

RENALTUMORS OF ADULTS

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URINARY TRACT TUMORS

Tumors of the <u>lower</u> urinary tract are twice as common as <u>kidney</u> tumors.

- The most common malignant tumor of the <u>kidney</u> is renal cell carcinoma.
- The most common <u>lower urinary tract</u> tumor is <u>urothelial carcinoma</u>.

RENAL CELL CARCINOMA (RCC)

- Origin: renal tubular epithelium.
- in cortex.
- 2%-3% of all cancers in adults.
- 6th-7th decades.
- •M:F 2:1

PREDISPOSING FACTORS

- smoking
- hypertension
- obesity
- occupational exposure to cadmium (nickel-cadmium batteries, etc).
- chronic dialysis (acquired polycystic disease)

NEW CLASSIFICATION BASED ON THE MOLECULAR ORIGINS OF THESE TUMORS

- I-Clear Cell Carcinomas
- 2-Papillary Renal Cell Carcinomas
- 3-Chromophobe Renal Carcinomas

I- CLEAR CELL CARCINOMAS

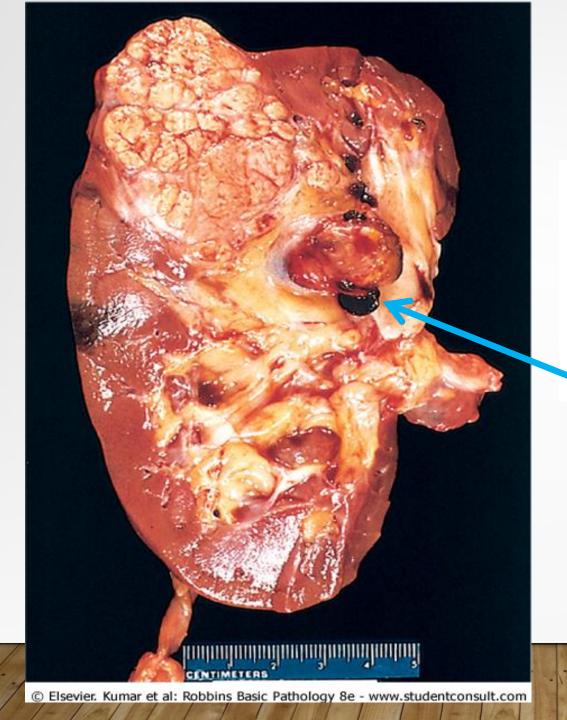
- most common type (70%-80% of RCC).
- cells with clear or granular cytoplasm.
- may be:
- **I-Sporadic**
- 2-Familial (including von Hippel-Lindau (VHL) disease)
- The VHL gene is involved in familial and also <u>sporadic</u> clear cell carcinomas (60%).

2- PAPILLARY RENAL CELL CARCINOMAS

- 10% to 15%.
- papillary growth pattern.
- multifocal and bilateral
- familial and sporadic forms.
- MET proto-oncogene on chromosome 7 →↑
 growth in proximal tubular epithelial cells

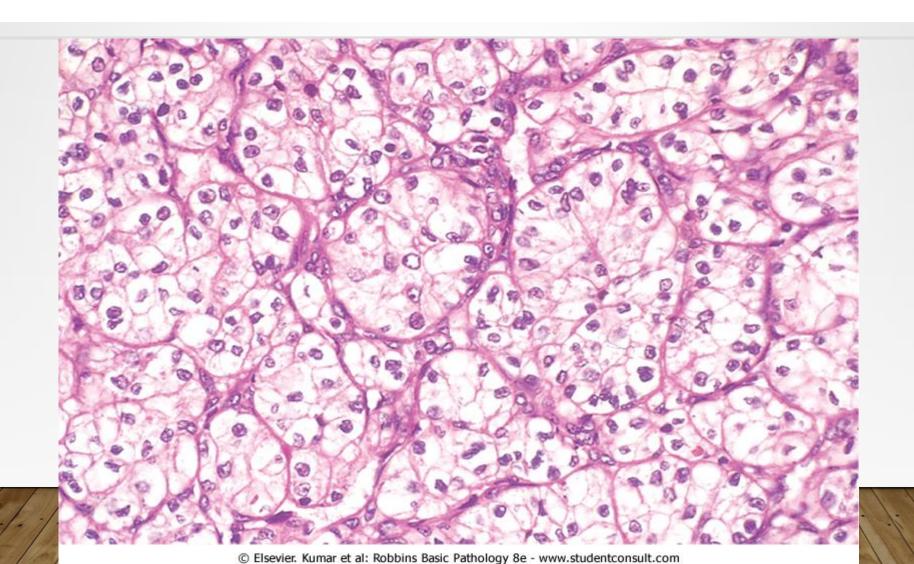
3- CHROMOPHOBE RENAL CARCINOMAS

- least common (5%)
- from intercalated cells of collecting ducts.
- tumor cells are "less clear" than cells in clear RCC
- multiple losses of entire chromosomes, including 1, 2, 6, 10, 13, 17, and 21.
- extreme hypodiploidy.
- good prognosis.



