

Pneumococcal Avidity Panel (12 Serotype)

Test Code: 401734P

Tests in this Panel

- Pneumo Ab Type 1 Avidity
- Pneumo Ab Type 3 Avidity
- Pneumo Ab Type 4 Avidity
- Pneumo Ab Type 8 Avidity
- Pneumo Ab Type 9 (9N) Avidity
- Pneumo Ab Type 12 (12F) Avidity
- Pneumo Ab Type 14 Avidity
- Pneumo Ab Type 19 (19F) Avidity
- Pneumo Ab Type 26 (6B) Avidity
- Pneumo Ab Type 51 (7F) Avidity
- Pneumo Ab Type 56 (18C) Avidity
- Pneumo Ab Type 23 (23F) Avidity

Clinical and Procedure

Clinical Utility

The avidity of antibody induced by a vaccine is an independent correlate of protection and this information is an important supplement to the measurement of antibody titer. The ability to generate higher avidity antibodies is a key aspect of a fully functional immune response. Avidity functions is an important determinant of protective efficacy against pneumococci. There is a strong correlation between avidity and opsonophagocytic activity.

Procedure

Multiplexed microsphere preparations, each coated with one Pne-PS serotype, are incubated with patient serum that has been preadsorbed with cell wall polysaccharide. After washing, varying concentrations of ammonium thiocyanate or PBS are added. Bound antibodies are detected with an antibody that recognizes human IgG. The IgG antibody concentration is calculated in samples from a standard curve referenced to WHO standard serum 89SF. The avidity is calculated based on binding with various concentrations of thiocyanate. This test has not been cleared or approved for diagnostic use by the U.S. Food and Drug Administration.

Turnaround Time

5-7 business days from receipt of specimen

Specimen Information

Specimen Type	Order Code	CPT Code	NY Approved	Volume	Assay Range
serum	401734P	86317	Yes	1 mL (min. 100 uL)	NA

Special Instructions

- Collect 1 mL, ambient, frozen, or refrigerated, so special shipping requirements.

Disclaimer

Specimens are approved for testing in New York only when indicated in the Specimen Information field above.

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References

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