do not bill any individual or other third-party payer for the cost of VFC-supplied or other vaccines purchased through CDC federal contracts.



Administration Fees

- Bill only Medicaid for the administration fee for VFC-eligible children enrolled in Medicaid.
 - o Administration fees are per vaccine and not per antigen.
- The vaccine administration fee for **non-Medicaid** VFC-eligible children **must not** exceed \$23.87 per dose.
- VFC providers may issue a single bill for the administration fee for **non-Medicaid** VFC-eligible children within 90 days of vaccine administration.
- Unpaid VFC vaccine administration fees may not be sent to collections and VFC providers may not refuse to vaccinate an eligible child whose parents have unpaid vaccine administration fees.

Vaccine Management

- Providers should follow VFC storage and handling requirements based on CDC's Vaccine Storage and Handling Toolkit including:
 - Ordering vaccines.
 - Utilizing required equipment.
 - Digital data loggers.
 - Vaccine cold chain.



Vaccine Management Plan

Procedures Procedures for for vaccine emergency Contact info ordering, Provider and situations Documented for current Proper receiving, vaccine (transport, training primary and storage and inventory coordinator related to equipment control, stock backup handling roles & malfunction, vaccine vaccine practices. rotation, and responsibilities power failure, management. coordinators. handling and natural vaccine loss disaster). and waste.

Plans must be updated annually or more frequently as needed.

Vaccine Management: Ordering Vaccine*

Vaccine ordering is completed through I-CARE.

Should be up-to-date before submitting an order:

- Patient immunization records.
- Temperature logs for all appliances.
- All data logger certificates of calibration are valid and not expired.
- All temperature excursions must have a vaccine incident report on file.
- No expired vaccines are showing in the clinic's inventory.
- The clinic's inventory in I-CARE matches the physical inventory.
- o The clinic's inventory in I-CARE is not showing any negative balances.
- Clinic must be open at least three days a week with at least four consecutive hours a day to be able to receive a delivery.
 - Delivery hours must be entered and updated in I-CARE, including specifying lunch hours or other closures, when placing orders.

Vaccine Management: Ordering Vaccine*

- Order and stock enough vaccine to meet patient demand for up to three months.
 - Consider patient numbers, patient age, vaccine uptake, etc.
 - o Smaller, more frequent orders help reduce the impact of incidents that may contribute to vaccine loss. **Providers may order as often as necessary**.
- Ensure each VFC vaccine administered is entered into I-CARE. Options are:
 - Direct entry
 - Electronic transmission to I-CARE from an electronic health record (EHR)

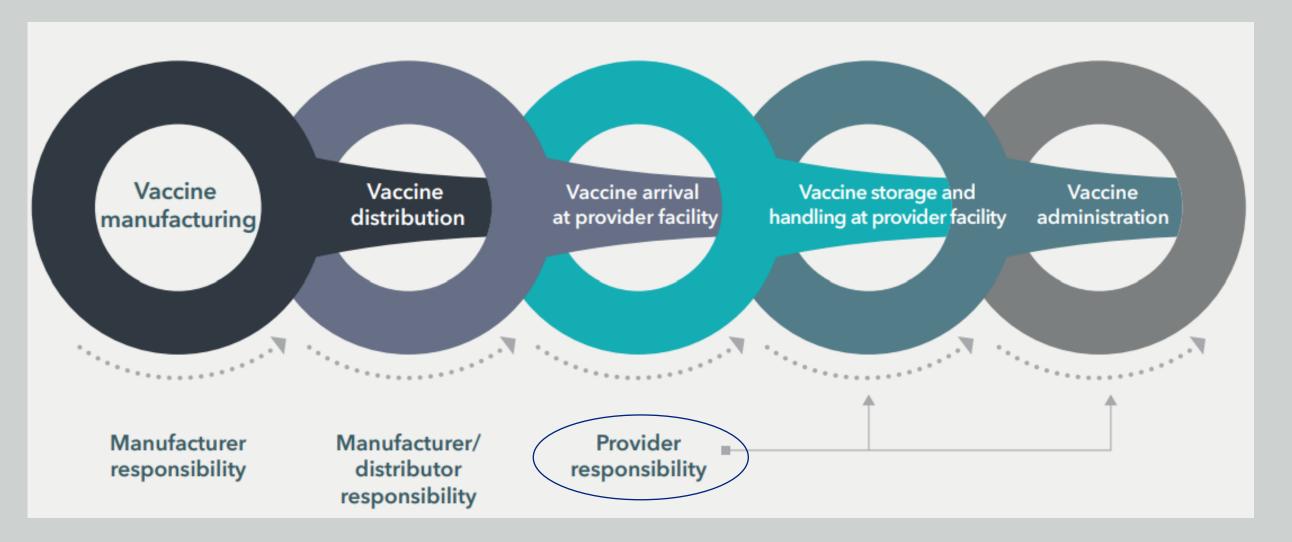
To set EHR to transmit data to I-CARE: DPH.HL7ICARE@Illinois.gov

Vaccine Management: Borrowing Vaccine

- Borrowing between public VFC & private vaccine inventory is not allowed.
- Transfers of VFC vaccine between VFC clinics are allowable with permission from IDPH VFC Program and proper transport storage equipment.
 - A transfer request form is available in I-CARE.

Vaccine Management: Cold Chain

- Vaccine cold chain must be maintained (ensures potency and useability).
 - Helps save money and avoid re-vaccination.



Vaccine
appearance is
NOT a reliable
indicator that
vaccines have
been stored in
appropriate
conditions.

Vaccine Management: Receiving Vaccine

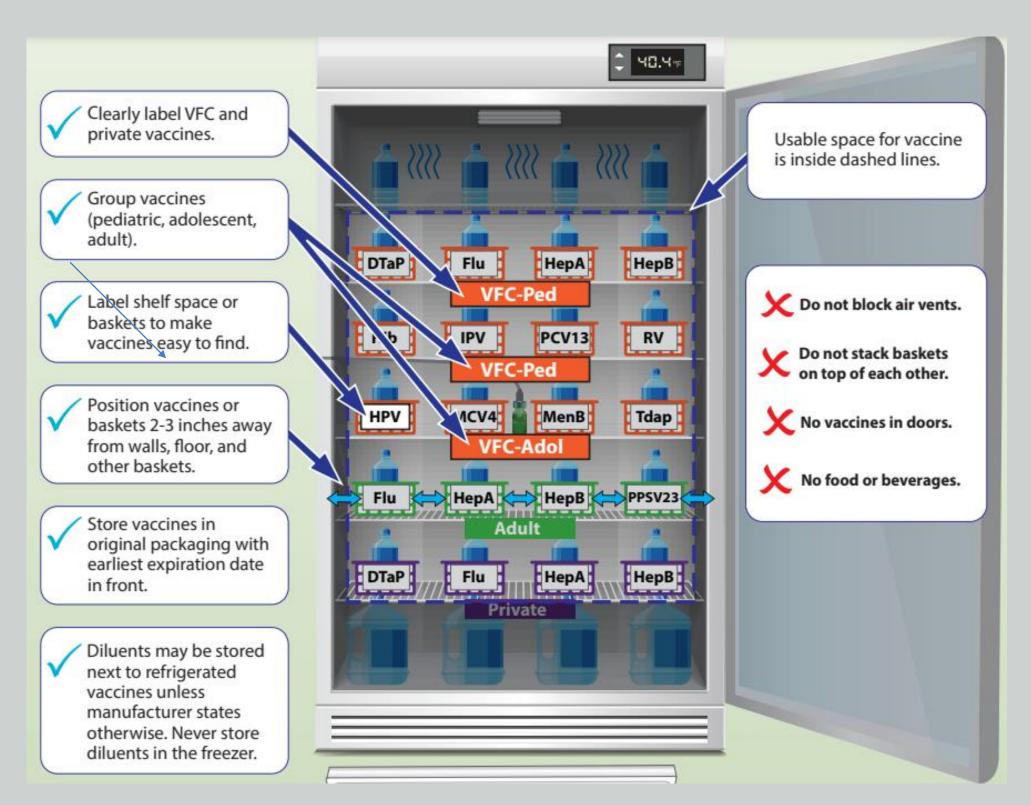
- Vaccine & diluent should be immediately unpacked, stored at recommended temperatures, and documented upon arrival.
 - Do not store shipment box in vaccine storage unit the combined storage methods may be too cold.
- Check immediately for:
 - Physical damage of shipping container.
 - Correct products were received
 - Diluent and Vaccine expiration dates.
 - Cold chain monitor, if included.

Any issues should be reported within two hours to the Illinois VFC Program Services Staff at 217-785-1455.



Vaccine Management: Storing Vaccine

- Store vaccine by funding type (check I-CARE or packing slips).
- Separate units are not required.
 - VFC: VFC eligible patients only.
 - 317: 317-eligible adults or approved outbreak response.



Vaccine Management: Storing Vaccine

- Stock rotation and removal:
 - Rotate vaccine stock so the vials with the soonest expiration date are at the front (used first).
 - Immediately remove expired vaccine from stock.

TIP: Determine regular intervals for rotation (i.e., weekly), including when there is a vaccine delivery.

Vaccine Management: Required Equipment*

- Purpose-built or pharmaceutical-grade refrigerators and freezers are preferred.
 - Still needs to be approved and met with the guidelines and re-certified by approved source.
 - Stand-alone refrigerator and freezer units may also be used.
 - The Department does not allow combination household refrigerator/freezer units for the storage of vaccines obtained through the VFC program.
 - Never store vaccine in a dorm-style or bar-style combined refrigerator/freezer unit.







Vaccine Management: Required Equipment*

- Some purpose-built units separate public & private vaccine stock electronically.
 - o If electronic, an inventory printout must be available upon request.
- Power Supply:
 - o Plug in only one storage unit per electrical outlet
 - Use a safety-lock plug or an outlet cover.
 - Post "DO NOT UNPLUG" warning signs at outlets and on storage units.
 - Label fuses and circuit breakers to alert others not to turn off these units.
 - Use caution when using power outlets that can be tripped or switched off and avoid using:
 - Built-in circuit switches (may have reset buttons).
 - Outlets that can be activated by a wall switch.
 - Multioutlet power strips.

Vaccine Management: Required Equipment Cont.*

- Storage units should be placed in a well-ventilated room, between 68°F 77°F, and without anything blocking them.
 - Refrigerators should maintain temps between 2° C 8° C (36°F 46°F).
 - Freezers should maintain temps between -50° C and -15° C (-58°F +5°F).
 - Recommended to set temps in Celsius and record to 1 decimal place.
 - Temperatures should be recorded any time staff are in the clinic, at least 3x/week.
 - Record the current temp 2x/day and the min/max temps at the start of every day.
 - Doors should always remain closed consider using locks or alarms.
- It can take multiple days to stabilize the temp in a new or repaired unit.
 - Min and max temps should be recorded 2x/day for 2 to 7 days.
 - Once two consecutive days of temperatures are recorded within the recommended range, the unit is stable and ready for use.

Vaccine Management: Required Equipment Cont.*

- Vaccines should be stored in their original packaging with lids closed.
 - Food and beverages should never be stored in the same unit as vaccines.
 - Water bottles can be used in vaccine storage units to help stabilize the temperature.
 - Water bottles are not recommended for use with certain pharmaceutical-grade and purpose-built units. For such units, follow the manufacturer's guidance.



Vaccine Management: Digital Data Loggers

- Digital Data Loggers (DDLs) are required to continually monitor the temperature of vaccine.
 - Must have a valid Certificate of Calibration Testing (some units have DDLs built in).
- Data from DDLs is retrieved using special software or a website.
 - Data should be downloaded and reviewed at least weekly.
 - Records should be kept for three years.
- A back-up DDL must be available in case another fails, or calibration testing is required.

Vaccine Management: Digital Data Loggers Cont.

- All data loggers must have a certificate of calibration that is current (based on the manufacturer's recommended re-testing timeline as indicated on the certificate of calibration).
- Some purpose-built units have built-in DDLs. The purpose-built unit DDLs must meet the same requirements as DDLs for other VFC storage units
- A back-up DDL must be available in case another fails or for emergency transportation.
 - Calibration testing is required.
 - Should have a different calibration testing date than other DDLs so they do not all go through testing at the same time.

Vaccine Management: Digital Data Loggers Cont.

The DDL must be equipped with:

- A temperature probe or sensor.
- An active temperature display outside the unit that can be easily read without opening the unit's door.
- Continuous temperature monitoring and recording capabilities and capacity to routinely download data.

Additional recommended DDL features:

- Alarm for out-of-range temperatures
- Temperature display showing current, minimum, and maximum temperatures
- Low battery indicator
- Accuracy of +/-1°F (0.5°C)
- User-programmable logging interval (or reading rate) recommended at a maximum time interval of no less frequently than every 30 minutes.

