



RENAL PATHOLOGY  
**INFECTION OF THE UPPER & LOWER URINARY  
TRACT**

April 2020

**Reference: Robbins & Cotran Pathology and Rubin's  
Pathology**

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# Objectives

## **Objectives:**

At the end of the two lectures the students will be able to:

- Recognize the predisposing factors for infections of the kidney and urinary tract.
- Describe the different types of infections in the kidney and urinary tract.
- Recognize acute and chronic pyelonephritis.
- Describe the causes of urinary tract obstruction.
- Recognize drug induced nephritis.

## **Key Outlines:**

- Urinary Tract Obstruction: causes and clinical manifestations in children and adults.
- Infections of the Urinary Tract: Predisposing Factors and Clinical Manifestations.
- Pathology of Acute and Chronic Pyelonephritis including causes and complications of urolithiasis.
- Drug induced interstitial nephritis and renal necrosis.

# Lecture outline for 2 lectures

- Upper urinary tract infection or inflammation
  - Tubulointerstitial nephritis
    - Acute pyelonephritis
    - Chronic pyelonephritis
    - Drug induced tubulointerstitial nephritis
- Urinary tract outflow obstruction
  - Causes
  - Urolithiasis
  - Hydronephrosis
- Lower urinary tract infection
  - Ureteritis
  - Cystitis

# Tubulointerstitial nephritis

Tubulointerstitial nephritis is inflammatory disease primarily involving the renal tubules and interstitium. It can be in the form of:

- Acute pyelonephritis
- Chronic pyelonephritis
- Drug induced tubulointerstitial nephritis



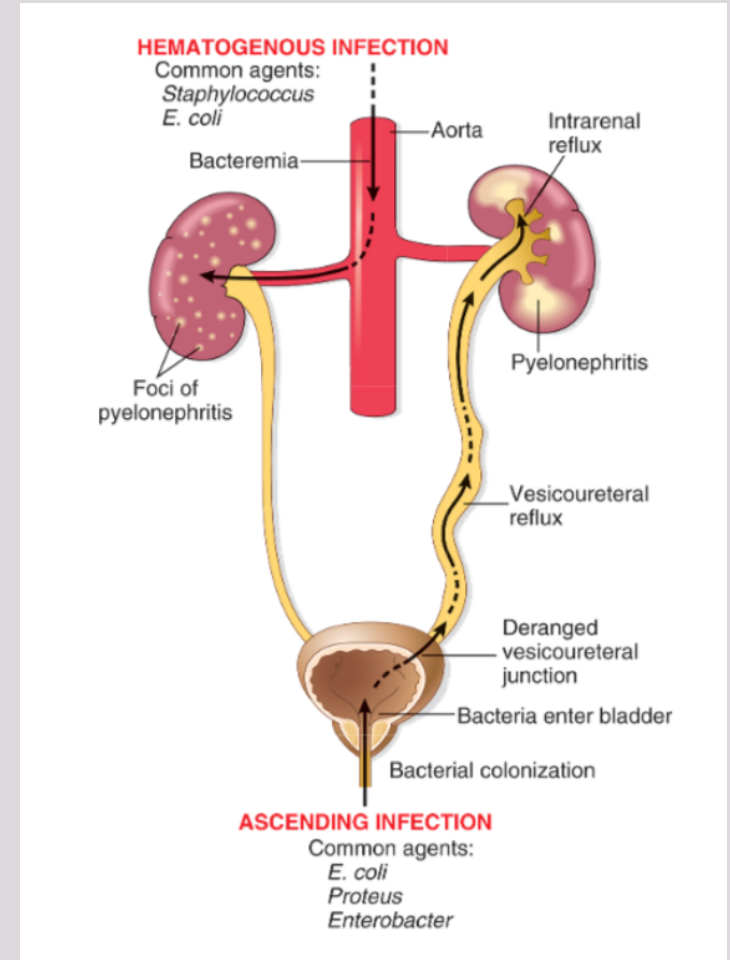
# **ACUTE PYELONEPHRITIS**

# Acute Pyelonephritis

- Acute pyelonephritis is an acute suppurative inflammation of the upper urinary tract (kidney & renal pelvis) caused by bacterial infection. It typically follows infection of the lower urinary tract.
- The most common causative organism is enteric gram-negative rods *Escherichia coli*. Other organisms include *Proteus*, *Klebsiella*, *Enterobacter* and *Pseudomonas*.

There are two routes by which bacteria can reach the kidneys:

1. **Ascending infection (more common) from the lower urinary tract:** bacteria ascends from the urethra into the urinary bladder and up the ureters to the kidneys.
2. **Through the bloodstream/hematogenous** (less common) e.g. from an infected heart valve in endocarditis, miliary tuberculosis etc.



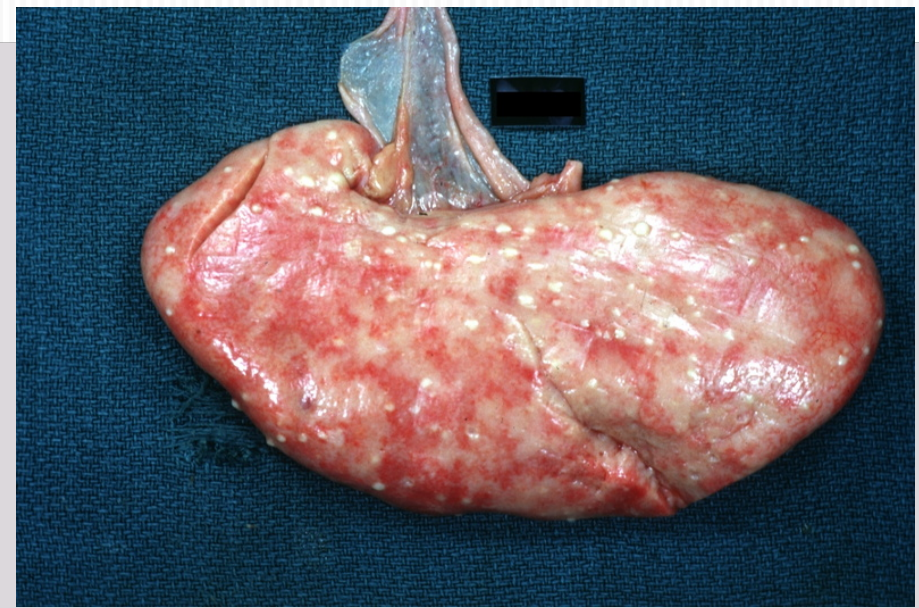
# Acute Pyelonephritis: predisposing factors

1. Urethral **instrumentation** e.g. catheterization and cystoscopy.
2. **Obstruction:** obstruction at the level of the urinary bladder (by stones, benign prostatic hypertrophy etc.)  
→ incomplete emptying of bladder and increased residual urine → stasis of urine in bladder → allows bacteria to multiply → bacteria ascend up the ureters and infect the kidneys
3. **Incompetence of the vesicoureteral orifice** → allows reflux of bladder urine to ascend up into the ureter (normally ureteral insertion into the bladder is a competent one-way valve that prevents retrograde flow of urine). About 1/3<sup>rd</sup> of children with UTI have vesicoureteral reflux (due to a congenital defect of the valve).
4. **Gender:** UTI most commonly affects females (anatomically predisposed because of the close proximity of the urethra to the rectum and the short urethra).
5. **Pregnancy:** Asymptomatic bacteriuria occurs in 10% of pregnant women, out of which some develop acute pyelonephritis.
6. **Preexisting renal lesions**
7. **Diabetes mellitus:** diabetic glycosuria predisposes to infection by providing a rich medium for bacterial growth. Diabetics also have increased risk of complications of pyelonephritis e.g. septicemia, necrotizing papillitis and recurrence of infection.
8. **Immunosuppression & immunodeficiency**

# Acute Pyelonephritis: morphology

## Macroscopic

- The kidney may be enlarged and swollen and show small yellow/ white subcapsular and cortical microabscesses (pus = suppurative necrosis).
- Rarely the kidney may become filled with large amounts of pus in the renal pelvis, calyces and ureter called pyonephrosis.



