

# NUP93

## Nephrotic Syndrome

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