National Center for Immunization and Respiratory Diseases



Meningococcal Disease and Meningococcal 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.

Continuing Education Information

- To claim continuing education (CE) for this course, please follow the steps below by July 1, 2026.
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- Pass the post-assessment at 80%.
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 during the previous 24 months as well as any use of unlabeled product(s) or products
 under investigational use.
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- CDC did not accept financial or in-kind support from any ineligible company for this continuing education activity.

Disclosure Statements

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

Meningococcal Disease

Meningococcal Disease (1)







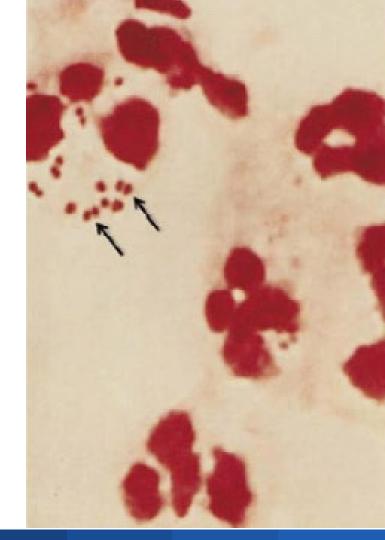
Meningococcal Disease (2)

- Clinical presentations primarily meningitis, bacteremia, or both
- Complications
 - 20% with long-term disabilities
 - 10%–15% fatality rate
 - Up to 40% in meningococcal bacteremia
- Less common presentations
 - Septic arthritis
 - Conjunctivitis
 - Urethritis



Neisseria meningitidis (1)

- Aerobic gram-negative bacteria
- Key antigens
 - Outer membrane protein
 - Polysaccharide capsule
- Spread by respiratory droplets or direct contact with respiratory secretions

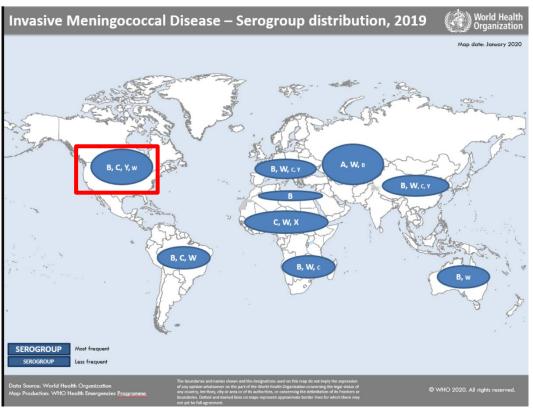


Neisseria meningitidis (2)

• Six serogroups cause most meningococcal disease

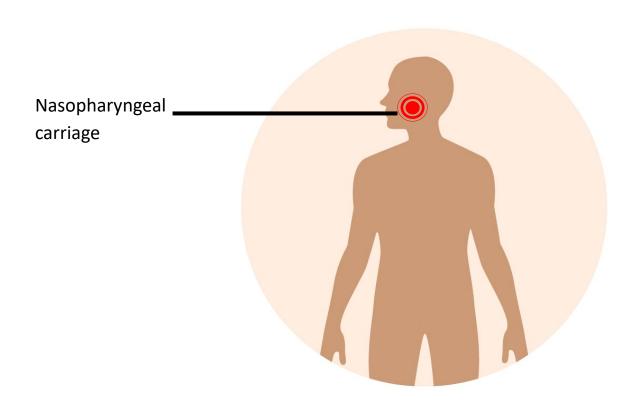


Neisseria meningitidis Serogroup Distribution



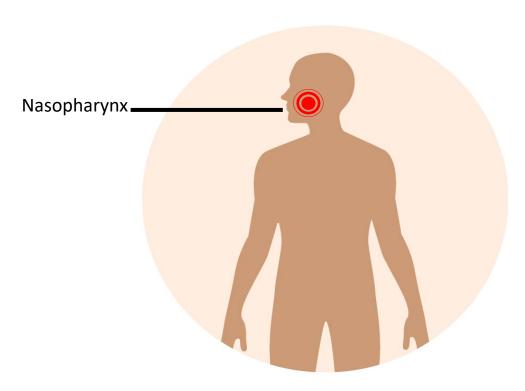
Information from: 9789240017481eng.pdf (who.int)

Transmission of *Neisseria meningitidis* (1)



About 10% of adolescents and adults are asymptomatic nasopharyngeal carriers.