Pathology Education Informational Resource (PEIR) Digital Library  
[http://peir.path.uab.edu/library/\\_data/i/upload/2013/08/01/20130801100330-254c35a8-me.jpg](http://peir.path.uab.edu/library/_data/i/upload/2013/08/01/20130801100330-254c35a8-me.jpg)



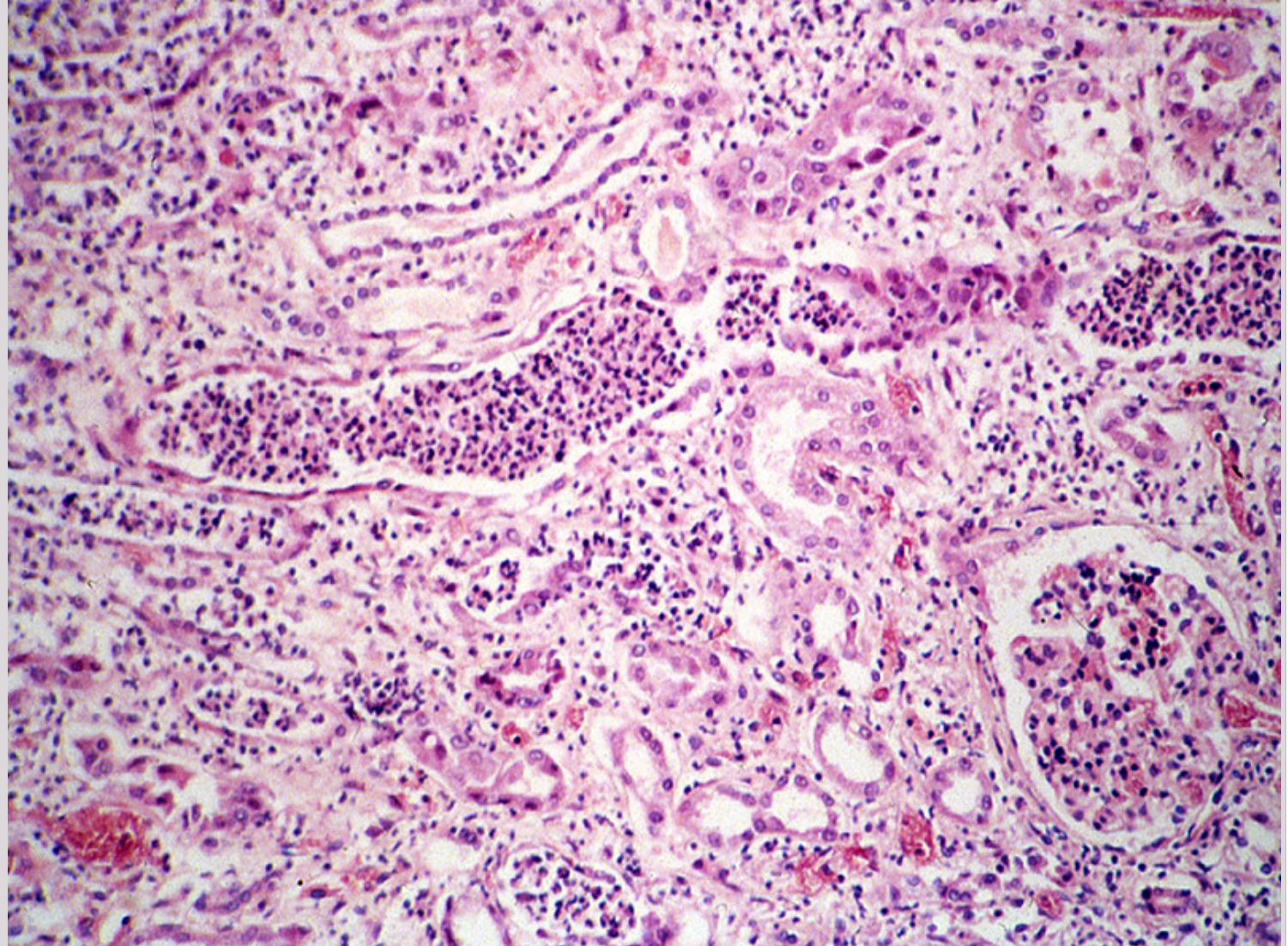
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# Acute Pyelonephritis: morphology