Vaccine Management: Digital Data Loggers Cont.

Certificates
of
Calibration
Testing
must
include:

- Model/device number.
- Serial number.
- Date of calibration (report or issue date).
- Confirmation the instrument passed testing (or instrument in tolerance).

Certificates
of
calibration
must
indicate at
least one of
the following
items:

- Conforms to ISO 17025.
- Testing was performed by an ILAC/MRS Signatory body accredited laboratory.
- Is traceable to the standards maintained by NIST.
- Meets specifications and testing requirements for the American Society for Testing and Materials (ASTM) Standard E2877 tolerance Class F (0.5 °C) or better.

Temperature Excursions

- Any temperature reading outside the recommended ranges in the manufacturers' package inserts.
 - o Manufacturers will help determine if vaccine is still viable after an excursion.
- Providers should immediately review their storage and handling policies and take the appropriate actions.
- Complete the Vaccine Incident Report in I-CARE.
- Unsure if an excursion occurred?
 - Mark vaccine "DO NOT USE!"
 - Do not use or discard until manufacturer determines viability and IDPH VFC is contacted.

Vaccine Emergency Response

- Onsite generators or backup batteries can be used to prevent transporting vaccines to another storage facility in the event of an emergency like a power outage.
 - If the unit breaks down, then it may be appropriate to transport.
- Generators and backups should be tested quarterly.

Expired, Spoiled, Wasted Vaccine

- Expired, spoiled, and wasted vaccine must be reported in I-CARE within one week of the expiration date.
- Expired and spoiled vaccines in unopened vials or unused manufacturer prefilled syringes should be returned to McKesson Specialty within 6 months of expiration date for Excise Tax Credit.
 - Must be unopened and in the original manufacturer vial or prefilled syringe.
- Wasted vaccines must be disposed of according to usual medical biosafety procedures, and according to your agency procedures.
 - May include open vials or prefilled syringes with or without the needles attached, vaccine that was drawn into a syringe, or vaccines compromised due to a dropped or broken container.

Transferring Vaccines

- Vaccine can occasionally be transferred between providers in these cases:
 - Vaccine is six months or less from outdate, and unable to be used by provider.
 - Area outbreak resulting in unexpected surge of walk-in patients.
 - Clinic closure requiring transfer of vaccines to other VFC providers.
 - Seasonal clinic needing to transfer vaccine to other VFC providers when closing.
- The following transfer requests will be reviewed on a case-by-case basis:
 - Vaccines are more than six months from the expiration date.
 - The provider has an immediate need for a couple of doses of vaccine before an order could be received.
- Cold chain must be maintained required to use DDL when transferring.
- Department must review & approve transfer requests.
- Influenza vaccine cannot be transferred.

Mobile Clinics

- Same VFC storage requirements with a permanently installed unit.
 - Mobile clinic should be plugged into a power source at home site location when not in use.
 - Will be inspected as part of compliance visit.
 - Vaccine cannot be transported to city of Chicago or out-of-state.
 - Vaccine must be delivered to home site.

Temporary Off-Site Vaccine Clinics

- Temporary off-site clinics can be held.
 - o Transportation, storage, and handling must meet VFC program guidelines.
 - Vaccine must be delivered to permanent location.
 - o Total time for travel + clinic day should not exceed 8 hours (e.g. if transport to an off-site clinic is 1 hour each way, the clinic may run for up to 6 hours).
 - When transporting vaccines, use a portable vaccine refrigerator or freezer unit or qualified container and packout that maintains appropriate storage temperatures.
 - Use of the manufacturer shipping container or a frozen water bottle transport system is not permitted for transport to temporary, mobile, or off-site clinics.
 - Vaccine must be returned to permanent location & DDLs must be reviewed to ensure proper temperatures were maintained.

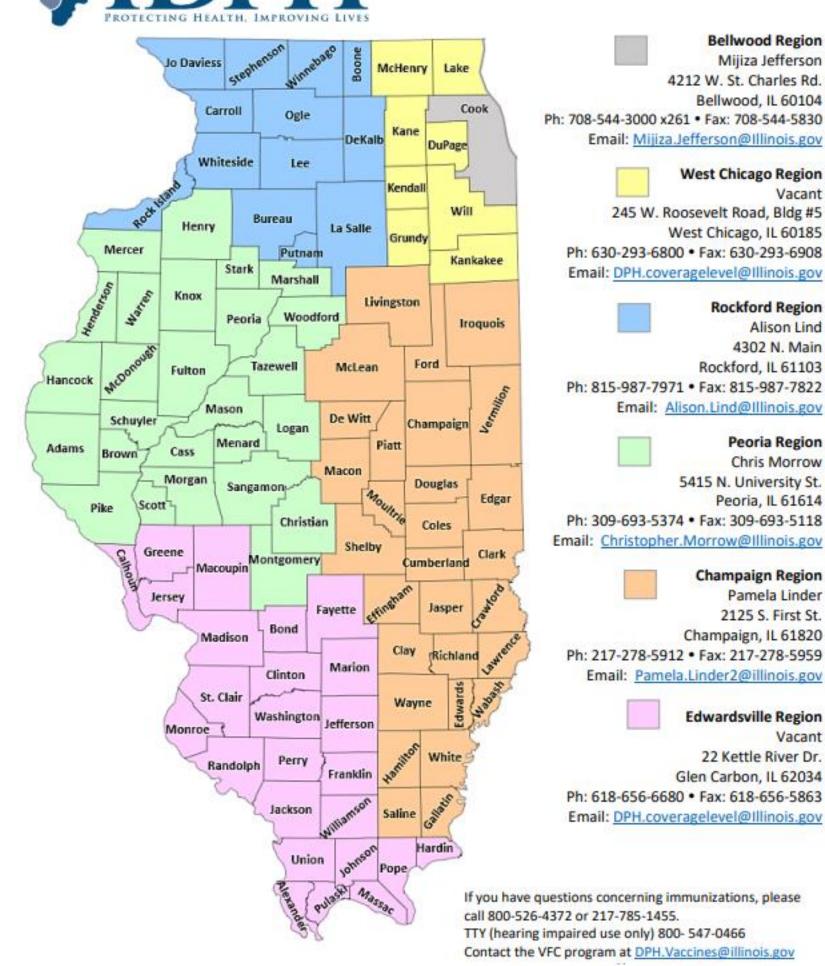
An off-site vaccine clinic notification form must be submitted in I-CARE 48 hours prior to the event.

Contacts

 Contact the VFC program at DPH.Vaccines@illinois.gov



IMMUNIZATION PROGRAM REGIONAL MAP



Break

Please return by 10:00

SESSION 2: VACCINE SCHEDULES

Jennifer Burns, APN



- Pediatric Family Travel
 Director
- Pediatric Infectious
 Diseases Section
- University of Chicago
 Comer Children's Hospital

After this Session

Outline new vaccine products and updates.

2

Apply the 2023
Advisory Committee
on Immunizations
Practices (ACIP)
pediatric vaccination
and catch-up
schedules.

Summarize current routine immunization rates.

Requirements

VFC providers must comply with:

- Current ACIP recommendations and VFC resolutions.
- Making vaccines identified in the Provider Profile based on the provider type and population served available including non-routine vaccines.
- Understanding state laws related to vaccination requirements and acceptable vaccine exemptions.
- Using ACIP recommendations and vaccine package inserts to understand contraindications for each vaccine type available through the VFC program.

State Vaccination Requirements

- State laws establish vaccination requirements for school children.
 - Laws often apply to public schools, private schools and day care facilities.
- Tools for maintaining high vaccination coverage rates.
- State laws also establish mechanisms for enforcement.
 - Documentation, signatures, etc.
- All states provide medical exemptions.
 - Some state offer religious and/or philosophical exemptions.

Exemptions Permitted for State Childcare and School (Kg-Gr 12) Immunization Requirements March 2023 ND WA MT MN SD WI ID OR WY IA CT NE PA OH IN NV UT CO KS MO DE. KY CA DC TN NC OK AR ΑZ NM SC GΑ MS AL TX LA Type of Exemption(s) Permitted Medical only AZ: Religious exemption for childcare only, personal belief Medical, religious exemption for school (Kg-Gr 12) only Medical, personal belief CO: Religious and personal belief exemptions combined under category of "nonmedical exemption" Medical, religious, personal belief DC and VA: Personal belief exemption for HPV only MO and NE: Personal belief exemption for childcare only WA: Personal belief exemption not allowed for MMR Immunize.org Source: state immunization programs For details, see data table: www.immunize.org/laws/exemptions.asp

IL Requirements

- Medical and religious exemptions.
- 105 III. Comp. Stat. Ann. 5/27-8.1; III. Adm. Code tit. 77, § 665.240; III. Adm. Code tit. 77, § 665.520; III. Adm. Code tit. 77, § 665.520; III. Adm. Code tit. 77, § Adm. Code 698.50.

ACIP

• Establishes:

- Age for vaccine recommendation.
- Number of doses and dosing interval.
- Precautions and contraindications.
- Technical recommendations on vaccine use and immunization practices
- Approves vaccines provided through VFC program.

Recommendations on the CDC's website.

ACIP Schedule - Approving Partners

Child Adolescent Schedule

American Academy of Pediatrics (AAP)

National Association of Pediatric Nurse Practitioners (NAPNAP) **Both Schedules**

American Academy of Family Physicians (AAFP)

American Academy of Physician Associates (AAPA)

American College of Obstetricians and Gynecologists (ACOG)

American College of Nurse-Midwives (ACNM) Adult Schedule

American College of Physicians (ACP)

Society for Healthcare Epidemiology of America (SHEA)

American Pharmacists Association (APhA)

VFC-ACIP Resolutions

ACIP votes on a resolution to include the vaccine change in the VFC program.

VFC resolutions passed by ACIP form the basis for VFC program policies on vaccine availability and use. Vaccines procured through the VFC program must be administered according to the guidelines outlined by ACIP in VFC resolutions (and in accordance with state laws).

CDC establishes contracts for VFC vaccines only after a VFC resolution is in place.

VFC-ACIP Vaccine Resolutions

rint

Vaccine procured through the VFC program must be administered according to the guidelines outlined by the ACIP in VFC resolutions. VFC vaccine also may be administered in accordance with State school attendance laws.

- 10/22-1 COVID-19 🔼 [1 page] Updated Oct 2022
- 6/21-1 <u>Dengue</u> [1 page] Updated Jun 2021
- 10/19-1 Diphtheria, Tetanus, & Pertussis 🚨 [4 pages, 508] Updated Oct 2019
- 6/19-2 <u>Haemophilus influenzae type b</u> [3 pages] Updated Jul 2019
- 6/19-6 Hepatitis A 🔼 [2 pages] Updated Jul 2019
- 6/19-3 Hepatitis B 🔼 [4 pages] Updated Jul 2019
- 10/16-2 <u>Human Papillomavirus (HPV)</u> [2 pages]
- 6/23-1 Influenza 🔼 [3 pages] Updated Jul 2023
- 6/20-1 Meningococcal 🔼 [4 pages] Updated Oct 2022
- 10/17-3 MMR & Varicella 🔼 [3 pages] Updated Jun 2022
- 06/94-16 Outbreak controls [1 page]
- 06/23-2 Pneumococcal 🔼 [4 pages] Updated Jul 2023
- 6/19-4 Polio 🔼 [2 pages] Updated Jul 2019
- 06/08-1 Rotavirus 🔼 [3 pages] Updated Feb 2023
- 08/23-2 Vaccines included in VFC Program [1] [1 page] Updated Aug 2023

PACKAGE INSERTS on FDA website

Immunization Schedules - Why They Matter

- There are no other alternative studied immunization schedule approve to provide to our patients.
- Protection against roughly 20 different life-threatening diseases.
- Prevention/control of infectious disease outbreaks.
- Gives children protection when they are most vulnerable.

Using the ACIP Schedule

To make vaccination recommendations, healthcare providers should:

- 1. Determine needed vaccines based on age (Table 1).
- 2. Determine appropriate intervals for catch-up, if needed (Table 2).
- 3. Assess for medical conditions and other indications (Table 3).
- 4. Review special situations (Vaccination Notes).
- 5. Review contraindications and precautions to vaccination (Appendix).

