National Center for Immunization and Respiratory Diseases



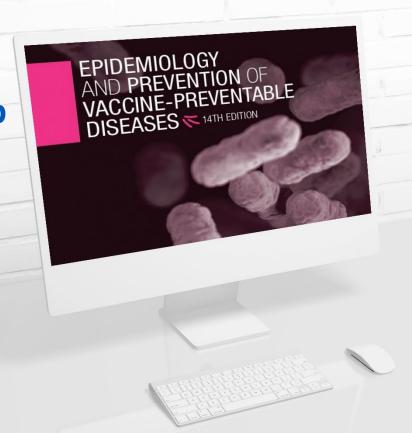
Polio and

Haemophilus influenzae type b

Vaccines

Pink Book Web-on-Demand Series

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Learning Objectives

- Describe the fundamental principles of the immune response.
- Describe immunization best practices.
- Describe an emerging immunization issue.
- For each vaccine-preventable disease, identify those for whom routine immunization is recommended.
- For each vaccine-preventable disease, describe characteristics of the vaccine used to prevent the disease.
- Locate current immunization resources to increase knowledge of team's role in program implementation for improved team performance.

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Disclosure Statements

- CDC did not accept financial or in-kind support from any ineligible company for this continuing education activity.
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Polio Disease

Poliomyelitis Disease

- Caused by poliovirus
- Epidemics starting in the late 19th century
- Polio epidemics were reported each summer and fall.
- 1952: More than 21,000 paralytic cases reported in the U.S.
- 1955: Vaccine introduction; polio incidence declined rapidly
- 1979: Last case of polio caused by wild poliovirus acquired in the U.S.
- 2022: Most recent case of polio caused by vaccine-derived poliovirus in the U.S. occurred in New York in an unvaccinated, immunocompetent young adult.

Poliovirus

- Enterovirus (RNA)
- Three serotypes: type 1, type 2, type 3
 - Immunity to one serotype does not produce significant immunity to other serotypes



Poliomyelitis Pathogenesis

- Enters through the mouth
- Replicates in oropharynx and gastrointestinal tract
 - Invades local lymphoid tissue, may enter bloodstream then infect cells of the central nervous system
- Viral spread along nerve fibers
- Destruction of motor neurons and brainstem cells

Poliovirus Epidemiology

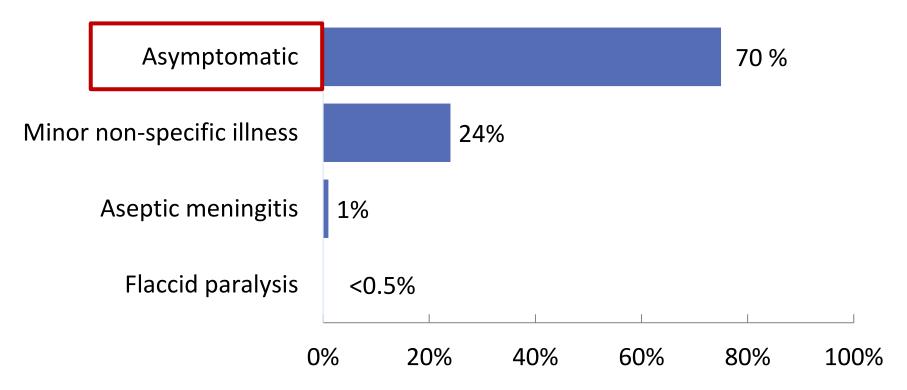
Reservoir	Human
Transmission	Fecal-oral Oral-oral possible
Communicability	Most infectious: 7–10 days before onset Virus present in stool 3–6 weeks

Poliomyelitis Clinical Features

- Incubation period
 - 3 to 6 days for nonparalytic poliomyelitis
 - 7 to 21 days for onset of paralysis in paralytic poliomyelitis

Risk of severe disease and death increases with age

Poliomyelitis Clinical Features in Children



Types of Paralytic Polio

Spinal polio

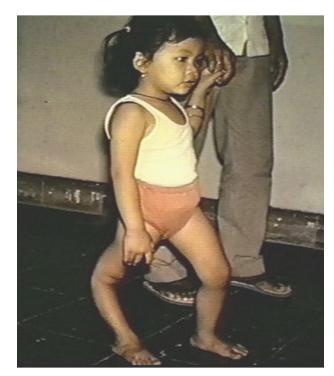
Asymmetric paralysis that most often involves the legs

Bulbar polio

 Presents with weakness of facial, oropharyngeal, and respiratory muscles innervated by cranial nerves

Bulbospinal polio

 Combination of bulbar and spinal paralysis, accounted for 19% of cases



Asymmetric Paralysis

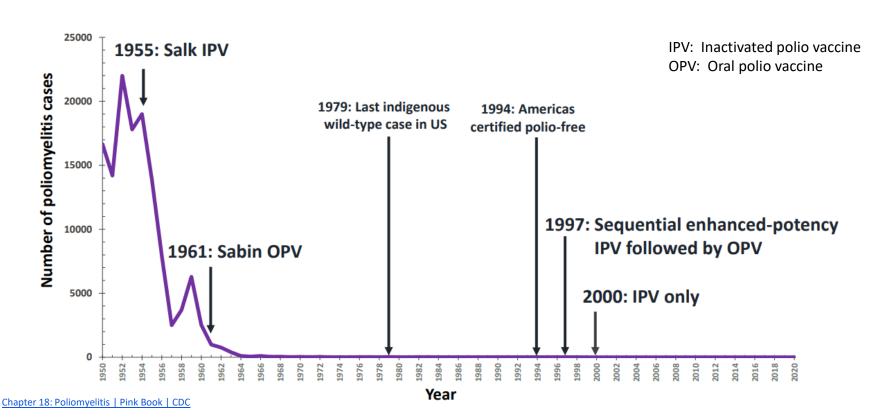
Poliomyelitis Clinical Features

- Historic case fatality rate for paralytic polio
 - 2% to 5% in children
 - 15% to 30% in adolescents and adults
 - 25% to 75% with bulbar involvement

Post-polio syndrome

- Slow, irreversible worsening of muscle weakness in the muscle groups involved during the original infection
- Occurs in 25% to 40% of persons who had paralytic polio in childhood
- Interval of 15 to 40 years after paralysis
- Not infectious, affected persons do not shed virus

Paralytic Polio Decreased Rapidly in the United States After Introduction of Polio Vaccine



Poliomyelitis in the United States, 2022–2024

- CDC was notified of a case of paralytic polio in an unvaccinated individual from Rockland County, New York.
 - The case was caused by vaccine-derived poliovirus type 2.
- Genetic sequencing indicated a linkage to polioviruses collected in wastewater in Israel, United Kingdom, and Canada.
- No additional paralytic cases were identified.