## Microscopic

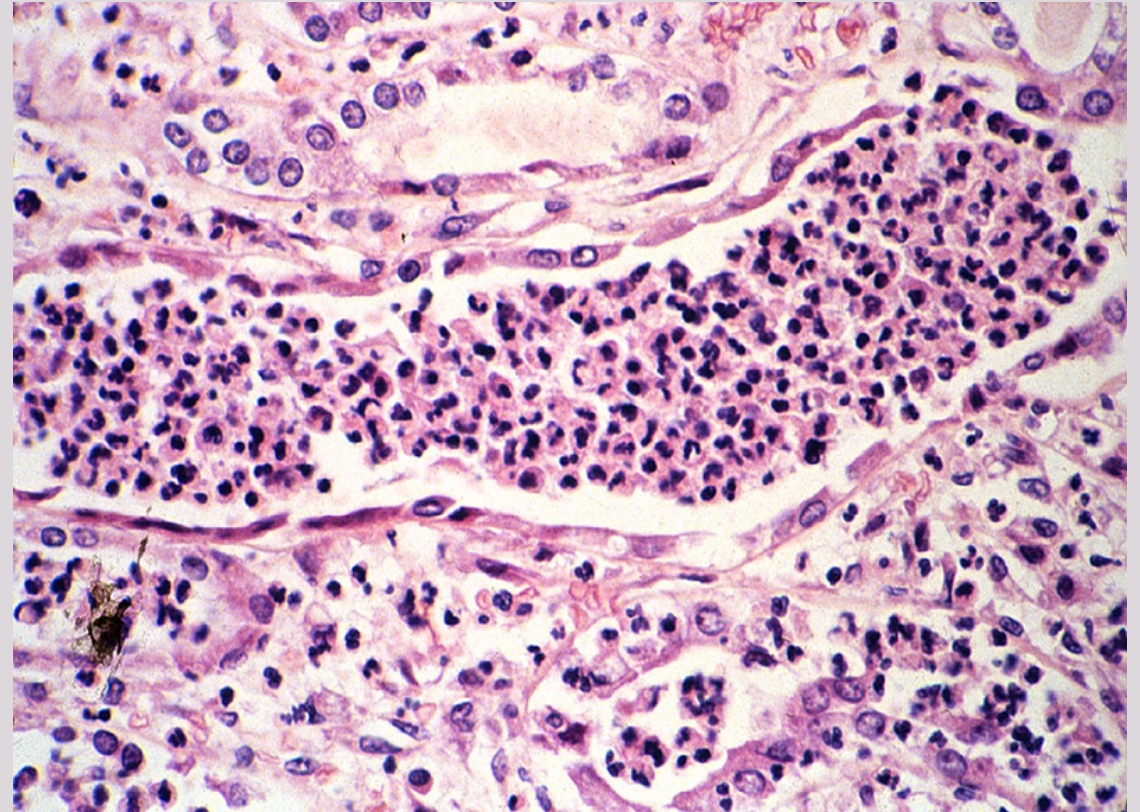
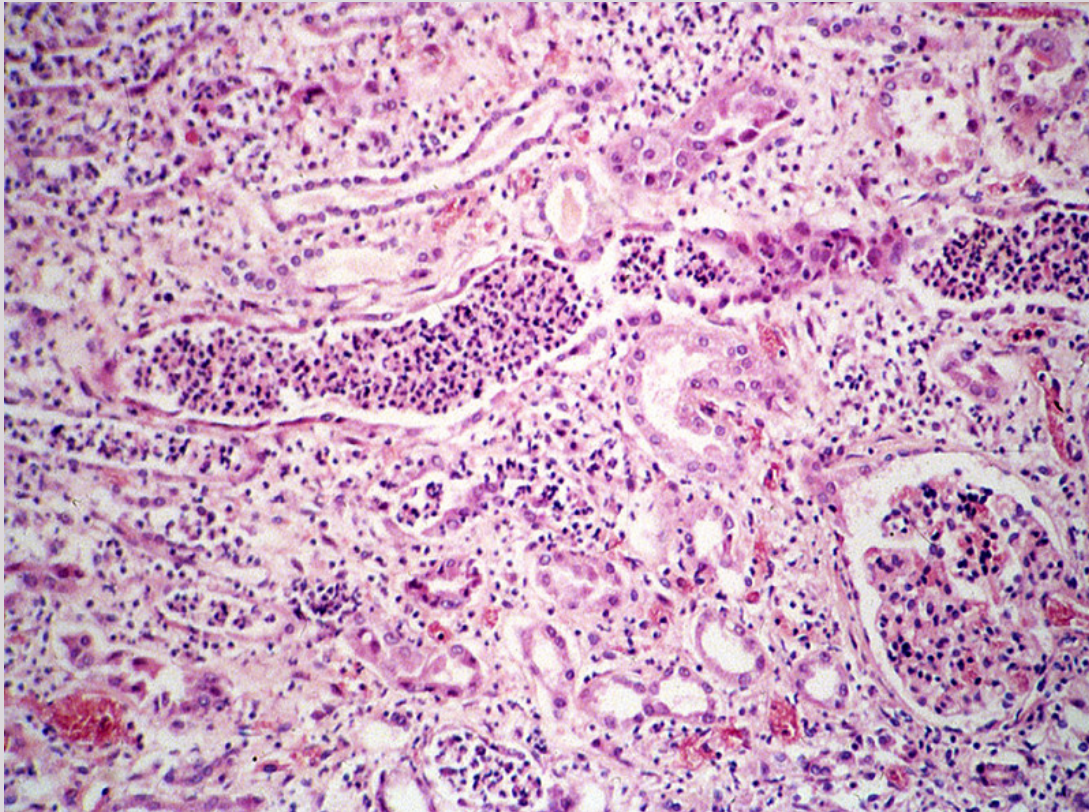
- There is a dense acute tubulointerstitial inflammation (neutrophils) with tubular destruction. The neutrophils fill the tubules and collecting ducts.
- The vessels and glomeruli often are preserved.
- In severe cases → perinephric abscess.



# Acute Pyelonephritis: morphology

## Microscopic

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# Acute Pyelonephritis:

## Clinical features

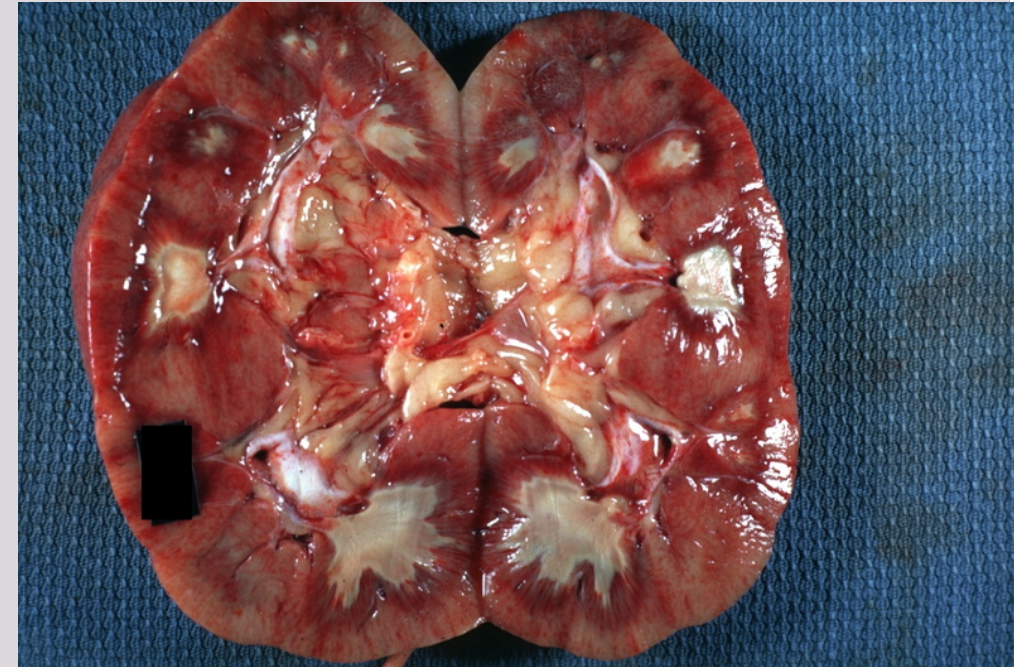
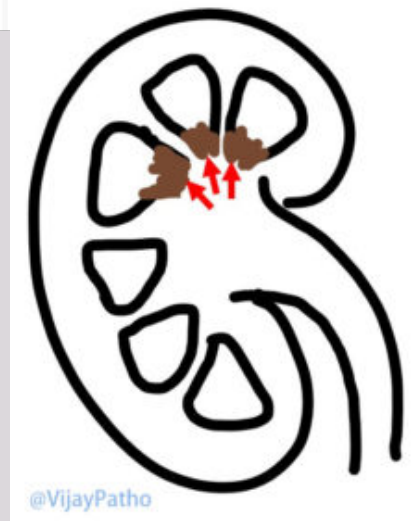
- Fever with chills and sweats.
- Flank pain with costovertebral angle tenderness.
- Dysuria, frequency and urgency.
- Pyuria, hematuria.
- Leukocytosis with neutrophilia.
- Positive urine culture and wbc casts in the urine.
- Differentiating upper from lower urinary tract infection is often clinically difficult.

## Complications

- Papillary necrosis
- Pyonephrosis
- Perinephric abscess
- Chronic pyelonephritis
- Septicemia

## Papillary necrosis (necrotizing papillitis)

- It is a type of pyelonephritis characterized by necrosis of the renal papillae (apex of renal pyramids).
- It is seen in
  - a) diabetics with acute pyelonephritis
  - b) acute pyelonephritis with urinary tract obstruction.
  - c) analgesic abuse associated interstitial nephritis
  - d) Chronic liver disease
  - e) Renal transplant rejection
  - f) Infections e.g. tuberculosis
  - g) Sickle cell disease
- Grossly: yellow white suppurative necrosis (pus) at the tips of renal papillae/ pyramids.
- Microscopically: coagulative necrosis & microabscess at the tips of renal papillae.





