#### 2024 CDPH VFC Training

#### March 6, 2024

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#### Agenda

| 8:00 AM  | Check - In and Networking  |
|----------|--|
| 8:30 AM  | CDPH VFC Program   |
| 9:30 AM  | Vaccine Hesitancy - <i>Virulent: The Vaccine</i><br><i>War</i> Screening |
| 10:30 AM | 15 Minute Networking Break   |
| 10:45 AM | Vaccine Schedules  |
| 11:45 AM | Questions and Closing  |
| 12:00 PM | Adjourn  |

## Session I: CDPH VFC Program

#### **Speaker: Victor Santiago**

#### **Learning Objectives**

After this session participants will be able to:

Objective 1:

Outline the VFC Program recommendations, requirements, and updates.

#### **Objective 2:**

Describe inventory reconciliation & vaccine ordering best practices and other Chicago VFC mandates.

Objective 3: Implement effective vaccine storage and handling practices and keep vaccine waste to a minimum.

Objective 4:

Overview of Immunization Information System (IIS) I-CARE

#### **Overview of the VFC Program**

- The Vaccines for Children (VFC) Program provides all routine vaccines recommended by the Advisory Committee on Immunization Practices (ACIP) at no cost to children who otherwise might not be vaccinated.
- Vaccines provided through the VFC Program must be administered <u>according to</u> the guidelines outlined by ACIP.

#### **Benefits of VFC**

- Many families cannot afford to pay for vaccines on their own, a common barrier in routine vaccination rates. VFC benefits include:
  - Reduces up-front costs of Chicago VFC providers because you will not have to pay to purchase vaccines for VFC program-eligible children.
  - Eliminating or reducing vaccine cost as a barrier to immunizing eligible children.
  - Enables patients to get the vaccines they need during routine appointments at their regular office.
  - Helps provide quality care to vulnerable children and adolescents.

#### **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.

#### **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).
    - Immunize.org: Vaccine Information Statements (available in 47 languages)
  - Reporting adverse reactions to VAERS.

#### Table 1. Guidance for Use of Vaccine Information 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<br>Compensation Program (VICP)-covered vaccine is<br>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<br>administering the vaccine*  |  |
| Must be the current version <sup>†</sup>  | Vaccine Information Statement version date and date<br>it is provided <sup>†</sup>  |  |
| Can provide (not substitute) other written or audio-<br>visual materials as necessary <sup>‡</sup>                      | Site (eg, deltoid area), route of administration<br>(eg, intramuscular), and expiration date of the<br>vaccine <sup>‡</sup> |  |

\*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 Cont.**

- The <u>National Childhood Vaccine Injury Act (NCVIA)</u> and/or <u>CDC</u> requires physicians to document the:
  - Name & Date of vaccine administered
  - 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).

#### **Provider Agreement**

- 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 Chicago 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 CDPH may terminate the provider agreement include:
  - Provider has not ordered vaccine in the past 12 months.
  - A provider on the List of Excluded Individual and Entities (LEIE) list maintained by Office of the Inspector General.
  - Failure to comply with requirements.

#### Vaccine Staff & Training – Vaccine Coordinators

- **Identify** a primary VFC vaccine coordinator and at least one backup VFC 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.

#### **Staff Training**

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



### **Training Opportunities**

- Some VFC site visits.
- This training  $\ensuremath{\textcircled{}}$
- CDC online training with both of the following modules:
  - You Call The Shots <u>Module 10 Storage and Handling</u>
  - You Call the Shots <u>Module 16 Vaccines for Children Program</u>



#### **VFC Enrollment Visits**

- VFC providers agree to VFC program site visits, which may include compliance visits, unannounced storage and handling visits, or educational site visits.
- The enrollment site visit is completed before a provider location can receive VFC vaccines. The goal of the enrollment site visit is to:
  - Educate providers about VFC program requirements.
  - Educate providers on proper vaccine storage and handling.
  - Certify provider locations have the appropriate resources to implement requirements.
  - Confirm providers know whom to contact if problems arise, especially with storage and handling issues.
  - Complete a Vaccine Management Plan.

#### **VFC Compliance Visits**

- Every year each VFC provider is required to have a comprehensive quality assurance review (QAR). This type of visit requires a thorough evaluation of the provider's compliance with all VFC program requirements including:
  - Verification of information in the provider profile
  - Review of VFC eligibility screening and documenting procedures
  - Review of vaccine storage and handling practices (including temperature logs and vaccine storage units)
  - Evaluation of provider's written procedures related to temperature monitoring, routine vaccine storage and handling and emergency vaccine storage and handling
  - Review of documentation of VIS given
  - Review of documentation for vaccine administration
  - Review of vaccine ordering and accountability
  - Verification that VFC Program policies are being properly implemented.

#### **VFC Storage & Handling Visits**

- Storage and Handling visits may be announced (scheduled) or unannounced.
- Reviewers assess individual storage units and DDLs, as well as overall storage and handling operations, based on VFC requirements and CDC's Vaccine Storage and Handling Toolkit.
- 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 dose-fordose basis with privately purchased vaccines.
- In order to replace each dose of VFC vaccine used on non-VFC eligible children, please submit a vaccine replacement request.
- Once the vaccine replacement request is approved, the provider's I-CARE inventory will be updated, and the provider will be notified on any changes.

#### Fraud & Abuse

- By enrolling in the VFC program, you agree to comply with all program requirements.
- It is your responsibility to read and understand our <u>Fraud and Abuse Policy</u>.
- 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.
  - Failing to properly maintain VFC records and requirements.
  - Failing to properly store and handle VFC vaccines, etc.

#### Fraud & Abuse Cont.

- The Department will investigate to determine intentional or unintentional fraud/misuse.
- The Chicago Department of Public Health Immunization Program may take the following actions when fraud and/or abuse may have occurred:
  - Determine if a situation requires immediate referral or if educational intervention and follow-up are adequate.
  - Make decisions to refer cases to the Medicaid Integrity Group (MIG) and any other state or city agencies that are required by law to refer suspect cases.
  - Make appropriate referrals and notify CDC of referral to MIG and any other appropriate agencies.

# Patient Eligibility Screening & Documentation

- Providers must screen and document patient eligibility screening in the patient's permanent medical record (paper-based or electronic medical record) using the VFC Patient Eligibility Screening Record or document the required elements in the electronic medical record.
- Patient eligibility screening records should be maintained on file for a minimum of three years.

### **Eligibility**

- All VFC Program providers must screen for a child's eligibility to receive vaccines through the VFC Program and record the screening results during each visit. A child is eligible for the VFC Program if they are 18 years of age or younger and are one of the following:
  - Uninsured.
  - Medicaid-eligible or Medicaid-enrolled
  - American Indian or Alaska Native.
  - Underinsured.
  - Underinsured VFC-eligible children can only receive VFC vaccine from a Federally Qualified Health Center (FQHC) or Rural Health Clinic (RHC).



# 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.
- A provider must select and document the VFC eligibility category that will require the least amount of out-of-pocket expenses to the parent/ guardian for the child to receive necessary immunizations.

#### **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 VFCsupplies or other vaccines purchased through CDC federal contracts.



#### **Administration Fees**

- Bill only Medicaid for the administration fee for VFC-eligible children enrolled in Medicaid.
  - 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.

## **Adolescent Vaccine Data**

# NIS – Teen: 2022 Summary

# Vaccination Coverage Estimates NIS – Teen Background

- The National Immunization Surveys (NIS) Teen is a telephone and mailed survey from the National Center for Immunization and Respiratory Diseases
  - Respondents include the parents/guardians of  $\sim$  45,000 adolescents aged 13 to 17 across the U.S. and their vaccination providers
- The surveys help estimate national and jurisdictional vaccination coverage for a selection of vaccines recommended for adolescents by the Advisory Committee on Immunization Practices (ACIP)
- For adolescents surveyed in the 2022 NIS Teen:
  - Interviews and mailed surveys were completed between January 2022 and February 2023
  - Respondents were born between January 2004 and January 2010

#### 🗚 Vaccination Coverage Among Adolescents Aged 13 - 17



■ City of Chicago ■ IL - Rest of State ■ United States

5

Source: National Immunization Survey - Teen 2022. Reported August 2023. Not all vaccine categories shown.

Hep A = hepatitis A vaccinate; Hep B = hepatitis B vaccine; HPV = human papilloma virus vaccine; Meningococcal = meningococcal conjugate vaccine; MMR = measles, mumps, and rubella vaccine; Tdap = tetanus toxoid, reduce diphtheria toxoid, and acellular pertussis. Varicella's estimates are inclusive of those who may have acquired chickenpox during childhood prior to being surveyed. HPV, Up-to-Date refers to receipt of 2 doses separated by 5 months for immunocompetent adolescents initiating the HPV vaccine series before their 15<sup>th</sup> birthday, and 3 doses for all others.

## Highlights

- Chicago's coverage estimates are roughly within ~ 5 points of both national and statewide estimates for virtually all adolescent vaccinations, although margins of error potentially mitigate these differences
- Overall, vaccine coverage across most vaccines has increased substantially since the NIS – Teen was first established in 2008
  - Tdap: 44.2% → 90.5%
  - Meningococcal:  $41.4\% \rightarrow 89.1\%$
- HPV remains the vaccine with the lowest coverage of all the adolescent vaccinations
  - While females have historically had higher coverage rates than males, recent changes have made it so that coverage for both are within one percentage point of each other


#### What is IQIP?

- IQIP is CDC's national, Vaccines for Children (VFC) provider-level immunization quality improvement (QI) program. IQIP serves to assist and support health care providers by identifying opportunities to improve vaccine uptake and to help providers be:
- Motivated to try new vaccination service delivery strategies and incorporate changes into their current practices
- Supported in sustaining changes and improvement to their vaccination service delivery
- Aware of and knowledgeable about vaccination coverage and missed opportunities to vaccinate
- Able to use available data from the IIS (I-CARE) to improve services and coverage

#### The IQIP Process

IQIP is a 12-month process during which public health representatives from CDPH and VFC providers collaborate to implement provider-level QI strategies to increase vaccine uptake by improving and enhancing vaccination workflow.

| Site Visit   | 2-Month and 6-Month Check-Ins  | 12-Month Follow-Up   |
|--|--|--|
| <ul> <li>Provider's vaccination workflow<br/>is observed, and initial<br/>coverage is reviewed</li> <li>QI strategies are selected</li> <li>Technical assistance is<br/>provided by the IQIP consultant</li> <li>Action items are chosen for<br/>strategy implementation plan</li> </ul> | <ul> <li>Progress toward strategy<br/>implementation is reviewed</li> <li>Technical assistance is<br/>provided by the IQIP consultant</li> <li>Strategy implementation plan is<br/>reviewed and updated</li> </ul> | <ul> <li>Progress toward strategy<br/>implementation is reviewed<br/>and updated</li> <li>Technical assistance is provided<br/>by the IQIP consultant</li> <li>Year-over-year coverage<br/>change is reviewed</li> </ul> |

#### **Benefits of Immunization QI Projects**

Quality improvement (QI) programs, such as IQIP, analyze processes and use a systematic approach to improve performance. Like other QI programs, the IQIP program is based on these basic steps:

- State the problem and desired result
- Use data to understand the problem
- Identify strategies for improvement
- Implement strategies and refine as needed
- Evaluate outcome

Beyond simply increasing vaccination coverage rates, conducting immunization quality improvement can provide additional benefits. Here are some broader effects that can result from immunization QI program like IQIP.

*Currently a part of a quality improvement project at your location? You may be able to get credit from CDC.* 

<u>Please think about participating in IQIP and therefore become eligible for next year's VFC</u> <u>Vaccine Coverage Awards!</u>



\* \* \* \*

#### Sign Up

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# Latest Alerts

• HAN Website: chicagohan.org

#### **VFC Program Website**



Welcome To The Chicago VFC Health Alert Network (HAN) Page!

This is your one-stop-shop for information, resources, VFC policies, and frequently-requested paperwork.