Transmission of *Neisseria meningitidis* (2)



- Transmission:
 Respiratory droplets
 or direct contact with
 respiratory secretions
- Limited communicability:
 2–4 cases per 1000
 household members at risk

Transmission of *Neisseria meningitidis* (3)



Can occur throughout the year, but incidence is highest in late winter and early spring



Meningococcal Disease Pathogenesis

Exposure to Neisseria meningitidis



Meningococci attach to and multiply in the nasopharynx and oropharynx mucosa



Penetrate mucosal cells and enter blood stream in less than 1% of persons

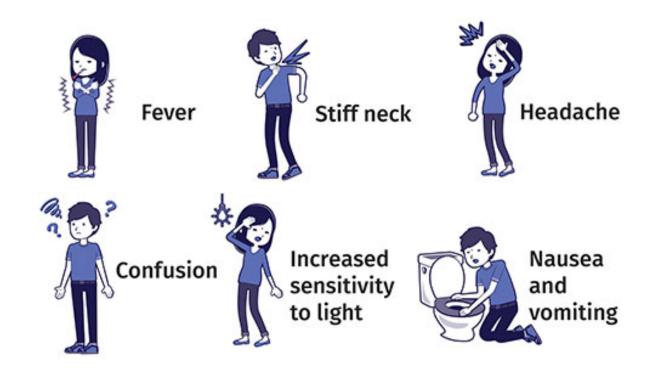


Spread through blood to cause systemic disease, and cross blood-brain barrier into cerebrospinal fluid to cause meningitis

Clinical Manifestations

- Incubation period 3–4 days (range 1–10 days)
- Common clinical manifestations:
 - Meningitis
 - Bacteremia/septicemia
- Other presentations:
 - Septic arthritis
 - Conjunctivitis
 - Urethritis

Meningococcal Meningitis



Meningococcal Bacteremia/Septicemia (1)



Meningococcal Bacteremia/Septicemia (2)



Risk Factors for Meningococcal Disease (1)



- Receipt of complement inhibitors (eculizumab, ravulizumab, etc.)
- Persistent complement component deficiency
- Functional or anatomic asplenia
- Human immunodeficiency virus (HIV)



Environmental Factors

- Active or passive smoking
- Antecedent viral infection
- Housing:
 - Military recruits
 - Unvaccinated or undervaccinated firstyear college students living in residence halls



Travel

 Traveling where meningococcal disease is hyperendemic or epidemic



Occupational Factors

Microbiologists



- Men who have sex with men
- People experiencing homelessness
- College campuses with outbreaks

Risk Factors for Meningococcal Disease (2)



- Receipt of complement inhibitors (eculizumab, ravulizumab, etc.)
- Persistent complement component deficiency
- Functional or anatomic asplenia
- Human immunodeficiency virus (HIV)



Environmental Factors

- Active or passive smoking
- Antecedent viral infection
- Housing:
 - Military recruits
 - Unvaccinated or undervaccinated firstyear college students living in residence halls



Trave

 Traveling where meningococcal disease is hyperendemic or epidemic



Occupational Factor

Microbiologists



- Men who have sex with men
- People experiencing homelessness
- College campuses with outbreaks

Risk Factors for Meningococcal Disease (3)



Host Factor

- Receipt of complement inhibitors (eculizumab, ravulizumab, etc.)
- Persistent complement component deficiency
- Functional or anatomic asplenia
- Human immunodeficiency virus (HIV)



Environmental Factors

- Active or passive smoking
- Antecedent viral infection
- Housing:
 - Military recruits
 - Unvaccinated or undervaccinated firstyear college students living in residence halls



Trave

 Traveling where meningococcal disease is hyperendemic or epidemic



Occupational Factor

Microbiologists



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Risk Factors for Meningococcal Disease (3)



Host Factor

- Receipt of complement inhibitors (eculizumab, ravulizumab, etc.)
- Persistent complement component deficiency
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- Human immunodeficiency virus (HIV)



Environmental Factors

- Active or passive smoking
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- Housing:
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Travel

 Traveling where meningococcal disease is hyperendemic or epidemic

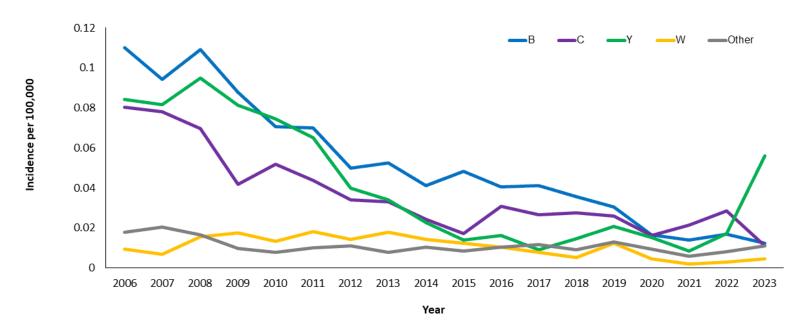


Microbiologists



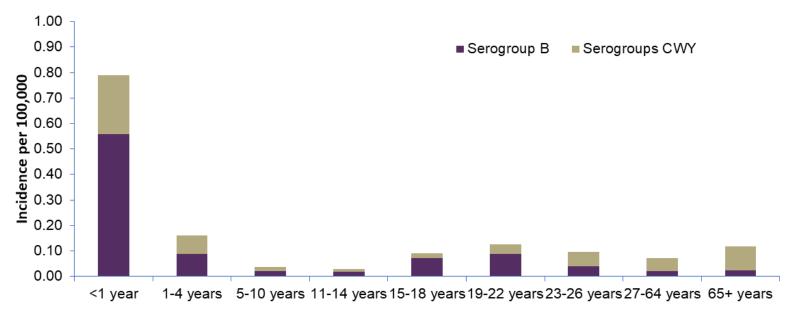
- Men who have sex with men
- People experiencing homelessness
- College campuses with outbreaks

