CASE DISCUSSIONS IN UROPATHOLOGY



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Clinical history:

 A 45 year old man presented with complaints of left loin pain which was non-radiating with intermittent fever for the past six months

Clinical examination:

• Smooth non-tender mass in the left flank

INVESTIGATIONS:

- Abdominal ultrasonography showed a multicystic lesion in the left kidney
- 2. CECT of the abdomen showed a heterogeneously hypodense multicystic lesion of 17 x 9cm with calcified thick and irregular walls, with small multiple daughter cysts within the lesion





Complete blood picture shows:

- 1. Mild eosinophilia
- 2. Raised ESR

Surgery:

Laparoscopic left sided nephrectomy was done



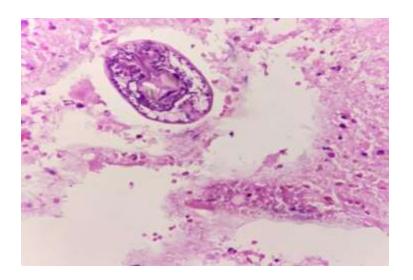
- Left nephrectomy specimen measuring 12x10x9 cms along with ureter measuring 3cms
- ➤Cut section shows multiple cysts noted in the renal parenchyma





MICROSCOPY FINDINGS

- Renal parenchyma with few tubules showing thyroidisation and chronic mononuclear inflammatory cell collection
- Thickened hyalinised cyst wall with lamellated membrane and brood capsules



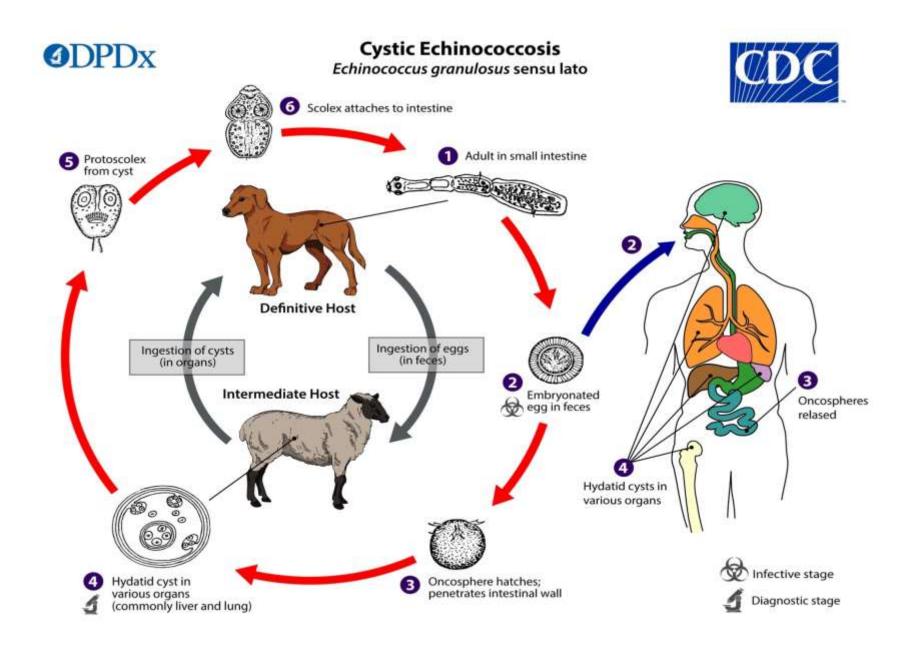




Left sided Renal hydatid cyst

RENAL HYDATID CYST

- Caused by Echinococcus granulosus
- > Definitive host is dog, intermediate host is sheep
- When transmitted to humans it can affect various organs like liver, lung, brain and urinary tract
- Kidney is the most commonly affected organ in the urinary tract



Benign disease remains asymptomatic for many years

Most common clinical presentation is hematuria and / hydatiduria

DIAGNOSIS:

