NURS 821 Alterations in Reproduction; Alterations in Elimination Lecture 10 Part 4 Intrarenal Causes of Renal Failure Renal Failure Classification ■ Intra-renal - those resulting from actual renal pathology - may also result from an unresolved prerenal or post-renal condition. - Cortical involvement may be vascular, infectious, or immunologic in origin. Causes of Renal Failure cont'd • Intra-Renal Amyloidosis - Pyelonephritis - Nephrocalcinosis - Acute glomerulonephritis Gout Rapidly progressive glomerulonephritis Hereditary nephropathy - Polycystic kidneys - Membranous Renal hypoplasia glomerulonephritis - Drugs - Goodpasture's syndrome Heavy metals - Systemic lupus Industrial solvents erythmatosis - Acute Tubular necrosis - Polyarteritis nodosa - Bilateral cortical necrosis - Scleroderma

- Sickle cell disease

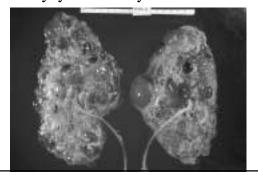
- Radiation Nephritis

- Diabetes mellitus

Polycystic Kidney Disease (PKD)

- Definition: Genetic disease in which numerous fluid-filled cysts fill the kidneys; affects other organs also
- Incidence: 500,000
- 4th leading cause of ESRD
- Affects all races, M=F
- Infantile form-rare, but rapidly progresses
- May have slight increase in women and Whites (NIDDK, 2000)

Polycystic 1	Kidney I	Disease
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PKD Pathophysiology

- Large fluid-filled cysts grow from nephrons
- Separate and enlarge
- Encroach upon total kidney mass and function-decreasing GFR
- Thousands may develop
- Kidney may weigh up to 22 lbs.!
- HTN usually develops before cysts appear (NIDDK, 2000)

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Calcium-Phosphate Metabolism

- Dietary intake is approximately 1 gm/day
- Kidney excretes 200 mg Ca and 400 mg PO₄ daily for homeostasis
- Hypocalcemia-
 - Parathyroid secretes parathormone causing renal excretion of PO₄
 - Bone release of calcium

PKD Forms

- Autosomal dominant
 - More common form
 - 90%
 - Symptomatic between age 30-40
- Autosomal recessive
 - Rare
 - May develop in womb or early infancy

PKD

- May have multi-organ system problemscysts in liver, pancreas; diverticula; aneurysms in brain; abnormal cardiac valves
- Associated renal problems-kidney stones, hematuria, UTI, HTN
- DX with ultrasound

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Hallmark Manifestations of Glomerular Disease

- Proteinuria
- Proteinemia
- Hematuria
- Edema-may be marked, including anasarca

Nephrotic Syndrome

- Vicious Cycle of Edema and Anasarca
 - Low oncotic pressure
 - Aldosterone secreted by adrenal cortex
 - Na reabsorbed
 - Increased plasma hydrostatic pressure
 - Inadequated serum oncotic pressure
 - edema

Glomerulonephritis

- Inflammation of membrane in kidney which filters wastes and fluid from blood
- Etiology
 - Primary-immune responses
 - Strept, staph, hepatitis
 - Secondary to systemic diseases
 - Autoimmune disorders such as SLE-lupus nephritis
 - Goodpasture's syndrome
 - Post-partum
 - (NIDDK, 2000)

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Glomerulonephritis

- Etiology
 - Acute post-streptococcal glomerulonephritis (PSGN)
 - Bacterial endocarditis
 - HIV infection

(NIDDK, 2000)

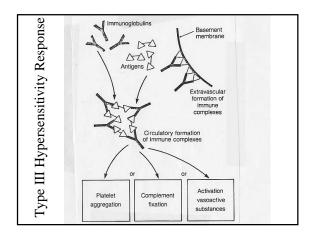


Acute Post-streptococcal Glomerulonephritis

- Etiology-streptococcal infection-throat or impetigo (rare)
- Increased incidence in boys, age 3-7
- One of most common causes of Acute Renal Failure (ARF)
 - occurs 2-3 weeks post-infection
- many have no long-term sequelae (NIDDK, 2000)

Acute Post-streptococcal Glomerulonephritis

- Pathophysiology-
 - Type III hypersensitivity response
 - Results in attack of glomeruli



HIV Glomerulonephritis

- Incidence-5-10% of PWA develop prior to full blown AIDS
- Manifestations-starts with heavy proteinuria and rapidly progresses to ESRD

(NIDDK, 2000)

Glomerulosclerosis

■ Definition-Scarring or hardening of the blood vessels in the kidneys

Glomerulosclerosis

- **Definition**-Scarring or hardening of the blood vessels in the kidneys
- Etiology
 - DM-leading cause of diabetic nephropathy
 - Scarring of kidney
 - Increases speed of processing of blood due to glycemia
 - Unknown-growth factors released by glomerulus or circulating to the glomerulus from elsewhere

 - (NIDDK, 2000)

Pyelonephritis

- Definition: Unilateral or bilateral infection of renal pelvis and interstitium
- Etiology: usually bacterial in origin, ascending from urethra
- Incidence: F>M
- Pathophysiology: Infection and inflammation ascend and worsen prognosis from renal medulla to cortex to necrosis of renal papilla
 - Glomerulus unaffected
 - Renal tubules usually affected

Pyelonephritis

- Pathophysiology:
 - Resolution of acute phase
 - Scarring
 - Tubular atrophy

Manifestations

- Fever and chills
- Flank pain, costovertebral tenderness
- Frequency, dysuria

Chronic Pyelonephritis (Interstitial Nephritis)

- Etiology:
 - Recurrent acute pyelonephritis
 - Ischemia
 - Radiation
 - Immune complex diseases
 - Drugs=aspirin, acetaminophen, phenacetin

Chronic Pyelonephritis

Pathophysiology

Initially, inflammation and fibrosis occurs in interstitium between tubules

- Gradual tubular atrophy, dilation, destruction and
- May affect renal concentrating ability

Manifestations

- Initially, none
 HTN
- Frequency, dysuria, flank pain

Pyelonephritis



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IgA Nephropathy

- Etiology-Deposition of IgA in glomeruliorigin unknown
- Increased incidence in males
- NIDDK studying impact of fish oil capsules and steroids

(NIDDK, 2000)

Goodpasture's Syndrome

- Etiology-
 - Unknown
 - Autoantibodies attack kidneys and lungs
- Lungs not as affected
- Increased incidence in young men (NIDDK, 2000)

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