#### Vaccines for Children - chicagohan.org/vfc

#### **Speaker: Kevin Hansen**

### Vaccine Management

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



#### Vaccine Management Plan

Contact info for current primary and backup vaccine coordinators.

Proper storage and handling practices. Procedures for vaccine ordering, receiving, inventory control, stock rotation, and handling vaccine loss and waste.

Procedures for emergency situations (transport, equipment malfunction, power failure, and natural disaster).

Documented training related to vaccine management. Provider and vaccine coordinator roles & responsibilitie s.

Plans must be updated annually or more frequently as needed.

# Vaccine Management: Ordering Vaccines

- Vaccine ordering is completed through I-CARE. Sites must complete these steps to place an order for vaccines in I-CARE:
  - Clear any errors from the Inventory Analysis Helper Report.
  - Run the Vaccine Accountability Report
  - Complete Temperature Log Report.
  - Review and Approve Delivery Hours for Their Site.
- COVID and Flu doses are ordered via a pre-book.
- After placing an order, allow at least **three business days** for order approval.
  - Orders may be delayed if steps for inventory accuracy are not followed.
- If you are in need of immediate vaccine supply, contact chicagovfc@cityofchicago.org.

# Vaccine Management: Ordering Vaccines

- Order and stock enough vaccine to be able to administer vaccine for their next period (monthly, bi-monthly, quarterly) plus a 5-week safety stock.
  - Consider patient numbers, patient age, vaccine uptake, etc.
  - Smaller, more frequent orders help reduce the impact of incidents that may contribute to vaccine loss.
- 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).

**Determining how much vaccine to order** In order to determine this amount, providers should use this formula:

**Doses Administered (previous period)** 

x 2.3 (monthly), 1.6 (bi-monthly), 1.4 (quarterly)

Inventory

=

Amount to Order

#### **Doses Administered** (previous period) 50 doses х 2.3 (monthly), 1.6 (bi- $50 \times 1.6 = 80$ (bi-monthly provider) monthly), 1.4 (quarterly) 80 - 24 = 56Inventory Amount to Order 56 (rounded up) = 60 doses

#### Vaccine Management: Blended Inventory & Transferring Vaccines

- Blended Inventory: CHIP and Vaccine vaccines merged into one inventory as of January 2023.
  - Benefiting eligible children and reducing administrative burden for providers.
- Vaccine Transfer Process
  - To transfer vaccine between Chicago VFC offices email <u>chicagovfc@cityofchicago.org</u> with site PINs, lot numbers, doses, and transfer date.
  - Ensure proper temperature monitoring during transport using digital data loggers and qualified cooler containers.

#### Vaccine Management: Cold Chains

- 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 Vaccines

- 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 the packing slip matches the vaccine received.
  - Ensure correct lot numbers are shown in I-CARE inventory.
  - Physical damage of shipping container.
  - Diluent and Vaccine expiration dates.
- Frozen vaccines, flu, and COVID ship separately from other refrigerated vaccines.

Any issues to the shipment (incorrect or missing), please email <a href="mailto:chicagovfc@cityofchicago.org">chicagovfc@cityofchicago.org</a> within one day of the delivery.



# Vaccine Management: Storing Vaccine

- To ensure viability of VFC vaccines, locations must have:
  - Storage units that maintain correct temperatures at all times.
  - Refrigerator temperature between 2°C and 8°C (36°F and 46°F).
  - Freezer temperature between -50°C and -15°C (-58°F and +5°F).
  - Digital data loggers (DDLs) with continuous monitoring capabilities and a current and valid Certificate of Calibration Testing for each unit, as well as at least one backup.



### Vaccine Management: Storing Vaccine

- Storage units must have enough room to store the largest inventory a provider location might have at the busiest point in the year without crowding.
- 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.

- Purpose-built or pharmaceutical-grade refrigerators and freezers are preferred.
  - Still needs to be approved and meet 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.



- Some purpose-built units separate public & private vaccine stock electronically.
  - If electronic, an inventory printout must be available upon request.
- Power Supply:
  - Plug in only one storage unit per electrical outlet.
  - Use a safety-lock plug or an outlet cover.
  - Post "DO NOT DISCONNECT" 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.

- 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, min/max temps, and the initials of the person recording the temps
  - 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.

- Vaccines should be stored in their original packaging with lids closed.
  - Never store food or beverage in a unit with vaccines.
  - Do not store vaccines in the deli, fruit, or vegetable bins, in the doors or on the floor of the unit, or under or near cooling vents.
  - Place water bottles throughout the units against walls, in the back, on the floor, and in the doors – to help stabilize temperatures.



- 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).
  - Review temperatures of VFC vaccine storage units twice daily. Record minimum, maximum, and current temperatures on paper log.
- Data from DDLs is retrieved using special software or a website.
  - CDPH recommends downloading data weekly, but at least once mon
  - Records should be kept for at least three years.
- A back-up DDL must be available in case another fails; calibration testing is required.



- 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.

- 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.

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.

The DDL must be equipped with:

- 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).

Certificate 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.
  - Manufacturers will help determine if vaccine is still viable after an excursion.
- Must notify <u>chicagovfc@cityofchicago.org</u> each time your unit goes out of range.
  - Email should include: site's VFC PIN, include DDL tag summary, fridge or freezer excursion, reason for being out of range, and if the unit is back in range.
- If unit's temperature goes out of range, pause administering affected vaccines until a VFC staff member confirms usability.

#### **Speaker: Danielle Belanger**



1. The Value of IIS

2. 2024 IIS Updates

#### The Value of IIS



#### Provides Consolidated Records

Comprehensive records containing immunizations administered at a previous provider office, hospital, pharmacy or school clinic give healthcare providers the full story, preventing patients from receiving too many or too few vaccines.

Manages Vaccine Inventory

Vaccine ordering, tracking, and administration are all managed in one tool.



#### Minimizes Waste

Ensures every vaccine is accounted for and prevents the administration of unnecessary doses of vaccines.

Forecasts

Helpful alerts notify providers to assist with clinical decisions and management of the complex immunization schedule.

#### **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 Chicago 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.

### Topics – 2024 IIS Updates

- 1. New I-CARE Enrollment
- 2. Quick Assist
- 3. Data Modernization
- 4. HL7 onboarding (Checking EMR messages to I-CARE)
- 5. Chicago HAN
- 6. I-CARE Training Videos

#### **New I-CARE Enrollment**

#### Welcome!

Welcome to the Illinois Comprehensive Automated Immunization Registry Exchange (I-CARE) Enrollment website.

On this website, you will find tools and resources to complete the following:

- New Organizations can complete site enrollments and designate a Portal Registration Authority (PRA)
- Current Organizations can report site updates and PRA changes
- · Individuals can request I-CARE Access

Please note that I-CARE will not enroll organizations outside of Illinois, enroll for the purpose of research, or human resource departments for employee immunization verification.

[No Title] New Individual Current **Organizations\*** Organizations **I-CARE Access** Provider \* New organizations are those with no prior enrollment in I-CARE Not sure if your organization is enrolled? Please contact us.

Access the new site here





Have Questions? **Contact Us** 

View the I-CARE Glossarv

#### **Quick Assist**

- Quick Assist is a new tab in I-CARE dedicated to provider support
- Password Reset
- Frequently used forms



HL7/EMR Updates – To be used when the site/organization is changing their EMR, adding HL7 or are having issues with their HL7 connection. I-CARE User Updates – Providers can request updates to user accounts. This includes a user's name change, email, site placement, access level, and delete user. Log in Procedures for I-CARE – Instructions for users to access the IDPH web portal and I-CARE. Password Reset/Log-In Issues – To be used if a password cannot be reset using the password reset help information. The login issues can be reported here. Report Organization Acquisition/ Merger – Providers should report an organizational acquisition or merger (change of ownership). Site Updates/ Organization – This form is used to request updates to site information (name, address). Submit/Contact Us – General questions about I-CARE can be submitted here.

#### Chicago Vaccine Management Resources

I-CARE

<u>Celsius Freezer</u> – Celsius Freezer Temperature Log <u>Celsius Refrigerator</u> – Celsius Refrigerator Temperature Log <u>Fahrenheit Freezer</u> – Fahrenheit Freezer Temperature Log <u>Fahrenheit Refrigerator</u> – Fahrenheit Refrigerator Temperature Log <u>Inventory Discrepancy</u> – Providers can utilize this form to submit their Inventory Discrepancy to ChicagoVFC <u>Vaccine Replacement Log</u> – Providers can use this form to submit their vaccine replacements to ChicagoVFC <u>Vaccine Return Form</u> – Chicago Providers to report any expired or wasted doses.

#### **Data Modernization – Data Quality**



#### Site Reports

Bad Address – Generate a list of active patients with an invalid primary address.

Invalid Doses – Generate a list of patients with invalid doses.

Missed Opportunities Detail – Generate a report of missed opportunites for a site. A missed opportunity for a patient is calculated by finding the most recent shot date for a patient, and then listing all forecasted shots where the forecasted date is before the most recent shot date.

Missed Opportunities Summary – Generate a summary report of missed opportunites for a site.

Missing Lots - Generate a list of patients with missing lot numbers on immunizations.

Monthly Statistics – Generate a monthly statistics report for patients aged 24 - 35 months or for patients aged 9 - 26 years. This report will always run as a background process.

Patient List Export - Generate a comma-delimited text file of patients that match the search criteria.

Shot Refusals – Generate a list of shots refusals reported by a site within a given time period.

HL7 Logs – Generate a HL7 Logs Report. View all HL7 upload errors and warnings.



# Submitting & 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</u> site. If you have questions, please contact
   <u>CDPH.HL7@illinois.gov</u> or <u>DPH.HL7@illinois.gov</u>.

# HL7 Onboarding and Ongoing Monitoring

- We launched a new process to onboard new providers to message their immunization data into the I-CARE Immunization Registry
- We provide guidance to your clinical, EMR, and integration team on messaging your immunization data into the registry
- Complete our survey on the Chicago HAN to sign up for review
- We can address any of your questions at <u>CDPH.HL7@cityofchicago.org</u>
# HL7 Onboarding and Ongoing Monitoring



# **Chicago HAN**

Highlighting some of the new and improved areas of the Chicago HAN.

https://www.chicagohan.org/vfc

## HAN Home > Programs > Vaccines for Children Overview + VFC Program Annual Re-Enrollment + **CHIP Vaccine Information** + VFC News Bulletins + Digital Data Loggers (DDL's) And Cloud Services + I-CARE Basics + VFC Tools And Policies + Training + Immunization Resources + **Data Quality Reviews And Onboarding** + I-CARE Training Videos +

## **Chicago HAN IIS Resources**

# **Chicago HAN I-CARE Trainings**

- We have created video training tutorials for a variety of I-CARE topics.
- This is a great resource if you need a refresher
- Also great for onboarding new staff who have not used I-CARE before.

https://www.chicagohan.org/vfc

## I-CARE Training Videos

- I-CARE Login : How to log into I-CARE and get started
- **Patient Module** : A comprehensive system for managing patient information related to immunization. It provides tools for adding new patients, searching for patient records, and updating patient information.
- Shots Add View Edit Delete : How to view, add, edit, and delete shots
- <u>Shots Immunizations and Contraindications</u> : How to add immunities, contraindications, and adverse events to a patient's profile
- Shots Overrides and Refusals : How to override a shot and what to do when a patient refuses a shot
- I-CARE Training VFC Tab Overview : A brief overview of the processes and systems within the VFC tab: VFC, Vaccine Requests, Staff, and Enrollment.
- VFC Vaccine Ordering : How to order VFC Vaccines via the VFC tab in I-CARE
- Bad Address Report : How to run a bad address report in I-CARE
- Immunizations Due/Given Reports : How to run the immunizations due and immunizations given reports in I-CARE
- <u>COVID Immunizations Activity/Due Reports</u>: How to run the COVID Immunization Activity report and COVID Immunizations Due report
- **<u>Reminder Recall</u>**: How to remind patients who are due or overdue for a vaccination to make an appointment with your office
- Patient Immunization History : How to access and understand a patient's immunization history.

