

NPHS2

Nephrotic Syndrome

Mode of Inheritance	<ul style="list-style-type: none">• Autosomal recessive
Renal Phenotype	<ul style="list-style-type: none">• Steroid resistant nephrotic syndrome• Age of onset: Childhood• Typical biopsy findings: MCD, FSGS, DMS
Extra-renal Manifestations	<ul style="list-style-type: none">• Sequelae from nephrotic syndrome:<ul style="list-style-type: none">• Hypoalbuminemia and edema• Hyperlipidemia• Hypothyroidism• Loss of immunoglobulins
Pre-Transplant Management	<ul style="list-style-type: none">• Avoidance of steroid and intensive immunosuppression therapy• Avoidance of renal biopsy
Transplant Considerations	<ul style="list-style-type: none">• Tailor immunosuppression given low risk of recurrence post-transplant
Post-Transplant Management	<ul style="list-style-type: none">• Lower risk of disease recurrence (4.5% vs 28.5%) (Trautmann <i>CJASN</i> 10:592, 2015)