Trends in Meningococcal Disease Incidence by Serogroup – United States, 2006–2023*



Source: NNDSS data with additional serogroup data from Active Bacterial Core surveillance (ABCs) and state health departments *2022 and 2023 data are preliminary

Meningococcal incidence by serogroup* and age-group, 2012–2021



^{*} Unknown serogroup (12%) and other serogroups (9%) excluded

SOURCE: CDC; National Notifiable Diseases Surveillance System with additional serogroup data from Active Bacterial Core surveillance and state health departments

National Notifiable Diseases Surveillance System Meningococcal Disease Surveillance and Trends | Meningococcal Disease | CDC



Knowledge Check

Meningitis with or without bacteremia is the primary clinical presentation of meningococcal disease.

- A. True
- B. False



Meningitis with or without bacteremia is the primary clinical presentation of meningococcal disease.

A. True

B. False

Meningococcal Vaccines

Meningococcal ACWY and Meningococcal B Vaccines, Unites States, 2024 (1)

Vaccine Product	Trade Name	Licensed Age Group*		
Quadrivalent meningococcal conjugate vaccines (MenACWY)				
MenACWY-CRM	Menveo	2 months-55 years		
MenACWY-TT	MenQuadfi	≥2 years		
Serogroup B meningococcal vaccines (MenB)				
MenB-FHbp	Trumenba	10–25 years		
MenB-4C	Bexsero	10–25 years		

^{*}ACIP recommends off-label use of vaccine products outside of the licensed maximum age

Information from: Chapter 14: Meningococcal Disease | Pink Book | CDC and https://www.cdc.gov/vaccines/vpd/mening/hcp/about-vaccine.html

Meningococcal ACWY and Meningococcal B Vaccines, Unites States, 2024 (2)

Vaccine Product	Trade Name	Licensed Age Group*		
Quadrivalent meningococcal conjugate vaccines (MenACWY)				
MenACWY-CRM	Menveo	2 months–55 years		
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Serogroup B meningococcal vaccines (MenB)				
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Meningococcal ACWY and Meningococcal B Vaccines, Unites States, 2024 (3)

Vaccine Product	Trade Name	Licensed Age Group*		
Quadrivalent meningococcal conjugate vaccines (MenACWY)				
MenACWY-CRM	Menveo	2 months-55 years		
MenACWY-TT	MenQuadfi	≥2 years		
Serogroup B meningococcal vaccines (MenB)				
MenB-FHbp	Trumenba	10–25 years		
MenB-4C	Bexsero	10–25 years		

^{*}ACIP recommends off-label use of vaccine products outside of the licensed maximum age

Information from Chapter 14: Meningococcal Disease | Pink Book | CDCl and https://www.cdc.gov/vaccines/vpd/mening/hcp/about-vaccine.html

Meningococcal Single-Component Vaccine Products (4)

MenACWY Vaccines



No adjuvants



No antibiotics



No preservatives

MenB Vaccines



Aluminum as an adjuvant



 Kanamycin as an antibiotic (MenB-4C/Bexsero only)



No preservatives

Vaccine Effectiveness

MenACWY – Wanes over time

- 79% effective within 1 year of vaccination
- 61% effective within 3–8 years after vaccination

MenB

- No data available on vaccine effectiveness against clinical disease among populations recommended for vaccination in the United States
- 84–88% immunogenicity in adolescents and college students
- Wanes 1–2 years after completion of primary series

Meningococcal Combination Vaccine Product

Vaccine Product	Trade Name	Licensed Age Group*		
Pentavalent meningococcal vaccine				
MenACWY-TT/MenB-FHbp	Penbraya	10 years-25 years		

^{*}ACIP recommends off-label use after age 25 years; the maximum licensed age

Vaccine Preparation: Menveo Two-Vial

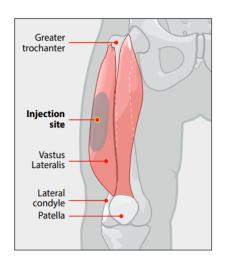
- Lyophilized formulation must be reconstituted (mixed) prior to administration
 - Vial containing liquid MenCYW (gray cap)
 - Vial containing lyophilized MenA (orange cap)
- Use only manufacturer-supplied vaccine diluent.