Table 1

*There is no other alternative studied immunization schedule

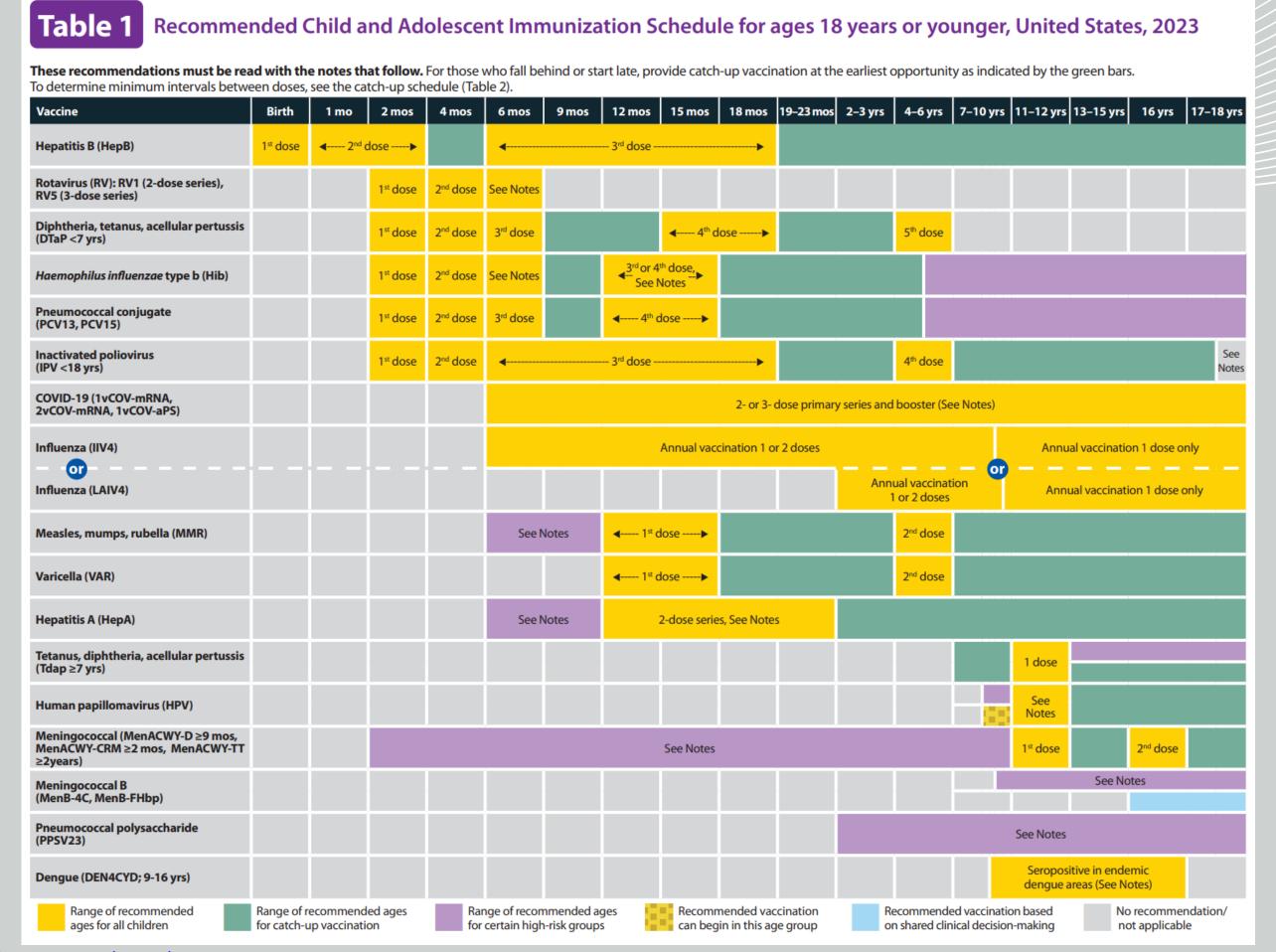


Table 2

Table 2

Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2023

The table below provides catch-up schedules and minimum intervals between doses for children whose vaccinations have been delayed. A vaccine series does not need to be restarted, regardless of the time that has elapsed between doses. Use the section appropriate for the child's age. **Always use this table in conjunction with Table 1 and the Notes that follow.**

			Children age 4 months through 6 years							
Vaccine	Minimum Age for	Minimum Interval Between Doses								
	Dose 1	Dose 1 to Dose 2	Dose 2 to Dose 3	Dose 3 to Dose 4	Dose 4 to Dose !					
Hepatitis B	Birth	4 weeks	8 weeks and at least 16 weeks after first dose minimum age for the final dose is 24 weeks							
Rotavirus	6 weeks Maximum age for first dose is 14 weeks, 6 days.	4 weeks	4 weeks maximum age for final dose is 8 months, 0 days							
Diphtheria, tetanus, and acellular pertussis	6 weeks	4 weeks	4 weeks	6 months	6 months					
Haemophilus influenzae type b	6 weeks	No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1st birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed if previous dose was administered at age 15 months or older 4 weeks if current age is younger than 12 months and first dose was administered at younger than age 7 months and at least 1 previous dose was PRP-T (ActHib®, Pentacel®, Hiberix®), Vaxelis® or unknown 8 weeks and age 12 through 59 months (as final dose) if current age is younger than 12 months and first dose was administered at age 7 through 11 months; OR if current age is 12 through 59 months and first dose was administered before the 1st birthday and second dose was administered at younger than 15 months; OR if both doses were PedvaxHIB® and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 doses before the 1 st birthday.						
Pneumococcal conjugate	6 weeks	No further doses needed for healthy children if first dose was administered at age 24 months or older 4 weeks if first dose was administered before the 1st birthday 8 weeks (as final dose for healthy children) if first dose was administered at the 1st birthday or after	No further doses needed for healthy children if previous dose was administered at age 24 months or older 4 weeks if current age is younger than 12 months and previous dose was administered at <7 months old 8 weeks (as final dose for healthy children) if previous dose was administered between 7–11 months (wait until at least 12 months old); OR if current age is 12 months or older and at least 1 dose was administered before age 12 months	8 weeks (as final dose) this dose is only necessary for children aged 12 through 59 months regardless of risk, or age 60 through 71 months with any risk, who received 3 doses before age 12 months.						
nactivated poliovirus	6 weeks	4 weeks	4 weeks if current age is <4 years 6 months (as final dose) if current age is 4 years or older	6 months (minimum age 4 years for final dose)						
Measles, mumps, rubella	12 months	4 weeks								
/aricella	12 months	3 months								
lepatitis A	12 months	6 months								
Meningococcal ACWY	2 months MenACWY-CRM 9 months MenACWY-D 2 years MenACWY-TT		See Notes	See Notes						
			Children and adolescents age 7 through 18 years							
Meningococcal ACWY	Not applicable (N/A)	8 weeks								
Tetanus, diphtheria; tetanus, diphtheria, and acellular pertussis	7 years	4 weeks	4 weeks if first dose of DTaP/DT was administered before the 1st birthday 6 months (as final dose) if first dose of DTaP/DT or Tdap/Td was administered at or after the 1st birthday	6 months if first dose of DTaP/DT was administered before the 1st birthday						
Human papillomavirus	9 years	Routine dosing intervals are recommended.								
Hepatitis A	N/A	6 months								
lepatitis B	N/A	4 weeks	8 weeks and at least 16 weeks after first dose							
nactivated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older and at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all previous doses were administered at <4 years or if the third dose was administered <6 months after the second dose.						
Measles, mumps, rubella	N/A	4 weeks								
Varicella	N/A	3 months if younger than age 13 years. 4 weeks if age 13 years or older								
		,								

Table 3

Table 3

Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2023

Always use this table in conjunction with Table 1 and the Notes that follow.

	INDICATION										
VACCINE	Pregnancy	Immunocom- promised status (excluding HIV infection)	HIV infection <15% or total CD4 cell count of <200/mm ³	>15% and total CD4 cell count of ≥200/mm³	Kidney failure, end-stage renal disease, or on hemodialysis	Heart disease or chronic lung disease	CSF leak or cochlear implant	Asplenia or persistent complement component deficiencies	Chronic liver disease	Diabetes	
Hepatitis B											
Rotavirus		SCID ^b									
Diphtheria, tetanus, and acellular pertussis (DTaP)											
Haemophilus influenzae ype b											
neumococcal conjugate											
nactivated poliovirus											
OVID-19		See Notes	See	Notes							
nfluenza (IIV4)											
nfluenza (LAIV4)						Asthma, wheezing: 2–4yrs ^c					
Measles, mumps, rubella	*										
Varicella	*										
Hepatitis A											
Tetanus, diphtheria, and acellular pertussis (Tdap)											
Human papillomavirus	*										
Meningococcal ACWY											
Meningococcal B											
Pneumococcal polysaccharide											
Dengue											
Vaccination according to routine schedule recommended		Recommended for persons with an addition factor for which the vact would be indicated	onal risk === a ccine n	/accination is recomr and additional doses necessary based on n condition or vaccine.	may be enedical	Precaution-vaccine night be indicated if benefit of protection outweighs risk of adverse reaction	be administ	led-vaccine should not	No recomme applicable	endation/no	

c. LAIV4 contraindicated for children 2-4 years of age with asthma or wheezing during the preceding 12 months

Notes

Notes

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2023

For vaccination recommendations for persons ages 19 years or older, see the Recommended Adult Immunization Schedule, 2023.

Additional information

 Consult relevant ACIP statements for detailed recommendations at www.cdc.gov/vaccines/hcp/acip-recs/ index html.

- For calculating intervals between doses, 4 weeks = 28 days. Intervals of ≥4 months are determined by calendar months.
- Within a number range (e.g., 12–18), a dash (–) should be read as "through."
- Vaccine doses administered ≤4 days before the minimum age or interval are considered valid. Doses of any vaccine administered ≥5 days earlier than the minimum age or minimum interval should not be counted as valid and should be repeated as age appropriate. The repeat dose should be spaced after the invalid dose by the recommended minimum interval. For further details, see Table 3-2, Recommended and minimum ages and intervals between vaccine doses, in *General Best Practice Guidelines for Immunization* at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/timing.html.
- Information on travel vaccination requirements and recommendations is available at www.cdc.gov/travel/.
- For vaccination of persons with immunodeficiencies, see
 Table 8-1, Vaccination of persons with primary and secondary
 immunodeficiencies, in General Best Practice Guidelines for
 Immunization at www.cdc.gov/vaccines/hcp/acip-recs/
 general-recs/immunocompetence.html, and Immunization in
 Special Clinical Circumstances (In: Kimberlin DW, Barnett ED,
 Lynfield Ruth, Sawyer MH, eds. Red Book: 2021–2024 Report
 of the Committee on Infectious Diseases. 32nd ed. Itasca, IL:
 American Academy of Pediatrics; 2021:72–86).
- For information about vaccination in the setting of a vaccine-preventable disease outbreak, contact your state or local health department.
- The National Vaccine Injury Compensation Program (VICP) is a no-fault alternative to the traditional legal system for resolving vaccine injury claims. All vaccines included in the child and adolescent vaccine schedule are covered by VICP except dengue, PPSV23, and COVID-19 vaccines. COVID-19 vaccines that are authorized or approved by the FDA are covered by the Countermeasures Injury Compensation Program (CICP). For more information, see www.hrsa.gov/vaccinecompensation or www.hrsa.gov/cicp.

COVID-19 vaccination

(minimum age: 6 months [Moderna and Pfizer-BioNTech COVID-19 vaccines], 12 years [Novavax COVID-19 Vaccine])

Routine vaccination

- Primary series:
- Age 6 months-4 years: 2-dose series at 0, 4-8 weeks (Moderna) or 3-dose series at 0, 3-8, 11-16 weeks (Pfizer-BioNTech)
- **Age 5–11 years:** 2-dose series at 0, 4-8 weeks (Moderna) or 2-dose series at 0, 3-8 weeks (Pfizer-BioNTech)
- **Age 12–18 years:** 2-dose series at 0, 4-8 weeks (Moderna) or 2-dose series at 0. 3-8 weeks (Novayax, Pfizer-BioNTech)
- For booster dose recommendations see www.cdc. gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html

Special situations

Persons who are moderately or severely immunocompromised

Primary series

- **Age 6 months-4 years:** 3-dose series at 0, 4, 8 weeks (Moderna) or 3-dose series at 0, 3, 11 weeks (Pfizer-RioNTech)
- **Age 5–11 years:** 3-dose series at 0, 4, 8 weeks (Moderna) or 3-dose series at 0, 3, 7 weeks (Pfizer-BioNTech)
- Age 12–18 years: 3-dose series at 0, 4, 8 weeks (Moderna) or 2-dose series at 0, 3 weeks (Novavax) or 3-dose series at 0, 3, 7 weeks (Pfizer-BioNTech)
- Booster dose: see www.cdc.gov/vaccines/covid-19/clinicalconsiderations/interim-considerations-us.html
- Pre-exposure prophylaxis (monoclonal antibodies) may be considered to complement COVID-19 vaccination. See www.cdc.gov/vaccines/covid-19/clinical-considerations/ interim-considerations-us.html#immunocompromised

For Janssen COVID-19 Vaccine recipients see COVID-19 schedule at www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html

Note: Administer an age-appropriate vaccine product for each dose. Current COVID-19 schedule and dosage formulation available at www.cdc.gov/vaccines/covid-19/downloads/COVID-19-immunization-schedule-ages-6months-older. pdf. For more information on Emergency Use Authorization (EUA) indications for COVID-19 vaccines, see www.fda.gov/emergency-preparedness-and-response/coronavirus-disease-2019-covid-19/covid-19-vaccines

Dengue vaccination (minimum age: 9 years)

Routine vaccination

- Age 9–16 years living in areas with endemic dengue AND have laboratory confirmation of previous dengue infection
 3-dose series administered at 0, 6, and 12 months
- Endemic areas include Puerto Rico, American Samoa, US Virgin Islands, Federated States of Micronesia, Republic of Marshall Islands, and the Republic of Palau. For updated guidance on dengue endemic areas and pre-vaccination laboratory testing see www.cdc.gov/mmwr/volumes/70/rr/r7006a1.htm?s.cid=rr7006a1 and www.cdc.gov/dengue/vaccine/hcp/index.html
- Dengue vaccine should not be administered to children traveling to or visiting endemic dengue areas.

