**Nephritic syndrome**

Glomerular diseases presenting with a nephritic syndrome are often *characterized by inflammation in the glomeruli*.

The nephritic patient usually presents with:

Hematuria.

Red cell casts in the urine.

Azotemia.

Oliguria.

Mild to moderate hypertension.

**Proteinuria and edema** but these are not as severe as those in the nephrotic syndrome.

Typically, it is characteristic of **poststreptococcal glomerulonephritis** and **crescentic glomerulonephritis**.

**Poststreptococcal glomerulonephritis**

This glomerular disease is a common disorder worldwide.

It usually appears 1 to 4 weeks after a streptococcal infection of the pharynx or skin.

Poststreptococcal glomerulonephritis occurs most frequently in children 6 to 10 years of age, but adults of any age can also be affected.

***Etiology and pathogenesis.***

Poststreptococcal glomerulonephritis is an immunologically mediated disease.

The latent period between infection and onset of nephritis is compatible with the time required for the production of antibodies and the formation of immune complexes.

Elevated titers of antibodies against streptococcal antigens are present in a great majority of patients. Serum complement levels are low, compatible with activation of the complement system and consumption of complement components.

***Clinical Course:***

In the classic case, a young child abruptly develops malaise, fever, nausea, oliguria, and hematuria (smoky or cola-colored urine) 1 to 2 weeks after recovery from a sore throat.

The patients have red cell casts in the urine, mild proteinuria (usually less than 1 gm/day), periorbital edema, and mild to moderate hypertension.

More than 95% of affected children eventually recover totally with conservative therapy.

***Morphology****.*

***Light microscope****:*

The changes are diffuse, that is, involving all lobules of all glomeruli, the changes include:

(1) Infiltration by leukocytes, neutrophils and monocytes.

(2) Proliferation of endothelial and mesangial cells.

(3) In **severe cases** there is crescent formation.

***Immunofluorescence microscopy***: granular deposits of IgG, IgM, and C3 in the mesangium and along the GBM.

***Electron microscopy*:** discrete, amorphous, electron-dense deposits on the epithelial side of the membrane, often having the appearance of “humps”, subendothelial intramembranous and mesangial deposits may be present.

***Rapidly progressive (crescentic) glomerulonephritis***

Rapidly progressive glomerulonephritis (RPGN) is a syndrome associated with severe glomerular injury and does not denote a specific etiologic form of glomerulonephritis.

It is characterized clinically by rapid and progressive loss of renal function associated with severe oliguria and signs of nephritic syndrome; if untreated, death from renal failure occurs within weeks to months.

*The most common histologic picture is the presence of crescents in most of the glomeruli* (crescentic glomerulonephritis).

**Classification of RPGN:**

RPGN is divided into three groups on the basis of immunological findings:

|  |
| --- |
| **TYPE I (ANTI-GBM ANTIBODY)** |
| |  |  |  | | --- | --- | --- | |  |  | Renal limited | |  |  | Goodpasture syndrome | |
| **TYPE II (IMMUNE COMPLEX)** |
| |  |  |  | | --- | --- | --- | |  |  | Idiopathic | |  |  | Post-infectious glomerulonephritis | |
| **TYPE III (PAUCI-IMMUNE)** |

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|  |

**Nephrotic syndrome**

The manifestations of the nephrotic syndrome include:

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| --- | --- | --- |
|  | **1.** | *Massive proteinuria*, with the daily loss of 3.5 gm or more of protein (less in children) |
|  | **2.** | *Hypoalbuminemia*, with plasma albumin levels less than 3 gm/dL |
|  | **3.** | *Generalized edema* |
|  | **4.** | *Hyperlipidemia and lipiduria* |

The lesion in glomerular capillary walls resulting in *increased permeability to plasma proteins, resulting in proteinuria*.

The proteinuria depletes serum albumin levels resulting in hypoalbuminemia

Generalized edema is the consequence of decreased colloid osmotic pressure of the blood with subsequent accumulation of fluid in the interstitial tissues.

The genesis of the *hyperlipidemia* seem to be due in part to *increased synthesis of lipoproteins in the liver, abnormal transport of circulating lipid particles, and decreased catabolism.*

Nephrotic patients are particularly vulnerable to *infection*, especially staphylococcal and pneumococcal, probably related to loss of immunoglobulins in the urine.

*Thrombotic and thromboembolic complications* are also common in nephrotic syndrome, due in part to loss of endogenous anticoagulants.

**Causes:**

The causes of the *nephrotic syndrome are either glomerular diseases or systemic diseases.*

*Glomerular diseases:*

The most frequent glomerular diseases causing nephrotic syndrome are:

*Minimal-change disease*: is most common in children.