## **CDPH VFC Manual**

- Updated VFC Manual for Chicago providers is now available!
- <u>https://illinoisaap.org/vaccines-for-</u> <u>children/</u>



Childhood vaccination information and resources as developed by the Chicago Department of Public Health in partnership with the Illinois Chapter of the American Academy of Pediatrics

# **Session II**

# Vaccine Misinformation and Hesitancy



Caroline Werenskjold, MPH

## **Learning Objectives**

After this session participants will be able to:

Objective 1:

Describe vaccine hesitancy, misinformation, and disinformation.

Objective 2:

Demonstrate strategies for combatting vaccine misinformation and disinformation.

Objective 3:

Outline ways to discuss vaccine hesitancy with patients.

# **Unfortunate Theme**

- Vaccination rates are still low post-pandemic
  - CDC report: kindergarten vaccination rates have not rebounded from the COVID-19 pandemic, and school exemptions reached an all-time high during the 2022-'23 school year.
- Many toddlers aged 19-35 months in the U.S. do not complete their full, recommended vaccine series.
  - Only 73% of toddlers finish all vaccine series. 10% of toddlers never initiate the vaccines, while 17% start the series, but never complete it.

# Implications

### Figure 3

Compared To 2019, More Adults Now Say Parents Should Be Able To Decide Not To Vaccinate Their Children For Measles, Mumps, And Rubella

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

Parents should be able to decide not to vaccinate their children, even if that may create health risks for other children and adults children are not vaccinated

| Total   |                            |                  |                                 |
|---|----------------------------|------------------|---------------------------------|
| Dec-22  | 28%                        | 71%              |                                 |
| Oct-19  | 16%                        | 82%              |                                 |
| Parents of children under age 18  |                            |                  |                                 |
| Dec-22  | 35%                        | 65%              |                                 |
| Oct-19  | 23%                        | 76%              |                                 |
| Democrat/Lean Democrat  |                            |                  |                                 |
| Dec-22  | 11%                        | 88%              |                                 |
| Oct-19  | 12%                        | 86%              |                                 |
| Republican/Lean Republican  |                            |                  |                                 |
| Dec-22  | 44%                        | 56%              |                                 |
| Oct-19  | 20%                        | 79%              |                                 |
| NOTE: See topline for full question wording.<br>SOURCE: KFF COVID-19 Vaccine Monitor (Nov 29-Dec 8, 2022) and | Pew Research Center (Oct 1 | -13, 2019) • PNG | KFF COVID-19<br>Vaccine Monitor |

## Figure 1

## Most Adults, Including Majorities Across Partisans, Say Benefits Of Childhood MMR Vaccines Outweigh Risks

Percent who say that the benefits of childhood vaccines for measles, mumps, and rubella outweigh the risks:

|                                    |                                 |                       |                       |                     | KFF 2022  | Pew 2019     |
|------------------------------------|---------------------------------|-----------------------|-----------------------|---------------------|-----------|--------------|
| Total                              |                                 |                       |                       |                     | 85% 🔶     | • 88%        |
| Parents of children under ag       | e 18                            |                       |                       |                     | 80% 🔷 🗢 8 | 3%           |
| Democrat/Lean Democrat             |                                 |                       |                       |                     | 88%       | • • 91%      |
| Republican/Lean Republicar         | 1                               |                       |                       |                     | 83% 🔵     | • 89%        |
|                                    | 0%                              | 20%                   | 40%                   | 60%                 | 80%       | 100%         |
| NOTE: See topline for full questio | n wording.<br>9 Monitor (Nov 29 | -Dec 8, 2022) and Pew | / Research Center (Or | rt 1-13 2019) • PNG | 1         | KFF COVID-19 |

## **Misinformation vs. Disinformation**

## Misinformation

• When people spread misinformation, they often **believe** the information they are sharing.

## Disinformation

• Disinformation is crafted and disseminated with the **intent to mislead** others.

# Misinformation & Myths

Figure 3

Adults Without A College Degree, Republicans, And Independents Are More Likely To Say COVID-19 And Vaccine Misinformation Is Definitely Or Probably True

Percent who say each of the following false claims is probably true or definitely true:

|                     | The COVID-19<br>vaccines have<br>caused<br>thousands of<br>sudden deaths<br>in otherwise<br>healthy people | lvermectin is<br>an effective<br>treatment for<br>COVID-19 | The COVID-<br>19 vaccines<br>have been<br>proven to<br>cause<br>infertility | The MMR<br>vaccines<br>have been<br>proven to<br>cause<br>autism in<br>children | More people<br>have died<br>from the<br>COVID-19<br>vaccines<br>than have<br>died from<br>the COVID-<br>19 virus |
|---------------------|--|--|---|---|--|
| Total               | 34%  | 31%  | 27%   | 24%   | 20%  |
| Education           |  |  |   |   |  |
| High school or less | 43%  | 32%  | 36%   | 31%   | 30%  |
| Some college        | 35%  | 40%  | 26%   | 23%   | 20%  |
| College degree      | 23%  | 24%  | 20%   | 16%   | 9%   |
| Race/Ethnicity      |  |  |   |   |  |
| Black               | 43%  | 32%  | 31%   | 35%   | 29%  |
| Hispanic            | 37%  | 33%  | 29%   | 25%   | 24%  |
| White               | 32%  | 31%  | 26%   | 21%   | 17%  |
| Gender              |  |  |   |   |  |
| Men                 | 33%  | 29%  | 26%   | 22%   | 18%  |
| Women               | 35%  | 34%  | 29%   | 25%   | 22%  |
| Age                 |  |  |   |   |  |
| 18-29               | 33%  | 25%  | 26%   | 23%   | 24%  |
| 30-49               | 39%  | 36%  | 34%   | 29%   | 23%  |
| 50-64               | 38%  | 34%  | 28%   | 25%   | 17%  |
| 65+                 | 23%  | 27%  | 17%   | 15%   | 13%  |
| Party ID            |  |  |   |   |  |
| Republicans         | 47%  | 46%  | 39%   | 29%   | 24%  |
| Independents        | 42%  | 32%  | 34%   | 34%   | 28%  |
| Democrats           | 18%  | 17%  | 13%   | 14%   | 12%  |
| Community type      |  |  |   |   |  |
| Rural               | 42%  | 34%  | 39%   | 25%   | 28%  |
| Suburban            | 34%  | 32%  | 28%   | 23%   | 19%  |
| Urban               | 32%  | 30%  | 22%   | 24%   | 18%  |
|                     |  |  |   |   |  |

https://www.kff.org/coronavirus-covid-19/poll-finding/kffhealth-misinformation-tracking-poll-pilot/

NOTE: Persons of Hispanic origin may be of any race but are categorized as Hispanic for this analysis; other groups are non-Hispanic. Partisans include independents who lean to either party. Independents are pure independents. See topline for full question wording. SOURCE: KF Health Misinformation Tracking Poll Pilot (May 23-June 12, 2023)

## 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.

yourlocalepidemiologist.substack.com/p/the-science-and-business-behind-covid

# **Strategies for Addressing Hesitancy**

- Give your strong recommendation
  - Healthcare professionals remain the most trusted source for vaccine-related information

## Presumptive approach

• Start each vaccine conversation with a strong, positive "You are due for x vaccine today"

## Motivational interviewing

Readiness scaling to learn what is preventing someone from getting vaccinated

## Show your vaccine confidence

- Display posters in clinic spaces with vaccine confident messages
- Share your own stories of why you got vaccinated and why you recommend vaccines

# **About the Film**



## VIRULENT THE VACCINE WAR

LAURA DAVIS PRODUCTIONS PRESENTS VIRULENT: THE VACCINE WAR & FILM BY TJARDUS GREIDANUS IN ASSOCIATION WITH WOED ORIGINAL MUSIC BY GARY LIONELLI MEDICAL ADVISOR DR. PAUL OFFIT EXECUTIVE PRODUCERS MARK JONATHAN HARRIS, DEB ACKLIN, ROB DENSEN, SRIDHAR TAYUR, ANDREW VAGELOS PRODUCED BY LAURA DAVIS & TJARDUS GREIDANUS WRITTEN & DIRECTED BY TJARDUS GREIDANUS

VIRULENTMOVIE.COM

## **BREAK** Please return at 10:45

# Session III Vaccine Schedules



Alexander Sloboda, MD, MPH

# **Learning Objectives**

After this session participants will be able to:

Objective 1:

Apply current pediatric and adolescent vaccine recommendations.

**Objective 2:** 

Outline new vaccine products and updates.

Objective 3:

Apply the 2024 Advisory Committee on Immunizations Practices (ACIP) pediatric vaccination and catch-up schedules.

Objective 4:

Summarize current routine immunization rates.

# Immunization Schedules – Why They Matter

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

# Value of the Immunization Program

- An AAP study demonstrated that routine childhood vaccines help prevent unnecessary morbidity and mortality, as well as have cost-saving impacts.
- From 2017-2021, the ACIP-recommended schedule for routine childhood immunization has targeted 14 vaccine-preventable disease.
  - Diphtheria, Hib, hepatitis A and B, flu, MMR, pertussis, invasive Streptococcus pneumoniae, polio, rotavirus, tetanus, varicella
- Using this recommended schedule and the 2017 birth cohort, it was demonstrated that immunizations prevented over 17 million cases of disease and 31,000 deaths.
- Estimated vaccines costs of \$8.5 billion were entirely offset by the avoided \$63.6 billion in disease-related costs.

# Using the ACIP Schedule

To make vaccination recommendations, healthcare providers should:

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



Centers for Disease Control and Prevention CDC 24/7: Saving Lives, Protecting People™

## Search

Vaccines site 🔻

## Immunization Schedules

## For Healthcare Providers

Child and Adolescent Schedule Recommended vaccination schedule for ages 18 years or younger



## **Clinical Vaccination Resources**

Download Schedule App for Healthcare Providers

Vaccination Resources for Healthcare Providers

Birth to 18 Years

## Adult Schedule

Recommended vaccination schedule for ages 19 years or older

19 Years or Older



## Interim COVID-19 Immunization Schedule for Ages 6 months and older

Guidance for COVID-19 vaccination schedules based on age and medical condition

**COVID-19 Vaccination Schedule** 



Scan me to access the schedules on your phone

https://www.cdc.gov/vaccines/schedules/index.html

## **Approving Partners**

| Child/Adolescent<br>Schedule   | Both Schedules   | Adult Schedule   |
|--|--|--|
| American Academy of<br>Pediatrics (AAP)                              | American Academy of<br>Family Physicians (AAFP)                  | American College of<br>Physicians (ACP)                      |
| National Association of<br>Pediatric Nurse<br>Practitioners (NAPNAP) | American Academy of<br>Physician Associations<br>(AAPA)          | Society for HealthCare<br>Epidemiology of American<br>(SHEA) |
|  | American College of<br>Obstetricians and<br>Gynecologists (ACOG) | American Pharmacists<br>Association (APhA)                   |
|  | American College of Nurse-<br>Midwives (ACNM)                    |  |

## Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger

Abbreviation(s) Trade name(s)

Vaccines and Other Immunizing Agents in the Child and Adolescent Immunization Schedule\*