Diphtheria, tetanus, and pertussis (DTaP) vaccination (minimum age: 6 weeks [4 years for Kinrix® or Quadracel®])

Routine vaccination

- 5-dose series at age 2, 4, 6, 15–18 months, 4–6 years
 Prospectively: Dose 4 may be administered as early as age
- 12 months if at least 6 months have elapsed since dose 3.

 Retrospectively: A 4th dose that was inadvertently administered as early as age 12 months may be counted if at least 4 months have elapsed since dose 3.

Catch-up vaccination

- Dose 5 is not necessary if dose 4 was administered at age
- 4 years or older and at least 6 months after dose 3.
- For other catch-up guidance, see Table 2.

Special situations

 Wound management in children less than age 7 years with history of 3 or more doses of tetanus-toxoid-containing vaccine: For all wounds except clean and minor wounds, administer DTaP if more than 5 years since last dose of tetanus-toxoid-containing vaccine. For detailed information, see www.cdc.gov/mmwr/volumes/67/rr/rr6702a1.htm.

Appendix

Appendix

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2023

Guide to Contraindications and Precautions to Commonly Used Vaccines

Adapted from Table 4-1 in Advisory Committee on Immunization Practices (ACIP) General Best Practice Guidelines for Immunization: Contraindication and Precautions available at www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html and ACIP's Recommendations for the Prevention and Control of 2022-23 seasonal influenza with Vaccines available at www.cdc.gov/mmwr/volumes/71/rr/rr7101a1.htm.

For COVID-19 vaccine contraindications and precautions see www.cdc.gov/vaccines/covid-19/clinical-considerations/interim-considerations-us.html#contraindications

Vaccine	Contraindicated or Not Recommended ¹	Precautions ²
Influenza, egg-based, inactivated injectable (IIV4)	 Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component³ (excluding egg) 	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Moderate or severe acute illness with or without fever
Influenza, cell culture-based inactivated injectable [(ccllV4), Flucelvax* Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) to any ccllV of any valency, or to any component ³ of ccllV4	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dos of any egg-based IIV, RIV, or LAIV of any valency. If using ccIV4, administer in medical setting under supervision of health care provider who can recognize and manage seve allergic reactions. May consult an allergist. Moderate or severe acute illness with or without fever
Influenza, recombinant injectable [(RIV4), Flublok* Quadrivalent]	• Severe allergic reaction (e.g., anaphylaxis) to any RIV of any valency, or to any component ³ of RIV4	 Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Persons with a history of severe allergic reaction (e.g., anaphylaxis) after a previous dos of any egg-based IIV, ccIIV, or LAIV of any valency. If using RIV4, administer in medical setting under supervision of health care provider who can recognize and manage seve allergic reactions. May consult an allergist. Moderate or severe acute illness with or without fever
Influenza, live attenuated [LAIV4, Flumist* Quadrivalent]	Severe allergic reaction (e.g., anaphylaxis) after previous dose of any influenza vaccine (i.e., any egg-based IIV, ccIIV, RIV, or LAIV of any valency) Severe allergic reaction (e.g., anaphylaxis) to any vaccine component³ (excluding egg) Children age 2 - 4 years with a history of asthma or wheezing Anatomic or functional asplenia Immunocompromised due to any cause including, but not limited to, medications and HIV infection Close contacts or caregivers of severely immunosuppressed persons who require a protected environment Pregnancy Cochlear implant Active communication between the cerebrospinal fluid (CSF) and the oropharynx, nasopharynx, nose, ear or any other cranial CSF leak Children and adolescents receiving aspirin or salicylate-containing medications Received influenza antiviral medications oseltamivir or zanamivir within the previous 48 hours, peramivir within the previous 5 days, or baloxavir within the previous 17 days	Guillain-Barré syndrome (GBS) within 6 weeks after a previous dose of any type of influenza vaccine Asthma in persons aged 5 years old or older Persons with underlying medical conditions (other than those listed under contraindications) that might predispose to complications after wild-type influenza virinfection [e.g., chronic pulmonary, cardiovascular (except isolated hypertension), renal, hepatic, neurologic, hematologic, or metabolic disorders (including diabetes mellitus)] Moderate or severe acute illness with or without fever

- 1. When a contraindication is present, a vaccine should NOT be administered. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html
- When a precaution is present, vaccination should generally be deferred but might be indicated if the benefit of protection from the vaccine outweighs the risk for an adverse reaction. Kroger A, Bahta L, Hunter P. ACIP General Best Practice Guidelines for Immunization. www.cdc.gov/vaccines/hcp/acip-recs/general-recs/contraindications.html
- 3. Vaccination providers should check FDA-approved prescribing information for the most complete and updated information, including contraindications, warnings, and precautions. Package inserts for U.S.-licensed vaccines are available at www.fda.gov/vaccines-blood-biologics/approved-products/vaccines-licensed-use-united-states

Hepatitis B Vaccine

- Recommended for infants born to mothers with positive test result for hepatitis B surface antigen (HBsAg) or unknown status.
- Catch up includes Heplisav-B and PreHevbrio for age 18 years and older.
- Heplisav-B not recommended during pregnancy.

Rotavirus Vaccine

- Rotavirus (Rotarix ™)
 - NO RECONSTITUTION NEEDED!
 - Oral-dosing applicator-only presentation.
 - FDA approved in Nov 2022.
 - Approved for use in infants 6 weeks old with the max age of the 1st dose at 18 weeks and the final dose no later than 8 months of age.
 - o Single 1.5ml dose.
 - Keep in original package to protect from light, store refrigerator 2-8°C.
 - New formulation contains disodium adipate & no longer contains sorbitol or phenol red.
 - Clinical trials comparing the 2 formulations showed similar safety profiles.
 - 2 variations of live vaccine Rotarix available until 2025 when older lyophilized formulation will retire.
 - Use up current 1ml lyophilized formulation (requires reconstitution) prior to using new liquid formulation.

DTaP

- 5-dose series at age 2, 4, 6, 15–18 months, 4–6 years.
- Prospectively: Dose 4 may be administered as early as age 12 months if at least 6 months have elapsed since dose 3.
- Retrospectively: A 4th dose that was inadvertently administered as early as age 12 months may be counted if at least 4 months have elapsed since dose 3.

Pneumococcal Vaccine

- Pneumococcal conjugate vaccines
 - o PCV13 (Prevnar 13, Pfizer) and PCV15 (Vaxneuvance, Merck), interchangeable.
 - o PCV20 (Prevnar 20, Pfizer).
 - Either PCV15 or PCV20 may be used for the full series or to complete the recommended schedule begun with PCV13.
 - o If only 13-valent PCV (PCV13) is available when the child is scheduled to receive a PCV, PCV13 may be given as previously recommended.
- Pneumococcal polysaccharide vaccine.
 - PPSV23 (Pneumovax23, Merck).
- Pneumococcal Vaccine added to the catch-up schedule, with 8 weeks marking the final dose. Age 12-59 months, regardless of risk.

Polio Vaccine

- Special situations -
 - Adolescents aged 18 years at increased risk of exposure.
 - No evidence of complete polio vaccination series (at least 3 doses)
 administer
 remaining doses to complete a 3-dose series.
 - Evidence of completed polio vaccination series (at least 3 doses)
 may administer 1

 lifetime IPV booster.

Influenza Vaccine

- Recommended for everyone 6 months+.
- Any licensed influenza vaccine appropriate by age and health status can be used.
- The AAP does not prefer any product over another for children and adolescents with no contraindications.
- Not given to individuals in close contact to immunocompromised people needing a protective environment.
- CAN be given to those with an egg allergy in medical setting where severe allergic reactions can be managed.

Influenza Vaccine

Influenza Vaccine Products for the 2023-2024 Influenza Season

Manufacturer	Trade Name	How Supplied	Mercury Content	Age Range	CVX Code	Vaccine Product Billing Code ² CPT	
	(vaccine abbreviation) ¹		(mcg Hg/0.5mL)		Code		
AstraZeneca FluMist (LAIV4)		0.2 mL (single-use nasal spray)	0	2 through 49 years	149	90672	
CCV	Fluarix (IIV4)	0.5 mL (single-dose syringe)	0	6 months & older ³	150	90686	
GSK	FluLaval (IIV4)	0.5 mL (single-dose syringe) 0		6 months & older ³	150	90686	
	Flublok (RIV4)	0.5 mL (single-dose syringe)	0	18 years & older	185	90682	
		0.5 mL (single-dose syringe)	0	6 months & older ³	150	90686	
	Fluzone (IIV4)	0.5 mL (single-dose vial)	0	6 months & older ³	150	90686	
Sanofi		5.0 mL multi-dose vial (0.25 mL dose)	25	6 through 35 months ³	158	90687	
		5.0 mL multi-dose vial (0.5 mL dose)	25	6 months & older	158	90688	
	Fluzone High-Dose (IIV4-HD)	0.7 mL (single-dose syringe)	0	65 years & older	197	90662	
	5.0 mL multi-dose vial (0.25 mL dose		24.5	6 through 35 months ³	158	90687	
	Afluria (IIV4)	5.0 mL multi-dose vial (0.5 mL dose)	24.5	3 years & older	158	90688	
Segirus		0.5 mL (single-dose syringe)	0	3 years & older ³	150	90686	
Seqirus	Fluad (alIV4)	0.5 mL (single-dose syringe)	0	65 years & older	205	90694	
	Elucolyay (ccll)/4)	0.5 mL (single-dose syringe)	0	6 months & older ³	171	90674	
	Flucelvax (ccIIV4)	5.0 mL multi-dose vial (0.5 mL dose)	25	6 months & older ³	186	90756	

NOTES

- IIV4 = egg-based quadrivalent inactivated influenza vaccine (injectable); where necessary to refer to cell culture-based vaccine, the prefix "cc" is used (e.g., ccIIV4); RIV4 = quadrivalent recombinant hemagglutinin influenza vaccine (injectable); aIIV4 = adjuvanted quadrivalent inactivated influenza vaccine.
- An administration code should always be reported in addition to the vaccine product code. Note: Third party payers may have specific policies and guidelines that might require providing additional information on their claim forms.
- Dosing for infants and children age 6 through 35 months:
- Afluria 0.25 mL
- Fluarix 0.5 mL
- Flucelvax 0.5 mL
- FluLaval 0.5 mL
- Fluzone 0.25 mL or 0.5 mL
- Afluria is approved by the Food and Drug Administration for intramuscular administration with the PharmaJet Stratis Needle-Free Injection System for persons age 18 through 64 years.

MMR

- 2-dose series at age 12–15 months, age 4–6 years.
- MMR or MMRV may be administered.
- Note: For dose 1 in children age 12–47 months, it is recommended to administer MMR and varicella vaccines separately. MMRV may be used if parents or caregivers express a preference.

Varicella

- 2-dose series at age 12–15 months, 4–6 years
- VAR or MMRV may be administered*
- Dose 2 may be administered as early as 3 months after dose 1 (a dose inadvertently administered after at least 4weeks may be counted as valid).
- *Note: For dose 1 in children age 12–47 months, it is recommended to administer MMR and varicella vaccines separately. MMRV may be used if parents or caregivers express a preference.

Hepatitis A

- 2-dose series (minimum interval: 6 months) at age 12–23 months.
- For persons traveling to or working in <u>countries with high or intermediate</u> endemic hepatitis A.
 - Infants age 6–11 months: 1 dose before departure; revaccinate with 2 doses (separated by at least 6 months) between age 12–23 months.
 - Unvaccinated age 12 months or older: Administer dose 1 as soon as travel is considered.