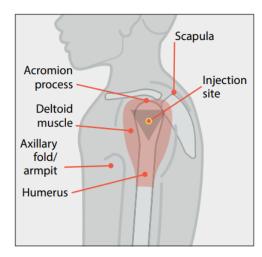
Vaccine Preparation: Penbraya

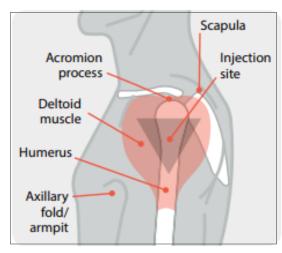
- Lyophilized formulation must be reconstituted (mixed) prior to administration
 - Vial containing lyophilized MenACYW (powder)
 - Prefilled syringe containing MenB (liquid)
 - Vial adapter
- Contains aluminum as an adjuvant but has neither antibiotic nor preservative

Vaccine Administration

All meningococcal vaccines are administered by intramuscular injection.







3

Clinical Considerations

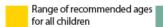
Meningococcal Vaccine Recommendations: Children and Adolescents

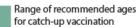


Recommended Child and Adolescent Immunization Schedule for Ages 18 Years or Younger, United States, 2025

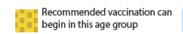
These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2).

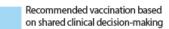
Vaccine and other immunizing agents	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 mos	18 mos	19–23 mos	2–3 yrs	4-6 yrs	7-10 yrs	11–12 yrs	13–15 yrs	16 yrs	17–18 yrs
$\label{eq:Menacwy-CRM} \begin{split} & \text{Meningococcal} (\text{MenACWY-CRM} \geq & 2 \text{mos}, \\ & \text{MenACWY-TT} \geq & 2 \text{years}) \end{split}$								See Notes						1st dose		2nd dose	
Meningococcal B (MenB-4C, MenB-FHbp)															See No	tes	













Meningococcal Vaccine Recommendations: Adults



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

Vaccine	19–26 years	27–49 years	50–64 years	≥65 years					
Meningococcal A, C, W, Y (MenACWY)	1 or 2 doses depending on indication (See Notes for booster recommendations)								
Meningococcal B (MenB)	19 through 23 years	2 or 3 doses depending on vaccine and indication (See Notes for booster recommendations)							
(19 tillough 25 years								

Recommended vaccination for adults who meet age requirement, lack documentation of vaccination, or lack evidence of immunity

Recommended vaccination for adults with an additional risk factor or another indication

Recommended vaccination based on shared clinical decision–making No Guidance/ Not Applicable

Routine MenACWY Recommendations for Children/Adolescents

Table 1

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

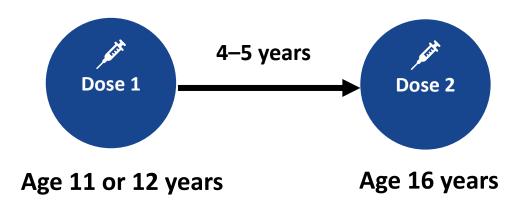
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Meningococcal (MenACWY-CRM ≥ 2 mos, MenACWY-TT ≥ 2 years)								See Notes						1st dose		2nd dose	
Meningococcal B (MenB-4C, MenB-FHbp)															See No	ites	

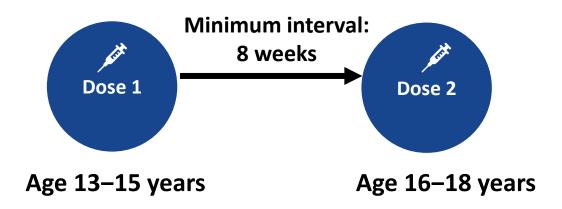
- Primary vaccination: 1 dose at age 11 or 12 years
- **Booster vaccination**: 1 dose at age 16 years
- Catch up vaccination
 - 1 dose at age 13–15 years
 - Single booster at age 16–18 years (minimum interval 8 weeks)
 - No booster if primary dose administered on or after 16th birthday
- Ages 19–21 years
 - Can receive 1 dose if have no dose administered after 16th birthday
 - After age 21 years, no booster dose is recommended for healthy persons

Information from https://www.cdc.gov/vaccines/hcp/imz-schedules/child-adolescent-age.html

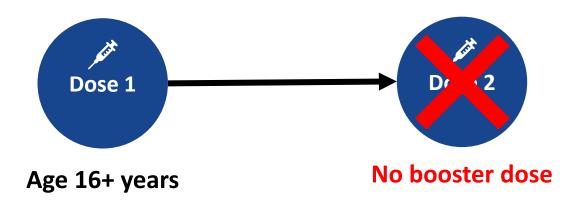
MenACWY Recommendations for Children and Adolescents: Routine Vaccination



Catch-up MenACWY Recommendations for Children and Adolescents (1)