Managlanal antibada

## ACIP Recommended Child & Adolescent Schedule 2024

| Respiratory syncytial virus monoclonal antibody (Nirseyimab)                            | BSV-mAb                        | Bevfortus <sup>IN</sup>   |
|---|--------------------------------|---|
| Vaccine   | Abbreviation(s)                | Trade name(s)   |
| COVID-19  | 1vCOV-mRNA                     | Comimaty*/Pfizer-<br>BioNTech COVID-19<br>Vaccine<br>Spikevax*/Moderna<br>COVID-19Vaccine |
|   | 1vCOV-aP5                      | Novavax COVID-19<br>Vaccine   |
| Dengue vaccine  | DEN4CYD                        | Dengvaxia*  |
| Diphtheria, tetanus, and acellular pertussis vaccine                                    | DTaP                           | Daptacel*<br>Infanrix*  |
| Haemophilus influenzae type b vaccine   | Hib (PRP-T)                    | ActHIB*<br>Hiberix*   |
| Recorder & constant   | Hib (PRP-OMP)                  | Pequexela   |
| Hepatitis A vaccine   | нера                           | Vaqta*  |
| Hepatitis B vaccine   | Нер8                           | Engerix-B*<br>Recombivax HB*  |
| Human papillomavirus vaccine  | HPV                            | Gardasil 9*   |
| Influenza vaccine (inactivated)   | BV4                            | Multiple  |
| Influenza vaccine (live, attenuated)  | LAIV4                          | FluMist* Quadrivalent   |
| Measles, mumps, and rubella vaccine   | MMR                            | M-M-R I*<br>Priorix*  |
| Meningococcal serogroups A, C, W, Y vaccine   | MenACWY-CRM<br>MenACWY-TT      | Menveo*<br>MenQuadfi*   |
| Meningococcal serogroup B vaccine   | MenB-4C<br>MenB-EMbn           | Bexsero*  |
| Meningococcal serogroup A, B, C, W, Y vaccine   | MenACWY-TT/<br>MenB-FHbp       | Penbraya'*  |
| Mpox vaccine  | Мрок                           | Jynneos*  |
| Pneumococcal conjugate vaccine  | PCV15<br>PCV20                 | Vaxneuvance <sup>34</sup><br>Prevnar 20*  |
| Pneumococcal polysaccharide vaccine   | PPSV23                         | Pneumovax 23*   |
| Poliovirus vaccine (inactivated)  | IPV                            | lpol*   |
| Respiratory syncytial virus vaccine   | RSV                            | Abrysvo <sup>m</sup>  |
| Rotavirus vaccine   | RV1<br>RV5                     | Rotarix*<br>RotaTecr*   |
| Tetanus, diphtheria, and acellular pertussis vaccine                                    | Tdap                           | Adacel <sup>®</sup><br>Brostrix <sup>®</sup>  |
| Tetanus and diphtheria vaccine  | Td                             | Tentvac*<br>Tdvax*  |
| Varicella vaccine   | VAR                            | Varivax*  |
| Combination vaccines (use combination vaccines instead of separate in                   | njections when appropriate)    |   |
| DTaP, hepatitis B, and inactivated poliovirus vaccine                                   | DTaP-HepB-IPV                  | Pediarbr*   |
| DTaP, inactivated poliovirus, and Haemophiks influenzae type b vaccin                   | e DTaP-IPV/Hib                 | Pentacel*   |
| DTaP and inactivated poliovirus vaccine   | DTaP-IPV                       | Kinrix*<br>Quadracel*   |
| DTaP, inactivated poliovirus, Haemophilus influenzae type b, and<br>hepatitis B vaccine | DTaP-IPV-Hib-<br>Hep8          | Vaxelis <sup>e</sup>  |
| Measles, mumps, rubella, and varicella vaccine  | MMRV                           | ProOuad*  |
| "Administer recommended vaccines if immunization history is incomplete or u             | inknown. Do not restart or add | doses to vaccine series for   |

"Administer recommended vaccines if immunization history is incomplete or unknown. Do not restart or add doses to vaccine series for extended intervals between doses. When a vaccine is not administered at the recommended age, administer at a subsequent visit. The use of trade names is for identification purposes only and does not imply endorsement by the ACIP or CDC. 11/16/2023

| How to<br>schedul                                       | use the cl<br>e  | hild and   | adolesce   | nt immun  | ization  |
|---|--|--|--|---|--|
| 1   | 2  | 3  | 4  | 5   | 6  |
| Determine<br>recommended<br>vaccine by age<br>(Table 1) | Determine<br>recommended<br>interval for catch-<br>up vaccination<br>(Table 2) | Assess need<br>for additional<br>recommended<br>vaccines<br>by medical | Review<br>vaccine types,<br>frequencies,<br>intervals, and<br>considerations | Review<br>contraindications<br>and precautions<br>for vaccine types<br>(Appendix) | Review new or<br>updated AOP<br>guidance<br>(Addendum) |

Recommended by the Advisory Committee on Immunization Practices (www.cdc.gov/vaccines/acip) and approved by the Centers for Disease Control and Prevention (www.cdc.gov), American Academy of Pediatrics (www.aap.org), American Academy of Family Physicians (www.aafp.org), American College of Obstetricians and Gynecologists (www.acog.org), American College of Nurse-Midwives (www.midwife.org), American Academy of Physician Associates (www.aapa.org), and National Association of Pediatric Nurse Practitioners (www.napnap.org).

situations

(Notes)

### Report

- Suspected cases of reportable vaccine-preventable diseases or outbreaks to your state or local health department
- Clinically significant adverse events to the Vaccine Adverse Event Reporting System (VAERS) at www.waers.hhs.gov or 800-822-7967

other indication

(Table 3)

## Questions or comments

Contact www.cdc.gov/cdc-info or 800-CDC-INFO (800-232-4636), in English or Spanish, 8 a.m.-8 p.m. ET, Monday through Friday, excluding holidays



Download the CDC Vaccine Schedules app for providers at www.cdc.gov/vaccines/schedules/hcp/schedule-app.html

## Helpful information

- Complete Advisory Committee on Immunization Practices (ACIP) recommendations:
- www.cdc.gov/vaccines/hcp/acip-recs/index.html
- ACIP Shared Clinical Decision-Making Recommendations:
- www.cdc.gov/vaccines/acip/acip-scdm-faqs.html
- General Best Practice Guidelines for Immunization (including contraindications and precautions):
- www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html
- Vaccine information statements:
- www.cdc.gov/vaccines/hcp/vis/index.html \* Manual for the Surveillance of Vaccine-Preventable Diseases (including case identification and outbreak response):
- (including case identification and outbreak response): www.cdc.gov/vaccines/pubs/surv-manual



U.S. Department of Health and Human Services Centers for Disease Control and Prevention Scan QR code for access to online schedule

UNITED STATES



C\$310020-D

## Major Updates: 2024 Child and Adolescent Immunization Schedules

- Changes to Format
  - Changed headers from "Vaccine" to "Vaccines and other Immunizing Agents"
  - Cover page: COVID-19 is added. As well as Trade name Priorix for MMR and Vaxneuvance (PCV15) for pneumococcal conjugate vaccine.

- Changes to Vaccination Notes
  - RSV-mAb (nirsevimab)
  - RSVPreF (Abrysvo)
  - Jynneos (mpox)
  - Influenza
  - COVID-19
  - Meningococcal A, C, W, Y
  - Meningococcal B
  - Pneumococcal
  - Polio

Changes to Appendix

- Column Header
- Influenza
- Hepatitis B
- HPV
- Measles, Mumps an d Rubella
- Varicella

## Table 1 Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, United States, 2024

These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2).

| Vaccine and other immunizing agents                         | Birth                | 1 mo                      | 2 mos                         | 4 mos                 | 6 mos                       | 9 mos                     | 12 mos 1                         | 5 mos               | 18 mos                     | 19-23 mos           | 2-3 yrs    | 4-6 yrs                      | 7-10 yrs      | 11–12 yrs            | 13-15 yrs                       | 16 yrs                     | 17–18 yrs    |
|---|----------------------|---------------------------|-------------------------------|-----------------------|-----------------------------|---------------------------|----------------------------------|---------------------|----------------------------|---------------------|------------|------------------------------|---------------|----------------------|---------------------------------|----------------------------|--------------|
| Respiratory syncytial virus<br>(RSV-mAb [Nirsevimab])       |                      | 1 dose dep<br>RSV vaccina | pending on i<br>ition status, | maternal<br>See Notes |                             | 1 dose (                  | 8 through 19 m                   | onths), S           | ee Notes                   |                     |            |                              |               |                      |                                 |                            |              |
| Hepatitis B (HepB)  | 1 <sup>st</sup> dose | 4 2 <sup>nd</sup>         | dose>                         |                       | •                           |                           | 3 <sup>rd</sup> dose             |                     | >                          |                     |            |                              |               |                      |                                 |                            |              |
| Rotavirus (RV): RV1 (2-dose series),<br>RV5 (3-dose series) |                      |                           | 1 <sup>st</sup> dose          | 2 <sup>nd</sup> dose  | See Notes                   |                           |                                  |                     |                            |                     |            |                              |               |                      |                                 |                            |              |
| Diphtheria, tetanus, acellular pertussis<br>(DTaP <7 yrs)   |                      |                           | 1 <sup>st</sup> dose          | 2 <sup>nd</sup> dose  | 3 <sup>rd</sup> dose        |                           | -                                | ۰ 4 <sup>th</sup> d | ose>                       |                     |            | 5 <sup>th</sup> dose         |               |                      |                                 |                            |              |
| Haemophilus influenzae type b (Hib)                         |                      |                           | 1 <sup>st</sup> dose          | 2 <sup>nd</sup> dose  | See Notes                   |                           | ▲ <u>3rd or 4th d</u><br>See Not | ose,                |                            |                     |            |                              |               |                      |                                 |                            |              |
| Pneumococcal conjugate<br>(PCV15, PCV20)                    |                      |                           | 1 <sup>st</sup> dose          | 2 <sup>nd</sup> dose  | 3 <sup>rd</sup> dose        |                           | <b>∢</b> 4 <sup>th</sup> dos     | e•                  |                            |                     |            |                              |               |                      |                                 |                            |              |
| Inactivated poliovirus<br>(IPV <18 yrs)                     |                      |                           | 1 <sup>st</sup> dose          | 2 <sup>nd</sup> dose  | 4                           |                           | 3 <sup>rd</sup> dose             |                     | >                          |                     |            | 4 <sup>th</sup> dose         |               |                      |                                 |                            | See<br>Notes |
| COVID-19 (1vCOV-mRNA, 1vCOV-aPS)                            |                      |                           |                               |                       |                             |                           |                                  | 1 or m              | ore doses                  | of updated          | (2023–2024 | Formula) v                   | accine (See I | Notes)               |                                 |                            |              |
| Influenza (IIV4)  |                      |                           |                               |                       |                             |                           | An                               | nual vaco           | ination 1 o                | r 2 doses           |            |                              |               | Annu                 | al vaccination                  | n 1 dose or                | nly          |
| Influenza (LAIV4)   |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     | Ann        | ual vaccinat<br>1 or 2 doses | tion          | Ann                  | ual vaccinatio                  | n 1 dose o                 | only         |
| Measles, mumps, rubella (MMR)                               |                      |                           |                               |                       | See                         | Notes                     | < 1 <sup>st</sup> dos            | e•                  |                            | 1                   |            | 2 <sup>nd</sup> dose         |               |                      |                                 |                            |              |
| Varicella (VAR)   |                      |                           |                               |                       |                             |                           | < 1 <sup>st</sup> dos            | e•                  |                            |                     |            | 2 <sup>nd</sup> dose         |               |                      |                                 |                            |              |
| Hepatitis A (HepA)  |                      |                           |                               |                       | See                         | Notes                     | 2-0                              | lose serie          | s, See Note                | s                   |            |                              |               |                      |                                 |                            |              |
| Tetanus, diphtheria, acellular pertussis<br>(Tdap ≥7 yrs)   |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              |               | 1 dose               |                                 |                            |              |
| Human papillomavirus (HPV)                                  |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              | (**)          | See<br>Notes         |                                 |                            |              |
| Meningococcal (MenACWY-CRM ≥2 mos,<br>MenACWY-TT ≥2years)   |                      |                           |                               |                       |                             |                           | Se                               | e Notes             |                            |                     |            |                              |               | 1ª dose              |                                 | 2 <sup>nd</sup> dose       |              |
| Meningococcal B<br>(MenB-4C, MenB-FHbp)                     |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              |               |                      | See Not                         | es                         |              |
| Respiratory syncytial virus vaccine<br>(RSV [Abrysvo])      |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              |               | du                   | Seasonal adn<br>Iring pregnan   | ninistration<br>cy, See No | n<br>tes     |
| Dengue (DEN4CYD; 9-16 yrs)                                  |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              |               | Seropos<br>dengue a  | itive in ender<br>areas (See No | nic<br>tes)                |              |
| Мрох  |                      |                           |                               |                       |                             |                           |                                  |                     |                            |                     |            |                              |               |                      |                                 |                            |              |
| Range of recommended  | Range of r           | ecommend<br>up vaccinati  | led ages<br>on                | Rar<br>for            | nge of recor<br>certain hig | mmended a<br>h-risk group | iges                             | Recomm<br>can begi  | nended vac<br>n in this ag | cination<br>e group | R          | ecommende<br>n shared clir   | ed vaccinatio | on based<br>n-making | No                              | recommer<br>applicable     | ndation/     |

