

HOTLINE: Effective December 21, 2020

New Test Available Now 3003477 Membranous Nephropathy Comprehensive Autoantibody Panel

**MNCA PAN** 



Additional Technical Information

Methodology: Semi-Quantitative Indirect Fluorescent Antibody

**Performed:** Tue **Reported:** 1-8 days

Specimen Required: Collect: Serum Separator Tube

Specimen Preparation: Separate serum from cells ASAP or within 2 hours of collection. Transfer 1 mL serum to an ARUP Standard

Transport Tube. (Min: 0.2 mL)

Storage/Transport Temperature: Refrigerated

Unacceptable Conditions: Hemolyzed, hyperlipemic, icteric, heat-treated or contaminated

Stability (collection to initiation of testing): Ambient: 48 hours; Refrigerated: 2 weeks; Frozen: 1 month

## **Reference Interval:**

2011828	Phospholipase A2 Receptor (PLA2R) Antibody, IgG with Reflex to Titer	Less than 1:10
3003480	Antithrombospondin Type-1 Domain-Containing 7A (THSD7A) Antibody, IgG with Reflex to Titer	Less than 1:10

## **Interpretive Data:**

Refer to individual components.

This test was developed and its performance characteristics determined by ARUP Laboratories. It has not been cleared or approved by the US Food and Drug Administration. This test was performed in a CLIA certified laboratory and is intended for clinical purposes.

**Note:** If Phospholipase A2 Receptor Antibody, IgG is positive, then a Phospholipase Receptor A2 Antibody, IgG titer will be added. Additional charges apply. If THSD7A Antibody, IgG is positive, then a THSD7A Antibody, IgG titer will be added. Additional charges apply.

**CPT Code(s):** 86255 x2; if reflexed, add 86256 x2

New York DOH approval pending. Call for status update.

HOTLINE NOTE: Refer to the Test Mix Addendum for interface build information.