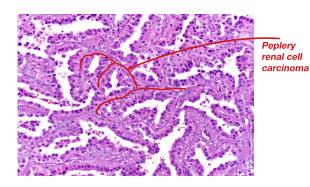


CLEAR CELL TYPE RENAL CELL CARCINO-MA

showing tumor cells with clear cytoplasm



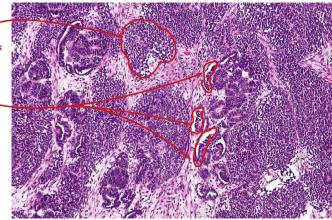
Blastemal cells

Immature epithelium tubules

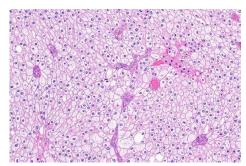
+ hale's colloidal iron stain

HALE'S COLLOIDAL IRON STAIN (SPECIAL STAINING)

Hale's colloidal iron stain is positive (blue) i.e. deep blue granular cytoplasmic positivity in the cytoplasm of Chromophobe RCC.



PEPLERY RENAL CELL CARCINOMA

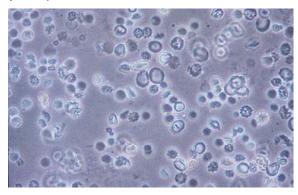


CHROMOPHOBE RENAL CELL CARCINO-MA (RCC)

Chromophobe RCC with admixture of classic (chromophobic) and eosinophilic cells. Characteristic features include perinuclear halos, nuclear "raisins", and distinct cytoplasmic borders and cytoplasm condenses around the edges, giving the appearance of thick prominent cell borders ("plant cell-like")

WILMS TUMOUR

- A) Malignant small round (blue) cells twice the size of resting lymphocyte (blastema component).
- B) Tubular structures/rosettes (epithelial component).
- C) Loose paucicellular stroma with spindle cells (stromal component)

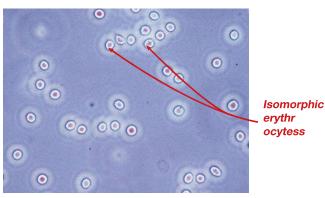


DYSMORPHIC RBC

These dysmorphic erythrocytes vary in size, shape, and hemoglobin content and reflect glomerular bleeding e.g. IgA nephropathy, Poststreptococcal glomerulonephritis

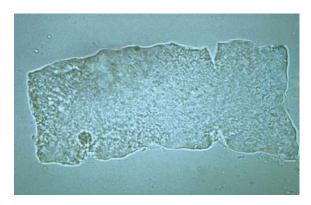
CONCEPTS IN PATHOLOGY

HEMATOLOGY



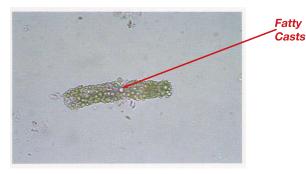
ISOMORPHIC ERYTHROCYTES

These erythrocytes are similar in size, shape, and hemoglobin content. Isomorphic cells reflect nonglomerular bleeding from lesions such as calculi and papillomas or hemorrhage from cysts in polycystic renal disease.



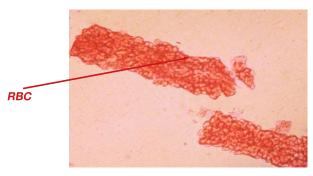
WAXY CAST

| WAXI OAGI | | |
|--|--|--|
| Waxy casts (renal fail- ure casts) | Edges are sharp and there are "cracks" i such casts. End product of cast evolution which is suggestive of very low urine flow associated with severe, longstanding kidney disease such as renal failure. | |



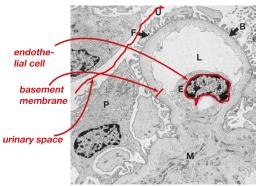
FATTY CASTS

| FALLY CASIS | | |
|--------------|---|--------------------|
| Waxy casts | | Tubular degenera- |
| (renal fail- | · Formed by the breakdown of lipid-rich epithelial | tion |
| ure casts) | re casts) cells, these are hyaline casts with fat globule in- | Nephrotic syndrome |
| a. 5 546t6) | clusions | Hypothyroidism |



RED CASTS

| Waxy casts | | Nephritic syndrome |
|----------------------------|----------------------------------|--|
| (renal fail- ure casts) | •Red Blood Cells within the cast | Pyelonephritis rinary tract injury |
| | | |



NORMAL GLOMERULAR CAPILLARY (ELECTRON MICROSCOPY)

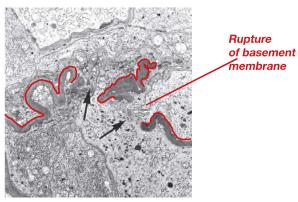
(ELECTRON MICROSCOPY)
Single capillary loop showing the capillary wall portion of the *lumen (L)* is lined by a thin layer of fenestrated endothelial cytoplasm that extends out from the *endothelial cell body (E)*. Endothelial cell body is in direct contact with the mesangium, which includes the *mesangial cell (M)* and adjacent matrix. The outer aspect of the *basement membrane (B)* is covered by *foot processes (F)* from the *podocyte (P)* that line the *urinary space (U)*.