Catch-up MenACWY Recommendations for Children and Adolescents (2)



Catch-up MenACWY Recommendations for Children and Adolescents (3)



If **no** dose was given after the 16th birthday

MenACWY may be given to persons ages 19–21 years

Catch-up MenACWY Recommendations for Children and Adolescents (4)

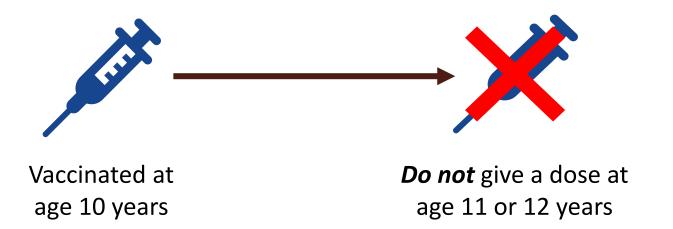


If **no** dose was given after the 16th birthday

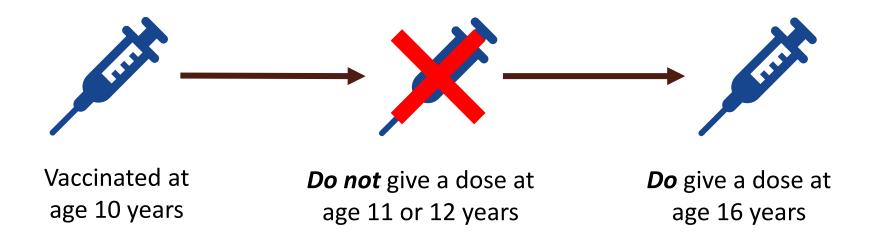
MenACWY may be given to persons ages 19–21 years

Not routinely recommended after age 21 years

Routine MenACWY Use in Children Before Age 11 Years (1)



Routine MenACWY Use in Children Before Age 11 Years (2)



Routine MenACWY Use in Children Before Age 11 Years (3)

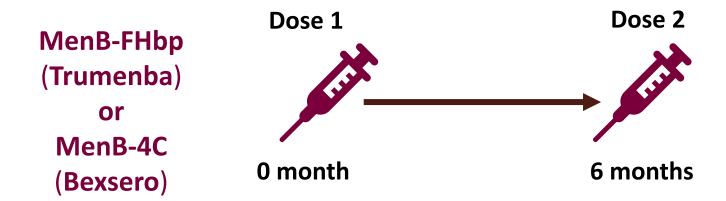


Routine MenB Recommendations for Adolescents

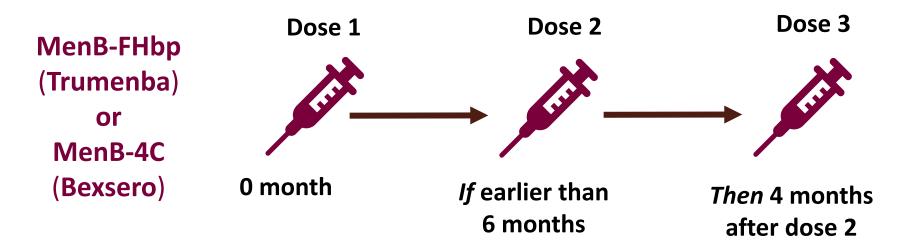
- Not routinely recommended for all adolescents
- Primary vaccination
 - 2 doses at ages 16–23 years based on *shared clinical decision-making*
 - Preferred age 16–18 years
- Booster vaccination not recommended*

^{*}Booster not recommended for healthy persons unless ongoing risk of exposure such as microbiologist exposed to *N. meningitidis* and persons exposed during an outbreak

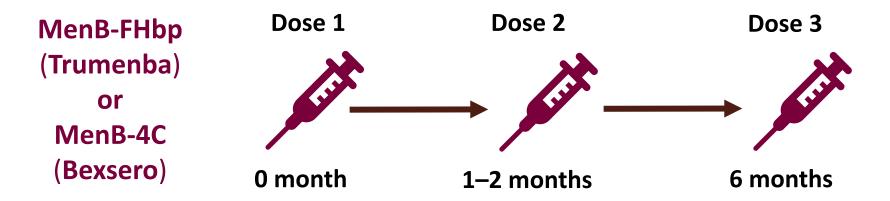
Routine MenB Recommendations for Adolescents: Dose Intervals (1)



Routine MenB Recommendations for Adolescents: Dose Intervals (2)



Routine MenB Recommendations for Adolescents: Dose Intervals if Rapid Protection Is Desired*



^{*} For example, for students starting college in less than 6 months.