HPV

- Routinely recommended at age 11–12 years (can start at age 9 years) and catch-up HPV vaccination recommended for all persons through age 18 years if not adequately vaccinated.
- 2- or 3-dose series depending on age at initial vaccination:
 - Age 9 –14 years at initial vaccination: 2-dose series at 0, 6–12 months (minimum interval: 5 months; repeat dose if administered too soon).
 - Age 15 years+ at initial vaccination: 3-dose series at 0, 1–2 months, 6 months (minimum intervals: dose 1 to dose 2: 4 weeks / dose 2 to dose 3: 12 weeks / dose 1 to dose 3: 5 months; repeat dose if administered too soon).
- If vaccination schedule is interrupted, the series does not need to be restarted.
- No additional doses recommended.

Meningococcal Vaccines

- In August 2022, Menactra was replaced with MenACWY-TT (MenQuadfi™).
 - Licensed down to 9 months of age.
 - Studies: To get approval down to 6 months.
- Menveo™ (GSK) will still be available.
 - o 2 vials-reconstitute, age 2mon-55yo (lyophilized).
 - 1 vial-no reconstitution, 10-55yo (liquid).
- MenQuadfi vs Menveo-immune non-inferiority based on seroresponse for all 4 serogroups.

Meningococcal Vaccines

- Reminders
 - Menactra® should be administered either before or at the same time as DTaP in infants.
 - Related to toxoid antigen in Menactra®.
 - DTaP may alter Menactra®.
- MenQuadfi® and Menveo can be given regardless of DTaP.
- MenACWY vaccines may be administered simultaneously with MenB vaccines if indicated, but at a different anatomic site, if feasible.
- In children under 10 years needing a meningitis vaccine for travel,
 Menveo liquid is not appropriate for use.

RSV

- Monoclonal antibody injection (Nirsevimab) approved by CDC in August, expected availability in fall.
 - Will be covered by VFC.
- For neonates and infants born during or entering their first RSV season.
- Dosage is based on body weight:
 - <5 kg should be administered a 50 mg dose.
 - o ≥5 kg should be administered a 100 mg dose.
- For children up to 24 months of age who remain vulnerable to severe RSV disease through their second RSV season:
 - Single 200 mg dose administered as 2 IM injections.
- Children who have received Nirsevimab should not receive Palivizumab for the same RSV season.

RSV

Timing

- For infants born just before or during the RSV season (October March):
 Administer 1 dose at birth or as soon as possible.
- For infants born outside of the RSV season (April September): Within the first months of the RSV season.
- Administration can occur during birth hospitalization or in the outpatient setting.
- Coadministration
 - Simultaneous administration of Nirsevimab with age-appropriate vaccines is recommended.
- Provides passive immunization that extends through 5 months regardless of infant birth month

Products in the Pipeline

- COVID-19 (more in the next session)
 - Monovalent vaccine with an XBB.1.5-lineage of the Omicron variant
- Meningococcal (pentavalent)
 - In June, ACIP reviewed the potential uses of a new pentavalent meningococcal vaccine that would protect against serotypes A, B, C, W, & Y
- PCV20 is now available for ordering.

Schedule Changes

- Provider sites should have a process established for informing staff of any changes to ACIP recommendations.
 - Have 1 person in charge of tracking updates.
 - Have monthly trainings to review clinical guidance updates.

Illinois Coverage

ABLE. Estimated* coverage† with measles, mumps, and rubella; diphtheria, tetanus, and acellular pertussis; poliovirus; and varicella vaccines; grace period or provisional enrollment§; and any exemption¶,** among kindergartners, by immunization program — United States,†† 2021–22 school year

Immunization program	Kindergarten population ^{§§}	Surveyed, ^{¶¶} %	2 Doses MMR,*** %	5 Doses DTaP,††† %	4 Doses polio, ^{§§§} %	2 Doses VAR,¶¶¶%	Grace period or provisional enrollment,%	Any exemption, %	Percentage point change in any exemption, 2020–2021
National estimate****	3,837,259	92.2	93.5	93.1	93.5	92.8	2.4	2.6	0.4
Median****		_	92.9	92.0	92.7	92.6	1.9	2.7	0.2
Illinois ++++,§§§§	137,699	100.0	92.1	91.9	91.9	91.8	NR	≥1.7	NA

^{††††} The proportion surveyed is reported as 100% but might be <100% if based on incomplete information about the actual current enrollment.

^{§§§§} Philosophical exemptions were not allowed.

Series Completion Coverage

- Recent study published in *Pediatrics* found that 17% of toddlers who started seven recommended vaccine series did not complete 1 or more:
 - 4 doses diphtheria, tetanus, and acellular pertussis (DTaP).
 - 4 doses pneumococcal conjugate vaccine (PCV).
 - 3 to 4 doses haemophilus type b (Hib).
 - 3 doses hepatitis B (HepB).
 - 3 doses polio (IPV).
 - 1 dose measles, mumps, rubella (MMR).
 - 1 dose varicella (VAR).

Strategies for Increasing Uptake

- Simplify Scheduling:
 - Alert patients when vaccines are coming.
 - Self-scheduling through patient portal.
 - Automatically schedule return visits.
 - Site options mass vaccination clinics, drive-thru vaccination, sites at local high school, or empty store fronts.

Strategies for Increasing Uptake

- Streamline Operations:
 - Pre-made vaccine card labels.
 - Only carry one vaccine manufacturer.
- Create a culture of vaccine confidence:
 - Staff townhalls so everyone stays up-to-date.
 - Empower vaccine champions.
 - Everyone in the office (clinical and non-clinical) should be trained on how to address patient concerns about vaccines.

Strategies for Increasing Uptake

- Celebrate Vaccination!
 - Show off "I'm vaccinated!" stickers, pins, or posters.
 - Marketing materials in waiting rooms and social media.
 - Offer/talk about vaccinations at every visit.
 - For hesitant people treat it as an ongoing conversation.

SESSION 3: COVID-19 VACCINES

After this Session

Summarize current data of COVID-19 vaccination rates.

2

Outline storage and expiration date information for each vaccine manufacturer.

Describe current COVID-19 vaccine administration guidelines.

Current Vaccination Rates

People with an Updated (Bivalent) Booster Dose‡	Count	Percent of U.S. Population
Total	56,478,510	17.0%
Population ≥ 5 Years of Age	56,352,709	18.0%
Population ≥ 12 Years of Age	54,974,636	19.4%
Population ≥ 18 Years of Age	52,996,306	20.5%
Population ≥ 65 Years of Age	23,699,191	43.3%

^{*}Vaccination coverage reporting is no longer required as of May 2023

Data as of 5/10/23

Commercialization

- Commercialization the transition from direct government purchase to purchase by public and private consumers.
- There might be upfront costs for providers who provide COVID-19 vaccines.
 - Moderna: \$130/dose.
 - Pfizer: \$110-\$130/dose.
- The AAP is advocating for:
 - Longer invoice and payment schedules so that pediatricians can be reimbursed before having to purchase additional vaccines.
 - Simplified and improved labeling of COVID-19 vials to minimize errors in administration.
 - An easier return process for expired doses.
 - Single doses vials for all products.

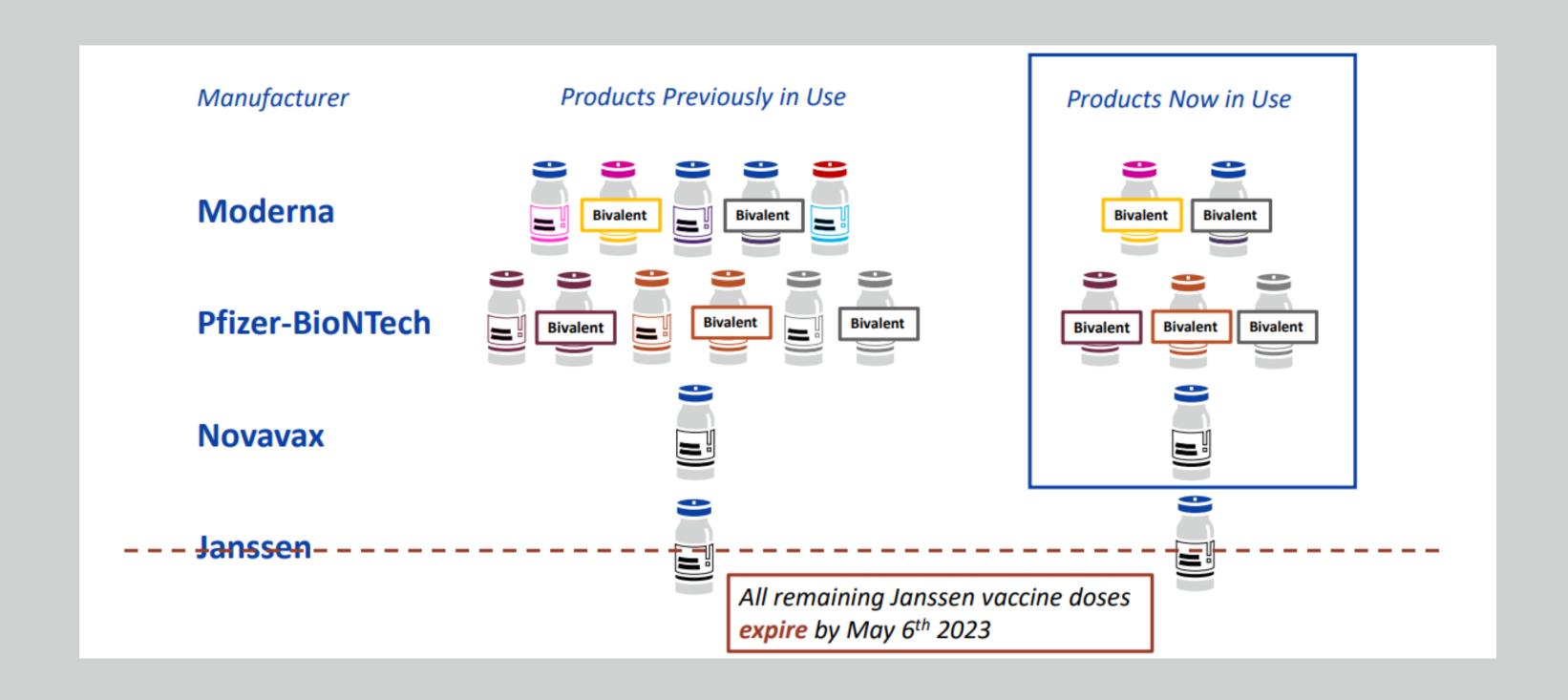
Commercialization Cont.

- HHS Commercialization Transition Guide: Sunsetting the US Government COVID-19
 Vaccine Distribution Program.
 - Issued July 6, 2023
- Access: VFC program, Bridge Access Program for adults without insurance, coverage by private insurance.
- Ordering:
 - VFC & Private Stock: ordered through the same process as other commercial vaccines
 - Bridge Access: separate ordering process.

Fall 2023 Vaccines

- Updating COVID-19 vaccines to XBB.1.5 strain: FDA recommendation 7/15/2023.
- Going back to a monovalent formula.
- Why?
 - WHO is not seeing any evidence that the earliest variant is still circulating.
 - We can drop that from the formula.
 - Including it could hurt as we keep teaching our immune system to recognize the old version of the virus rather than the new one.
- Novavax and Moderna found that a monovalent vaccine may be more advantageous to mice's immune systems than a bivalent vaccine.

Products Currently in Use



New Products Timeline

Ordering for bivalent COVID-19 vaccine closed in early August

Continue vaccinating with bivalent vaccines/stock available*

Prepare for commercialization

Communicate with patients

Pre-order monovalent vaccines

August 2023

Continue to vaccinate!

"We are likely to see this as a recommendation as an annual COVID shot, just as we have an annual flu shot." - Mandy Cohen

After September

September 2023

Commercialization

Fall COVID-19 vaccines are expected to be approved and available

ACIP Meeting 9/12

"We anticipate that they are going to be available for most folks by the third or fourth week of September" – CDC Director Mandy Cohen, MD, MPH

Expected Ordering

Expected

XBB.1.5 ordering *subject to change*

FALL 2023 VACCINE ORDERING

ILLINOIS VACCINATES
AGAINST COVID-19
A project led by the Illinois Chapter
of the American Academy of Pediatrics

Last updated August 17, 2023. Information is subject to change.