Poliovirus Vaccine

2 Types of Polio Vaccines

Inactivated polio vaccine (IPV)

Protects against all 3 serotypes

Oral polio vaccine (OPV)

- Trivalent OPV (tOPV): Protects against all 3 serotypes
- Bivalent OPV (bOPV): Protects against serotypes 1 and 3
- Monovalent OPV (mOPV): Protects against only 1 serotype

Not available in U.S.

Polio Vaccine Products

Vaccine (Manufacturer)	Vaccine Components	Age Indication	Doses in Polio Vaccine Series	Injection Route
Ipol (SP)	IPV	6 weeks and older	Any	IM or SC
Pentacel (SP)	DTaP-IPV/Hib	6 weeks–4 years	1, 2, 3, 4	IM
Vaxelis (Merck)	DTap-IPV-Hib-HepB	6 weeks–4 years	1, 2, 3	IM
Pediarix (GSK)	DTaP-HepB-IPV	6 weeks–6 years	1, 2, 3	IM
Kinrix (GSK)	DTaP-IPV	4–6 years	4	IM
Quadracel (SP)	DTaP-IPV	4–6 years	4, 5	IM

IM = Intramuscular; SC = Subcutaneous; All vaccines in the table above are non-live.

Chapter 18: Poliomyelitis | Pink Book | CDC | POL, Kinrix, Pediarix, Pentacel, VAXELIS, Quadracel.

Inactivated Polio Vaccine Efficacy

- Seroconversion rates and antibody titers after vaccination vary depending on:
 - Age at receipt of the first dose
 - Vaccination schedule
- IPV highly effective in producing immunity to poliovirus
 - 95% or more of recipients develop protective antibodies after 3 doses
- Duration of immunity not known with certainty

Preparation and Administration for IPV-Containing Vaccines

- Preparation: Prepare the vaccine just prior to administration.
 - Pentacel requires reconstitution.
 - Reconstitute the lyophilized Haemophilus influenzae type b (Hib) vaccine with the DTaP-IPV liquid diluent supplied by the manufacturer.
 - Do not use Kinrix or Quadracel to reconstitute the Hib component of Pentacel.

Route:

- Intramuscular injection for all products
- Ipol may be administered by subcutaneous injection.

3

Clinical Considerations

Recommended Polio Vaccination Schedule for Children and Adolescents



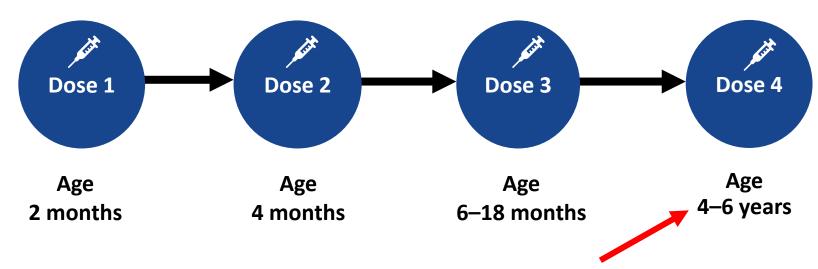
Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, United States, 2025

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	15 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
Inactivated poliovirus (IPV)			1st dose	2nd dose	←		3rd dose					4th dose					See Notes



Polio Vaccination in Children: Routine Schedule



Final dose on or after age 4 years <u>and</u> at least 6 months after the previous dose.

Recommended Polio Vaccination Schedule for Children and Adolescents: Catch-up Vaccination

Table 2

Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2025

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						
Inactivated poliovirus	6 weeks	4 weeks	4 weeks if current age is <4 years 6 months (as final dose) if current age is 4 years or older	6 months (minimum age 4 years for final dose)							
			Children and adolescents age 7 through 18 years								
Inactivated poliovirus	N/A	4 weeks	6 months A fourth dose is not necessary if the third dose was administered at age 4 years or older <i>and</i> at least 6 months after the previous dose.	A fourth dose of IPV is indicated if all previous doses were administered at <4 years OR if the third dose was administered <6 months after the second dose.							

A fourth dose is not necessary if the third dose was administered at age 4 years or older <u>and</u> at least 6 months after the previous dose.

Accelerated Routine Polio Vaccination Schedule for Children and Adolescents Ages 17 Years and Younger

• If accelerated protection is needed (e.g., travel to polio-endemic area), minimum age and intervals may be used as follows:

Dose	Minimum Age	Minimum Interval to the Next Dose
Dose 1	6 weeks	4 weeks
Dose 2	10 weeks	4 weeks
Dose 3	14 weeks	6 months
Dose 4	4 years	

Recommended Polio Vaccination Schedule: People Ages 18 Years and Older



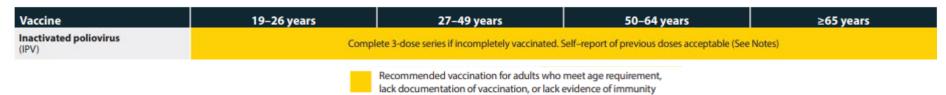
Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, United States, 2025

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	15 mos	18 mos	19–23 mos	2-3 yrs	4-6 yrs	7–10 yrs	11–12 yrs	13–15 yrs	16 yrs	
Inactivated poliovirus (IPV)			1st dose	2nd dose	∢		3rd dose					4th dose					See Notes
					_	ge of recor Il children	nmended a	ages		of recomme	_	es					

Table 1

Recommended Adult Immunization Schedule by Age Group, United States, 2025



Routine Polio Vaccination Schedule for People Ages 18 Years and Older

- Known or suspected to be incompletely vaccinated:
 - Complete 3-dose series
 - Dose 1 at any time
 - Dose 2; 1 to 2 months later
 - Dose 3; 6 to 12 months after dose 2
- Most adults born and raised in the United States can assume they were vaccinated as children.
 - Unless there are specific reasons to believe otherwise



Accelerated Polio Vaccination Schedule for People Ages 18 Years and Older

- If accelerated protection is needed (e.g., travel to polio-endemic area), use the minimum intervals.
- If cannot complete accelerated schedule: give remaining doses as soon as possible, with dose 3 given at least 6 months after dose 2.