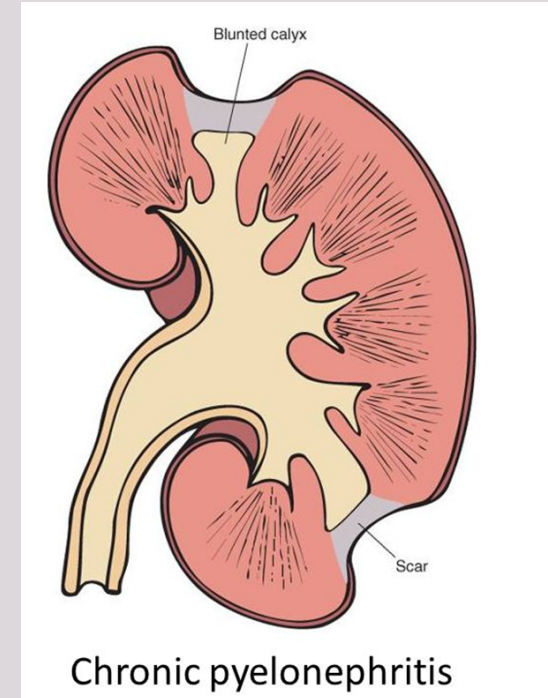
# **CHRONIC PYELONEPHRITIS**

# Chronic Pyelonephritis

- It is a chronic tubulointerstitial inflammation of the kidney caused by repeated bouts of inflammation and healing → resulting in scarring of the involved kidney with deformed renal pelvis & calyces and gradual renal insufficiency.
- It is caused by:
  - Chronic urinary obstruction (e.g. obstruction of the ureter by calculi (stones), tumor within the ureter, or extrinsic compression etc.)
  - Chronic reflux.

## Macroscopy

- The kidneys are small and contracted.
- The surface of the kidney is irregularly scarred with areas of depression.
- The cortex is thinned out and there is deformity and blunting of the pelvicalyceal system.



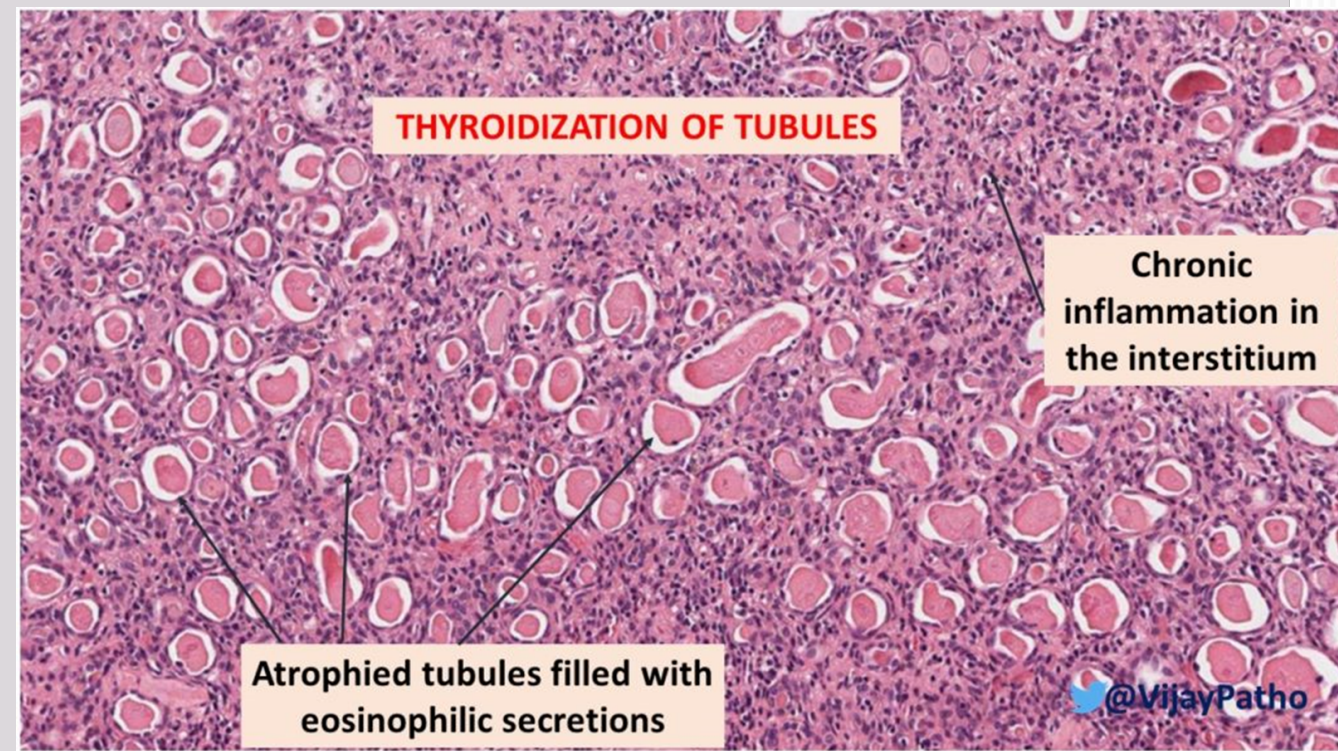
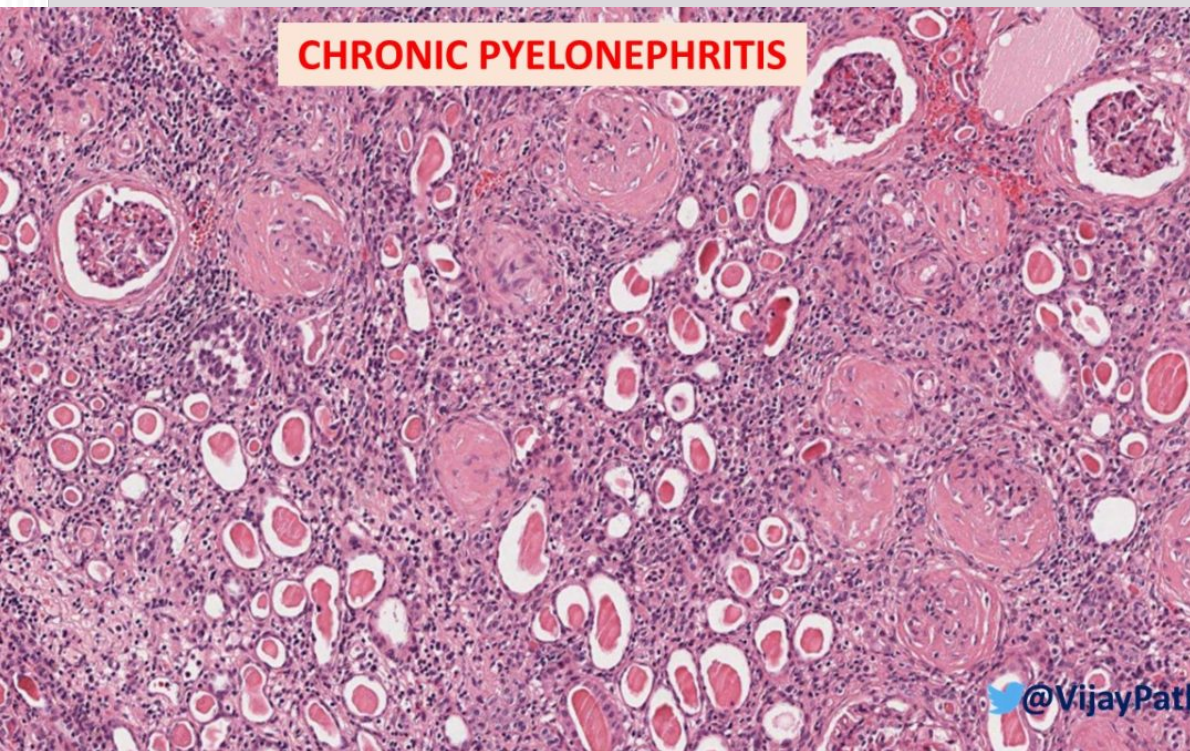
Robbins basic pathology, 9<sup>th</sup> Ed Elsevier 2013