Renal cell carcinoma:
yellowish, spherical neoplasm
in one pole of kidney.
Note the tumor in the
dilated, thrombosed renal
vein.

RENAL CELL CARCINOMA (CLEAR CELL TYPE)



CLINICAL COURSE OF ALL RCC

- 1- Painless hematuria (50%)
- 2- palpable abdominal mass
- 3- dull flank pain
- 4-Fever
- 5-Polycythemia (5% 10%): elaboration of erythropoietin by tumor.

CLINICAL COURSE OF ALL RCC

6- other Paraneoplastic syndromes:

- I-hypercalcemia
- 2-Hypertension
- 3-Cushing syndrome
- 4-feminization or masculinization
- 7- Metastasis: most commonly to lungs and bones.
- 8- may invade the renal vein

UROTHELIAL TUMORS (TRANSITIONAL CELL CARCINOMA)

classified into:

- I -benign papilloma: rare
- 2-papillary urothelial neoplasms of low grade: most frequent
- 3-papillary urothelial carcinoma of high grade

TRANSITIONAL CELL CARCINOMA OF BLADDER



UROTHELIAL (TRANSITIONAL) CELL CARCINOMAS

- Low-grade carcinomas are rarely invasive.
- may recur after removal.
- staging at the time of initial diagnosis is the most important prognostic factor

PAPILLARY UROTHELIAL (TRANSITIONAL) CARCINOMA-LOW GRADE







SQUAMOUS CELL CARCINOMAS

- only 5% of bladder cancers
- Associated with:
- **Schistosomiasis** infection
- chronic inflammation
- stone formation

CLINICAL COURSE OF BLADDER CANCERS

- Painless hematuria.
- M:F 3:1
- 50 to 70 years.
- Prognosis
- low-grade + shallow → good prognosis.
- High grade lesions + deep → bad

- Predisposing factors of bladder cancers:
- not familial.
- I β-naphthylamine (paints; cigarettes)
- 2- Cigarette smoking.
- 3- Chronic cystitis.
- 4- Schistosomiasis.
- 5- drugs as cyclophosphamide.

Treatment:

- transurethral resection in cystoscopy
- (BCG) injections → granulomatous reaction (immune response against cancer)
- Radical cystectomy and chemotherapy for advanced cases
- Follow-up for recurrence with cystoscopy and urine cytologic studies for the rest of life.

Renal tumors of childhood

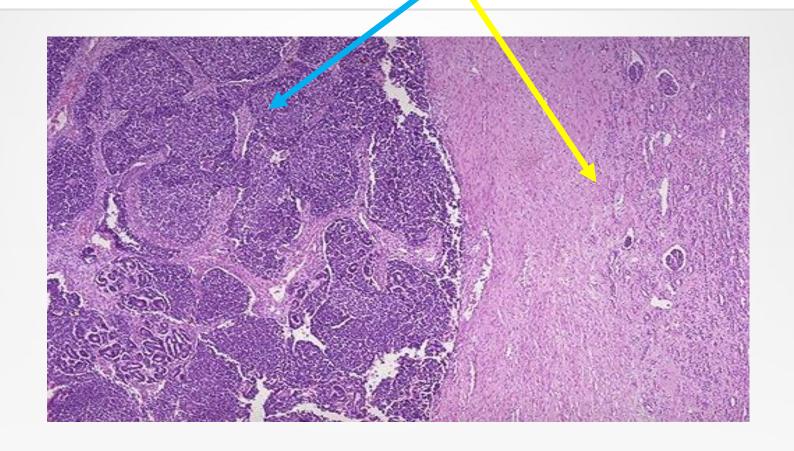
WILMSTUMOR

- 3rd most common solid cancer < 10 years.
- derived from the mesoderm.
- sporadic or familial (autosomal dominant).
- Mutations: WT-land 2 genes.
- primitive glomerular and tubular structures
- Treatment: surgery & chemotherapy



WILM'S TUMOR

NESTS AND SHEETS OF DARK **BLUE** CELLS OF <u>WILMS TUMOR</u> AT THE LEFT WITH COMPRESSED <u>NORMAL RENAL PARENCHYMA</u> AT THE RIGHT.



WILMSTUMOR: THE TUMOR SHOWS ATTEMPTS TO FORM PRIMITIVE GLOMERULAR AND TUBULAR STRUCTURES