FORMULATION	CPT CODE	PRESENTATION	MINIMUM ORDER QTY HOW TO ORDER		соѕт	DELIVERY TIME	
Novavax							
Patient age 12 years and older	91304	Multi-dose vials containing 5 doses	50 doses	 Through VFC and other state programs Wholesaler/distribution channels 	Unknown - Prices will depend on the contract or group affiliation	Varies	
Pfizer							
Patient age 6 months-4 years	91318	Multi-dose vials containing 3 doses	30 doses	Through V/EQ and other state			
Patient age 5 years-11 years	91319	Single-dose vials			Through VFC and other state programs Pfizer Prime: 15% of an order that is received can be returned	Unknown - Prices will depend on the contract or group affiliation	Varies
Patient age 12 years and older	91320	Single-dose vials or single-dose prefilled syringes*	10 doses	Wholesaler/distribution channels			
Moderna							
Patient age 6 months-11 years	91321	Single-dose vials that come in 10-count cartons		Through VFC and other state programs	Unknown - Prices	Varies, Moderna would begin	
Patient age 12 years and older	91322	Single-dose prefilled syringes sold in 10-count cartons with a blister pack option	10 doses	 Moderna Direct: 10% of an order that is received can be returned Wholesaler/distribution channels 	will depend on the contract or group affiliation	shipping the vaccines after FDA approval	

^{*}There will be limited quantities of prefilled syringes and they will not be available for initial stocking orders.

Sources: www.pfizer.com/news/announcements/pfizer-and-biontech-submit-applications-us-fda-omicron-xbb15-adapted-monovalent https://publications.aap.org/aapnews/news/25384?autologincheck=redirected

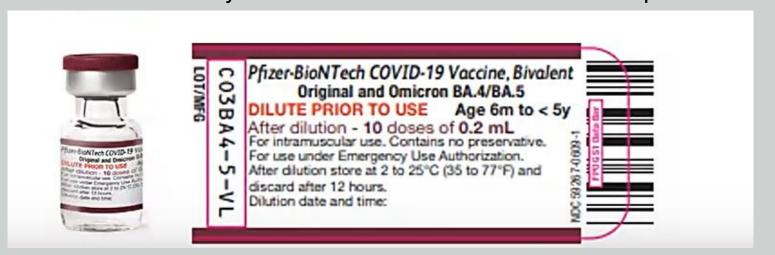
illinoisvaccinates.com

Preordering

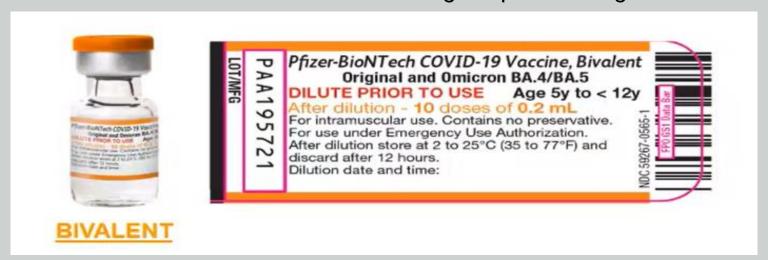
- Illinois Department of Public Health:
 - VFC COVID-19 Vaccine order link
 - COVID-19 Vaccine Bridge Access Program
 - Due to limited quantities, Bridge Access Program vaccine will be available to a limited group of providers. Target providers include LHDs, FQHCs, RHCs.
 - Link will be sent to target group providers.

Current Packaging-Pfizer

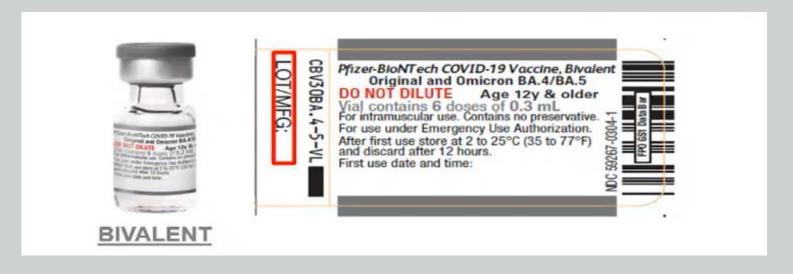
Pfizer: months – 4 years bivalent vaccine has a maroon cap and label.



Pfizer: 5-11 bivalent vaccine has an orange cap and orange label.



Pfizer: 12+ bivalent vaccine has a gray cap and gray label.

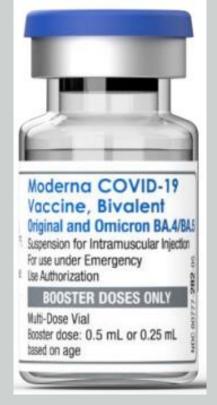


Current Packaging- Moderna

Moderna: Bivalent vaccine for ages 6 months – 5 years has a magenta cap and a yellow label border.



Moderna: Bivalent vaccine for ages 6 months and older has a blue cap and a grey label border.



Current Storage and Handling - Pfizer

AGE INDICATIONS	6 months through 4 years	5 years through 11 years	12 years and older		
FORMULATION	Bivalent	Bivalent	Bivalent		
VIAL CAP COLOR/LABEL WITH COLOR BORDER	Maroon/ Bivalent label	Orange/ Bivalent label	Gray/ Bivalent label		
PREPARATION	Dilute Before Use	Dilute Before Use			
AMOUNT OF DILUENT NEEDED PER VIAL®	2.2 mL	1.3 mL	Do Not Dilute		
DOSE VOLUME/DOSE	0.2mL/3mcg 0.2 mL/10 mcg		0.3 mL/30 mcg		
DOSES PER VIAL	10 doses (after dilution)	10 doses (after dilution)	6 doses or 1 dose		
ULT FREEZER (-90°C TO -60°C) ^b	18 months				
FREEZER (-25°C TO -15°C)	DO NOT STORE				
REFRIGERATOR (2°C TO 8°C)	10 weeks				
ROOM TEMPERATURE (8°C TO 25°C) including any thaw time	12 hours prior to first puncture				
AFTER FIRST PUNCTURE (2°C TO 25°C)	Discard after 12 hours				

- Diluent: Sterile 0.9% Sodium Chloride Injection, USP. Do not use bacteriostatic 0.9% Sodium Chloride Injection or any other diluent.
- Regardless of storage condition, vaccines should not be used after 12 months from the date of manufacture printed on the vial and cartons.

Current Storage and Handling - Moderna

AGE INDICATIONS AND FORMULATION	6 months through 5 years 6 months and older (Bivalent)		
VIAL CAP COLOR	Dark Pink	Dark Blue	
VIAL LABEL BORDER COLOR	Yellow	Gray	
PREPARATION	Do Not Dilute	Do Not Dilute	
DOSE VOLUME/DOSE	0.2 mL/10 mcg	6 months to 5 years old: 0.25 mL/25 mcg 6 to 11 years old: 0.25 mL/25 mcg 12 years and older: 0.5 mL/50 mcg	
DOSES PER VIAL	2 doses	Varies by age	
ULT FREEZER (-90°C TO -60°C)	DO NOT STORE		
FREEZER (-50°C TO -15°C)°	Until expiration		
REFRIGERATOR (2°C TO 8°C)	30 days		
ROOM TEMPERATURE (8°C TO 25°C) including any thaw time	24 hours		
AFTER FIRST PUNCTURE (2°C TO 25°C) ^b	Discard after 8 hours Discard after 12 hours		

- Regardless of storage condition, vaccines should not be used after the expiration date.
- After a vial is punctured 20 times it should be discarded even if the 12-hour time limit has not been met.

Storage and Handling - XBB.1.5

XBB.1.5 VACCINE STORAGE & HANDLING

I-VAC

ILLINOIS VACCINATES
AGAINST COVID-19
Agroject is d by the linois Chepter
of the American Academy of Pediatrics

Last updated August 31, 2023

	MODI	ERNA	PFIZER			NOVAVAX	
AGE INDICATIONS AND FORMULATION	6 Months- 11 Years	12+ Years	6 Months- 4 Years	5-11 Years	12+ Years	12+ Years Prefilled Syringe	12+ years
VIAL CAP COLOR	Unknown	Unknown	Yellow	Blue	Gray	N/A	Unknown
VIAL LABEL BORDER COLOR	Unknown	Unknown	Yellow	Blue	Gray	N/A	Unknown
PREPARATION	Do Not Dilute	Do Not Dilute	Dilute	Do Not Dilute	Do Not Dilute	N/A	Do Not Dilute
DOSE	0.25 mL	0.5 mL	3 mcg/ 0.3 mL dosage	10 mcg/ 0.3 mL dosage	30 mcg/ 0.3 mL dosage	30 mcg/ 0.3 mL dosage	5 mcg/ 0.5 mL dosage
DOSES PER VIAL	1	1	3	1	1	1 – Prefilled Syringe	5
ULT FREEZER (-90°C TO -60°C)	DO NOT STORE	DO NOT STORE	12 Months	12 Months	18 Months	9 Months	DO NOT STORE
FREEZER (-50°C TO -15°C)	Until Expiration	Until Expiration	DO NOT STORE	DO NOT STORE	DO NOT STORE	DO NOT STORE	DO NOT STORE
REFRIGERATOR (2°C TO 8°C)	30 Days	30 Days	10 Weeks	10 Weeks	10 Weeks	10 Weeks	Until Expiration
ROOM TEMPERATURE (8°C TO 25°C)	24 Hours	24 Hours	12 Hours Prior to First Puncture	12 Hours Prior to Use	12 Hours Prior to Use	4-12 Hours*	Unknown
AFTER FIRST PUNCTURE (2°C TO 25°C)	N/A	N/A	Discard After 12 Hours	N/A	N/A	N/A	12 Hours

Last updated September 1, 2023

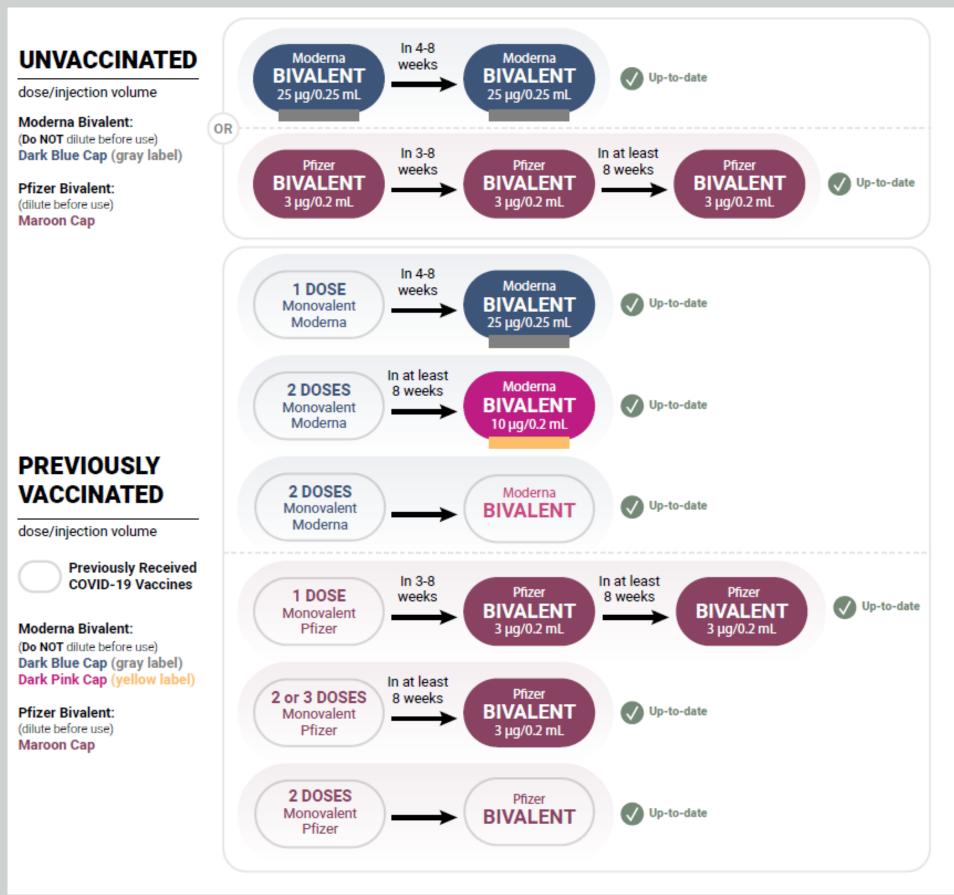
^{*}Thawed in carton: 12 hours prior to use. Thawed outside of carton: Use within 4 hours of thawing.

Moderna Carton Dimensions-XBB.1.5

TYPE	DESCRIPTION	ITEM AND PACKING INFORMATION					
Prefilled Syringe Blister Pack • 0.5 mL single-dose prefilled syringe • 10-count carton (blister pack) • 12+ years of age	prefilled syringe		Weight (lbs)	Depth (in)	Width (in)	Height (in)	Volume (in³)
	(blister pack)	Carton	0.24 lbs	2.25 in	5.25 in	3.43 in	38.84 in
Prefilled Syringe • 0.5 mL single-dose prefilled syringe • 10-count carton • 12+ years of age		ITEM AND PACKING INFORMATION					
		Weight (lbs)	Depth (in)	Width (in)	Height (in)	Volume (in³)	
	12 v yours or ago	Carton	0.20 lbs	4.10 in	4:14 in	1.42 in	24.10 in ³
		ITEM AND PACKING INFORMATION					
O.5 mL single- dose vial 10-count carton 12+ years of age		Weight (lbs)	Depth (in)	Width (in)	Height (in)	Volume (in³)	
	• 12+ years of age	Carton	0.21 lbs	3.68 in	1.54 in	2.20 in	11.76 in ³

6mo-<5yrs

Additional Considerations An 8-week interval between the first and second doses of Moderna and Pfizer-BioNTech COVID-19 vaccines might be optimal for some people ages 6mo-64yrs especially for males ages 12-39yrs, as data suggests it reduces the risk of myocarditis and pericarditis associated with these vaccines.