Protection needed in	Accelerated Schedule
8 weeks or more	3 doses at least 4 weeks apart
More than 4 weeks, but less than 8 weeks	2 doses at least 4 weeks apart
Less than 4 weeks	1 dose

Polio Vaccine Recommendations | Polio | CDC, Use of Inactivated Polio Vaccine Among U.S. Adults: Updated Recommendations of the Advisory Committee on Immunization Practices — United States, 2023 | MMWR

Adults at Increased Risk of Poliovirus Exposure



Laboratory workers handling specimens that may contain polioviruses



Health care personnel treating patients who could have polio or have close contact with a person who could be infected with poliovirus



Travelers to areas where poliomyelitis is endemic or epidemic



Adults identified by public health authorities as part of a group or population at increased risk for exposure to poliovirus because of an outbreak

Polio Vaccination in Adults with Increased Risk of Exposure

- Has not completed polio vaccination series (at least 3 doses):
 - Administer remaining doses to complete a 3-dose series
- Has completed polio vaccination series (at least 3 doses):
 - With increased risk, may administer one lifetime IPV booster



OPV Administered Outside the United States

- Persons with doses of OPV that do not count towards the U.S. vaccination requirements should receive IPV.
- Use the date of administration to make a presumptive determination of what type of OPV was received.
- Trivalent OPV was used throughout the world prior to April 1, 2016.
 - OPV prior to April 1, 2016: count as valid (except campaign doses)
 - OPV on or after April 1, 2016: invalid dose

Polio Vaccination Schedules that Include Both IPV and OPV

- Mixed-product series containing both OPV and IPV is acceptable
 - Only trivalent OPV (tOPV) counts toward completing the series
- Children with an incomplete series:
 - Administer IPV to complete a series that includes doses of tOPV
 - Ensure doses meet minimum ages and intervals
- Administer 1 dose of IPV to children who received 4 doses of tOPV (or more) before 4 years of age
 - Should be at least 6 months between the last dose of OPV and the IPV dose

Fractional IPV Clinical Considerations



- For persons who received fractional (1/5 full dose) IPV administered intradermally outside of the United States
 - 2 fractional doses of IPV (fIPV) *should* be considered valid and counted as 1 full intramuscular dose of IPV towards the US vaccination schedule.



 If a person received only 1 dose of fIPV, this dose should <u>not</u> be considered valid or counted towards the US vaccination schedule.

Unknown or Uncertain Polio Vaccination Status

- Children and adolescents ages 17 years or younger
 - Only accept written, dated records of tOPV or IPV as evidence of vaccination.
 - No or questionable documentation: administer IPV according to U.S. schedule.
- People ages 18 years and older
 - In general, self-reports are acceptable unless the clinician has specific reasons to believe the patient was not vaccinated such as:
 - Did not receive consistent medical care as an infant
 - Parents were against vaccination
 - Person has other reason to doubt their vaccination status.
 - Specific immigration programs might have additional polio documentation requirements.

Safety

Contraindications to IPV

- Severe allergic reaction (e.g., anaphylaxis) after a previous dose or to a vaccine component
- The following ingredients are used in the production of this vaccine:
 - Neomycin
 - Formaldehyde
 - Polymyxin B
 - Streptomycin
 - 2-phenoxyethanol



Precautions to IPV

- Pregnancy: can vaccinate if at increased risk of polio infection
- Moderate or severe acute illness with or without fever



IPV Adverse Reactions

- Transient local reactions at the site of injection
- Erythema (redness), induration or swelling, and pain
- Elevated temperature, irritability, sleepiness, fussiness, crying
- Serious adverse reactions rarely occur

Polio Vaccine Administration Errors (1)

Schedule errors:

- Doses administered 5 or more days before the minimum age and/or interval do not count and should be repeated when age-appropriate.
- Wait the minimum interval from the invalid dose before giving the repeat dose.

Polio Vaccine Administration Errors (2)

- Age and dose errors: Kinrix or Quadracel for doses 1–3
 - If the minimum age and minimum interval from the last dose of polio vaccine have been met, the dose can count and does not need to be repeated.

5

Storage and Handling

Polio Vaccine Storage and Handling

For all IPV-containing vaccines:

- Store in the refrigerator between 2°C and 8°C (36°F and 46°F)
- Should not be frozen
- Refrigerate on arrival
- Protect from light (Vaxelis and Ipol)

For Pentacel:

- Use immediately after reconstitution.



Knowledge Check

When DTaP-IPV/Hib (Pentacel) is used, 4 doses of IPV are given at ages 2, 4, 6, and 15–18 months. This results in 4 doses of IPV by the age of 18 months. The series is complete at 18 months.

- A. True
- B. False



When DTaP-IPV/Hib (Pentacel) is used, 4 doses of IPV are given at ages 2, 4, 6, and 15–18 months. This results in 4 doses of IPV by the age of 18 months. The series is complete at 18 months.

A. True

B. False



Knowledge Check

The vaccination record of a 14-year-old from Uzbekistan shows receipt of four OPV vaccines between 6/2/2011 and 2/20/2016. All were given using U.S. minimum age and intervals. This series is considered valid and complete for the U.S. vaccination schedule.

- A. True
- B. False



The vaccination record of a 14-year-old from Uzbekistan shows receipt of four OPV vaccines between 6/2/11 and 2/20/2016. All were given using U.S. minimum age and intervals. This series is considered valid and complete for the U.S. vaccination schedule.

A. True

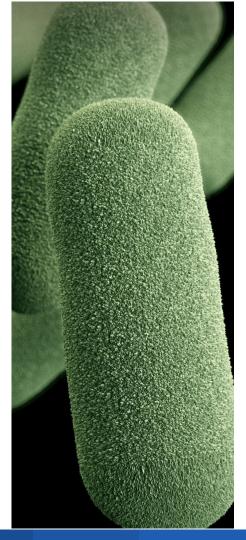
B. False

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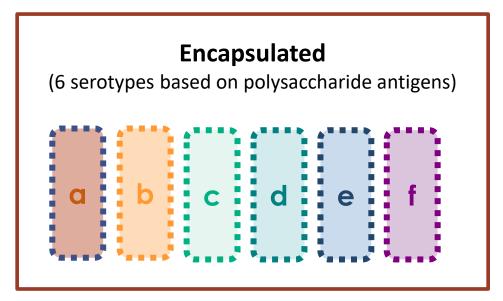
Haemophilus influenzae Disease

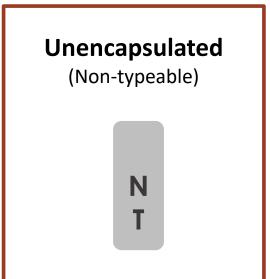
Haemophilus influenzae

- Gram-negative bacilli
- Originally thought to be the cause of influenza
- Abbreviated "H. flu" or Hi
- Infections range from mild to severe invasive disease



