



Let's talk about pneumococcal pneumonia.

Take this guide to your next doctor's appointment or pharmacy visit. Here are some questions to ask:

Am I at risk because of my age or health conditions?

- How does my age put me at risk for pneumococcal pneumonia?
- How do my health conditions put me at risk for pneumococcal pneumonia?
- How could pneumococcal pneumonia make my health condition worse?
- Do the medications I take suppress my immune system, putting me at risk for pneumococcal pneumonia? (Bring a list of all your medications to show your doctor.)

Is Pevnar 20® right for me?

- Can Pevnar 20 help provide additional protection if I have already been vaccinated with another pneumonia vaccine?
- Where can I get Pevnar 20?

Have questions about pneumococcal pneumonia for your doctor or pharmacist? Write them here:

IMPORTANT SAFETY INFORMATION

- Pevnar 20® should not be given to anyone who has had a severe allergic reaction to any component of Pevnar 20 or to diphtheria toxoid
- Individuals with weakened immune systems may have a lower immune response. Safety data are not available for these groups
- In individuals 18 years of age and older, the most common side effects were pain at the injection site, muscle pain, fatigue, headache, and joint pain. Additionally, injection site swelling was also common in individuals 18 through 59 years of age
- Ask your healthcare provider about the risks and benefits of Pevnar 20. Only a healthcare provider can decide if Pevnar 20 is right for you

INDICATION FOR PREVNAR 20

Pevnar 20 is a vaccine approved for:

- the prevention of invasive disease caused by *Streptococcus pneumoniae* serotypes 1, 3, 4, 5, 6A, 6B, 7F, 8, 9V, 10A, 11A, 12F, 14, 15B, 18C, 19A, 19F, 22F, 23F, and 33F in individuals 6 weeks of age and older
- the prevention of pneumonia caused by *S. pneumoniae* serotypes 1, 3, 4, 5, 6A, 6B, 7F, 8, 9V, 10A, 11A, 12F, 14, 15B, 18C, 19A, 19F, 22F, 23F, and 33F in individuals 18 years of age and older
- The indication for the prevention of pneumonia caused by *S. pneumoniae* serotypes 8, 10A, 11A, 12F, 15B, 22F, and 33F in individuals 18 years of age and older is approved based on immune responses. Continued approval may depend on a supportive study

Patients should always ask their healthcare providers for medical advice about adverse events. You are encouraged to report negative side effects of vaccines to the US Food and Drug Administration (FDA) and the Centers for Disease Control and Prevention (CDC). Visit <https://www.vaers.hhs.gov/reportevent.html> or call 1-800-822-7967.

Please see Full Prescribing Information [here](#).



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Manufactured by Wyeth Pharmaceuticals LLC. Marketed by Pfizer Inc.

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