Ages 5mo to <5yrs moderately to severely immunocompromised

For individuals 6mo-4yrs of age with certain types of immunocompromise who have previously received three 0.2mL doses (Pfizer-BioNTech COVID-19 Vaccine or Pfizer-BioNTech COVID-19 Vaccine, Bivalent:

- A fourth dose may be administered at least 1 month following the most recent dose.
- Additional doses may be administered at the discretion of the healthcare provider, taking into consideration the individual's clinical circumstances.

UNVACCINATED

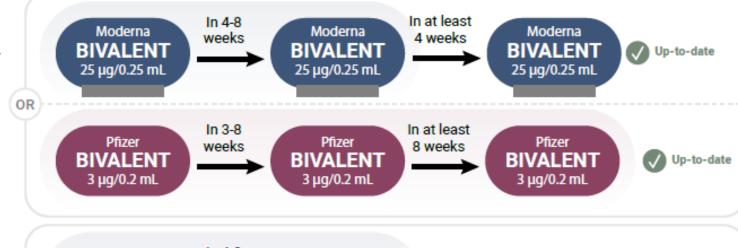
dose/injection volume

Moderna Bivalent:

(Do NOT dilute before use) Dark Blue Cap (gray label)

Pfizer Bivalent:

(dilute before use) Maroon Cap



PREVIOUSLY VACCINATED

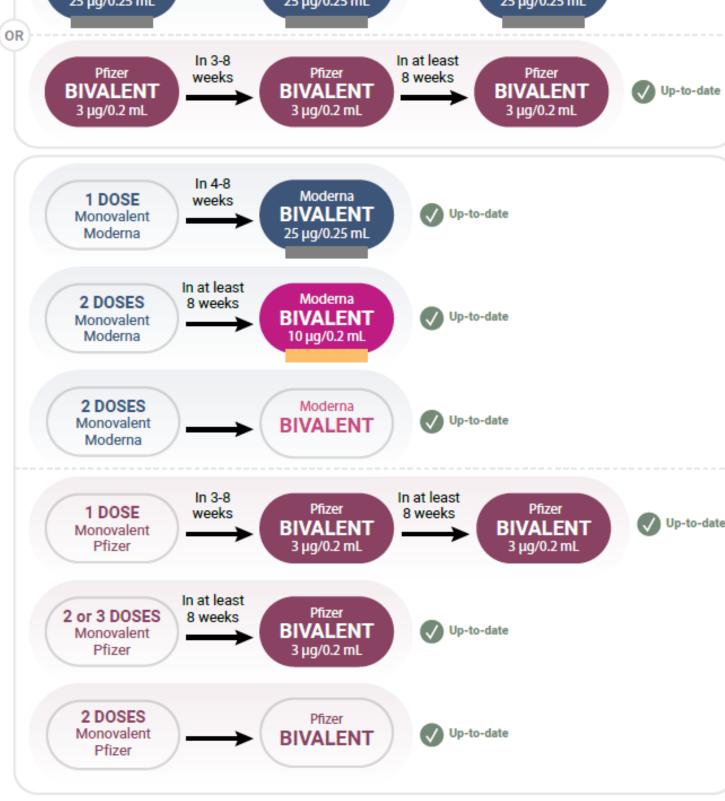
dose/injection volume

Previously Received **COVID-19 Vaccines**

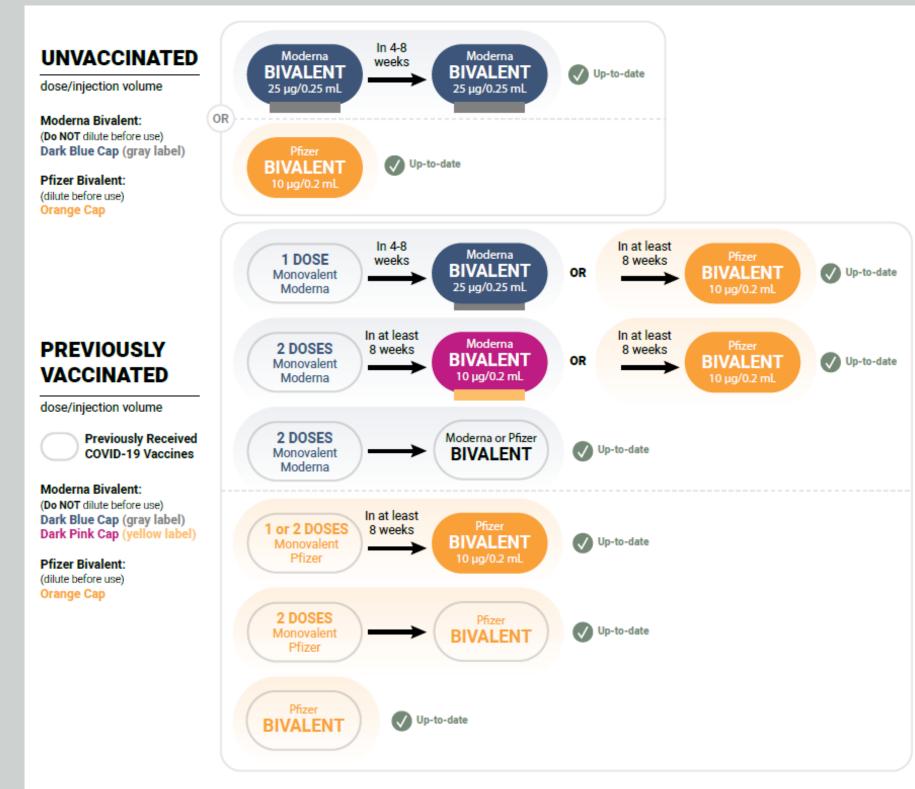
Moderna Bivalent:

(Do NOT dilute before use) Dark Blue Cap (gray label) Dark Pink Cap (yellow label)

Pfizer Bivalent: (dilute before use) Maroon Cap



5yrs-<6yrs



Ages 6 to <12yrs



dose/injection volume

Moderna Bivalent:

(Do NOT dilute before use)
Dark Blue Cap (gray label)

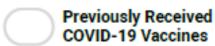
Pfizer Bivalent:

(dilute before use)
Orange Cap



PREVIOUSLY VACCINATED

dose/injection volume

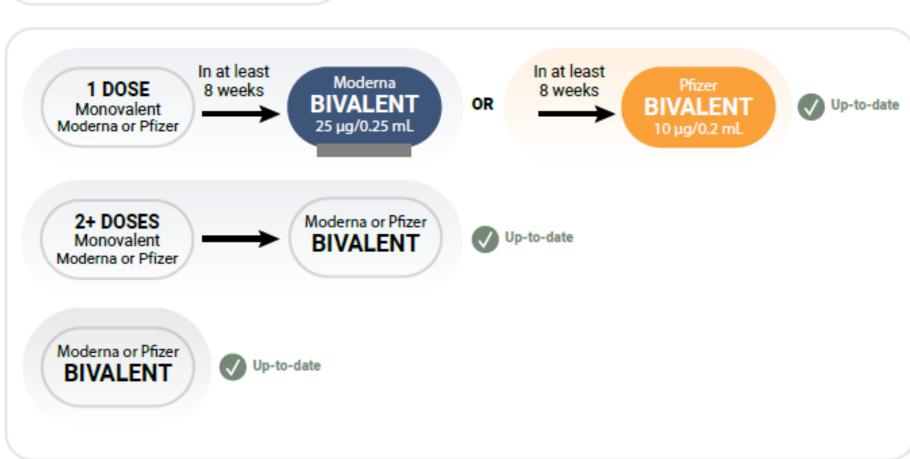


Moderna Bivalent:

(Do NOT dilute before use)
Dark Blue Cap (gray label)
Dark Pink Cap (yellow label)

Pfizer Bivalent:

(dilute before use) Orange Cap



Ages 12yrs+

UNVACCINATED

dose/injection volume

Moderna Bivalent:

(Do NOT dilute before use) Dark Blue Cap (gray label)

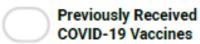
Pfizer Bivalent:

(Do NOT dilute before use) Gray Cap

Moderna BIVALENT Up-to-date 50 μg/0.5 mL OR Pfizer Up-to-date BIVALENT 30 µg/0.3 ml

PREVIOUSLY VACCINATED

dose/injection volume



Moderna Bivalent: (Do NOT dilute before use)

Dark Blue Cap (gray label)

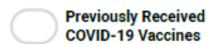
Pfizer Bivalent:

(Do NOT dilute before use) Gray Cap



ADDITIONAL DOSES

dose/injection volume



Moderna Bivalent: (Do NOT dilute before use)

Dark Blue Cap (gray label)

Pfizer Bivalent: (Do NOT dilute before use)

Gray Cap

65 YEARS AND OLDER

People ages 65 years and older have the option to receive 1 additional bivalent mRNA vaccine dose at least 4 months after the first dose of a bivalent mRNA vaccine.







Expiration Dates

Manufacturer	How to find it
Moderna (Spikevax)	 Scan QR code on the vial or carton. Website: <u>Look up tool.</u>
Pfizer	 Website: Look up tool. See EUA fact sheets for providers.
Novavax	 Scan QR code on outer carton. Website: <u>Look up tool</u>.

Dosing and Administration XBB.1.5

- Clinical guidance for fall Omicron XBB.1.5 COVID-19 vaccines is not yet issued.
- We are awaiting sign-offs from the FDA and the CDC.
- ACIP Meeting September 12

How to Guide Patients Now: Not Official Guidance

Get the available COVID-19 vaccine or wait until the new one comes out?

It depends on a person's individual risk:

- People who are 65 or older or who have multiple health conditions should get the currently available shot if it's been more than 6 to 8 months since their last dose.
- Children who are not vaccinated and entering their first season.
- Pregnant persons.
- For others: it is okay to wait for the new version.

Reminders: Clinical guidance for the new formulation has not yet been issued. We don't know yet which population groups will be authorized for an annual shot.

Reporting to VAERS

Required by law to report the following to VAERS:

- 1. Vaccine administration errors, whether or not associated with an adverse event (AE)
 - If the incorrect mRNA COVID-19 vaccine product was inadvertently administered for a second dose in a 2-dose series, VAERS reporting is required.
- 2. Cases of myocarditis after a Pfizer-BioNTech, Moderna, or Novavax vaccine.
- 3. Cases of pericarditis after a Pfizer-BioNTech, Moderna, or Novavax vaccine.

VAERS reporting is not required for:

Mixed series is given intentionally (e.g., due to hypersensitivity to a vaccine ingredient).

Reporting to VAERS

- 4. Serious AEs regardless of whether the reporter thinks the vaccine caused the AE:
 - Death
 - A life-threatening AE
 - Inpatient hospitalization or prolongation of existing hospitalization
 - A persistent or significant incapacity or substantial disruption of the ability to conduct normal life functions
 - A congenital anomaly/birth defect
 - An important medical event that based on appropriate medical judgement may require medical or surgical intervention to prevent one of the outcomes listed above
- 5. Cases of Multisystem Inflammatory Syndrome in adults or children.
- 6. Cases of COVID-19 that result in hospitalization or death.

Report to VAERS any clinically significant AEs following vaccination, even if you are not sure whether the vaccine caused the event.

Reporting to VAERS

- Fill out a separate form for each patient.
- List any medicines, supplements, and remedies the patient is taking or using.
- List any allergies.
- List acute illnesses and chronic conditions the patient may have.
- Fill out the information of the person filling out the form and contact information of the provider to contact, correctly and completely.
- Call 1-800-822-7967 or email info@vaers.org for help or information.

Combating Hesitancy and Misinformation

- Misinformation vs. disinformation
 - When people spread misinformation, they often believe what they are sharing
 - Disinformation is intended to mislead others.
- Address hesitancy and misinformation by increasing vaccine confidence:
 - Create vaccine champions of your staff
 - Help simplify scheduling for patients and alert them when vaccines are available.
 - Streamline operations for your staff.
 - Be empathetic, compassionate, and sensitive to culture, family dynamics, and circumstances that may influence how patients view vaccines.
- Motivational interviewing is an evidence-based and culturally sensitive way of speaking with unvaccinated patients about getting vaccinated.