MenB Recommendations: Considerations for Shared Clinical Decision-Making



Clinical factors



Risk factors



Vaccine effectiveness

Considerations for Shared Clinical Decision-Making (1)

Clinical factors

- Serious nature of meningococcal infections
- High death rate
- Severe complications
 - e.g., loss of limb(s), neurologic disability, hearing loss



Considerations for Shared Clinical Decision-Making (2)

Considerations:

- Increased risk among college students
- Rapid course of severe illness and risk of death even with prompt recognition and antibiotic treatment



Considerations for Shared Clinical Decision-Making (3)

Vaccine effectiveness

- Protection against most strains of serogroup
 B N. meningitidis
- Estimated relatively short duration of MenB protection
- Evidence to date suggests MenB vaccination has no effect on meningococcal carriage





Knowledge Check

A healthy 20-year-old college freshman has previously received two doses of MenACWY vaccine.

- Dose 1 at age 13 years
- Dose 2 at age 15 years

Her school is requesting an additional dose of MenACWY because she will be living in college residential housing. Can she get a 3rd dose today?

- A. Yes
- B. No

Answer

A healthy 20-year-old college freshman has previously received two doses of MenACWY vaccine.

- Dose 1 at age 13 years
- Dose 2 at age 15 years

Her school is requesting an additional dose of MenACWY because she will be living in college residential housing. Can she get a 3rd dose today?

- A. Yes
- B. No

Meningococcal Vaccination for Persons at *Increased Risk* (1)

Table 1

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
Meningococcal (MenACWY-CRM ≥2 mos, MenACWY-TT ≥2 years)								See Notes						1st dose		2nd dose	
Meningococcal B (MenB-4C, MenB-FHbp)															See No	otes	

Table 1

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

Vaccine	19–26 years	27–49 years	50–64 years	≥65 years					
Meningococcal A, C, W, Y (MenACWY)		1 or 2 doses depending on indication (See Notes for booster recommendations)							
Meningococcal B		2 - 2 days days dispersion of indication (Co. Natural Co. Natural							
(MenB)	19 through 23 years	2 or 3 doses depending on vaccine and indication (See Notes for booster recommenda							

Meningococcal Vaccination for Persons at Increased Risk (2)

Risk Group	MenACWY	MenB
Persons with complement component deficiency (including patients using a complement inhibitor)	Age ≥2 months	Age ≥10 years
Persons with functional or anatomic asplenia (including sickle cell disease)	Age ≥2 months	Age ≥10 years
Persons with HIV infection	Age ≥2 months	No recommendation
Microbiologists routinely exposed to Neisseria meningitidis	Age appropriate*	Age appropriate [†]
Persons exposed during an outbreak of meningococcal disease due to a vaccine-preventable serogroup	Age ≥2 months	Age ≥10 years
Persons who travel to or live in countries where meningococcal disease is hyperendemic or epidemic	Age ≥2 months	No recommendation
College freshmen living in residence halls	Age appropriate*	No recommendation
Military recruits	Age appropriate*	No recommendation

[•] Persons aged ≥2 months in these risk groups are recommended to receive MenACWY vaccination.

[†] Persons aged ≥10 years in this risk group are recommended to receive MenB vaccination

^{*}Shared clinical decision-making recommendation for all persons ages 16-23 years

Meningococcal Vaccination for Persons at Increased Risk (3)

Risk Group	MenACWY	MenB
Persons with complement component deficiency (including patients using a complement inhibitor)	Age ≥2 months	Age ≥10 years
Persons with functional or anatomic asplenia (including sickle cell disease)	Age ≥2 months	Age ≥10 years
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Military recruits	Age appropriate*	No recommendation

[•] Persons aged ≥2 months in these risk groups are recommended to receive MenACWY vaccination.

[†] Persons aged ≥10 years in this risk group are recommended to receive MenB vaccination.

^{*}Shared clinical decision-making recommendation for all persons ages 16-23 years

Meningococcal Vaccination for Persons at Increased Risk (4)

Risk Group	MenACWY	MenB
Persons with complement component deficiency (including patients using a complement inhibitor)	Age ≥2 months	Age ≥10 years
Persons with functional or anatomic asplenia (including sickle cell disease)	Age ≥2 months	Age ≥10 years
Persons with HIV infection	Age ≥2 months	No recommendation
Microbiologists routinely exposed to Neisseria meningitidis	Age appropriate*	Age appropriate [†]
Persons exposed during an outbreak of meningococcal disease due to a vaccine-preventable serogroup	Age ≥2 months	Age ≥10 years
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Military recruits	Age appropriate*	No recommendation

[•] Persons aged ≥2 months in these risk groups are recommended to receive MenACWY vaccination.

[†] Persons aged ≥10 years in this risk group are recommended to receive MenB vaccination.