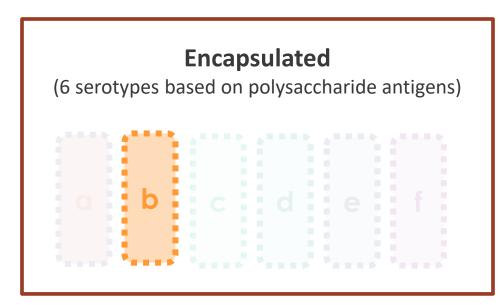
Classification of *H. influenzae*





H. influenzae serotype b (Hib)

- The most virulent serotype
- The only Hi serotype preventable through vaccination
- Before the introduction of effective vaccines, the leading cause of bacterial meningitis and other invasive bacterial disease among children aged less than 5 years in the United States



H. influenzae serotype b (Hib) pathogenesis

- Enters body through nasopharynx
- Can colonize the nasopharynx
- Invasive infection: Bacteria spread in bloodstream to distant sites in the body

Hib Epidemiology

Reservoir	Human asymptomatic carriers
Transmission	Airborne respiratory droplets or direct contact with respiratory secretions
Temporal pattern	Pre-vaccine era: Peaks in Sept. through Dec. and March through May
Communicability	Generally limited but higher in some circumstances (e.g., household, childcare)

Risk Factors for Invasive Hib Disease in the Pre-vaccine Era

Demographic factors

- Male sex
- Race/Ethnicity
 - American Indian
 - Alaska Native
 - Black

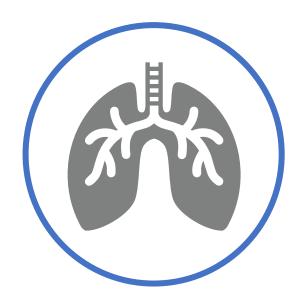
Social factors

- Household crowding
- Large household size
- Low socioeconomic status
- School-aged siblings
- Daycare attendance

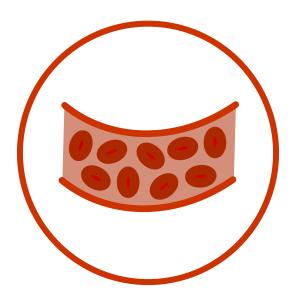
Immunocompromising conditions

- HIV infection
- Asplenia or Sickle cell disease
- IgG deficiency
- Early component complement deficiency
- Hematopoietic stem cell transplantation
- Chemotherapy

Common Clinical Syndromes of Invasive Hib Disease



Bacteremic pneumonia



Bacteremia without a focus



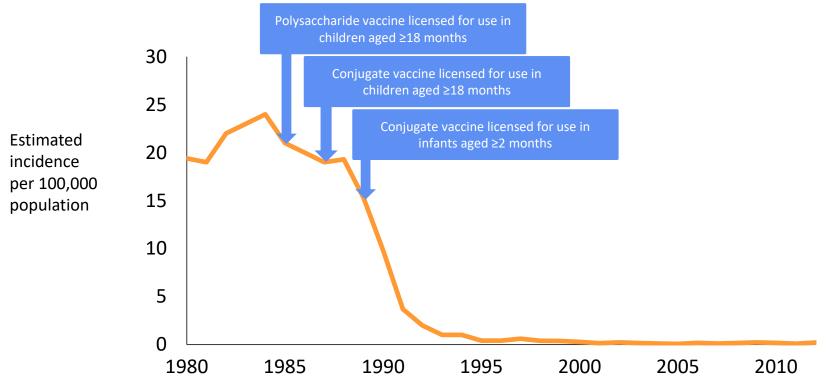
Meningitis

Other Clinical Syndromes of Hib Disease

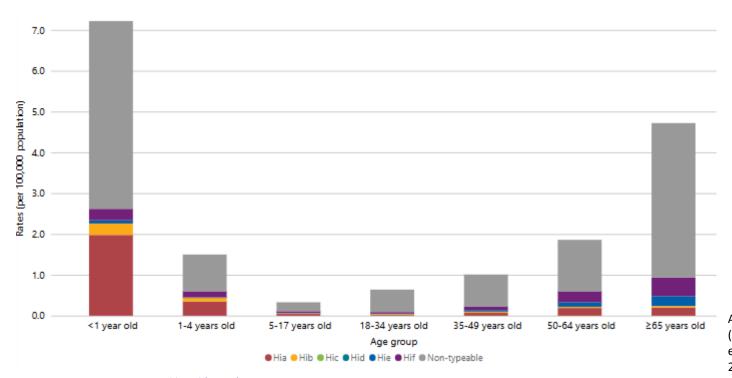
- Invasive disease:
 - Epiglottitis
 - Septic arthritis
 - Cellulitis
 - Purulent pericarditis
 - Endocarditis
 - Osteomyelitis

- Non-invasive disease:
 - Otitis media
 - Sinusitis
 - Bronchitis

Invasive Hib Disease in Children Decreased Dramatically After Hib Vaccine Introduction



Estimated U.S. Incidence of Invasive H. influenzae Disease by Age Group and Serotype



Active Bacterial Core Surveillance (ABCs) cases from 2018-2023 and estimated to the U.S. population. 2023 data are preliminary.

Hib Vaccine

Hib Polysaccharide Conjugate Vaccines

- Capsular polysaccharide (PRP) conjugated to carrier proteins
 - Tetanus toxoid (PRP-T)
 - Outer membrane protein of meningococcal serogroup B (PRP-OMP)
- Highly immunogenic via activation of T-cell dependent immunity
 - 95% of infants develop protective antibody levels after a primary series.
 - No cross protection against non-b/non-typeable serotypes
- Estimated clinical efficacy 95%–100%
- Invasive Hib disease is uncommon in children who are fully vaccinated.

Hib-containing Vaccine Products

Vaccine Product	Age Indications	Dose in Series						
PRP-T (polysaccharide, tetanus toxoid)								
ActHIB	2 months-5 years	1, 2, 3, booster						
Pentacel (SP)	6 weeks–4 years	1, 2, 3, booster						
Hiberix (GSK)	6 weeks–4 years	1, 2, 3, booster						
PRP-OMP (polysaccharide, meningococcal outer membrane protein)								
PedvaxHIB	2–71 months	1, 2, booster						
Vaxelis (Merck)	6 weeks–4 years	1, 2, 3*						

Chapter 8: Haemophilus influenzae | Pink Book | CDC

^{*} Vaxelis should not be used as a booster dose, but a booster dose is still recommended.