# Chronic Pyelonephritis:

## Microscopy

- Tubules → tubular atrophy with thyroidization of tubules (tubules are filled with eosinophilic hyaline casts resembling colloid of thyroid gland).
- Interstitium → interstitial fibrosis and chronic interstitial inflammation (lymphocytes and plasma cells)
- Glomeruli → periglomerular fibrosis and glomerulosclerosis

Chronic pyelonephritis can ultimately lead to renal failure (end stage renal disease)



# Chronic Pyelonephritis: clinical features

- Most patients have episodic symptoms of urinary tract infection or acute pyelonephritis.
- Hypertension.
- Some patients have a silent course until end-stage renal disease develops.
- Imaging studies show deformed pelvicalyceal system and cortical scarring.





**DRUG INDUCED TUBULOINTERSTITIAL  
NEPHRITIS**

# Drug induced tubulointerstitial nephritis

- Drugs are an important cause of renal injury.
- Drug-induced interstitial nephritis is an IgE (hypersensitivity) and T cell–mediated immune reaction to a drug. It is characterized by interstitial inflammation with many eosinophils.
- It can be acute or chronic.
- Implicated therapeutic drugs: penicillins (e.g. ampicillin), rifampicin, diuretics (thiazides), nonsteroidal anti-inflammatory agents, etc.

## Microscopy

- Interstitial infiltration by chronic inflammatory cells (lymphocytes and macrophages), typically with increased eosinophils.
- Interstitial non-necrotizing granulomas with multinucleated giant cells +/-.
- Tubular atrophy +/-.
- The glomeruli are usually normal.

## Clinical features

- Abnormal renal function test after few days or weeks after exposure to the drug.
- Urine: hematuria and eosinophils.
- Can present as acute kidney injury (rising serum creatinine and oliguria).
- It is important to diagnose this condition, because if remove the offending drug on time the injury maybe reversible.



# URINARY TRACT OBSTRUCTION (URINARY OUTFLOW OBSTRUCTION)

# Obstruction in the outflow of urine

## Causes of urinary tract outflow obstruction

- Congenital anomalies
- Urolithiasis
- Benign prostate hyperplasia
- Tumors (of prostate, cervix, urinary bladder etc.)
- Inflammations (prostatitis, urethritis etc.)
- Pregnancy
- Others (paralysis of urinary bladder)



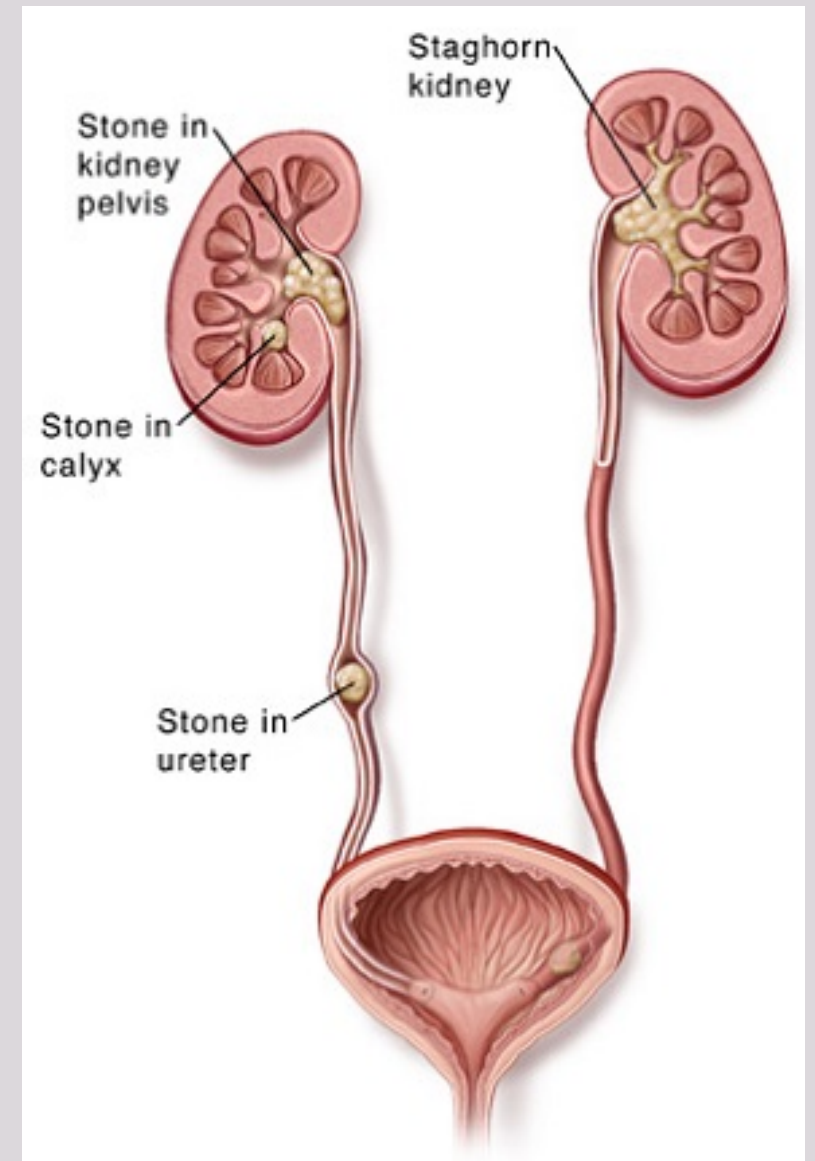
# UROLITHIASIS

# Urolithiasis/nephrolithiasis

- **Urolithiasis** is formation of urinary calculi/stone along the urinary system e.g. kidney, bladder, ureters etc.
- Renal pelvis and calyces are common sites for calculi formation (called nephrolithiasis).

## Predisposing factors:

- Metabolic factors: hypercalciuria, hyperphosphaturia, oxaluria, gout etc.
- Persistently alkaline urine favors formation of calcium phosphate stones.
- Persistently acidic urine favors formation of oxalate or uric acid stones.
- Stasis of urine facilitates precipitation of salts and stone formation.
- Chronic dehydration: concentrates urine and favors stone formation
- For unknown reasons, renal stones are more common in men than in women.
- There may be a familial tendency toward stone formation.



