

## Pneumococcal Antibody Panel (23 Serotype)

Order Name: **PNEUMO 23** Test Number: 5575605 Revision Date: 01/01/2025

TEST NAME Pneumococcal Antibody Panel (23 Serotype)			METHODOLOGY	LOINC CODE
			Fluoroimmunoassay	
SPECIMEN REQU	REMENTS			
Specimen	Specimen Volume (min)	Specimen Type	Specimen Container	Transport Environment
Preferred	1.0 mL (0.5 mL)	Serum	Clot Activator SST	Room Temperature
Instructions GENERAL INFORM	Notes: 0.5 mL (Note: This volume Does NOT allow for repeat testing).   Specimen Type: Gel-barrier tube   Specimen Storage: Maintain specimen at room temperature.   Specimen Collection: Not Available   Special Instructions: Testing referred to Eurofins Viracor LLC EURKS#401835   Specimen Stability: Ambient: 1 week, Refrigerated : 1 week, Frozen: 5 months			
Expected TAT	7-10 Days			
Clinical Use Includes Serotypes 1, 2, 3, 4, 5, 6B, 7F, 8, 9N, 9V, 10A, 11A, 12F, 14, 15B, 17F, 18C   Serotypes: 1, 3, 4, 5, 6B, 7F, 9V, 14, 18C, 19A, 19F, and 23F.).   A pre-and post-vaccination sample comparison is required in order to assess the hum   Streptococcus pneumoniae vaccine. Post vaccination samples should be obtained at   only general immune status of an individual to various serotypes of pneumococci. Imm   individuals based on the age, vaccination status, immunologic competence and the segenerally considered to be a serotype antibody level of 1.3 ug/ mL or greater.   Following vaccination, an antibody ratio of less than twofold is considered a non-respondence.				nune response to vaccination with post-immunization. A single sample provides sponse to pneumococci may vary in different of the organism. Long-term protection is

Following vaccination, an antibody ratio of less than twofold is considered a non-responder; a ratio of two-to fourfold is a weak responder; a ratio of fourfold or greater is a good responder. The higher the pre-vaccination antibody level for a specific pneumococcal serotype, the less likely the response will increase significantly after vaccination.

An increased antibody level to 50-70 percent or more of the serotypes is thought to represent a normal humoral response. In the case of pure polysaccharide vaccine, indication of immune system competence is further delineated as an adequate response to at least 70 percent of the serotypes in the vaccine challenge for those 6-65 years of age, or to at least 50 percent of the serotypes in the vaccine challenge for those 2-5 years of age.

CPT Code(s)

Service Provided By



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