Hib-containing Vaccine Products: Single Antigen Vaccines

Vaccine Product	Age Indications	Dose in Series							
PRP-T (polysaccharide, tetanus toxoid)									
ActHIB	2, 4, and 6 months	1, 2, 3, booster							
Pentacel (SP)	6 weeks–4 years	1, 2, 3, booster							
Hiberix (GSK)	2, 4, and 6 months	1, 2, 3, booster							
PRP-OMP (polysaccharide, meningococcal outer membrane protein)									
PedvaxHIB	2 and 4 months	1, 2, booster							
Vaxelis (Merck)	6 weeks–4 years	1, 2, 3							

Hib-containing Vaccine Products: Combination Vaccines

Vaccine Product	Age Indications	Dose in Series							
PRP-T (polysaccharide, tetanus toxoid)									
ActHIB	2, 4, and 6 months	1, 2, 3, booster							
Pentacel (SP)	6 weeks–4 years	1, 2, 3, booster							
Hiberix (GSK)	2, 4, and 6 months	1, 2, 3, booster							
PRP-OMP (polysaccharide, meningococcal outer membrane protein)									
PedvaxHIB	2 and 4 months	1, 2, booster							
Vaxelis (Merck)	6 weeks-4 years	1, 2, 3							

Preparation and Administration of Hib-containing Vaccines

- Preparation: just prior to administration
 - ActHIB, Hiberix, and Pentacel require reconstitution.
 - Reconstitute the lyophilized vaccine with manufacturer-supplied diluent.
- Route: intramuscular injection

Vaccines with Diluents: How to Use Them

Be sure to reconstitute (mix) the following vaccines correctly before administering them! Reconstitution means that the lyonhilized (freeze-dried) vaccine powder in one vial must be mixed with the diluent (liquid) in another.

- . Only use the diluent provided by the manufacturer for that vaccine as indicated on the chart.
- . ALWAYS check the expiration date on the diluent and vaccine. NEVER use expired diluent or vaccine.
- . Never freeze diluents.

Vaccine product name	Manufacturer	Lyophilized (powder) vaccine	Liquid diluent (may contain vaccine)	Time allowed between mixing and use ^a	Diluent storage environment
Abrysvo	Pfizer	RSV	Sterile water	4 hrs	Refrigerator or room temp
ActHIB (Hib)	Sanofi	Hib	Sodium chloride 0.4%	24 hrs	Refrigerator
Arexvy	GSK	RSV	ASO1E adjuvant suspension	4 hrs	Refrigerator
COVID-19, Pfizer-BioNTech, 6 mos through 4 yrs formulation	Pfizer-BioNTech	see footnote ^b	Sodium chloride 0.9%	12 hrs	Refrigerator or room temp
Dengvaxia (DEN4CYD)	Sanofi	Dengue	Sodium chloride 0.4%	30 min	Refrigerator
Hiberix (Hib)	GSK	Hib	Sodium chloride 0.9%	Immediately ^c	Refrigerator or room temp
Imovax (RAB _{HDCV})	Sanofi	Rabies	Sterile water	Immediately ^c	Refrigerator
bxchiq	Valneva	Chikungunya	Sterile water	Immediately ^c	Refrigerator
M-M-R II (MMR)	Merck	MMR	Sterile water	8 hrs	Refrigerator or room temp
Menveo ^d (MenACWY)	GSK	MenA	MenCWY component	8 hrs	Refrigerator
Penbraya (MenABCWY)	Pfizer	MenACWY	MenB component	4 hrs	Refrigerator
Pentacel (DTaP-IPV/Hib)	Sanofi	Hib	DTaP-IPV component	Immediately ^c	Refrigerator
Priorix (MMR)	GSK	MMR	Sterile water	8 hrs	Refrigerator or room temp
ProQuad (MMRV)	Merck	MMRV	Sterile water	30 min	Refrigerator or room temp
RabAvert (RAB _{PCECV})	GSK	Rabies	Sterile water	Immediately ^c	Refrigerator
Rotarix ^d (RV1)	GSK	RV1	Sterile water, calcium carbonate, and xanthan	24 hrs	Refrigerator or room temp
Shingrix (RZV)	GSK	RZV	ASO1B adjuvant suspension	6 hrs	Refrigerator
Varivax (VAR)	Merck	VAR	Sterile water	30 min	Refrigerator or room temp
Vaxchora (CVD 103-HgR)	Bavarian Nordic	Cholera	Buffer solution plus bottled water	see footnote ^e	Refrigerator
YF-VAX (YF)	Sanofi	YF	Sodium chloride 0.9%	60 min	Refrigerator or room temp

Always refer to package inserts for detailed instructions on reconstituting specific vaccines. In general, follow the steps below. 1 For single-dose vaccine products (exceptions: Rotarix. 3 Reconstitute (i.e. mix) vaccine before use by: Vaxchora), select a syringe and needle of proper length removing the protective caps and wiping each

to be used for both reconstitution and administration of the vaccine. For Rotarix and Vaxchora, see the package insert.

- lyophilized vaccine vial and the diluent to verify that . they are the correct two products to mix together. - neither the vaccine nor the diluent has evalued
- stopper with an alcohol swab. . inserting needle of syringe into diluent vial and withdrawing contents, and injecting diluent into lyophilized vaccine vial and rotating or inverting to thoroughly dissolve the
- 4 Check the appearance of the reconstituted vaccing . Reconstituted vaccine may be used if the color and appearance match the description on the package
- . If there is discoloration, particulate matter, obvious
- USE," return it to proper storage conditions, and immunization program or the vaccine manufacture 5. If reconstituted vaccine is not used immediately
 - . clearly mark the vial with the date and time the · maintain the product at 2°-8°C (36°-46°F); do no
 - · use only within the time indicated on chart above
- a. If the reconstituted vaccine is not used within this time period, it must be discarded.
- b. The Pfizer-BioNTech COVID-19 formulation for children age 6 mos through 4 yrs is a liquid concentrate that requires dilution c. For purposes of this guidance, Immunize.org defines "Immediately" as within 30 minutes or less.
- d States and Manuacy varyings are available as either a liquid formulation that does not require dilution or as a lyophilized vaccine that requires reconstitution. Both formulations of the Rotarix vaccine are administered by mouth, they should not be administered as an injection.
- e. Vaxchora dilution: 30 minutes if sucrose or unflavored stevia added; 4 hours if sucrose or unflavored stevia have