^{*}Shared clinical decision-making recommendation for all persons ages 16-23 years

MenACWY-CRM (Menveo): Anatomic or Functional Asplenia, HIV Infection, Persistent Complement Component Deficiency, Complement Inhibitor Use (1)

- 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
 - Then 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

MenACWY-TT (MenQuadfi): Anatomic or Functional Asplenia, HIV Infection, Persistent Complement Component Deficiency, Complement Inhibitor Use (2)

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

MenACWY: Travel to Countries With Hyperendemic or Epidemic Meningococcal Disease

- Ages 2–23 months: MenACWY-CRM (Menveo)
 - 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
 - Then 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
- Ages 2 years or older: 1 dose Menveo or MenQuadfi

MenACWY: First-Year College Students Who Live in Residential Housing, or Military Recruits

 1 dose MenACWY-CRM (Menveo) or MenACWY-TT (MenQuadfi) if not previously vaccinated at age 16 years or older

MenACWY Booster Recommendations for Persons at *Increased Risk*

- For persons who remain at increased risk and completed the primary vaccination at age:
 - Younger than 7 years
 - 1 dose 3 years after primary series
 - Boosters every 5 years
 - 7 years and older
 - 1 dose 5 years after primary series
 - Boosters every 5 years

MenB Vaccination Schedule for Persons at Increased Risk

- MenB-4C (Bexsero) or MenB-FHbp (Trumenba): 3-dose series at 0, 1–2, 6 months
 - If dose 2 administered at least 6 months after dose 1, dose 3 not needed
 - If dose 3 administered earlier than 4 months after dose 2, a 4th dose should be given at least 4 months after dose 3

MenB Booster Recommendations for Persons at Increased Risk

- Persons with complement deficiency, complement inhibitor use, or functional or anatomic asplenia, or who are routinely exposed to Neisseria meningitidis at work
 - Booster dose at least 1 year since primary series
 - Repeat every 2–3 years as long as risk remains
- At risk due to Serogroup B outbreak
 - Booster dose at least 1 year since primary series
 - If recommended by public health officials, booster dose may be given if it has been at least 6 months since primary series

Use of Penbraya

 Combination MenACWY-TT/MenB-FHbp (Penbraya) is an option when both MenACWY and MenB are recommended (inclusive of SCDM) for same visit.

- Minimum 6-month interval between:
 - Any 2 doses of Penbraya
 - Penbraya and Trumenba



Knowledge Check

A 16-year-old recently began treatment with eculizumab—a complement inhibitor—and is recommended to begin MenB vaccination series. Your clinic has Trumenba in stock. How many doses of Trumenba should she receive?

- A. Trumenba 2 doses at 0 and 6 months
- B. Trumenba 3 doses (0, 1–2, and 6 months)



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Interchangeability of Meningococcal Vaccine Products (1)



MenACWY vaccines are interchangeable



MenB vaccines are not interchangeable

Interchangeability of Meningococcal Vaccine Products (2)



MenACWY vaccines are interchangeable

Same vaccine product is recommended but not required for all doses.



MenB vaccines

are not

interchangeable

Interchangeability of Meningococcal Vaccine Products (3)



MenACWY vaccines

Are interchangeable



MenB vaccines *are not* interchangeable

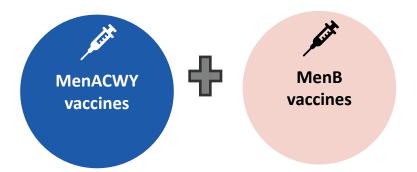
Use same product for all doses.

If 2 different products are administered, pick one and invalidate the dose of the other.

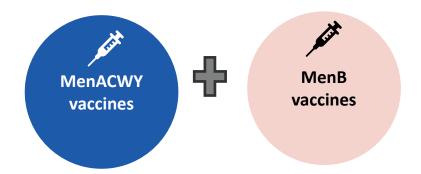
Space repeat dose at a minimum interval of 4 weeks.