CRESCENTIC GLOMERULONEPHRITIS (PAS STAIN) Showing collapsed glomerular tufts and the crescent-shaped

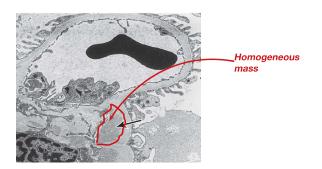
Showing collapsed glomerular tufts and the crescent-shaped mass of proliferating parietal epithelial cells and leukocytes internal to Bowman capsule.



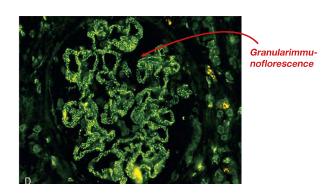


CRESCENTIC GLOMERULONEPHRITIS

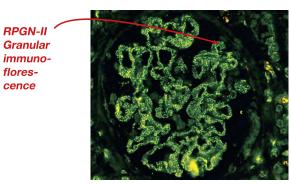
Electron micrograph showing characteristic wrinkling of glomerular basement membrane with focal disruptions (arrows).



HOMOGENEOUS AND LINEAR PATTERN IN IMMUNOFLUORESCENCE SEEN IN GOOD PASUTURE SYNDROME

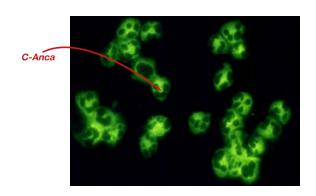


WIRE LOOP LESION

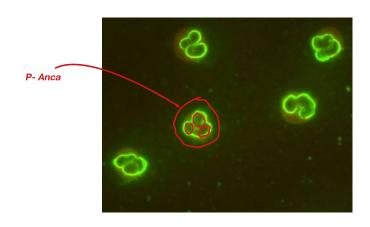


GRANULAR IMMUNOFLORESCENCE

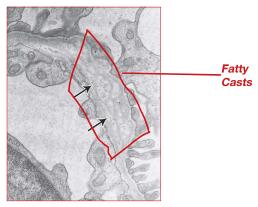
| <u> </u> | | |
|------------|--|---|
| casts (Im- | Formed by Tamm-Horsfall mucoprotein. Hyaline cast has faintly visible outlines. The contours are smooth and the matrix is amorphous. | Normal individuals Heavy exercise Dehydration |



C-ANCA & P-ANCA IN GRANULARIMMU-NOFLORESCENCE

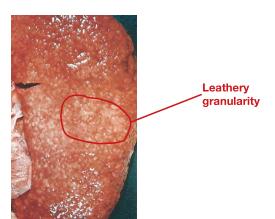






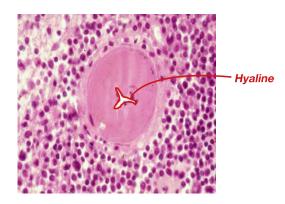
ALPORT SYNDROME

The lamina densa of the glomerular basement membrane is laminated (arrows) rather than forming a single dense band.



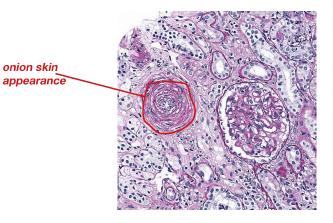
BENIGN NEPHROSCLEROSIS (GROSS APPEARANCE)

Showing cortical surface with fine and leathery granularity of the surface



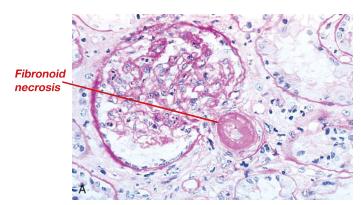
HYALINE

MALIGNANT HTN (FLEA BITTEN KIDNEY)



MALIGNANT HTN (-ONION-SKIN)

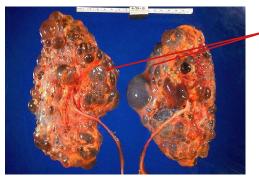
Thickening of the renal arteriolar wall associated with a hyperplastic arteriolosclerosis (hyperplastic arteriolitis). This arteriole has an "onion skin" appearance due to endothelial and muscular hyperplasia.