## Table 2Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More<br/>than 1 Month Behind, United States, 2024

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 5  |  |  |  |  |  |  |
| Hepatitis B   | Birth   | 4 weeks  | 8 weeks and at least 16 weeks after first dose<br>minimum age for the final dose is 24 weeks  |   |   |  |  |  |  |  |  |
| Rotavirus   | 6 weeks<br>Maximum age for first<br>dose is 14 weeks, 6 days. | 4 weeks  | 4 weeks<br>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<br>A fifth dose is not necessary<br>if the fourth dose was<br>administered at age 4 years or<br>older <i>and</i> at least 6 months<br>after dose 3 |  |  |  |  |  |  |
| Haemophilus influenzae<br>type b  | 6 weeks   | No further doses needed<br>if first dose was administered at age 15<br>months or older.<br>4 weeks<br>if first dose was administered before the<br>1 <sup>s</sup> birthday.<br>8 weeks (as final dose)<br>if first dose was administered at age<br>12 through 14 months.   | No further doses needed<br>if previous dose was administered at age 15 months or older<br>4 weeks<br>if current age is younger than 12 months and first dose was administered at younger than age 7 months and<br>at least 1 previous dose was PRP-T (ActHib's, Pentacel*, Hiberix*), Vaxelis* or unknown<br>8 weeks and age 12 through 59 months (an final dose)<br>if current age is younger than 12 months and first dose was administered at age 7 through 11 months; OR<br>if current age is 12 through 59 months and first dose was administered before the 1 <sup>st</sup> birthday and second<br>dose was administered at younger than 15 months; OR<br>if both doses were PedvaxHIB* and were administered before the 1st birthday | 8 weeks (as final dose)<br>This dose only necessary<br>for children age 12 through<br>59 months who received<br>3 doses before the<br>1" birthday.  |   |  |  |  |  |  |  |
| Pneumococcal conjugate  | 6 weeks   | No further doses needed for healthy<br>children if first dose was administered at<br>age 24 months or older<br>4 weeks<br>if first dose was administered before the<br>1 <sup>st</sup> birthday<br>8 weeks (as final dose for healthy<br>children)<br>if first dose was administered at the<br>1 <sup>st</sup> birthday or after | No further doses needed<br>for healthy children if previous dose was administered at age 24 months or older<br>4 weeks<br>if current age is younger than 12 months <i>and</i> previous dose was administered at <7 months old<br>8 weeks (as final dose for healthy children)<br>if previous dose was administered between 7–11 months (wait until at least 12 months old); OR<br>if current age is 12 months or older <i>and</i> at least 1 dose was administered before age 12 months   | 8 weeks (as final dose)<br>This dose is only necessary<br>for children age 12 through<br>59 months regardless of risk,<br>or age 60 through 71 months<br>with any risk, who received 3<br>doses before age 12 months. |   |  |  |  |  |  |  |
| Inactivated poliovirus  | 6 weeks   | 4 weeks  | 4 weeks<br>if current age is <4 years<br>6 months (as final dose)<br>if current age is 4 years or older   | 6 months (minimum age 4<br>years for final dose)  |   |  |  |  |  |  |  |
| Measles, mumps, rubella   | 12 months   | 4 weeks  |   |   |   |  |  |  |  |  |  |
| Varicella   | 12 months   | 3 months   |   |   |   |  |  |  |  |  |  |
| Hepatitis A   | 12 months   | 6 months   |   |   |   |  |  |  |  |  |  |
| Meningococcal ACWY  | 2 months MenACWY-CRM<br>2 years MenACWY-TT                    | 8 weeks  | See Notes   | See Notes   |   |  |  |  |  |  |  |
|   |   |  | Children and adolescents age 7 through 18 years   |   |   |  |  |  |  |  |  |
| Meningococcal ACWY  | Not applicable (N/A)  | 8 weeks  |   |   |   |  |  |  |  |  |  |
| Tetanus, diphtheria;<br>tetanus, diphtheria, and<br>acellular pertussis | 7 years   | 4 weeks  | 4 weeks<br>if first dose of DTaP/DT was administered before the 1 <sup>st</sup> birthday<br>6 months (as final dose)<br>if first dose of DTaP/DT or Tdap/Td was administered at or after the 1 <sup>st</sup> birthday   | 6 months<br>if first dose of DTaP/DT was<br>administered before the 1 <sup>st</sup><br>birthday   |   |  |  |  |  |  |  |
| Human papillomavirus  | 9 years   | Routine dosing intervals are<br>recommended.   |   |   |   |  |  |  |  |  |  |
| Hepatitis A   | N/A   | 6 months   |   |   |   |  |  |  |  |  |  |
| Hepatitis B   | N/A   | 4 weeks  | 8 weeks and at least 16 weeks after first dose  |   |   |  |  |  |  |  |  |
| Inactivated poliovirus  | N/A   | 4 weeks  | <b>6 months</b><br>A fourth dose is not necessary if the third dose was administered at age 4 years or older <b>and</b> at least 6 months<br>after the previous dose.   | A fourth dose of IPV is<br>indicated if all previous doses<br>were administered at <4<br>years <b>OR</b> if the third dose was<br>administered <6 months after<br>the second dose.                                    |   |  |  |  |  |  |  |
| Measles, mumps, rubella   | N/A   | 4 weeks  |   |   |   |  |  |  |  |  |  |
| Varicella   | N/A   | <b>3 months</b> if younger than age 13 years.<br><b>4 weeks</b> if age 13 years or older   |   |   |   |  |  |  |  |  |  |
| Dengue  | 9 years   | 6 months   | 6 months  |   |   |  |  |  |  |  |  |

## Table 3 Recommended Child and Adolescent Immunization Schedule by Medical Indication, United States, 2024

Always use this table in conjunction with Table 1 and the Notes that follow. Medical conditions are often not mutually exclusive. If multiple conditions are present, refer to guidance in all relevant columns. See Notes for medical conditions not listed.



## Notes Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2024

For vaccination recommendations for persons ages 19 years or older, see the Recommended Adult Immunization Schedule, 2024.

### Additional Information

 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. 32<sup>nd</sup> 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, RSV, 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 Pfize BioNTech COVID-19 vaccines], 12 years [Nova COVID-19 Vaccine])

### Routine vaccination

ge 6 months-4 years

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Unvaccinated:
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- 2-dose series of updated (2023–2024 Formula) Moderna at 0, 4-8 weeks
- 3-dose series of updated (2023–2024 Formula) Pfizer-BioNTech at 0, 3-8, 11-16 weeks
- Previously vaccinated\* with 1 dose of any Moderna: 1 dose of updated (2023–2024 Formula) Moderna 4-8 weeks after the most recent dose.
- Previously vaccinated<sup>+</sup> with 2 or more doses of any Moderna: 1 dose of updated (2023–2024 Formula) Moderna at least 8 weeks after the most recent dose.
- Previously vaccinated: with 1 dose of any Pfizer-BioNTech: 2-dose series of updated (2023–2024 Formula: Pfizer-BioNTech at 0, 8 weeks iminimum interval between previous Pfizer-BioNTech and dose 1: 3-8 weeks)
- Previously vaccinated: with 2 or more doses of any Pfizer-BioNTech: 1 dose of updated (2023–2024 Formula) Pfizer-BioNTech at least 8 weeks after the most recent dose.

### ge 5-11 years

- Invaccinated: 1 dose of updated (2023–2024 Formular Iodema or Pfizer-BioNTech vaccine
- Previously vaccinated with 1 or more doses of Moderna

### Special situations

Persons who are moderately or severely immunocompromised\*\* Age 6 months=4 years

- Unvaccinated:
- 3-dose series of updated (2023–2024 Formula) Moderna at 0, 4, 8 weeks
- 3-dose series of updated (2023–2024 Formula) Pfizer BioNTech at 0:3:11 weeks
- Previously vaccinated: with 1 dose of any Moderna:
   2-dose series of updated (2023–2024 Formula) Moderna at 0 4 weeks (minimum interval between previous Moderna and dose 1; 4 weeks).
- Previously vaccinated with 2 doses of any Moderna: 1 dose of updated (2023-2024 Formula) Moderna at least 4 weeks after the most recent dose
- Previously vaccinated: with 3 or more doses of any Moderna: 1 dose of updated (2023–2024 Formula) Moderna at least 8 weeks after the most recent dose
- Previously vaccinated: with 1 dose of any Pfizer-BioNTech: 2-dose series of updated (2023–2024 Formula) Pfizer-BioNTech at 0, 8 weeks (minimum interval between previous Pfizer-BioNTech and dose 1: 3 weeks).
- Previously vaccinated: with 2 or more doses of any Pfizer-BioNTech: 1 dose of updated (2023–2024 Formula) Pfizer-BioNTech at least 8 weeks after the most recent dose

### Age 5-11 years

 Unvaccinated:
 - 3-dose series of updated (2023–2024 Formula) Moderna at 0. 4, 8 weeks

The National Vaccine Injury Compensation Program (VICP) is a nofault 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, **RSV**, 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.

### Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2024 Notes

### Influenza vaccination

(minimum age: 6 months [IIV], 2 years [LAIV4], 18 years [recombinant influenza vaccine, RIV4])

### **Routine vaccination**

 Use any influenza vaccine appropriate for age and health status annually:

Age 6 months-8 years who have received fewer than 2 influenza vaccine doses before July 1, 2023, or whose influenza vaccination history is unknown: 2 doses, separated additional doses of MMR including 3rd dose of MMR, see by at least 4 weeks. Administer dose 2 even if the child turns 9 years between receipt of dose 1 and dose 2.

Age 6 months-8 years who have received at least 2 influenza vaccine doses before July 1, 2023: 1 dose - Age 9 years or older: 1 dose

### For the 2023-2024 season, see www.cdc.gov/mmwr/ volumes/72/rr/rr7202a1.htm.

 For the 2024–25 season, see the 2024–25 ACIP influenza vaccine recommendations.

### Special situations

 Close contacts (e.g., household contacts) of severely immunosuppressed persons who require a protected environment: these persons should not receive LAIV4. If LAIV4 is given, they should avoid contact with for such immunosuppressed persons for 7 days after vaccination

Note: Persons with an egg allergy can receive any influenza vaccine (egg-based and non-egg-based) appropriate for age and health status.

## Added information for vaccinating persons with a history of egg allergy.

## Notes

## es Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2024

### Special situations

- Revaccination is not generally recommended for persons with a normal immune status who were vaccinated as infants, children, adolescents, or adults
- Post-vaccination serology testing and revaccination of anti-HBs < 10mlU/mLi is recommended for certain populations inclusion.
- Infants born to HRs40-nositive mothers
- -Persons who are predialysis or on maintenance dialysis

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- uther minunocompromised perso
- gov/vaccines/http/acip-rectiva

Note: Heplisav-B and PreHevbrio ar pregnancy due to lack of safety dat

### Human papillomavirus va (minimum age: 9 years)

### **Routine and catch-up vaccin**

- HPV vaccination routinely recommended as age 11 12 ye (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 or older 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)
- No additional dose recommended when any HPV vaccine series of any valency has been completed using recommended dosing intervals.