Co-Administration of Meningococcal Vaccine Products

Same day administration

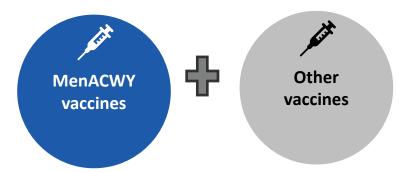


Any interval



Co-Administration of MenACWY and Other Vaccines

Same day administration

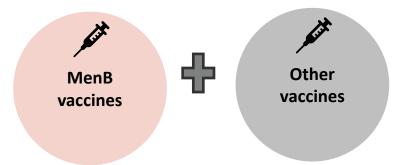


Any interval

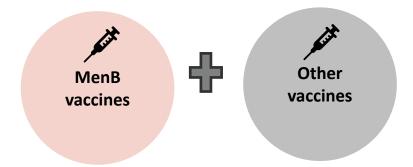


Co-Administration of MenB and Other Vaccines

Same day administration



Any interval



Off-Label Meningococcal Vaccination Recommendations for Persons at Risk

Age Group	Indication
2 years and older	Administration of a 2-dose MenACWY primary series in persons at increased risk for serogroups A, C, W, or Y meningococcal disease
	Repeated booster doses of MenACWY for certain persons who remain at increased risk for serogroups A, C, W, or Y meningococcal disease
	(MenACWY-CRM is licensed for a single booster dose for persons aged 15–55 yrs if at least 4 yrs have elapsed since the last dose. MenACWY-TT is licensed for a single booster dose for persons aged ≥15 yrs if at least 4 yrs have elapsed since the last dose of MenACWY)
10 years and older	MenB booster doses in certain persons who remain at increased risk for serogroup B meningococcal disease
26 years and older	MenB primary series administration in persons at increased risk for serogroup B meningococcal disease
56 years and older	Administration of MenACWY-CRM in persons at increased risk for serogroups A, C, W, or Y meningococcal disease

Abbreviations

MenACWY = quadrivalent meningococcal conjugate vaccine; MenACWY-CRM = meningococcal groups A, C, W, and Y oligosaccharide diphtheria CRM₁₉₇ conjugate vaccine; MenACWY-D = meningococcal groups A, C, W, and Y polysaccharide diphtheria toxoid conjugate vaccine; MenACWY-TT = meningococcal groups A, C, W, and Y polysaccharide tetanus toxoid conjugate vaccine; MenB = serogroup B meningococcal vaccine

Mbaeyi SA, Bozio CH, Duffy J, et al. Meningococcal Vaccination: Recommendations of the Advisory Committee on Immunization Practices, United States, 2020. MMWR Recomm Rep 2020;69(No. RR-9):1–41. DOI: http://dx.doi.org/10.15585/mmwr.rr6909a1external icon.

Safety

Common Adverse Reactions: MenACWY Vaccines



Fever



Irritability



Fatigue, drowsiness



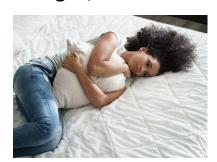
Injection site reactions: pain and erythema



Headache



Myalgia



Malaise

Common Adverse Reactions: MenB Vaccines



Headache



Fever



Fatigue



Myalgia



Arthralgia



Injection site reactions: pain, induration, erythema

Information from: Chapter 14: Meningococcal Disease | Pink Book | CDC

Contraindications: MenACWY and MenB

- Severe allergic reaction (e.g., anaphylaxis) after a previous dose
- Severe allergic reaction (e.g., anaphylaxis) to a vaccine component:
 - For MenACWY-CRM (Menveo) only: severe allergic reaction to any diphtheria toxoid- or CRM197-containing vaccine
 - For MenACWY-TT (MenQuadfi) only: severe allergic reaction to a tetanus toxoidcontaining vaccine

Precautions to MenACWY and MenB

- Moderate or severe acute illness, with or without fever
- For MenACWY-CRM (Menveo) only: infants born preterm if younger than age 9 months
- For MenB only: pregnancy
- For MenB-4C (Bexsero) only: anaphylactic allergy to latex



5

Storage & Handling

Meningococcal Vaccine Storage and Handling

- Store meningococcal vaccines refrigerated between
 2°C and 8°C (36°F and 46°F).
- Do not freeze vaccine or diluents, or expose to freezing temperatures.
- Store meningococcal vaccines in original packaging.



Vaccine Storage and Handling: MenACWY-CRM (Menveo)

- MenACWY-CRM (Menveo) two-vial presentation requires reconstitution.
- The MenA (lyophilized) component should be reconstituted only with the liquid C-W-Y component of Menveo.
- Reconstituted vaccine should be used immediately but may be stored between 36°F and 77°F (2°C and 25°C) for up to 8 hours.
- Do not use if the reconstituted vaccine cannot be resuspended with thorough agitation.



Improper Storage and Handling of Meningococcal Vaccines

- If the vaccine product is exposed to inappropriate temperatures or conditions:
 - Store the vaccine at the appropriate temperature.
 - Isolate from other vaccines.
 - Mark "Do NOT Use."
 - Consult the vaccine manufacturer or your state or local immunization program for guidance.

6

Resources

Meningococcal Resources

Meningococcal disease

Meningococcal Disease | Meningococcal | CDC

ACIP's Meningococcal Recommendations

ACIP Recommendations: Meningococcal Vaccine | ACIP Recommendations
 CDC

Meningococcal Vaccination for healthcare providers

Meningococcal Vaccination | For Providers | CDC

Immunize.org

https://www.immunize.org/meningococcal-acwy/

Children's Hospital of Philadelphia Vaccine Education Center Meningococcal web page

• https://www.chop.edu/centers-programs/vaccine-education-center/vaccine-details/meningococcal-vaccine

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 WD4810-080124 in CDC TRAIN.
- Pass the post-assessment at 80%.
- Complete the evaluation.
- Visit "Your Learning" to access your certificates and transcript.
- If you have any questions, contact CDC TRAIN at train@cdc.gov 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.