Vaccine abbreviations in column 3: Hib »Homophiks influenter type b MenA = Meningopoccal serogroup A MenACWY » Meningopoccal serogroups A, C, W, Y MMR » Measles, rumps, & rubella MMRV » MMR » unicula

RZV - Zoster vaccine, recombinary



FOR PROFESSIONALS www.immunize.org / FOR THE PUBLIC www.vaccineinformation.org



8

Clinical Considerations

Recommended Hib Vaccination Schedule for Children and Adolescents

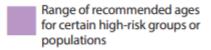


Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, United States, 2025

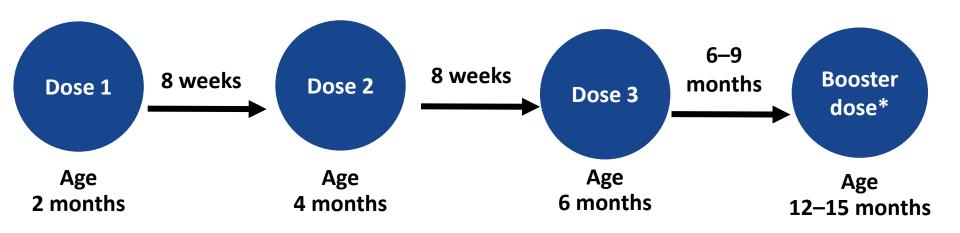
These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2).

Vaccine and other immunizing agents	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 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
Haemophilus influenzae type b (Hib)			1st dose	2nd dose	See Notes		3rd or 4tl (See N	h dose lotes)▶									



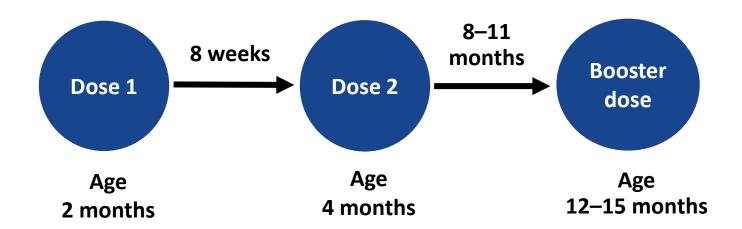


Hib Vaccine Recommendations: Routine Schedule for ActHib, Pentacel, Hiberix, and Vaxelis*

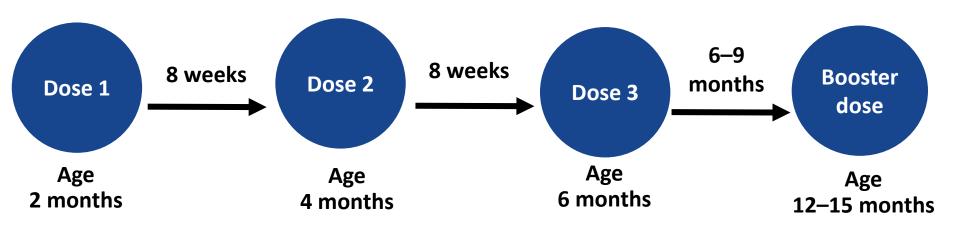


^{*}Vaxelis is not recommended for use as a booster dose. A different Hib-containing vaccine should be used for the booster dose.

Hib Vaccine Recommendations: Routine Schedule for PedvaxHIB



Hib Vaccine Recommendations: Routine Schedule for Mixed or Unknown Products*



^{*}If any dose in the series is ActHIB, Pentacel, Hiberix, Vaxelis or the product is not known, follow the 4-dose schedule.

Hib Vaccine Recommendations: Catch-up Schedule

Recommended Catch-up Immunization Schedule for Children and Adolescents Who Start Late or Who Are More than 1 Month Behind, United States, 2025

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						
Haemophilus influenzae type b	6 weeks	No further doses needed if first dose was administered at age 15 months or older. 4 weeks if first dose was administered before the 1st birthday. 8 weeks (as final dose) if first dose was administered at age 12 through 14 months.	No further doses needed if previous dose was administered at age 15 months or older 4 weeks if current age is younger than 12 months and first dose was administered at younger than age 7 months and at least 1 previous dose was PRP-T (ActHib, Pentacel, Hiberix), Vaxelis or unknown 8 weeks and age 12 through 59 months (as final dose) if current age is younger than 12 months and first dose was administered at age 7 through 11 months; OR if current age is 12 through 59 months and first dose was administered before the 1st birthday and second dose was administered at younger than 15 months; OR if both doses were PedvaxHIB and were administered before the 1st birthday	8 weeks (as final dose) This dose only necessary for children age 12 through 59 months who received 3 doses before the 1st birthday.							

Hib Immunization Recommendations: Catch-up Schedule Resources

- Children starting late might not need entire 3- or 4-dose series.
 - Number of doses child requires depends on current age

Resources

Catch-up guidance for healthy children

Catch-Up Guidance for Healthy¹ Children 4 Months through 4 Years of Age

Haemophilus influenzae type b Vaccines: ActHIB, Hiberix, Pentacel, Vaxelis, or Unknown²

The table below provides guidance for children whose vaccinations have been delayed. Start with the child's age and information on previous doses previous doses must be documented and must meet minimum age requirements and minimum intervals between doses. Use this table in conjunction with table 2 of the Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, found at www.cdc.gov/vaccines/schedules/hcp/child-doses

	AND # of previous doses is							
	Unknown or 0	-	→	Give Dose 1 today	Give Dose 2 at least 4 weeks after Dose 1			
	1		en at least ince Dose 1	Give Dose 2 today	Give Dose 3 at least 4 weeks after Dose 2			
4 through 6 months	'		peen at least ince Dose 1	No dose today	Give Dose 2 at least 4 weeks after Dose 1			
			en at least ince Dose 2	Give Dose 3 today	Give Dose 4 (Final Dose) at 12 months of age or older ⁴			
	2		peen at least ince Dose 2	No dose today	Give Dose 3 at least 4 weeks after Dose 2			
7 through 11 months	Unknown or 0	→	→	Give Dose 1 today	Give Dose 2 at least 4 weeks after Dose 1			
	1	It has been at least 4 weeks since Dose 1	→		IF Dose 1 was given before 7 months of age, give Dose 3 at least 4 weeks after Dose 2			
					IF Dose 1 was given at 7 months of ago or older, give Dose 3 (Final Dose) at least 8 weeks after Dose 2 and no earling than 12 months of age			
		It has not been at least 4 weeks since Dose 1	→	No dose today	Give Dose 2 at least 4 weeks after Dose 1			
	2				Dose 1 was given	It has been at least 4 weeks since Dose 2	Give Dose 3 today	Give Dose 4 (Final Dose) at least 8 weeks after Dose 3 and no earlier that 12 months of age ⁴
		7 months of age	It has not been at least 4 weeks since Dose 2	No dose today	Give Dose 3 at least 4 weeks after Dose 2			
	Dose 1 was given at 7 months of age or older		→	No dose today	Give Dose 3 (Final Dose) at least 8 weeks after Dose 2 and no earlier that 12 months of age			