*Membranous glomerulopathy*: is most common in older adults.

*Focal segmental glomerulosclerosis*: occurs at all ages.

*Systemic diseases:*

The most frequent systemic causes of the nephrotic syndrome are:

Diabetes mellitus.

Amyloidosis.

SLE.

Prevalence of causes of nephrotic syndrome

|  | **Prevalence (%)[\*]** | |
| --- | --- | --- |
| **Causes** | ***Children*** | ***Adults*** |
|  | | |
| Membranous glomerulopathy | 5 | 30 |
| Minimal-change disease | 65 | 10 |
| Focal segmental glomerulosclerosis | 10 | 35 |
| Membranoproliferative | 10 | 10 |
| glomerulonephritis[†] | 10 | 15 |
| Other proliferative glomerulonephritides (focal, “pure mesangial,” IgA nephropathy)[†] |

**Membranous Nephropathy**

*Membranous nephropathy is a common cause of the nephrotic syndrome in adults.*

*It is characterized by diffuse thickening of the glomerular capillary wall due to the accumulation of electron-dense deposits along the subepithelial side of the basement membrane*.

**Etiology**

Membranous glomerulopathy is divided into:

Primary idiopathic membranous glomerulopathy in 85% of cases: no associated condition can be identified.

Secondary membranous glomerulopathy in 15% of cases: occurring in association with other systemic diseases and a variety of identifiable etiologic agents.

The most notable such associations in Secondary membranous glomerulopathy as follows:

|  |  |  |
| --- | --- | --- |
|  | **•** | ***Drugs*** : penicillamine, captopril, gold, nonsteroidal anti-inflammatory drugs NSAIDs. |
|  | **•** | ***Underlying malignant tumors***, particularly carcinomas of the lung and colon, and melanoma. |
|  | **•** | ***SLE*.** About 10% to 15% of glomerulonephritis in SLE is of the membranous type. |
|  | **•** | ***Infections*** (chronic hepatitis B, hepatitis C, syphilis, schistosomiasis, malaria). |
|  |  |  |

**Clinical Features.**

This disorder usually begins with the insidious onset of the nephrotic syndrome or, in 15% of patients, with non-nephrotic proteinuria.

Hematuria and mild hypertension are present in 15% to 35% of cases.

The course of the disease is variable but generally indolent.

The proteinuria is nonselective and usually does not respond well to corticosteroid therapy.

Progression is associated with increasing sclerosis of glomeruli, rising serum creatinine reflecting renal insufficiency, and development of hypertension.

About 10% die or progress to renal failure within 10 years.

It is necessary in any patient to first rule out the secondary causes, since treatment of the underlying condition (malignant neoplasm, infection, or SLE) or discontinuance of the offending drug can reverse the injury.

**Morphology.**

**By light microscopy:**

1-The glomeruli either appear normal in the early stages of the disease or exhibit uniform, diffuse thickening of the glomerular capillary wall.

2-As the disease advances glomerular sclerosis may occur; with time glomeruli may become totally sclerosed.

**By electron microscopy**

1-Irregular dense deposits of immune complexes between the basement membrane and the overlying epithelial cells (subepithelial deposit).

2-Basement membrane has irregular spikes.

3-Visceral epithelial cells having effaced foot processes.

**Immunofluorescence microscopy**:

Granular deposits of both immunoglobulins and complement.

**Minimal-Change Disease**

It is the *most frequent cause of nephrotic syndrome in children*, but it is less common in adults.

*It is characterized by diffuse effacement of foot processes of visceral epithelial cells (podocytes) in glomeruli that appear virtually normal by light microscopy*.

**Etiology and Pathogenesis.**

Two hypotheses:

1. Immune dysfunction, eventually resulting in the elaboration of a cytokine that damages visceral epithelial cells and causes proteinuria. *(The disease sometimes follows a respiratory infection or routine prophylactic immunization. Its most characteristic feature is its usually dramatic response to corticosteroid therapy.)*
2. Mutations in several podocytes proteins, including *nephrin and podocin*, discussed in the section on focal glomerulosclerosis below. These structural proteins are localized to the slit diaphragm

In adults, it has been noted, that minimal-change disease can be associated with Hodgkin lymphoma and, less frequently, other lymphomas and leukemias. In addition, secondary minimal-change disease may follow NSAID therapy.

**Clinical Features.**

The peak incidence is between 2 and 6 years of age.

Patients present with massive proteinuria, renal function remains good, and there is commonly no hypertension or hematuria.

