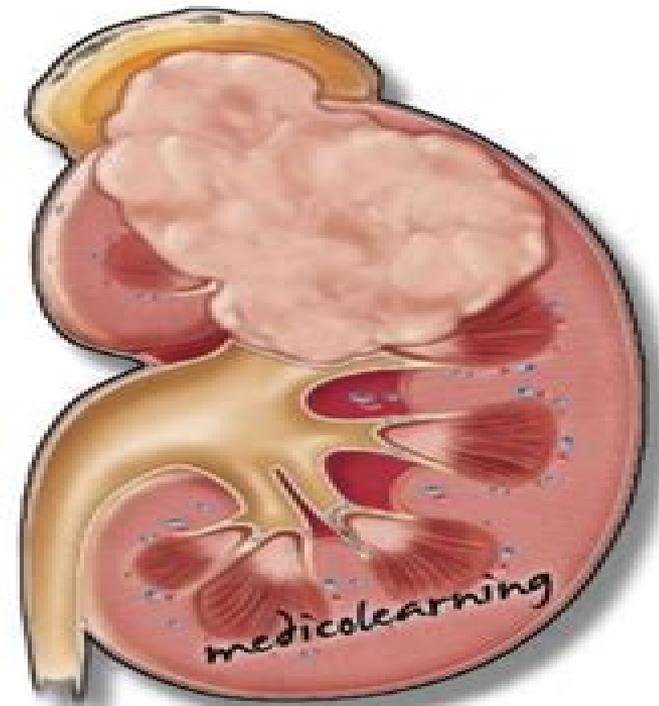


# Wilms' Tumor

- 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

**RM Cervero**



## **Wilms Tumour:**

## Enjoying the Melody of Appearance: An Mental Symphony within **Wilms Tumour**

In a world used by monitors and the ceaseless chatter of quick interaction, the melodic elegance and psychological symphony developed by the published word often diminish in to the background, eclipsed by the constant sound and disturbances that permeate our lives. But, situated within the pages of **Wilms Tumour** a marvelous fictional prize filled with organic thoughts, lies an immersive symphony waiting to be embraced. Constructed by a masterful musician of language, that interesting masterpiece conducts viewers on a mental journey, skillfully unraveling the hidden songs and profound impact resonating within each cautiously crafted phrase. Within the depths of the moving review, we can explore the book is main harmonies, analyze its enthralling writing type, and submit ourselves to the profound resonance that echoes in the depths of readers souls.

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