Sectioned Kidneys, Ureters, and Bladder showing location of Kidney Stones. SOURCE: pickup from 5A11872  
Referenced from [www@  
http://graphics8.nytimes.com/images/2007/08/01/health/adam/17091.jpg](http://graphics8.nytimes.com/images/2007/08/01/health/adam/17091.jpg)  
Anatomy text:  
Clemente Anatomy, 4th Ed. Carmine D. Clemente. Lippincott Williams & Wilkins. plate 233 figure 356

# Urolithiasis

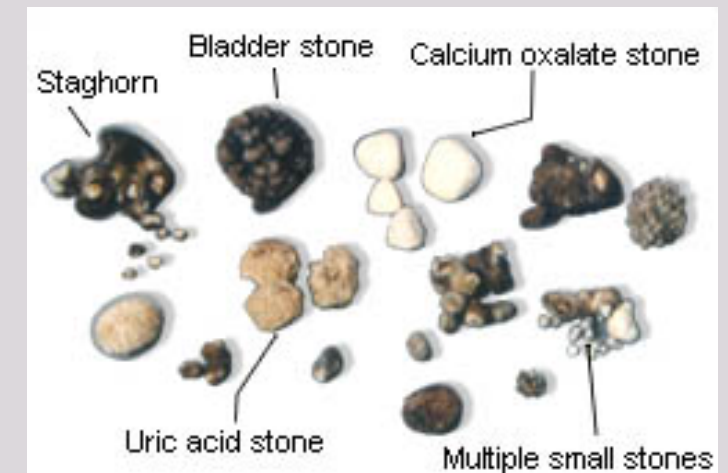
## Morphology

Stones vary in composition.

Types of stones seen are:

1. **Calcium stones:** 75% of kidney stones are calcium oxalate or phosphate (radio-opaque).
2. **Infection stones:** 10% of stones are caused by infection → infection results in alkaline urine → leads to formation of **magnesium ammonium phosphate stones**. Infection stones can occasionally fill the pelvis and calyces to form **large staghorn calculi**.
3. **Uric acid stones:** are formed in acidic pH. Patients with hyperuricemia and gout are predisposed to uric acid stones, but it can also be seen in people with no hyperuricemia or gout. Pure uric acid stones are radiolucent.
4. **Cystine stones:** are uncommon.

- Majority (80%) of the stones are unilateral
- Common sites: are renal pelvis and calyces and urinary bladder
- Commonly multiple stones are found in one kidney
- Stones vary in size from few mm to large stones that dilate the entire renal pelvis.
- They range from hard to soft, from smooth to rough
- The staghorn calculi take the shape of the pelvicalyceal system. They are large stones are usually composed of magnesium ammonium phosphate



# Urolithiasis

## Clinical features

- May be asymptomatic esp. stones lodged in the renal pelvis
- Stone can erode the mucosa (ulceration) → hematuria.
- Smaller stones can pass into the ureter, where they cause obstruction → intense episodes of flank pain radiating toward the groin known as renal or ureteral colic. They may pass out in urine → painful.
- Stones obstruct urine flow → predispose to bacterial infection

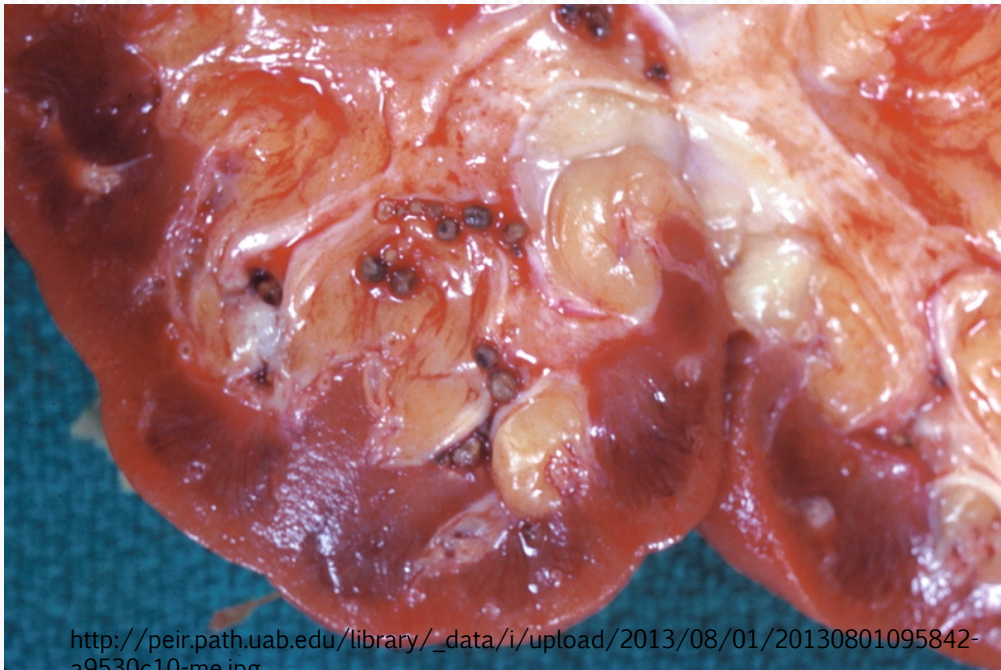
## Complications

- Recurrent infection
- Hydronephrosis
- Pyelonephritis
- Pyonephrosis
- Acute urinary retention
- Renal failure.

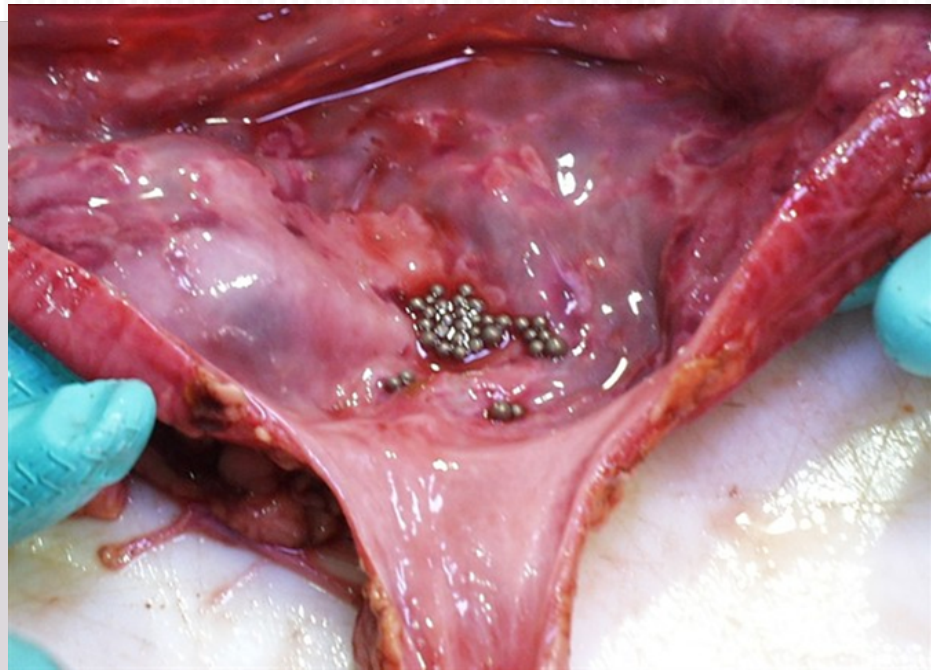
## Diagnosis and management

- Majority, the diagnosis → made radiologically.
- In the past most kidney stones required surgical removal, but now ultrasonic disintegration (lithotripsy) and endoscopic removal are now effective.