The proteinuria usually is highly selective, most of the protein being albumin.

Most children (>90%) with minimal-change disease respond rapidly to corticosteroid therapy.

However, proteinuria may recur, and some patients may become steroid-dependent or resistant.

The long-term prognosis for patients is excellent, and even steroid-dependent disease resolves when children reach puberty. Although adults are slower to respond, their long-term prognosis is also excellent.

**Morphology**.

**Light microscopy:**

The glomeruli are normal.

**Electron microscopy:**

The GBM appears normal, and no electron-dense material is deposited. The principal lesion is in the visceral epithelial cells, which show a uniform and diffuse effacement of foot processes.

Foot process effacement is also present in other proteinuric states (e.g., membranous glomerulopathy, diabetic nephropathy); it is only when effacement is associated with normal glomeruli by light microscopy that the diagnosis of minimal-change disease can be made.

**Immunofluorescent microscopy:**

Show no immunoglobulin or complement deposits.

**Focal Segmental Glomerulosclerosis**

This lesion is characterized by sclerosis of some, but not all, glomeruli (thus, it is focal); and in the affected glomeruli, only a portion of the capillary tuft is involved (thus, it is segmental). Idiopathic focal segmental glomerulosclerosis accounts for as many as 10% (In children) and 35% (In adults) of cases of nephrotic syndrome.

**Classification:**

Focal segmental glomerulosclerosis (FSGS) occurs in the following settings:

|  |  |  |
| --- | --- | --- |
|  | • | primary disease (idiopathic focal segmental glomerulosclerosis) |
|  | • | Secondary :   1. In association with other known conditions, such as HIV infection (HIV-associated nephropathy), heroin addiction (heroin nephropathy), sickle-cell disease, and massive obesity |
|  |  | 1. As a secondary event, in previously active glomerular disease e.g., IgA nephropathy. |
|  |  | 1. As adaptive response to loss of renal tissue in advanced stages of other renal disorders, such as reflux nephropathy, hypertensive nephropathy. |
|  |  |  |

**Clinical features:**

The clinical features of FSGS differ from those of minimal-change disease in the following respects:

(1) There is a higher incidence of hematuria, reduced GFR, and hypertension.

(2) Proteinuria is more often nonselective.

(3) There is poor response to corticosteroid therapy.

(4) There is progression to chronic kidney disease, with at least 50% developing end-stage renal disease within 10 years.

There is little tendency for spontaneous remission in idiopathic FSGS, and responses to corticosteroid therapy are variable. In general, children have a better prognosis than adults do.

**Morphology**:

**Light microscopy:**

The focal and segmental lesions may involve only a minority of the glomeruli and may be missed if the biopsy specimen contains an insufficient number of glomeruli.

Glomeruli that do not show segmental lesions usually appear normal on light microscopy.

**Electron microscopy:**

Both sclerotic and nonsclerotic areas show diffuse effacement of foot processes, focal detachment of the epithelial cells and denudation of the GBM.

**Immunofluorescence microscopy**:

IgM and C3 may be present in the sclerotic areas and/or in the mesangium.

Glomerular lesions associated with systemic diseases

**Diabetic Nephropathy**

Diabetes mellitus is a major cause of renal morbidity and mortality, and diabetic nephropathy is one of the leading causes of chronic kidney failure

Advanced or end-stage kidney disease occurs in as many as 40% of both insulin-dependent type 1 diabetics and type 2 diabetics.

By far the most common lesions involve the glomeruli and are associated clinically with three glomerular syndromes: non-nephrotic proteinuria, nephrotic syndrome, and chronic renal failure.

**Pathogenesis:**

The pathogenesis of diabetic glomerulosclerosis involved:

|  |  |  |
| --- | --- | --- |
|  | **•** | *The metabolic defect*s, are responsible for biochemical alterations in the GBM. |
|  | **•** | *Non enzymatic glycosylation* of glomerular proteins. |
|  | **•** | *Hemodynamic changes* involved in the initiation and progression of diabetic glomerulosclerosis. |

To sum up, two processes seem to play a role in the fully developed diabetic glomerular lesions:

1. A metabolic defect, probably linked to advanced glycosylation end products, that accounts for the thickened GBM and increased mesangial matrix
2. Hemodynamic effects, associated with glomerular hypertrophy, which also contribute to the development of glomerulosclerosis.

**Morphology**

The morphologic changes in the glomeruli include:

(1) Capillary basement membrane thickening.

(2) Diffuse mesangial sclerosis.

(3) Nodular glomerulosclerosis.

The morphologic manifestations of diabetic nephropathy are identical in type 1 and type 2 diabetes.