FIBRINOID NECROSIS (MALIGNANT HYPERTENSION)

Fibrinoid necrosis in the arteriole showing normal muscle layer of the media has been replaced by the fibrinoid material.



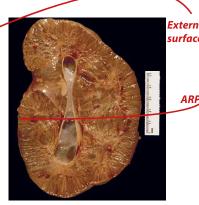


Autosomal dominant polycystic kidney disease.

AUTOSOMAL DOMINANT POLYCYSTIC KIDNEY DISEASE.

The kidneys are enlarged and studded with multiple fluid-filled structures. Renal parenchyma is almost entirely replaced by cysts of varying size.

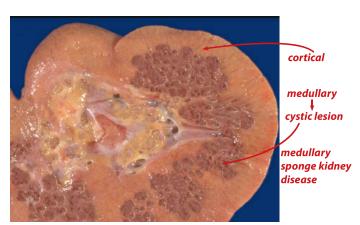




External surface ARPKD

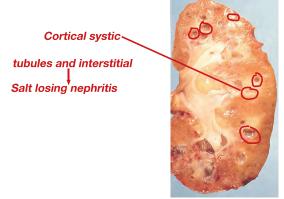
AUTOSOMAL RECESSIVE POLYCYSTIC KIDNEY DISEASE.

The dilated collecting ducts (Both cortical and medullary) are arranged radially and the external surface is smooth.



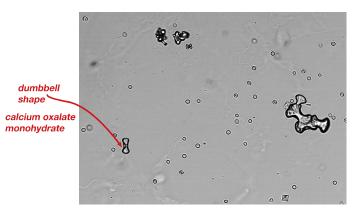
MEDULLARY SPONGE KIDNEY

Showing cysts involving the inner medullary and papillary regions in this kidney with normal cortex



MEDULLARY CYSTIC DISEASE

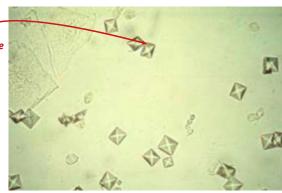
Cut section showing cysts at the corticomedullary junction and in the medulla.



GRANULAR IMMUNOFLORESCENCE

| Calcium oxalate monohy- drate | crystals vary in size and may have a spindle, oval, or dumbbell shape | Urolithiasis. Hypercalciuric or hyperoxaluric disorders Ethylene glycol poisoning. |
|--|---|--|
|--|---|--|

calcium oxalate dihydrate

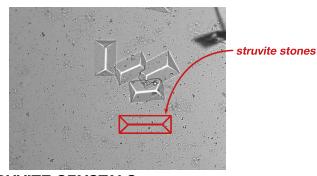


ALCIUM OXALATE DIHYDRATE

Calcium oxhydrate

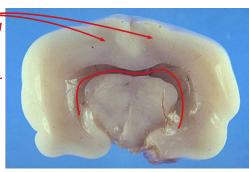
alate monocross-striations, resembling an "EN-

High dietary oxalate ngestion. Ethylene glycol poi-Nephrolithiasis.



No L/R cerebral hemisphere

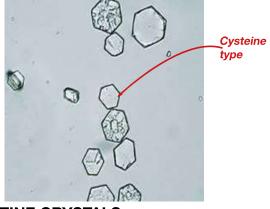
monoventricular cavity



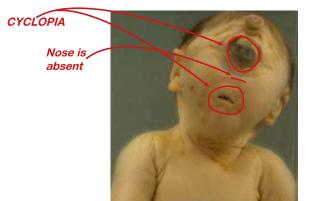
STRUVITE CRYSTALS

| Waxy casts | | |
|------------|---|---|
| • | , | UTIs caused by urea-splitting bac- teria. |

A) HOLOPROSENCEPHALY
A coronal section of the brain showing Forebrain of the embryo fails to develop into two cerebral hemispheres (holoprosencephaly) with monoventricular cavity





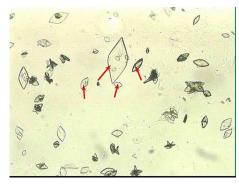


CYSTINE CRYSTALS

| CYSTINE CRYSTALS Hexagonal shape with irregular sides | Cystinuria |
|---|------------|
|---|------------|

B) CYCLOPIA

Affected fetuses and neonates typically have severe facial defects like CYCLOPIA. Associated with trisomy 13 and maternal diabetes mellitus.



Uric acid crystals





agyria

N= Brain

URIC ACID CRYSTALS

URIC ACID **CRYSTALS**

Usually lozenges but varying shape, yellow-tinged

Acute uric acid nephropathy as part of tumor lysis syndrome, and

A) LISSENCEPHALY VS B) NORMAL BRAIN Absence of cortical gyri in brain .