### Special situations

- Immunocompromising conditions, including HIV Infection: 3-dose series, even for those who initiate vaccination at age 9 through 14 years
- + History of sexual abuse or assault: Start at age 9 years
- Pregnancy: Pregnancy testing not needed before vaccination; HPV vaccination not recommended until after pregnancy; no intervention needed if vaccinated while pregnant

Close contacts (e.g., household contacts) of severely mmunosuppressed persons who require a protected environment: these persons should not receive LAV4. FLAIV4 is given, they should avoid contact with for such mmunosuppressed persons for 7 days after vaccination.

Note: Persons with an egg allergy can receive any influenz; raccine legg-based and non-egg-based; appropriate for ac and health status;

### Aeasies, mumps, and rubella vaccination minimum age: 12 months for routine vaccinati

### **Routine vaccination**

- + 2-dose series at age 12–15 months, age 4–6 years
- MMR or MMRV may be administered

Deleted MenACWY-D (Menactra)

Added MenABCWY (Penbraya)

recommendations from all sections.

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.

### Catch-up vaccination

- Unvaccinated children and adolescents: 2-dose series at least 4 weeks apart
- The maximum age for use of MMRV is 12 years
- \* Minimum interval between MMRV doses: 3 month

### Special situation

### International travel

- Infants age 6–11 months: 1 dose before departure, revaccinate with 2-dose series at age 12–15 months (12 months for children in high-risk areas) and dose 2 as early as 4 weeks later
- Unvaccinated children age 12 months or older:
   2-dose series at least 4 weeks apart before departure
- In mumps outbreak settings, for information about additional doses of MMR lincluding 3rd dose of MMR, see www.cdc.adv.immwr.ixdlumes/67.vvi.ntine/20167.htm

### Meningococcal serogroup A,C,W,Y vaccination (minimum age: 2 months [MenACWY-CRM, Menveo], 2 years [MenACWY-TT, MenQuadfi]), 10 years [MenACWY-TT/MenB-FHbp, Penbraya])

### **Routine vaccination**

2-dose series at age 11–12 years; 16 years

### Catch-up vaccination

- Age 13–15 years: 1 dose now and booster at age 16–18 years (minimum interval: 8 weeks)
- Age 16–18 years: 1 dose

## Special situations

Anatomic or functional asplenia (including sickle cell disease), HIV infection, persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use:

### Menveo<sup>®\*</sup>

- Dose 1 at age 2 months: 4-dose series (additional 3 doses at age 4, 6, and 12 months)
- Dose 1 at age 3–6 months: 3- or 4-dose series (dose 2 [and dose 3 if applicable] at least 8 weeks after previous dose until a dose is received at age 7 months or older, followed by an additional dose at least 12 weeks later and after age 12 months)
- Dose 1 at age 7–23 months: 2-dose series (dose 2 at least 12 weeks after dose 1 and after age 12 months)
- Dose 1 at age 24 months or older: 2-dose series at least 8 weeks apart

### MenQuadfi<sup>®</sup>

 Dose 1 at age 24 months or older: 2-dose series at least 8 weeks apart

Travel to countries with hyperendemic or epidemic meningococcal disease, including countries in the African meningitis belt or during the Haii (www.cdc.gov/travel/):

## Notes

## Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2024

### Children less than age 24 months.

-Menveo \* (age 2-23 months

Dose 1 at age 2 months: 4-dose series (additional 3 doses at age 4, 6, and 12 months)

Dose 1 at age 3–6 months: 3- or 4-dose series (dose 2 [and dose 3 if applicable] at least 8 weeks after previous dose until a dose is received at age 7 months or older, followed by an additional dose at least 12 weeks later and after age 12 months:

Dose 1 at age 7–23 months. 2-dose series (dose 2 at least 12 weeks after dose 1 and after age 12 months)

 Children age 2 years or older: 1 dose Menveol\*\* or MenQuadfi\*

First-year college students who live in residential housing (If not previously vaccinated at age 16 years or older) or military recruits:

+ 1 dose Menveo - or MenQuadfi

Adolescent vaccination of children who received MenACWY prior to age 10 years:

Children for whom boosters are recommended because of an ongoing increased risk of meningococcal disease (e.g., those with complement component deficiency, HIV, or asplenial: Follow the booster schedule for persons at increased risk.

Children for whom boosters are not recommended is g, a healthy child who received a single dose for travel to a country where meningococcal disease is endemic: Administer MenACWY according to the recommended adolescent schedule with dose 1 at age 11–12 years and dose 2 at age 16 years

Menveo has two formulations: lyophilized and liquid. The liquid formulation should not be used before age 10 years. See when ada gov vocomes ypd. mening-downloads menueo-single-matpresentation pdf.

Note: For MeriACWY booster dose recommendations for groups listed under "Special situations" and in an outbreak setting and additional meningococcal vaccination information, sea www.cdc.gov/mmwcvolumes/69/m/m5909a1 htm.

Children age 10 years or older may receive a single dose of Penbraya \*\* as an alternative to separate administration of MenACWY and MenB when both vaccines would be given on the same clinic day, and a single injection with Penbraya \* is preferred (see "Meningococcal serogroup B vaccination" section below for more information. Meningococcal serogroup 8 vaccination (minimum age: 10 years (Men8-4C, Bexsero"; Men8-FHbp, Trumenba"])

## Shared clinical decision-making

Adolescents not at increased risk age 16–23 years (preferred age 16–18 years) based on shared clinical decision-making

Bexsero 1: 2-dose series at least 1 month apart

Trumenba 1: 2-dose series at least 6 months apart (If dose 2 is administered earlier than 6 months, administer a 31º dose at least 4 months after dose 2.

## Special situation

Anatomic or functional asplenia (including sickle cell disease), persistent complement component deficiency, complement inhibitor (e.g., eculizumab, ravulizumab) use

Bexsero 1:2-dose series at least 1 month apart

Trumenball: 3-dose series at 0, 1–2, 6 months lif dose 2 was administered at least 6 months after dose 1, dose 3 not needed; if dose 3 is administered earlier than 4 months after dose 2, a 4<sup>ll</sup> dose should be administered at least 4 months after dose 3)

Note: Bexsero and Trumenba are not interchangeable, the same product should be used for all doses in a series.

For Men8 booster dose recommendations for groups listed under "Special situations" and in an outbreak setting and additional meningococcal vaccination information, see www.cdc.gov.mmvr.volumes.69/mmc908a1.htm

Children age 10 years or older may receive a dose of Penbraya as an alternative to separate administration of MenACWY and MenB when both vaccines would be given on the same clinic day, and a single injection with Penbraya" is preferred. If usin Penbraya" for dose 1 MenB subsequent MenB doses must be either MenB-FHbp (Trumenbai or Penbraya" (minimum interval between Penbraya" doses 6 months). Children age 10 years or older recommended to receive booster doses of MenACWY and MenB less than 6 months after a dose of Penbraya" should receive MenACWY and MenB-FHbp (Trumenbai separately.

## Apox vaccination minimum age: 18 years [Jynneos\*

## Special situations

 Age 18 years and at risk for Mpox infection: 2-dose series, 28 days apart.

Risk factors for Mpox infection include:

 Persons who are gay, bisexual and other MSM, transgender or nonbinary people who in the past 6 months have had;

At least 1 sexually transmitted dis

More than 1 sex partner

Sex at a commercial sex venue.

Sex in association with a large public event in a geographic area where Mpox transmission is occurring.

Persons who are sexual contacts of the persons described above

Persons who anticipate experiencing any of the situations described above.

 Persons deemed at risk by public health authorities in mpoxoutbreak settings

 Pregnancy: There is currently no ACIP recommendation for Jynneos use in pregnancy due to lack of safety data in pregnant persons. Pregnant persons with any risk factor described above may receive Jynneos.

For detailed information, see, www.cdc.gov/pow/ros/mpox/ Interim-considerations/wineos-vaccine.html

## Pneumococcal vaccination (minimum age: 6 weeks [PCV15], [PCV 20]; 2 years [PPSV23])

## **Routine vaccination with PCV**

4-dose series at 2, 4, 6, 12–15 months

## Catch-up vaccination with PCV

 Healthy children ages 2–4 years with any incomplete\* PCV series: 1 dose PCV

For other catch-up guidance, see Table 2.

Note: Either PCV15 or PCV20 can be used when PCV is indicated. PCV20 is not indicated for children who have received 4 doses of PCV13 or PCV15 or another age appropriate complete PCV series.



## Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2024

Mpox vaccination (minimum age: 18 years [Jynneos\*]) Special situations **Special situations**  Age 18 years and at risk for Mpox infection: 2-dose series, 8 days apart. Risk factors for Mpox infection include: Age 18 years and at risk for Mpox - Persons who are gay, bisexual, and other MSM, transgender or nonbinary people who in the past 6 months have had: infection: 2-dose series, 28 days apart. A new diagnosis of at least 1 sexually transmitted disease. More than 1 sex partner. Risk factors for Mpox infection include: Sex at a commercial sex venue. Sex in association with a large public event in a geographic area where Mpox transmission is occurring. - Persons who are sexual partners of the persons described above. - Persons who anticipate experiencing any of the situations described above. Pregnancy: There is currently no ACIP recommendation for Jynneos use in pregnancy due to lack of safety data in Added bullet on use of Jynneos in Children for whom boosters are red . pregnant persons. Pregnant persons with any risk factor described above may receive Jynneos. pregnant persons For detailed information, see: https://www.cdc.gov/ vaccines/acip/meetings/downloads/ Children for whom boosters are no slides-2023-10-25-26/04-MPOX-Rao-508.pdf

# February ACIP Meeting Updates

- COVID-19: ACIP recommends that persons ≥ 65 years of age should receive an additional dose of 2023-2024 Formula COVID-19 vaccine.
- Chikungunya: Chikungunya vaccine is recommended for persons aged ≥18 years traveling to a country or territory where there is an outbreak and for laboratory workers with potential for exposure to chikungunya virus.
- Diphtheria, Tetanus, and Pertussis: Approve the VFC resolution for diphtheria, tetanus, and pertussis to (1) add Td vaccine for use in children < 7 years of age for whom receipt of the pertussis component is contraindicated and to (2) update the language regarding the Tdap booster to align with ACIP recommendations.
- Updates to the meningococcal vaccine schedule may come later this year.

## **RSV Prevention Products**

- Nirsevimab (Beyfortus)
- Abrysvo
- Arexvy

## **RSV** Products Visual Guide

FOR PATIENTS WHO ARE:

## FOR PATIENTS WHO ARE: Pregnant

Administer ABRYSVO. Abrysvo is supplied in a kit that includes a vial of Lyophilized Antigen Component, a prefilled syringe containing diluent, and a vial adapter.

Timing: a single dose during weeks 32 through 36 of pregnancy during September through January

## A newborn or infant

Administer **BEYFOTRUS**. Beyfortus is supplied in a prefilled syringe. 50mg doses of Beyfortus are light blue and 100mg doses are purple.

Timing: One dose just before or during RSV season.



## FOR PATIENTS WHO ARE: 60 years or older

Administer AREXVY. Arexvy is supplied in two vials that must be combined prior to administration.

Timing: One dose just before RSV season.





Premature/at high risk






### Niservimab (Beyfortus) - RSV Monoclonal Antibody)

- Infants born shortly before or during RSV season (October March)
  - Administer 1 dose nirsevimab within 1 week of birth if mother did not receive Abrysvo, mother's RSV vaccination status is unknown, or if mother received Abrysvo less than 14 days prior to delivery
  - May be given in birth hospital or PCP office
- Infants born April–September entering their first RSV season
  - Mother did not receive RSV vaccine OR mother's RSV vaccination status is unknown: administer 1 dose nirsevimab shortly before start of RSV season
- May consider giving nirsevimab to children 8 19 months who are entering their 2nd RSV season who are at increased for severe disease related to RSV infection.

