# Standing Order for Administering Meningococcal ACWY Vaccine (Adult)

**Purpose:** To reduce morbidity and mortality from meningococcal disease by vaccinating all individuals who meet the criteria established by the Centers for Disease Control and Prevention's Advisory Committee on Immunization Practices (ACIP), the Food and Drug Administration (FDA) product labeling, and the Department of Defense (DoD).

**Policy:** Under this standing order, eligible health care professionals working within their scope of practice may vaccinate patients who meet the criteria below.

#### **Procedure:**

- 1. Identify individuals ≥ 19 years of age in need of vaccination against meningococcal serogroups A, C, W, and Y based on the <u>following criteria</u>:
  - No documented receipt of a complete routine series of meningococcal ACWY vaccine (MenACWY) at the appropriate ages and intervals.
  - At increased risk due to:
    - Asplenia (anatomic or functional) or sickle cell disease (SCD)
    - HIV infection
    - Microbiologists routinely exposed to Neisseria meningitidis
    - Men who have sex with men (MSM)
    - Military recruits
    - Persistent (e.g., genetic) complement deficiency or using a complement inhibitor medication
    - o Travel to or living in countries where meningococcal disease is hyperendemic or epidemic
    - Unvaccinated or undervaccinated 1<sup>st</sup> year college students living in residence halls
    - Meningococcal outbreaks
- 2. Using DD Form 3111, screen all patients for contraindications and precautions to MenACWY:

### **Contraindications:**

- History of a serious reaction (e.g., anaphylaxis) after a previous dose of meningococcal vaccine or to a vaccine component
- MenACWY-CRM (Menveo): severe allergic reaction to a diphtheria toxoid
   or CRM<sub>197</sub>
   containing vaccine
- MenACWY-TT (MenQuadfi) and MenABCWY (Penbraya): severe allergic reaction to a tetanus toxoid-containing vaccine
- Penbraya: severe allergic reaction to yeast
- For information on vaccine components, refer to the package inserts for MenQuadfi, Menveo, and Penbraya, and The CDC Pink Book Appendix B.

## **Precautions:**

- Moderate or severe acute illness with or without fever
- For questions or concerns, consider consulting the DHA Immunization Healthcare Support Center at (877) 438-8222, Option 1 or DSN 312-761-4245.

## **Special Populations:**

- Pregnancy and Lactation: Pregnant and lactating women should receive MenACWY vaccine if indicated.
- 3. Prior to vaccine administration, provide all patients (or their parent/legal representative) with a copy of the current federal <u>Vaccine Information Statement</u> (VIS). Provide non-English speaking patients with a copy of the VIS in their native language, if available and preferred (<u>www.immunize.org/vis</u>).
- 4. Provide MenACWY as follows:
  - Administer the appropriate vaccine intramuscularly (IM) according to Tables 1 3.

- Off-label ACIP recommendations covered under this standing order:
  - A 2-dose primary series in persons at increased risk due to certain underlying medical conditions
  - o Repeated booster doses for persons who remain at increased risk
  - ≥ 56 years: administration of MenACWY-D (Menactra) or Menveo in persons at increased risk
    - Production of Menactra was discontinued in 2022. Remaining stock may be used according to previous schedules through the expiry date or until it is no longer FDAlicensed, whichever is earlier.
- MenACWY vaccines are interchangeable; the same product is recommended, but not required, for all doses (primary and booster).
- MenACWY and meningococcal B vaccine (MenB) may be administered simultaneously (at different anatomic sites) if indicated.
- Penbraya may only be used when both MenACWY and MenB are indicated at the same visit.
  Consult the age appropriate MenACWY and MenB standing orders for indications and dosing.
  Vaccination of healthy individuals aged 19–23 years (preferred at 16-18 years) with a 2-dose
  MenB primary series is based on shared clinical decision-making (SCDM) and is not covered
  under this standing order. These individuals must obtain a written order from a privileged
  provider.

TABLE				
	MenQuadfi (MenACYW-TT)	Menveo / 1-vial (MenACWY-CRM)	Menveo / 2-vial (MenACWY-CRM)	Penbraya (MenABCWY)
Age	≥ 2 years	10 – 55 years	2 mo - 55 years	10 – 25 years
Dilute	No: single-dose vial	No: single-dose vial (pink cap)	Yes: MenA vial (orange cap) & MenCWY vial (gray cap)	Yes: MenACWY vial & MenB syringe

TABLE 2. IM Needle Length and Injection Site Guide, Adult ≥ 19 years				
<ul> <li>Use a 22 – 25-gauge needle</li> <li>Choose needle gauge and length appropriate to the patient's age, sex, and weight</li> </ul>				
Patient Group	Needle Length	Injection Site		
Men and women, <60 kg (130 lbs)	5/8* - 1 inch (16-25 mm)			
Men and women, 60-70 kg (130-152 lbs)	1 inch (25 mm)	Deltoid muscle of arm		
Men, 70-118 kg (152-260 lbs)	1.1.5 in shop (25.20 mms)			
Women, 70-90 kg (152-200 lbs)	1-1.5 inches (25-38 mm)			
Men, >118 kg (260 lbs)	4.5 in also a (00 mm)			
Women, >90 kg (200 lbs)	1.5 inches (38 mm)			
Men and women, any weight	1 inch* - 1.5 inches (25-38 mm)	Anterolateral thigh		

Adapted from the CDC General Best Practice Guidelines: https://www.cdc.gov/vaccines/hcp/acip-recs/general-recs/index.html.

<sup>\*</sup> If skin is stretched tightly and subcutaneous tissues are not bunched.

Not recommended unless person becomes at increased risk due another indication
<ul> <li>Every 5 years base on exposure risk</li> </ul>
Single dose 5 years after primary vaccination and every 5 years thereafter

- 5 location of administration, lot number, manufacturer, VIS date, and the identification of the person administering the vaccine. If vaccine was not given, record the reason for non-receipt.
- 6. Be prepared to manage a medical emergency related to the administration of vaccines by having a written emergency medical protocol available, as well as equipment and medications.
- 7. Adverse events occurring after administration of any vaccine should be reported to the Vaccine Adverse Event Reporting System (VAERS) online at <a href="https://vaers.hhs.gov">https://vaers.hhs.gov</a>. Additional VAERS information is available by telephone at (800) 822-7967.

8.	This standing order shall remain in effect for all patients of theuntil rescinded and/or upon a change in the Medical Director, whichever is earlier.				
	Medical Director's Signature	Date			