Primary membranous glomerulonephritis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Primary membranous glomerulonephritis. ORPHA:97560

A rare glomerular disease, histologically characterized by thickening of the capillary wall, with immune deposits predominantly containing IgG4 and C3 on the sub-epithelial side, and typically manifesting with nephrotic syndrome.