### RSV Vaccines: Abrysvo (Pfizer), Arexvy (GSK)

### Pregnancy:

- All pregnant people should only receive Abrysvo (Pfizer)
- One dose should be administered during RSV season (September-January) for those between 32 – 36 weeks gestation
- Currently, there is no data on revaccinating with every pregnancy like Tdap – studies are ongoing

### Older Adults:

- Arexvy and Abrysvo are recommended for adults 60 years and older
- One dose recommended before onset of fall/winter RSV season
- Currenly, the recommendation is one dose Studies are ongoing to assess if boosters are needed in older adults

## **RSV Season**

- Abrysvo ordering through the CDPH VFC program is no longer available,
- All remaining VFC doses of Abrysvo and nirsevimab have expiration dates that will allow them to be used for parts or all of the next RSV season.
- The CDC recommends administering nirsevimab to eligible newborns through March 31, 2024.
- Pediatricians can also now reserve private doses of Beyfortus for the 2024-2025 respiratory virus season through Sanofi's new reservation program.
  - Does not apply to VFC program Sanofi continues to engage with CDC to ensure supply for the VFC program

## **Hepatitis B Vaccine**

- Routine vaccination:
  - 3-dose series at age 0, 1–2, 6–18 months (use monovalent HepB vaccine for birth dose and any doses administered before age 6 weeks).
  - Additional steps dependent on mother's HBsAg status.
- Heplisav-B and PreHevbrio may be used for catch up for age 18 years and older.
- Heplisav-B not recommended during pregnancy.

## Hepatitis B Vaccine cont.

- Providers are required to refer pregnant women who are HBsAg-positive within 7 days after receipt of the test result to the local health department for case management.
- Chicago providers can fulfill this reporting requirement by providing contact information for the patient, along with demographics, and HBsAg test date via <u>CDPH's secure</u> <u>online reporting form on REDCap</u>. This is especially important if providers use a commercial lab.

## **Rotavirus Vaccine**

- Products
  - RotaTeq® (RV5)
  - Rotarix® (RV1)
- Routine Vaccination:
  - Rotarix®: 2-dose series at age 2 and 4 months
  - RotaTeq®: 3-dose series at age 2, 4, and 6 months
  - If any dose in the series is either RotaTeq® or unknown, default to 3-dose series.
- Catch-Up Vaccination:
  - Do not start the series on or after age 15 weeks, 0 days.
  - The maximum age for the final dose is 8 months, 0 days.

## **Rotavirus Vaccine**

- Rotavirus (Rotarix ™)
  - <u>NO RECONSTITUTION NEEDED!</u>
  - Oral-dosing applicator-only presentation.
  - FDA approved in Nov 2022.
  - There are 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/Tdap

### DTaP:

- 5-dose series (3-dose primary series at age 2, 4, and 6 months, followed by a booster doses at ages 15–18 months and 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

### Tdap:

• Age 11–12 years: 1 dose Tdap (adolescent booster)

### **Catch-Up Vaccination:**

- DTaP: Dose 5 is not necessary if dose 4 was administered at age 4 years or older and at least 6 months after dose 3
- Tdap: Dependent on age and DTaP vaccination history

## Hib

# **Products:** ActHIB®, Hiberix®, Pentacel®, PedvaxHIB® or Vaxelis®

### **Routine Vaccination**

- 4-dose series (3-dose primary series at age 2, 4, and 6 months, followed by a booster dose\* at age 12–15 months)
  - Vaxelis® is not recommended for use as a booster dose. A different Hib-containing vaccine should be used for the booster dose.
  - PedvaxHIB®: 3-dose series (2-dose primary series at age 2 and 4 months, followed by a booster dose at age 12–15 months)

### **Catch-Up Vaccination**

Refer to catch up schedule: Dependent on age and vaccination history

## **Pneumococcal Vaccine**

- Products:
  - Pneumococcal conjugate vaccines (PCVs, specifically PCV15 and PCV20) ACIP has not made a preferential statement
  - Pneumococcal polysaccharide vaccine (PPSV23)
- Routine Vaccination with PCV:
  - 4-dose series at 2, 4, 6, 12–15 months
- Catch-Up Vaccination with PCV:
  - Healthy children ages 2–4 years with any incomplete\* PCV series: 1 dose PCV
  - Note: Either PCV15 or PCV20 can be used when PCV is indicated. For children without risk conditions, PCV20 is not indicated if they have received 4 doses of PCV13 or PCV15 or another age appropriate complete PCV series

## Pneumococcal Vaccine cont.

- PCV13 is no longer distributed or recommended for use in the U.S
- Pneumococcal polysaccharide vaccine PPSV23 (Pneumovax23, Merck)
  - No longer routinely recommended for all children and adolescents aged ≥2 years at increased risk for invasive pneumococcal disease. It is still recommended in certain circumstances

## **Polio Vaccine**

- Routine Vaccination:
  - 4-dose series at ages 2, 4, 6–18 months, 4–6 years
  - 4 or more doses of IPV can be administered before age 4 years when a combination vaccine containing IPV is used

### Catch-Up Vaccination:

- In the first 6 months of life, use minimum ages and intervals only for travel to a polio-endemic region or during an outbreak
- Adolescents age 18 years known or suspected to be unvaccinated or incompletely vaccinated: administer remaining doses (1, 2, or 3 IPV doses) to complete a 3-dose primary series

## COVID-19

- 2023 2024 Pfizer, Moderna, and Novavax COVID-19 vaccines were authorized and recommended in September 2023.
- Everyone 6 months and older should receive a COVID-19 vaccine
  - Most people only require one dose
  - Children 6 months to 4 years will need multiple doses if they are starting a series or having not completed a primary series.
- All VFC providers are required to stock and recommend COVID vaccines
  - VFC providers are required to meet the private inventory requirement no later than March 31, 2024.

### **COVID-19 VACCINATION SCHEDULE AND DOSING**

#### **AGES 6 MONTHS TO 4 YEARS**





### **COVID-19 VACCINATION SCHEDULE AND DOSING**

#### AGES 6 MONTHS TO 4 YEARS IMMUNOCOMPROMISED



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



**Complete at least a three-dose series with a COVID-19 vaccine, each dose one month apart.** At least one dose should be with a COVID-19 vaccine (2023-2024 Formula).

Additional age-appropriate doses may be administered <u>at the discretion</u> of the healthcare provider, taking into consideration the individual's clinical circumstances. The timing of the additional doses may be based on the individual's clinical circumstances.

### **COVID-19 VACCINATION SCHEDULE AND DOSING** AGES 5 TO 11 YEARS





#### PREVIOUSLY VACCINATED

dose/injection volume

Previously Received COVID-19 Vaccines

Moderna 2023-2024: (Do NOT dilute before use) Dark Blue Cap (green label)

Pfizer 2023-2024: (Do NOT dilute before use) Blue Cap



### **COVID-19 VACCINATION SCHEDULE AND DOSING**

#### **AGES 5 TO 11 YEARS IMMUNOCOMPROMISED**





#### PLEASE NOTE

Complete at least a three-dose series with a COVID-19 vaccine, each dose one month apart. At least one dose should be with a COVID-19 vaccine (2023-2024 Formula).

Additional age-appropriate doses may be administered <u>at the discretion</u> of the healthcare provider, taking into consideration the individual's clinical circumstances. The timing of the additional doses may be based on the individual's clinical circumstances.

### **COVID-19 VACCINATION SCHEDULE AND DOSING** AGES 12 YEARS AND OLDER





#### PREVIOUSLY VACCINATED

dose/injection volume

Previously Received COVID-19 Vaccines

Moderna 2023-2024: (Do NOT dilute before use) Dark Blue Cap (dark blue label)

Pfizer 2023-2024: (Do NOT dilute before use) Gray Cap



### **COVID-19 VACCINATION SCHEDULE AND DOSING**

#### **AGES 12 YEARS AND OLDER IMMUNOCOMPROMISED**





#### PLEASE NOTE

**Complete at least a three-dose series with a COVID-19 vaccine, each dose one month apart.** At least one dose should be with a COVID-19 vaccine (2023-2024 Formula).

Additional age-appropriate doses may be administered at the discretion of the healthcare provider, taking into consideration the individual's clinical circumstances. The timing of the additional doses may be based on the individual's clinical circumstances.

### COVID-19 VACCINATION SCHEDULE AND DOSING AGES 12 YEARS AND OLDER



#### UNVACCINATED

dose/injection volume

Novavax 2023-2024: (Do NOT dilute before use) Royal Blue Cap



#### PREVIOUSLY VACCINATED

dose/injection volume

Previously Received COVID-19 Vaccines

Novavax 2023-2024: (Do NOT dilute before use) Royal Blue Cap



### **COVID-19 VACCINATION SCHEDULE AND DOSING**

### **AGES 12 YEARS AND OLDER IMMUNOCOMPROMISED**



Royal Blue Cap

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

PLEASE NOTE

Additional doses may be administered at the discretion of the healthcare provider, taking into consideration the individual's clinical circumstances. The timing of the additional doses may be based on the individual's clinical circumstances.

## Influenza Vaccine

• Annual vaccination of 1 or 2 doses recommended for everyone 6 months and older.



- People with a history of egg allergy of any severity can be vaccinated with any influenza vaccine indicated for the recipient's age and health status with no additional safety considerations.
  - If using egg-based IIV4 or LAIV4, administer in medical setting under supervision of health care provider who can recognize and manage severe allergic reactions.

## **MMR Vaccine**

### • Products:

- M-M-R II® (MMR) vaccine Merck & Co, Inc.
- PRIORIX® (GSK).
- ProQuad® (MMRV) vaccine.

### Routine Vaccination:

- 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.
- \*Note: If MMRV is used, the minimum interval between MMRV doses is 3 months.
- Recommendation for the MMRV to be separated for the 1st dose is because there was a slight increase in febrile seizures seen in the combination vaccine.

## **MMR Vaccine**

- Catch Up Vaccination:
  - Unvaccinated children and adolescents: 2-dose series at least 4 weeks apart.\*
  - The maximum age for use of MMRV\* is 12 years.
  - \*Note: If MMRV is used, the minimum interval between MMRV doses is 3 months.
- Providers should ensure everyone is up-to-date with MMR vaccine
  - Especially tin the context of international travel and local outbreaks



January 25, 2024

**Stay Alert for Measles Cases** 

### **Measles Reminder**





DEDICATED TO THE HEALTH OF ALL CHILDREN®

### Think Measles

Consider measles in any patient presenting with a febrile rash illness, especially if unvaccinated for measles or traveled internationally in the last 21 days.

#### Measles Symptoms

- High Fever
- Cough
- Coryza (runny nose)
- Conjunctivitis (red, watery eyes)
- Maculopapular Rash
  - Typically appears 2-4 days after symptoms begin
  - Begins at hairline, spreads downward, to face, neck, and trunk
  - Rash appears red on light complexions, but may be harder to see or appear as purple or darker than surrounding skin on dark complexions.

#### **2** Pre-Visit Telephone Triage

- For those reporting measles symptoms, assess the risk of exposure:
  - · Are measles cases present in your community?
  - Did the patient spend time out of the country in the 21 days before symptom onset?
  - Has the patient ever received the MMR vaccine?
- Triage should only be completed by a clinically trained person.
- If patient will be seen in the office, provide instructions on face masks for patient (2 years of age and older) and family.
- Instruct to arrive to a side or back entrance instead of the main entrance.

## Varicella

### Routine Vaccination:

- 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 4 weeks 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.
- Catch-Up Vaccination:
  - Ensure persons age 7–18 years without <u>evidence of immunity</u> have a 2-dose series.