Refer to notes of the Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger-United States, 2024 for immunization guidance for children at increased risk for Haemophilix influenzae type b disease

Reference: Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger-United States, 2024. www.cdc.gov/vaccines/sched



U.S. Department of Health and Human Services Centers for Disease Control and Prevention

Catch-Up Guidance for Healthy1 Children 4 Months 4 years of age-Haemophilus influenzae type b Vaccines: ActHIB, Hiberix, Pentacel, Vaxelis, or Unknown2- Revised December 2023

Catch-up Guidance for Children 4 months through 4 years of age-Haemophilus Influenzae type b-PedvaxHIB - December 2023

Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger; 2025 U.S.

² See separate job aid for HIB vaccination with PedvaxHIB.
³ Next dose due is not the final dose in the series unless explicitly stated.
⁴ Vaxelis should not be used for Dose 4.

Hib Vaccine Recommendations: American Indian/Alaska Native Children

Vaccine Product	Age Indications	Dose in Series						
PRP-T (polysaccharide, tetanus toxoid)								
ActHIB	2 months–5 years	1, 2, 3, booster						
Pentacel (SP)	6 weeks–4 years	1, 2, 3, booster						
Hiberix (GSK)	6 weeks–4 years	1, 2, 3, booster						
PRP-OMP (polysaccharide, meningococcal outer membrane protein)								
PedvaxHIB	2–71 months	1, 2, booster						
Vaxelis (Merck)	6 weeks–4 years	1, 2, 3						

<u>Use of Haemophilus influenzae Type b–Containing Vaccines Among American Indian and Alaska Native Infants: Updated Recommendations of the Advisory Committee on Immunization Practices — United States, 2024 | MMWR</u>

Hib Vaccine Recommendations: Special Populations (1)

- Recommendations for children and adolescents with conditions that increase the risk of invasive Hib are based on age, vaccination history, and condition.
 - Functional or anatomic asplenia (including sickle cell disease)
 - Immunoglobulin deficiency or early complement component deficiency
 - HIV infection
 - Receipt of chemotherapy or radiation therapy
 - Hematopoietic stem cell transplant

Hib Vaccine Recommendations: Special Populations (2)

· Chemotherapy or radiation treatment:

Age 12-59 months

- · Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart
- o 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Doses administered within 14 days of starting therapy or during therapy should be repeated at least 3 months after therapy completion.

- Hematopoietic stem cell transplant (HSCT):
 - 3-dose series 4 weeks apart starting 6 to 12 months after successful transplant regardless of Hib vaccination history
- · Anatomic or functional asplenia (including sickle cell disease):

Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart
- o 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

Unvaccinated* persons age 5 years or older

- 1 dose
- Elective splenectomy:

Unvaccinated* persons age 15 months or older

- 1 dose (preferably at least 14 days before procedure)
- HIV infection:

Age 12-59 months

- · Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart
- 2 or more doses before age 12 months; 1 dose at least 8 weeks after previous dose

Unvaccinated* persons age 5-18 years

- 1 dose
- Immunoglobulin deficiency, early component complement deficiency:

Age 12-59 months

- Unvaccinated or only 1 dose before age 12 months: 2 doses, 8 weeks apart
- o 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

^{*}Unvaccinated = Less than routine series (through age 14 months) OR no doses (age 15 months or older)

Hib Vaccine Recommendations: Special Populations (3)

- Children 12–59 months of age at increased risk of invasive Hib disease*
 - 0 or 1 dose before age 12 months: 2 doses, 8 weeks apart
 - 2 or more doses before age 12 months: 1 dose at least 8 weeks after previous dose

^{*}Chemotherapy, radiation therapy, anatomic or functional asplenia, HIV infection, immunoglobulin deficiency, or early component complement deficiency

Hib Vaccine Recommendations: Special Populations (4)

- Unvaccinated persons ages 15 months or older undergoing elective splenectomy:
 - 1 dose, preferably at least 14 days before procedure
- Unvaccinated persons ages 5 years or older with anatomic or functional asplenia:
 - 1 dose
- Unvaccinated persons ages 5–18 years with HIV infection:
 - 1 dose
- Hematopoietic stem cell transplant recipients (any age):
 - 3 doses, 4 or more weeks apart, beginning 6–12 months post-transplant
- Note: "Unvaccinated" includes partially vaccinated.

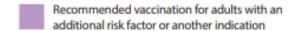
Hib Vaccine Recommendations: Special Populations (5)

- Children younger than 24 months of age with invasive Hib disease:
 - Administer complete series as recommended for child's age.
 - Vaccinate during the convalescent phase of the illness.

Recommended Hib Vaccination Schedule for Adults

Table 1 Recommended Adult Immunization Schedule by Age Group, United States, 2025

Vaccine	19-26 years	27-49 years	50-64 years	≥65 years
Haemophilus influenzae type b (Hib)	1 or 3 doses depending on indication			



Recommended Hib Vaccination Schedule for Adults Based on Indication

- Anatomical or functional asplenia (including sickle cell disease):
 - 1 dose if previously did not receive Hib vaccine
 - For elective splenectomy: 1 dose preferably at least 14 days before splenectomy
- Hematopoietic stem cell transplant:
 - 3-dose series 4 weeks apart starting 6–12 months after successful transplant; regardless of Hib vaccination history

Use of Hib-containing Vaccine Products is Offlabel in Older Children and Adults

Vaccine Product	Age Indications	
ActHIB	2 months–5 years	
Pentacel (SP)	6 weeks–4 years	
Hiberix (GSK)	6 weeks–4 years	
PedvaxHIB	2–71 months	
Vaxelis (Merck)	6 weeks–4 years	

Hib Vaccine Interchangeability

- Single-component conjugate Hib vaccines are interchangeable.
 - 3-dose primary series (4 doses total) if more than one brand of vaccine used at 2 or 4 months of age
- Whenever feasible, use same combination vaccine for subsequent doses
 - If the vaccine used for earlier doses is unknown or not available, use any brand to complete the primary series.



