



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 Prevnar 20™ right for me?

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

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

IMPORTANT SAFETY INFORMATION

Prevnar 20™ should not be given to anyone with a history of severe allergic reaction to any component of Prevnar 20™ or to diphtheria toxoid

Adults with weakened immune systems may have a lower response to Prevnar 20™. Safety data are not available for these groups. Your healthcare provider can tell you if Prevnar 20™ is right for you

In adults 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 adults 18 through 59 years of age

Ask your healthcare provider about the risks and benefits of Prevnar 20™. Only a healthcare provider can decide if Prevnar 20™ is right for you

Please [click here](#) for Prevnar 20™ Full Prescribing Information. Full Prescribing Information is also available at <https://www.prevnar20.com>.

INDICATION FOR PREVNAV 20™

Prevnar 20™ is a vaccine indicated for active immunization for the prevention of pneumonia and 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 adults 18 years of age and older

The indication for preventing pneumonia caused by *S. pneumoniae* serotypes 8, 10A, 11A, 12F, 15B, 22F, and 33F 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 <http://www.vaers.hhs.gov> or call 1-800-822-7967.



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