## **Hepatitis A**

- Routine Vaccination:
  - 2-dose series (minimum interval: 6 months) at age 12–23 months.
  - Never need to restart the series you pick up where you left off (same with Hep B).
- Catch-Up Vaccination:
  - Unvaccinated persons through age 18 years should complete a 2-dose series (minimum interval: 6 months).
  - Persons who previously received 1 dose at age 12 months or older should receive dose 2 at least 6 months after dose 1.
  - Twinrix® (HepA and HepB vaccine): 18 years or older.
    - 3-dose series (0, 1, and 6 months) or
    - 4-dose series (3 doses at 0, 7, and 21–30 days, followed by a booster dose at 12 months.

\*May give Hep A for travel to infants as young as 6 months but it does not count in the series if traveling internationally

## HPV

- HPV vaccination is cancer prevention
  - Cervical, oropharyngeal, anal, and others
- Recommended for both girls and boys
- Routinely recommended starting at 11 but administration can start at 9
  - Starting HPV vaccine discussions at age 9 gives more time for parents to make the decision to vaccinate
- View the HPV vaccine schedule on the next slide



### **HPV**

**If your child is 9- to 14-years-old,** your child's doctor will determine whether your child needs a 2-dose or 3-dose schedule of GARDASIL 9.



<sup>a</sup>If the second shot is given earlier than 5 months after the first shot, your child will need to get a third shot at least 4 months after the second shot was given.



## **Meningococcal Vaccines**

- Products:
  - Meningococcal conjugate or MenACWY vaccines (Menveo® and MenQuadfi®).
  - Serogroup B meningococcal or MenB vaccines (Bexsero® and Trumenba®).
  - Pentavalent meningococcal or MenABCWY vaccine (PenbrayaTM).
- Routine Vaccination:
  - 2-dose series at age 11-12 years; 16 years.
- Catch-Up Vaccination:
  - Age 13-15 years; 1 dose now and booster at age 16-18 years (minimum interval: 8 weeks).
  - Age 16-18 years: 1 dose.

## Meningococcal Vaccines Cont.

- MenACWY (Menactra) not recommended or distributed.
- Pfizer's pentavalent meningococcal vaccine (Penbraya) approved by ACIP in October 2023 for use in adolescents and young adults ages 10 to 25 years. The vaccine includes serogroups A, B, C, W, and Y.
  - Approved for the VFC program.
- 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.

## **Mpox Vaccine**

- Since December 1, 2023, 22 mpox cases have been reported in Chicago; of which 17 (77%) have been reported since January 14th, 2024.
- Age 18 years and at risk for Mpox infection: 2-dose series, 28 days apart.
- Risk factors for Mpox infection include:
  - Persons who are gay, bisexual, and other MSM, transgender or nonbinary people who in the past 6 months have had:
  - A new diagnosis of at least 1 sexually transmitted disease.
  - More than 1 sex partner.
  - Sex at a commercial sex venue.
  - Sex in association with a large public event in a geographic area where Mpox transmission is occurring.
  - Persons who are sexual partners of the persons described above.
  - Persons who anticipate experiencing any of the situations described above.
- This vaccine has not yet been studied in pregnancy.

### Start vaccinating now, before spring and summer festivals!

## **VAERS Reminder**

- Healthcare providers are <u>required by law</u> to report to VAERS:
- Any adverse event listed in the VAERS Table of Reportable Events Following Vaccination that occurs within the specified time period after vaccinations.
- An adverse event listed by the vaccine manufacturer as a contraindication to further doses of the vaccine.
- Healthcare providers are <u>strongly encouraged</u> to report to VAERS:
- Any adverse event that occurs after the administration of a vaccine licensed in the United States, whether it is or is not clear that a vaccine caused the adverse event.
- Vaccine administration errors.

| /AERS Home                               |   |   |  |  |
|--|---|---|--|--|
| VAERS Home<br>About VAERS                | Home / Report an Adverse Event  | /_en Españ  |  |  |
| Report an Adverse Event –                | Report an Adverse Event to VAERS  |   |  |  |
| Report Online<br>Report Using a PDF Form | VAERS Reporting Information for COVID-19 Vaccines   |   |  |  |
| /AERS Data +                             | VAERS Reporting Requirements for Beyfortus (nirsevimab)   |   |  |  |
| Resources +                              | VAERS Reporting Requirements for Monkeypox vaccines   |   |  |  |
| ubmit Follow-Up Information              | Knowingly filing a false VAERS report is a violation of Federal law (18 U.S.  | Click here for information on reporting   |  |  |
| requently Asked Questions                | Code § 1001) punishable by fine and imprisonment.   | to VAERS after COVID-19 vaccination   |  |  |
| Contact Us                               | Two Ways to Submit an Online Report to VAERS  |   |  |  |
| Privacy                                  | Option 1 - Report Online to VAERS         Submit a VAERS report online. The report must         be completed online and submitted in one sitting         and canot be saved and returned to at a later         time. Your information will be erased if you are         inactive for 20 minutes; you will receive a         warning after 15 minutes. | Checklist<br>What will I need to fill out the<br>report?<br>• Patient information (age, date of<br>birth, sex)<br>• Vaccine information (brand name,<br>dosage)   |  |  |
|  | Option 2 - Report using a Writable PDF<br>Form<br>Download the Writable PDF Form to a computer.<br>Complete the VAERS report offline if you do not<br>have time to complete it all at once. Return to<br>this page to upload the completed Writable PDF<br>form by clicking here.   | <ul> <li>Date, time, and location<br/>administered</li> <li>Date and time when adverse<br/>event(s) started</li> <li>Symptoms and outcome of the<br/>adverse event(s)</li> <li>Medical tests and laboratory<br/>results (if applicable)</li> <li>Physician's contact information (if<br/>applicable)</li> </ul> |  |  |

## **Chicago Vaccine Disparities**

100%

100%

Influenza and COVID-19 Vaccination Coverage, 0-17 Years



Influenza and COVID-19 Vaccination Coverage, 65+ Years



Influenza and COVID-19 Vaccination Coverage, 18-64 Years



Influenza and COVID-19 Vaccination Coverage, All Ages



100%

## **HCEZ Racial Vaccine Disparities**

#### **Percent of People Up to Date with COVID-19 Vaccinations by Region**

Welcome to the COVID-19 Vaccination Healthy Chicago Equity Zones (HCEZ) dashboard. To view coverage and insights for a specific region, age group, and race-ethnicity, click on the cell in the table for the group you are interested in.

| Age Group and<br>Race/Ethnicity | Citywide | Far South | Near South | North/Central | Northwest | Southwest | West |
|---------------------------------|----------|-----------|------------|---------------|-----------|-----------|------|
| 0-17 yrs                        |          |           |            |               |           |           |      |
| All Race/Ethnicities            | 8%       | 2%        | 4%         | 17%           | 9%        | 4%        | 6%   |
| Asian, non-Latinx               | 11%      | 8%        | 20%        | 13%           | 10%       | 5%        | 22%  |
| Black, non-Latinx               | 2%       | 2%        | 2%         | 5%            | 7%        | 2%        | 2%   |
| Latinx                          | 6%       | 3%        | 7%         | 13%           | 5%        | 4%        | 6%   |
| White, non-Latinx               | 16%      | 4%        | 31%        | 21%           | 13%       | 6%        | 18%  |
| 18–64 yrs                       |          |           |            |               |           |           |      |
| All Race/Ethnicities            | 13%      | 7%        | 8%         | 21%           | 13%       | 7%        | 11%  |
| Asian, non-Latinx               | 14%      | 15%       | 12%        | 16%           | 13%       | 8%        | 16%  |
| Black, non-Latinx               | 6%       | 6%        | 6%         | 12%           | 11%       | 6%        | 5%   |
| Latinx                          | 8%       | 5%        | 9%         | 15%           | 7%        | 5%        | 7%   |
| White, non-Latinx               | 19%      | 9%        | 21%        | 21%           | 16%       | 9%        | 17%  |
| 65+ yrs                         |          |           |            |               |           |           |      |
| All Race/Ethnicities            | 32%      | 29%       | 27%        | 47%           | 30%       |           | 25%  |
| Asian, non-Latinx               | 28%      |           | 31%        | 30%           | 25%       | 25%       | 27%  |
| Black, non-Latinx               | 26%      | 26%       | 25%        | 35%           | 25%       | 24%       | 21%  |
| Latinx                          | 24%      | 22%       | 16%        | 30%           | 24%       | 23%       | 21%  |
| White, non-Latinx               | 38%      | 31%       | 51%        | 48%           | 29%       | 24%       | 36%  |
| All Ages                        |          |           |            |               |           |           |      |
| All Race/Ethnicities            | 14%      | 10%       | 10%        | 23%           | 14%       | 8%        | 11%  |
| Asian, non-Latinx               | 15%      | 15%       | 13%        | 17%           | 15%       | 10%       | 17%  |
| Black, non-Latinx               | 9%       | 9%        | 8%         | 14%           | 12%       | 8%        | 7%   |
| Latinx                          | 8%       | 6%        | 9%         | 16%           | 8%        | 6%        | 8%   |
| White, non-Latinx               | 21%      | 12%       | 27%        | 25%           | 18%       | 13%       | 18%  |



Data reported through: January 27, 2024.

Data are updated the first Wednesday of the month at 3:30 p.m., except for City holidays. All data are provisional and subject to change.

## What Can Be Done To Increase Vaccine Uptake?

### AAP suggests:

- Use the presumptive approach.
- Implementing reminder recall strategies.
- Promoting the VFC program to ensure un/under-insured have access to all recommended vaccines.
- Make every visit a vaccine visit by making a strong recommendation at **all** visits.

**CDC's** <u>Let's RISE initiative</u>: Routine vaccination is rebounding but unevenly and not among all groups. Many are still behind schedule.

- Send reminders to families whose children are behind on or due for vaccination,
- Improve vaccine related communications
- Offer vaccination-only appointments or hold vaccination clinics
- Have standing orders
- Be prepared to answer questions and address specific concerns
## **Presumptive Approach**

It's so great to see you back, I missed you guys!

...

Mike is going to get Tdap, HPV, COVID and meningococcal vaccine today.

> I'm so excited to protect Mike from pertussis, cancer, COVID and meningitis today. Let's get this done!



He is so sick of school half on the computer and not seeing his friends. We're ready!

## **Provider Recommendations Make a Difference to Their**

### **Patients**



# **Helpful Links**

- The <u>Recommended Child and Adolescent</u>
  <u>Immunization Schedule, United States, 2024</u> is now
  available on *Red Book Online* (RBO).
- AAP Policy Statement: <u>Recommended Childhood</u> <u>and Adolescent Immunization Schedule: United</u> <u>States, 2024.</u>
- CDC: <u>2024 Child and Adolescent Immunization</u> <u>Schedule.</u>

## **Thank you!** We will now distribute awards and raffle prizes, share important last steps, and take questions.

## Vaccine Coverage Awards

# If you see your clinic, be ready to come up to the stage for an **AWARD**

**High Pediatric Vaccine Coverage** 80% or greater for 4DTaP, 3Polio, 1MMR, Hib UTD, HBV UTD, 1 VAR, PCV UTD (≥ 15 records)

C01890 ADVOCATE MED GRP EVERGEEN

Congratulations and thank you for your hard work!

#### High Adolescent Vaccine Coverage

Tdap ≥ 90%, MCV ≥ 80%, HPV UTD ≥ 85% (>1000 records)

C01720 ESPERANZA MARQUETTE CLINIC

C01581 FRIEND FAMILY HC PULASKI

Congratulations and thank you for your hard work!

## **High HPV Vaccine Coverage** 85% or greater for HPV UTD (≥ 10 records)

C01720 ESPERANZA MARQUETTE

Congratulations and thank you for your hard work!

# **Raffle Winners!**

## **Upcoming Events 2024**

- March 19 at 12PM Adolescent Immunization Action Week: A National Movement to Get Adolescents Up to Date on Immunizations
- April 17 at 12PM HPV Vaccinations (Trends and Updates)
- Find and register for all events at: <u>https://illinoisaap.org/upcoming-events/</u>



# **Evaluation**

• You must complete the evaluation by 3/13/24 to receive CME and/or a completion certificate.



# **Questions?**