

Renal

1. GLOMERULAR

R DISEASE:

Clinical Syndromes of Glomerular Disease:

- ➔ Nephritic:
 - Haematuria (macro- or microscopic)
 - Oliguria (sometimes anuria)
 - Hypertension
 - Proteinuria (mild if present)
- ➔ Nephrotic:
 - Proteinuria (often massive) – causes hypoproteinaemic related lipid and lipoprotein changes result in lipid and cholesterol derangements
 - Oedema
- ➔ Renal failure: (see Chemical Pathology lectures/notes)

Nomenclature of Glomerular Injury:

- *Diffuse* – a lesion affecting > 50% of the glomeruli
- *Focal* – a lesion involving < 50% of the glomeruli
- *Global* – affecting the whole glomerulus
- *Segmental* – affecting only part of the glomerulus

Additional terms are used to describe the character of the light microscopic changes within the glomeruli:

- *Proliferative*:
Increased numbers of cells within the glomerulus; due to proliferation of indigenous cells and to recruitment of polymorphs and macrophages from the circulation as a consequence of activation of the complement cascade
- *Membranous change*:
The peripheral loops are thickened due to basement membrane expansion
- *Membrano-proliferative*:
A combination of *Proliferative* and *Membranous change*, often with accentuation of the lobular architecture
- *Crescentic*:
Florid proliferation of cells including macrophages lining Bowman's capsule, often compressing the glomerulus

Primary Glomerulopathies:

Disorders in which the kidney is the only or predominant organ involved

1. Acute diffuse proliferative glomerulonephritis
 - 1.1. Poststreptococcal
 - 1.2. Non-poststreptococcal
2. Rapidly progressive (crescentic) glomerulonephritis
3. Membranous glomerulopathy
4. Focal segmental glomerulosclerosis

Secondary Glomerulopathies:

Systemic diseases with glomerular involvement

1. Systemic lupus erythematosus
2. Diabetes mellitus
3. Amyloidosis
4. Goodpasture syndrome
5. Microscopic polyarteritis/polyangitis

Hereditary Disorders:

1. Alport syndrome
2. Thin basement membrane disease
3. Fabry disease

Mechanisms of Immune-mediated Glomerular Disease:

Both the clinical manifestations and glomerular histologic changes in primary and secondary forms can be altered immune mechanisms:

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