



PENMENVY

Meningococcal Groups
A, B, C, W, and Y Vaccine

You are invited to the following presentation:

Topic

Presented by

Date/Time

Location

*Featuring the personal story of a
meningococcal disease patient advocate*



RSVP/Questions

Program code

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Indication

PENMENVY is a vaccine indicated for active immunization to prevent invasive disease caused by *Neisseria meningitidis* serogroups A, B, C, W, and Y in individuals 10 through 25 years of age.

Important Safety Information for PENMENVY

- Do not administer PENMENVY to individuals with a severe allergic reaction (e.g., anaphylaxis) to a previous dose of PENMENVY, to any component of this vaccine, or to any other diphtheria toxoid-containing vaccine
- Appropriate medical treatment must be immediately available to manage potential anaphylactic reactions following the administration of PENMENVY

Please see additional Important Safety Information on the next page, along with the accompanying [Prescribing Information](#) for PENMENVY.

Important Safety Information for PENMENVY (Meningococcal Groups A, B, C, W, and Y Vaccine) (cont'd)

- Syncope (fainting) has occurred in association with administration of PENMENVY. Procedures should be in place to avoid injury from fainting
- Vaccination with PENMENVY may not protect all vaccine recipients and may not provide protection against all meningococcal serogroup B strains
- Immunocompromised persons, including those receiving immunosuppressive therapy, may have reduced immune responses to PENMENVY
- Individuals with certain complement deficiencies and individuals receiving treatment that inhibits terminal complement activation are at increased risk for invasive disease caused by *N. meningitidis*, including disease caused by serogroups A, B, C, W, and Y, even if they develop antibodies following vaccination with PENMENVY
- Guillain-Barré syndrome (GBS) has been reported in temporal relationship following administration of a U.S.-licensed meningococcal quadrivalent polysaccharide conjugate vaccine. The decision by the healthcare professional to administer PENMENVY to persons with a history of GBS should take into account the expected benefits and potential risks
- The most commonly reported ($\geq 10\%$) solicited adverse reactions in individuals aged 10 through 25 years after Dose 1 and Dose 2, respectively, were pain at the injection site (92% and 88%), fatigue (51% and 42%), headache (42% and 36%), myalgia (15% and 12%), nausea (15% and 10%), erythema (13% and 12%), and swelling (13% and 12%). The most commonly reported ($\geq 10\%$) solicited adverse reactions in MenACWY conjugate vaccine-experienced individuals aged 15 through 25 years after Dose 1 and Dose 2, respectively, were pain at the injection site (80% and 74%), headache (41% and 33%), fatigue (40% and 33%), myalgia (15% and 13%), and nausea (15% and 12%)

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