

## Summary of Nephrotic Syndrome

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<b>Nephrotic Syndrome</b>	<b>Minimal change disease.MCD (known as lipoid nephrosis)</b>	<b>Focal segmental GS Similar to MCD</b>	<b>Membranous glomerulopathy/GN</b>
<b>Occurs in</b>	children	Older children and adult	adult
<b>Character but not specific</b>	Effacement of epithelial cell (podocyte) foot processes	Effacement of epithelial cell (podocyte) foot processes	Deposit of immune complex in subepithelium zone of glomerular capillary (immune complex disease)
<b>LM</b>	<ol style="list-style-type: none"> <li>1) Normal glomeruli.</li> <li>2) Cytoplasm of PCT cells are heavily laden with protein droplet and lipid.</li> <li>3) No tubular atrophy or interstitial fibrosis.</li> </ol>	<ol style="list-style-type: none"> <li>1) Glomeruli of juxtamedullary area are affected.</li> <li>2) Some glomeruli are involved (focal).</li> <li>3) Focal and segmental sclerosis of glomeruli.</li> <li>4) Adhesion and hyalinosis.</li> </ol>	<ol style="list-style-type: none"> <li>1) Thickened of capillary wall.</li> <li>2) Deposit separated by glomerular basement membrane matrix (spikes).</li> <li>3) At progresses there is fibrosis and sclerosis.</li> </ol>
<b>IF</b>	Negative for Ig and C.	Negative, sometimes IgM is positive.	IgG and C3 is positive.
<b>EM</b>	Diffuse/fusion/effacement of epithelial cell for (podocyte) foot processes.	Patchy effacement of epithelial cell (podocyte) for processes.	<ol style="list-style-type: none"> <li>1) The immune complex deposits in the subepithelial space with intervening basement membrane spikes.</li> <li>2) Effacement of epithelial cell podocyte.</li> <li>3) Microvillous formation.</li> </ol>

Name of disease	Diabetic nephropathy	Renal amyloidosis	Lipus nephropathy
<b>LM</b>	1) Diffuse glomerulosclerosis <ol style="list-style-type: none"> <li>a) Thickness of capillary wall</li> <li>b) Diffuse mesangial matrix &amp; cells</li> </ol> 2) Nodular sclerosis (kimmelstiel Wilson nodules) <ol style="list-style-type: none"> <li>a) Nodule formation in mesengium.</li> <li>b) Spherical eosinophil.</li> <li>c) Central cellular area.</li> </ol>	It can: <ol style="list-style-type: none"> <li>1) Primary → idiopathic</li> <li>2) Secondary → associated with:               <ol style="list-style-type: none"> <li>a) Inflammatory disease (rheumatoid arthritis)</li> <li>b) Plasma cell disorder (myeloma)</li> </ol> </li> </ol>	It is show: <ol style="list-style-type: none"> <li>1) Active lesion:               <ol style="list-style-type: none"> <li>a) Endo/extra capillary proliferative (crescent).</li> <li>b) Inflammation.</li> <li>c) Fibrinoid necrosis.</li> <li>d) Subendothelial deposit.</li> </ol> </li> <li>2) Chronic:               <ol style="list-style-type: none"> <li>a) Glomerular sclerosis.</li> <li>b) Tubular atrophy.</li> <li>c) Interstitial fibrosis.</li> </ol> </li> </ol>
<b>EM</b>	<ol style="list-style-type: none"> <li>1) Thickness of GBM.</li> <li>2) Arterial hyaline deposits.</li> </ol>	Criss-crossing non-branching fibrils.	
<b>General Notes</b>	End result id scarred non-functioning kidney.		Immune complex mediated disease. Deposit of anti-DNA in glomeruli → lead to glomerular necrosis.