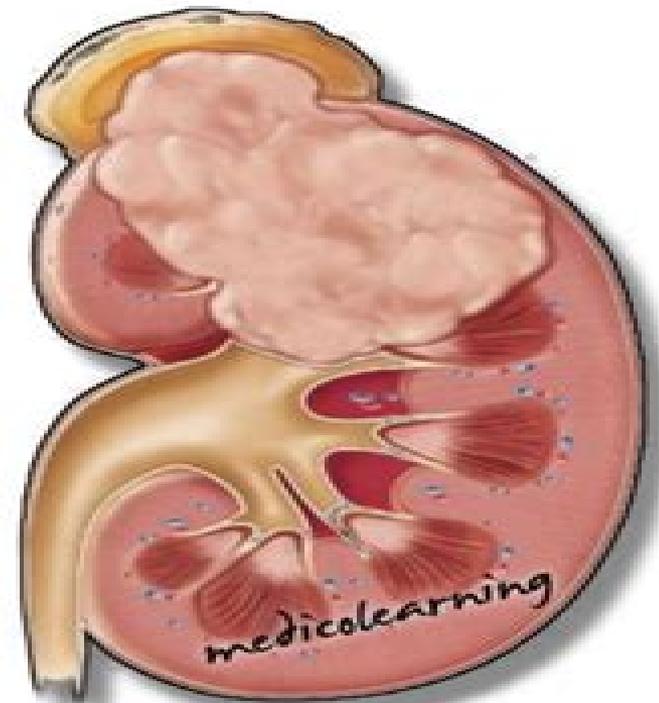


Wilms' Tumor

- Wilms tumor is most common primary renal tumor of childhood,
- Highly malignant primary embryonal tumor.
- Most common between 2 - 5 years of age & more than 95% occur below 10 Years of age.
- WT1 gene: Tumor suppressor gene located in the chromosome 11p13.
- Wilm tumor: Deletion of short arm of chromosome 11p13.
- Wilms' tumor: Associated congenital syndromes
 1. WAGR syndrome
 2. Denys-Drash syndrome
 3. Blackwith-Wiedmann syndrome.
- WT1 and PAX6 genes are located at chromosome 11p13.
- Wilms tumor microscopy: Three components:
 1. Blastemal
 2. Immature stromal
 3. Immature epithelial.
- Anaplasia in Wilms tumor: Associated with:
 - TP53 mutations
 - Resistance to chemotherapy.
- Wilms tumor:
 1. Abdominal mass
 2. Microscopic hematuria.

Clinical Presentation :

- Palpable smooth abdominal mass
- Abdominal pain
- Gross Hematuria
- Tumor rupture with hemorrhage in peritoneal cavity
- Extension into renal vein and IVC
- Hypertension



Wilms Tumour

Rosina Ehmann



Wilms Tumour:

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