[http://peir.path.uab.edu/library/\\_data/i/upload/2013/08/01/20130801095842-a9530c10-me.jpg](http://peir.path.uab.edu/library/_data/i/upload/2013/08/01/20130801095842-a9530c10-me.jpg)



**Staghorn calculi**

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**Staghorn calculi**

Qiao's Pathology: Kidney - Staghorn Calculus  
[https://www.flickr.com/photos/jian-hua\\_qiao\\_md/5648360864/in/album-72157622371115615/](https://www.flickr.com/photos/jian-hua_qiao_md/5648360864/in/album-72157622371115615/)



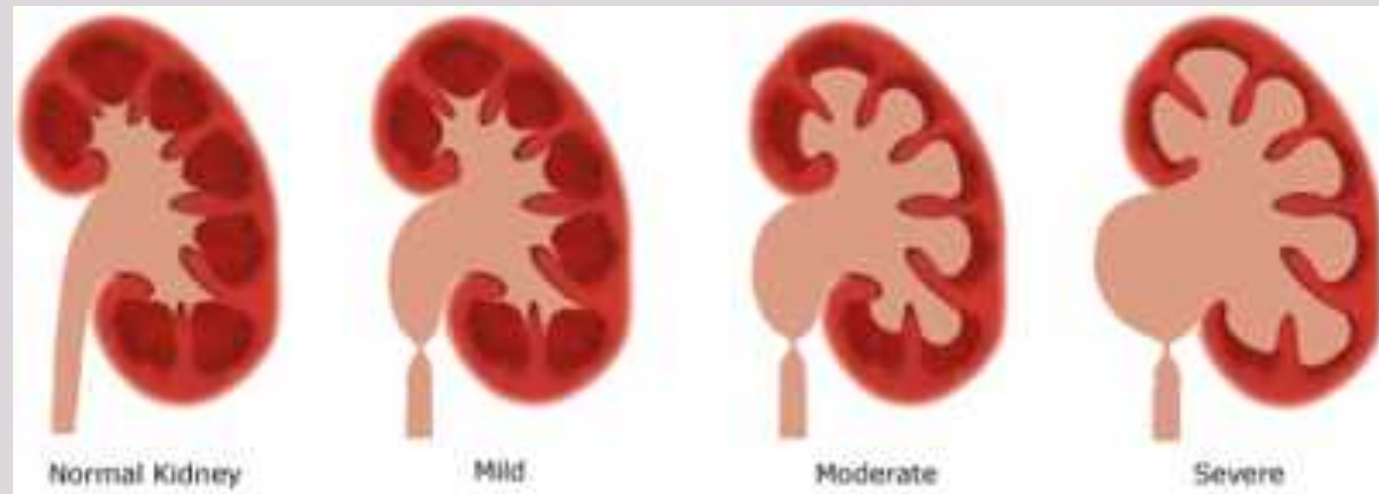
# **HYDRONEPHROSIS**

# Hydronephrosis

- It is the dilation of the renal pelvicalyceal system with resultant renal parenchymal atrophy.
- It is caused by (or is a complication of) the obstruction to the outflow of urine.
- The obstruction can be acute or chronic and be at any level of the urinary tract.
- Obstruction below the level of ureters causes bilateral hydronephrosis.
- The obstruction can be:
  - a) Congenital: e.g. atresia of urethra.
  - b) Acquired:
    - Calculi
    - Benign prostatic hyperplasia
    - Tumors: of prostate, bladder, cervix tumors etc.
    - Inflammation: Prostatitis, ureteritis, urethritis
    - Neurogenic: Spinal cord damage with bladder paralysis
    - Pregnancy

# Hydronephrosis: pathogenesis

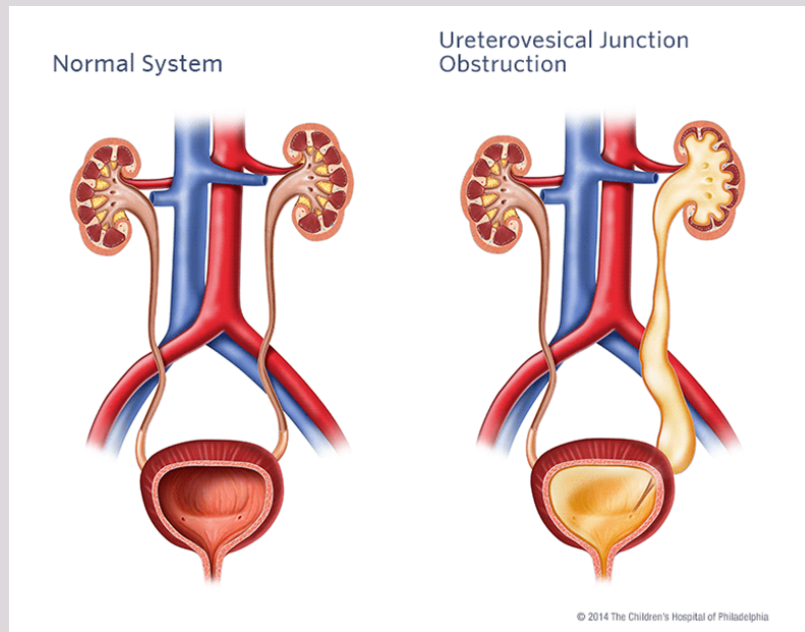
- There is backflow of the urine into the kidney → the calyces and pelvis become markedly dilated → back flow in collecting ducts of kidney → ultimately back flow in renal cortex → renal cortical atrophy.
- The obstruction also triggers an interstitial inflammatory reaction leading eventually to tubular atrophy and interstitial fibrosis.
- Initially the glomeruli are spared but eventually they become sclerotic.



# Hydronephrosis:

## Gross morphology:

- Initially enlarged kidney due to **dilation of the renal pelvis and calyces**
- Followed by atrophy or compression of the renal parenchyma.
- Depending on the level of the obstruction, one or both ureters may also be dilated (hydroureter)



# Hydronephrosis: clinical features

- Unilateral hydronephrosis may be silent/ asymptomatic for long periods
- Bilateral hydronephrosis can lead to oliguria, anuria and renal failure
- In hydronephrosis, the kidney is more susceptible to pyelonephritis, which causes additional injury
- With time the changes become irreversible
- Early diagnosis and timely removal of obstruction within a few weeks usually permits full return of function



