

Pathology - Renal Cancer

🕒 Published: 15 May 2006



- Clear cell carcinoma: comprises >70% of renal lesions
 - VHL gene mutation principle event. Recent association between VHL protein and hypoxia inducing factor [HIF] protein ties pathology into angiogenesis cascade pathway.
- Papillary carcinoma: comprises 10-15% of renal lesions
 - Sporadic and hereditary forms
 - Associated with alterations in chromosomes 7, 17, and Y
 - Generally better survival

Read more on "[A Molecular Classification of Papillary Renal Cell Carcinoma](#)"

- Chromophobe tumors: 5 % of cases
 - Loss on chromosome 1
- Collecting duct carcinoma: one percent or less of cases
 - Can mimic transitional cell Ca
 - Generally poor outcome
- Oncocytoma: 5 % of renal tumors
 - Generally localized and encapsulated. 5% bilaterality
 - Mahogany brown color, acidophilic cells secondary to dense mitochondrial hyperplasia
 - Distinction from renal cell cancer difficult on imaging or needle biopsy. Best treated with surgical removal
- Angiomyolipoma: Renal Hamartomas comprised of fat, muscle and blood vessels. Tissue signature on CT by demonstration of negative Hounsfield units.
 - Sporadic, isolated lesions present age 35-50 with a 4:1 female ratio
 - Tuberous Sclerosis patients demonstrate multiple and bilateral lesions. 80% of patients will develop AML.
 - Treatment based on tumor size: those <4 cm are observed, those >4cm undergo selective angioembolization or partial nephrectomy
- Renal Sarcoma
 - Pure sarcoma is rare and usually leiomyosarcoma
 - All tumor types can degenerate towards sarcoma
 - Generally poorer outcome
- Rare Renal lesions
 - Adult Wilms tumor
 - Lymphoma
 - Xanthogranulomatous Pyelonephritis [XPG]
 - Haemangiopericytoma

From the BJU International Mini Reviews: [Renal Haemangiopericytoma: The Characteristics of a Rare Tumour.](#)

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