Discussion

- Case scenario: You are carrying both the Moderna and Pfizer products for all ages at your clinic. What are some ways you can minimize vaccine administration errors?
- 5 minutes to discuss different strategies at your tables.
- What are some ways you minimize vaccine administration errors in your own practice?
- 5 minutes to present one strategy to the group.

Lunch

Please return by 12:45

SESSION 4: OTHER TOPICS

After this Session

•

Describe vaccine hesitancy, misinformation, and disinformation.

Demonstrate strategies for combatting vaccine misinformation and disinformation.

Outline ways to discuss vaccine hesitancy with patients.

Unfortunate Theme

- 23 million children missed out on basic childhood vaccines through routine health services in 2020.
 - The highest number since 2009 and 3.7 million more than in 2019.

Children were on track to miss an estimated 9 million vaccination doses in 2020, a
decrease of up to 26% in childhood vaccination doses compared to 2019.

Unfortunate Theme

Vaccination	BCBS 2020 estimated vax rate	BCBS % decrease from 2019	CDC herd immunity thresholds	BCBS 2020 est. vax rate vs herd immunity thresholds
Measles	88.2%	-26%	94%	-4.8
Pertussis	79.3%	-26%	92%	-12.7
Polio	88.9%	-16%	86%	+2.9

Unfortunate Theme

- Vaccination coverage among kindergartners nationwide:
 - Lower during the 2020-21 school year compared with 2019-20 school year (95% to ~94%).
 - Lower during the 2021-22 school year compared with the 2020-21 school year (~94% to ~93% for all state-required vaccines).
- Non-exempt, undervaccinated students often attend school while in a grace period or are provisionally enrolled.
- In many states, exemption policies were expanded either formally or informally during the 2020–21 school year.

Importance of Influenza, COVID-19 & Routine Vaccines

Since we do not have vaccines to protect against every disease, it is even more important to use the vaccines that we do have to keep children out of the hospital.

Preliminary interim estimates—NVSN

- Through January 25, 2023, influenza vaccination significantly reduced laboratory confirmed medically attended influenza
 - 68% (95% CI: 46, 81) against pediatric hospitalizations
 - 42% (95% CI: 25, 56) against pediatric ED visits
- Important protection against both A/H3N2 and A/H1N1 associated illness

School Exemptions



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Measles Case Summary: Central Ohio Outbreak



(As of: Wednesday, January 25, 2023 9:56 AM)

Data Source: Ohio Disease Reporting System (ODRS). All data are preliminary and subject to change.

PUBLIC HEALTH

Case Information for <u>confirmed measles cases</u> who are part of the current central Ohio outbreak.

Since the start of the outbreak in November 2022, we have seen:

 $85\,\mathrm{cases}$ (of which $36\,\mathrm{were}$ hospitalized and $0\,\mathrm{have}$ died)

Vaccination status

Unvaccinated	80
Partially vaccinated (1 dose)	4
Fully vaccinated (2 doses)	0
Unknown vaccination status	1

Vaccination status definitions:

Unvaccinated: Case has had 0 doses of the MMR (Measles, Mumps, and Rubella) vaccine - some cases may not have been eligible for any doses

Partially vaccinated: Case has had 1 dose of the MMR vaccine - some cases may not have been eligible yet for the 2nd dose due to age Fully vaccinated: Case has had 2 doses of the MMR vaccine

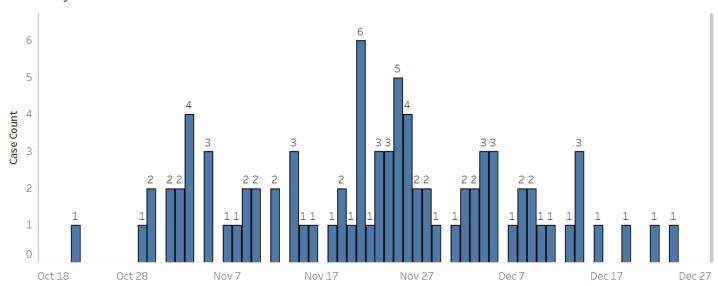
65% of cases are 1-5 years old.

<1 year old	25
1-2 years old	36
3-5 years old	19
6-17 years old	5
18+ years old	C

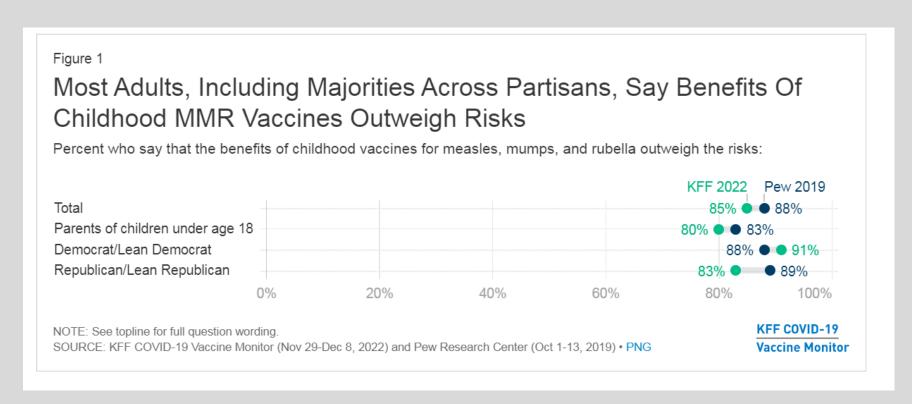
48% of cases are female and 52% of cases are male.

Female	41
Male	44
Unknown sex	(

Cases by Date of Rash Onset



Implications



Dec-22
Oct-19
Democrat/Lean Democrat

Figure 3

Rubella

Total

Dec-22

Oct-19

Oct-19

Dec-22

Oct-19

Parents of children under age 18

Republican/Lean Republican

NOTE: See topline for full question wording.



20% **79%**

SOURCE: KFF COVID-19 Vaccine Monitor (Nov 29-Dec 8, 2022) and Pew Research Center (Oct 1-13, 2019) • PNG

Compared To 2019, More Adults Now Say Parents Should Be Able To

Which comes closer to your views about childhood vaccines for measles, mumps, and rubella, even if neither is exactly

Parents should be able to decide not to vaccinate their children, even if that may create health risks for other children and because of the potential risk for others when

adults children are not vaccinated

Decide Not To Vaccinate Their Children For Measles, Mumps, And

KFF COVID-19 Vaccine Monito

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Misinformation vs. Disinformation

- When people spread misinformation, they often believe the information they are sharing.
- Disinformation is crafted and disseminated with the intent to mislead others.

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Dictionary
Definitions from Oxford Languages

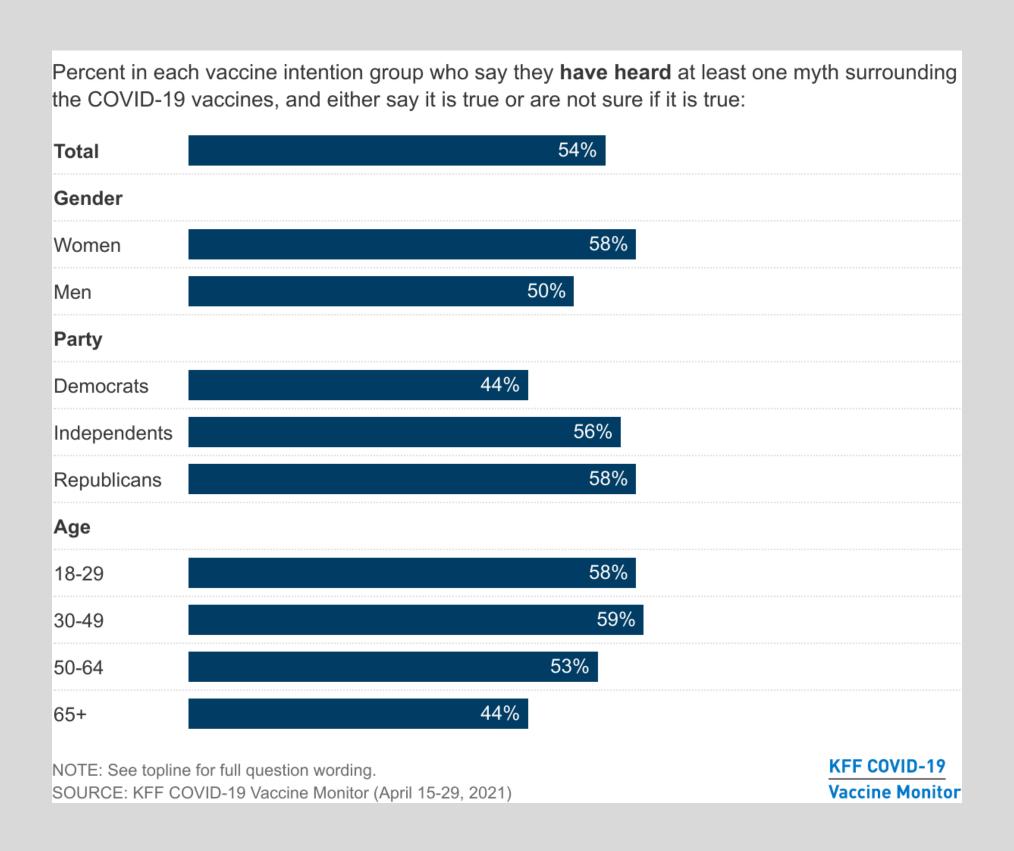
mis·in·for·ma·tion

/misinfərˈmāSH(ə)n/

1.false or inaccurate information, especially that which is deliberately intended to deceive.

"nuclear matters are often entangled in a web of secrecy and misinformation"

Misinformation & Myths



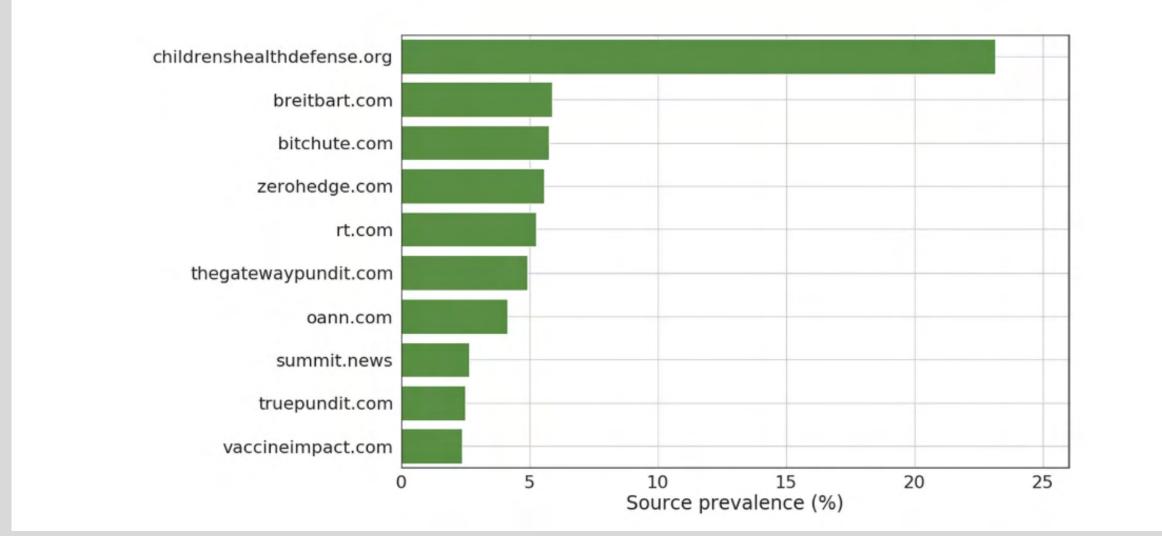
Pandemic + Overload of Information = Infodemic



Vaccine Landscape

"Disinformation campaigns are deliberate, often orchestrated, and highly effective in confusing people enough to change behaviors, like not getting the COVID-19 vaccine."

Top low-credibility Sources



Tweets shared by users geolocated in the U.S. that link to a low-credibility source. Sources are ranked by percentage of the tweets considered.

Documentary Discussion Questions

- What are your reactions to this documentary?
- What has been the most challenging part about vaccinating since 2020?
- What vaccine misinformation/disinformation patterns have you seen in the past?
 - What are the sources for this information?
 - How have these patterns changed?
- What has your clinic/organization done to respond to misinformation/disinformation?
 - Among staff (internally)
 - Among patients/families (externally)
- How do you initiate conversations about vaccines with patients?
 - How do you address misinformation/myths with a patient?
 - How do you address vaccine hesitancy?
- What strategies to address vaccine hesitancy have worked?
 - What hasn't worked?

Next Steps

- Make sure you signed in
- Fill out your evaluation (needed for CME)
- Let us know if you want additional outreach materials









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 &

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Thank You