# **INFECTIONS OF THE LOWER URINARY TRACT**

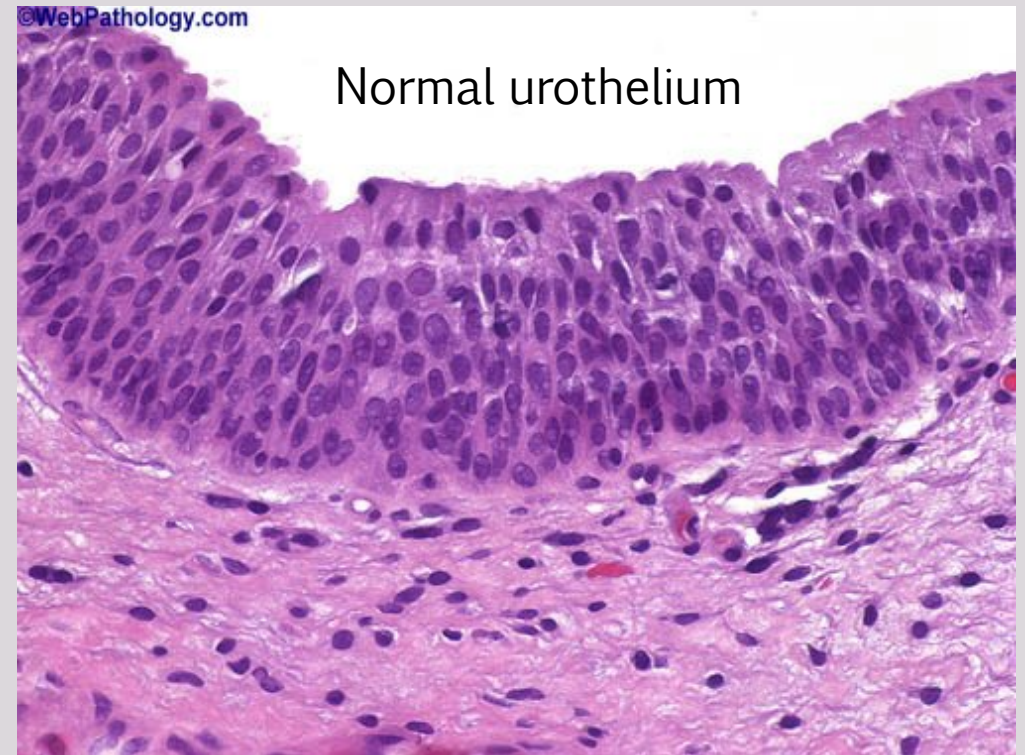
# Lower urinary tract

- Lower urinary tract = the ureters + urinary bladder + urethra.
- The ureters, urinary bladder and urethra are lined by transitional epithelium (urothelium) except the terminal urethra which is lined by squamous epithelium.

## Inflammation in lower urinary tract:

- Ureteritis
- Cystitis
- Urethritis

Infection of the lower urinary tract is referred to as UTI (urinary tract infection).





# Pathology

## Ureteritis

- Ureteritis is an inflammation of the ureters. It is a complication of descending or ascending infections; ureteritis is often associated with ureteral obstruction (e.g. calculi or extrinsic compression of ureter by an adjacent tumor/lymph node or pregnant uterus).

## Cystitis

- It is inflammation of the urinary bladder. It may be acute or chronic.
- It is the most common site of urinary tract infection. Bacterial cystitis is most common form of cystitis. It is caused mainly by coliform bacteria e.g. E coli, Proteus vulgaris, Pseudomonas and Enterobacter spp.
- Predisposing factors: bladder calculi, bladder outlet obstruction (e.g. prostatic hyperplasia), diabetes mellitus, immunodeficiency, radiation therapy, and chemotherapy, prior instrumentation or catheterization (often seen as a nosocomial infection in hospitalized patients, common in patients with indwelling catheters for prolonged periods).
- The risk of cystitis is increased in females because of a short urethra, especially during pregnancy.

# Cystitis: pathology

- Acute cystitis : edema, hemorrhage and a neutrophilic infiltrate.
- Chronic cystitis: a predominance of lymphocytes and fibrosis. Occasionally there are numerous lymphoid follicles → called follicular cystitis or dense infiltrates of eosinophils → eosinophilic cystitis.

## Special forms of cystitis:

- **Tuberculous cystitis:** there is granulomatous cystitis with or without necrosis.
- **Hemorrhagic cystitis:** shows mucosal hemorrhages. Is often seen in acute bacterial infection, adenovirus infection, bleeding diatheses (e.g., leukemia, treatment with cytotoxic drugs and disseminated intravascular coagulation).
- **Chronic interstitial cystitis (chronic pelvic pain syndrome):** is a persistent, painful form of chronic cystitis typically affects middle-aged women. It is characterized by intermittent, suprapubic pain, urinary frequency, urgency, hematuria and dysuria without evidence of bacterial infection; cause is unknown; urine cultures are usually negative. The disease is refractory to all forms of therapy. Eventually → bladder fibrosis → contracted bladder.

# Cystitis: pathology

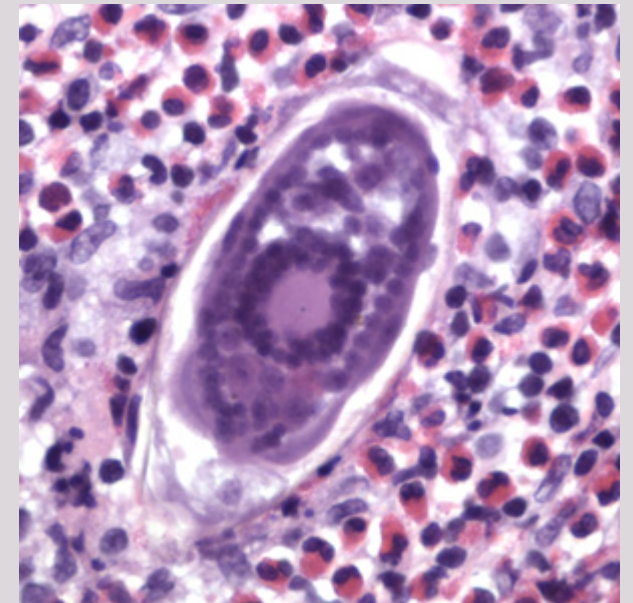
- Acute cystitis : edema, hemorrhage and a neutrophilic infiltrate.
- Chronic cystitis: lymphocytic, histiocytic and plasma cell infiltrate and fibrosis.

## Special forms of cystitis:

- › **Follicular cystitis:** chronic inflammation with many lymphoid follicles
- › **Eosinophilic cystitis:** inflammation with prominent eosinophilic infiltrate.
- › **Tuberculous cystitis:** shows granulomatous cystitis with or without necrosis.
- › **Hemorrhagic cystitis:** shows mucosal hemorrhages. Seen in acute bacterial infection, adenovirus infection and bleeding diatheses (e.g., leukemia, treatment with cytotoxic drugs and disseminated intravascular coagulation).
- › **Chronic interstitial cystitis** (chronic pelvic pain syndrome): persistent, painful chronic cystitis typically affects middle-aged women; characterized by intermittent, suprapubic pain, urinary frequency, urgency, hematuria and dysuria without evidence of bacterial infection; urine cultures are usually negative; cause is unknown; refractory to treatment. Eventually → bladder fibrosis → contracted bladder.

# Special forms of cystitis contd...

- › **Malakoplakia:** uncommon; inflammatory disorder of unknown etiology. Seen in the bladder and other sites within and outside the urinary tract; characterized by plaques on the mucosal surface of the bladder. Histology shows a chronic inflammation with numerous macrophages. Some macrophages contain laminated, basophilic calcified bodies called **Michaelis-Gutmann bodies**.
- › **Polypoid cystitis:** bladder inflammation in which there is marked mucosal edema (resulting in polypoidal elevations).
- › **Schistosomiasis:** infection by parasitic flatworms called Schistosoma (*Schistosoma hematobium*) in which the worms may lay eggs in the wall of urinary bladder. These eggs elicit a granulomatous reaction and an eosinophilic infiltrate. The ova can calcify and appear as grains of sand in bladder wall. Occasionally the entire urinary bladder becomes calcified → known as calcific cystitis. Chronic Schistosomiasis can predispose to squamous cell carcinoma of the urinary bladder.



**Schistosoma Egg and eosinophils**

## Acute and chronic cystitis: clinical features

- Fever with chills
- Excessive urinary frequency, painful burning urination (dysuria) and lower abdominal or pelvic discomfort.
- Examination of urine usually reveals inflammatory cells and causative organism can be identified by urine culture.
- Most cases of cystitis respond well to treatment with antimicrobial agents.