Knowledge Check

Vaxelis can be used for all 4 recommended doses in the Hib series.

- A. True
- B. False



Vaxelis can be used for all 4 recommended doses in the Hib series.

A. True

B. False



Knowledge Check

PedvaxHIB can be administered to a 26-year-old who has sickle cell disease and no history of previous Hib vaccination.

- A. True
- B. False



PedvaxHIB can be administered to a 26-year-old who has sickle cell disease and no history of previous Hib vaccination.

A. True

B. False

Safety

Hib Vaccine Contraindications

- Severe allergic reaction (e.g. anaphylaxis)
 - After a previous dose

or

- To a vaccine component
- Age younger than 6 weeks
 - Potential for development of immunologic tolerance



Hib Vaccine Precautions

 Moderate or severe acute illness with or without fever



Hib Vaccine Adverse Reactions



Injection site pain, redness, swelling



Fever



Crying



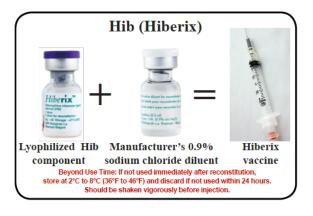
Irritability

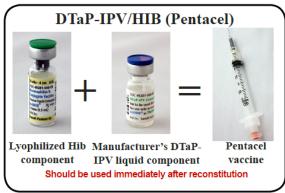


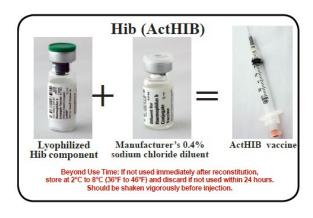
Rash

Hib Vaccine Administration Errors (1)

- Using wrong diluent to reconstitute the lyophilized component
 - Doses using the wrong diluent are not valid.
 - Dose should be repeated after reconstitution with correct diluent.
 - The repeat dose may be given as soon as possible; there is no waiting period.







Hib Vaccine Administration Errors (2)

Schedule errors:

- Doses administered 5 or more days before the minimum age and/or interval do not count and should be repeated when age-appropriate.
- Wait the minimum interval from the invalid dose before giving the repeat dose.

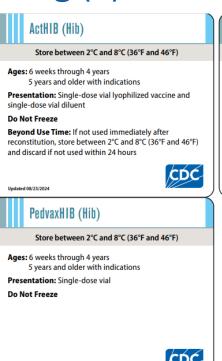
Hib Vaccine Administration Errors

- For all Hib vaccine administration errors:
 - Inform the recipient of the vaccine administration error.
 - Determine how the error occurred and implement strategies to prevent it from happening again.
 - Providers are encouraged to report the error to Vaccine Adverse Event Reporting System (VAERS), even for errors not associated with an adverse event.

Storage and Handling

Hib Vaccine Storage and Handling (1)

- All Hib-containing vaccines: store refrigerated between 2°C and 8°C (36°F and 46°F).
- Store in the original packaging.
 - Hib vaccines requiring reconstitution (Pentacel, ActHIB, Hiberix) should be stored together with their diluent in the refrigerator.



Hiberix (Hib)

Store between 2°C and 8°C (36°F and 46°F)

Ages: 6 weeks through 4 years 5 years and older with indications

Presentation: Single-dose vial lyophilized vaccine and single-dose vial or manufacturer-filled syringe diluent

Protect From Light

Do Not Freeze

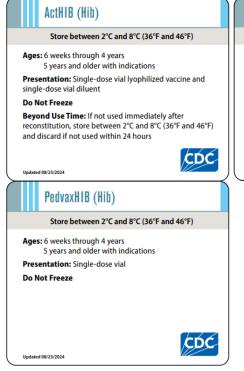
Updated 08/23/2024



Updated 08/23/20

Hib Vaccine Storage and Handling (2)

- Do not freeze vaccine or diluents; do not expose to freezing temperatures.
- If vaccines are exposed to inappropriate temperatures or handled improperly:
 - Store at the appropriate temperature.
 - Isolate from other vaccines.
 - Mark "Do NOT Use."
 - Consult the vaccine manufacturer and/or state or local immunization program for guidance.



Hiberix (Hib)

Store between 2°C and 8°C (36°F and 46°F)

Ages: 6 weeks through 4 years
5 years and older with indications

Presentation: Single-dose vial lyophilized vaccine and single-dose vial or manufacturer-filled syringe diluent

Protect From Light
Do Not Freeze

111

Resources

Vaccine Information Statements

- Provide the polio and Hib vaccine information statement (VIS) when a combination vaccine is administered.
 - There are no VISs specific for Kinrix,
 Pediarix, Pentacel, Vaxelis, or Quadracel.
- Other option: Multiple Vaccines VIS
 - May be used in place of the individual VISs for DTaP,
 Hib, hepatitis B, polio, and PCV when two or more of these vaccines are administered during the same visit
 - May be used for infants through children receiving their routine 4- to 6-year vaccines





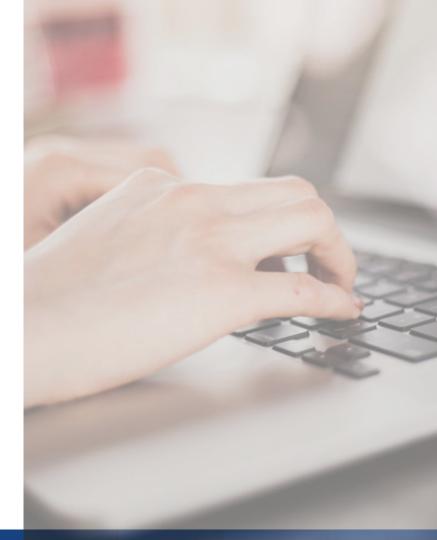


CDC Clinical Resources

- www.cdc.gov/vaccines/
 - Advisory Committee on Immunization Practices
 (ACIP) Vaccine Recommendations and Guidelines
 - Recommended Immunization Schedules
 - Vaccine Storage and Handling Toolkit
 - Vaccine Information Statements

Pink Book Training Materials





Continuing Education Information

- To claim continuing education (CE) for this course, please follow the steps below by July 1, 2026.
- Search and register for course WD4810-080824 in CDC TRAIN.
- Pass the post-assessment at 80%.
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 or CE Coordinator, Melissa Barnett, at MBarnett2@cdc.gov



Email Us Your Immunization Questions



nipinfo@cdc.gov

Thank You From Atlanta!

For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.