Requires high index of suspicion

> Peripheral eosinophilia

> Positive ecchinococcal antigen immunofluorescence

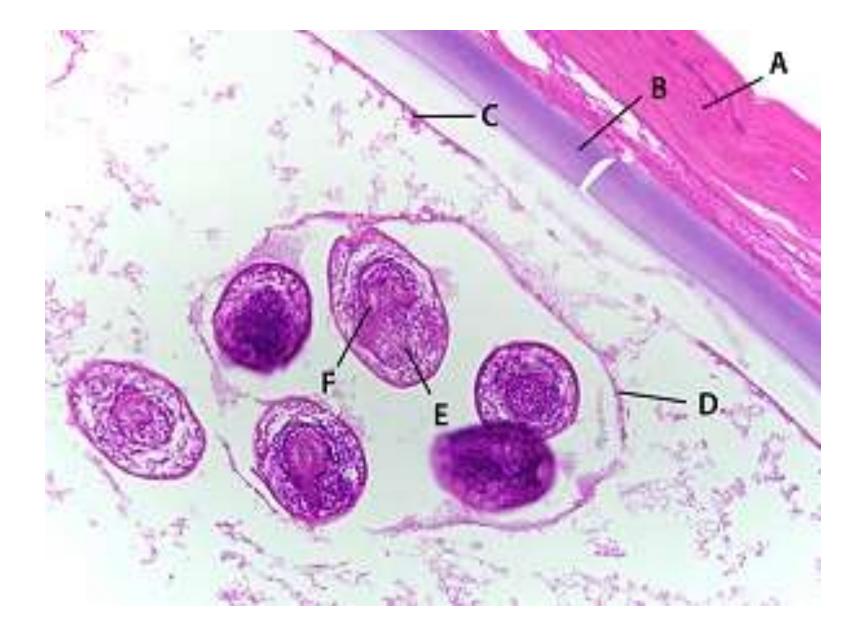
Hemagglutination tests

> H/O exposure in an endemic areas

MORPHOLOGY

Cyst is composed of 3 layers

- 1. Outermost *pericyst* is fibrous
- 2. Middle *ectocyst* which is laminated, hyaline, acellular
- Inner *endocyst* is germinative layer consists of daughter cyst and brood capsules with scolices



DIFFERENTIAL DIAGNOSIS

1.Multicystic renal cell carcinoma:

- Variably sized cyst with thin septae lined by clear cells with low nuclear grade
- 2. Multicystic nephroma:
- Cysts are lined by epithelium of variable morphology with hypo/hypercellular stroma
- Stroma composed of closely packed spindle cells with areas of calcifications, multinucleated giant cells



Case history:

• A 35 year old *female* presented with swelling in the region of external urethral orifice, gradually increasing in size since 6 months along with hematuria, dysuria

Clinical examination:

• Firm, mobile, nontender mass sized 3cmx2.5cm located just proximal to the urethral meatus *anteriorly*

Investigations:

- Ultrasound abdomen and pelvis revealed a normal genitourinary system
- Uroflometry was normal
- Cystourethroscopy showed a mass arising from the anterior wall of the urethra, extending 2cms proximally from the meatus

Surgery:

The mass was completely excised under spinal anaesthesia

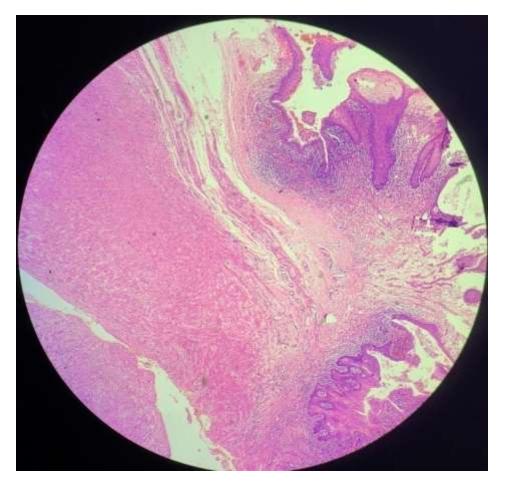
Gross findings:

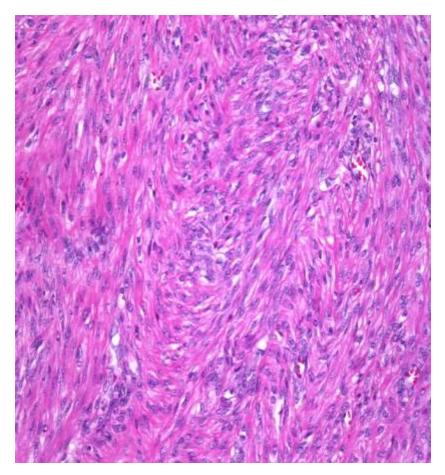
 Pedunculated growth of urethral wall measuring 3cmx1.5cmx1cm













URETHRAL LEIOMYOMA

URETHRAL LEIOMYOMA

- Leiomyoma of urethra is a rare benign mesenchymal tumor
- > Occur in 1 among 1000 women of reproductive age
- > They constitute 5% of all paraurethral masses
- > Most common site of presentation is *proximal urethra*
- > Distal urethral involvement is very rare

➤The tumor has been reported to enlarge during pregnancy and shrink after delivery

Diagnosis of such tumors is primarily based on clinical history, physical examination and imaging techniques

➤The final diagnosis should made based on histopathological report

Types of leiomyoma:

- 1. Conventional
- 2. Subtypes:
- Cellular
- Leiomyoma with bizarre nucleus
- Lipoleiomyoma
- Epitheloid
- Myxoid
- Dissecting
- Diffuse leioyomatosis



> A 72 year male complaints of lower abdominal pain

since 2 months, associated with gross hematuria

> On examination:

P/A: soft, non tender, bowel sounds present, external genitalia: meatus -normal

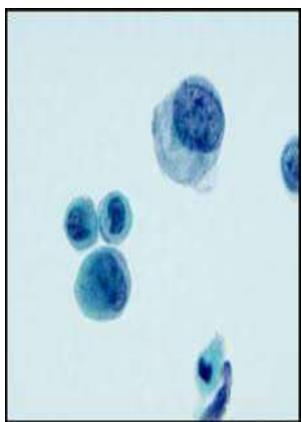
INVESTIGATIONS:

Urine cytology:

• Positive for atypical urothelial cells

Ultrasound abdomen and pelvis:

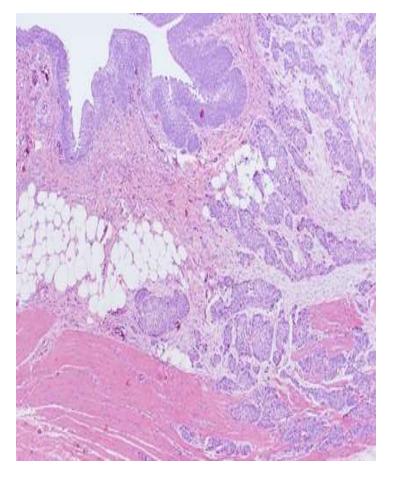
• Gross hydroureteronephrosis

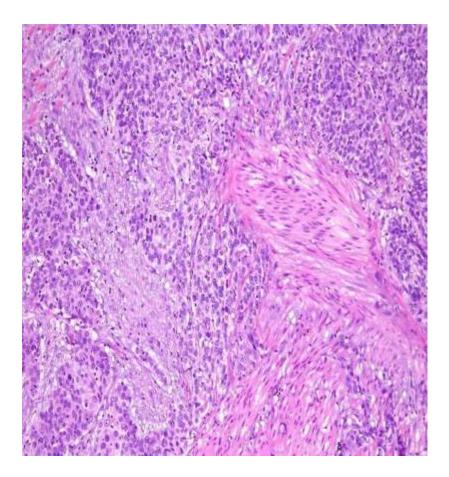


Gross:

Bladder and ureter biopsy

Microscopy:







INFILTRATING UROTHELIAL CARCINOMA

UROTHELIAL NEOPLASMS

RISK FACTORS:

- 1. Smoking
- 2. Environmental exposure
- 3. Occupational exposure
- 4. Schistosoma hematobium
- 5. Congenital bladder exostrophy

Localization:

- > 90% arises in bladder
- Among bladder lateral wall involvement is most common, anterior wall involvement is rare

Urethral involvement is rare

Can cause hydronephrosis

≻<u>IMAGING:</u>

- ➤ cystoscopy
- Bimanual examination under anaesthesia
- ➢ Biopsy
- Transurethral resection

Non invasive urothelial lesions

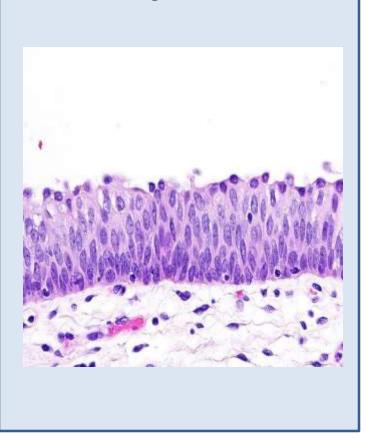
- Urothelial dysplasia
- Urothelial papilloma
- Inverted urothelial papilloma
- Urothelial carcinoma in situ
- Papillary urothelial neoplasm of low malignant potential
- Non invasive papillary urothelial carcinomalow grade and high grade

INFILTRATING UROTHELIAL CARCINOMA

- Nested, including large nested
- Microcystic
- Micropapillary
- Lymphoepithelioma like
- Plasmacytoid/ signet ring cell/ diffuse
- Sarcomatoid
- Giant cell
- Lipid rich
- Clear cell
- Poorly differentiated

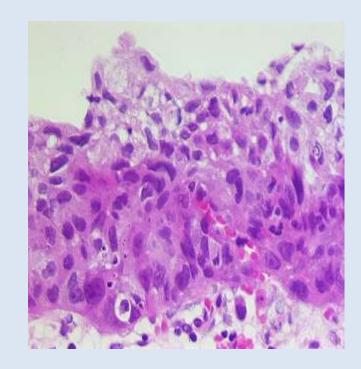
1.UROTHELIAL DYSPLASIA

Defined as degree of atypia believed to be definitely preneoplastic yet not sufficient for diagnosis of CIS



2. Urothelial carcinoma in situ

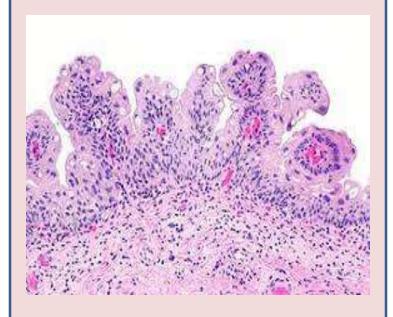
Flat lesion in which surface epithelium contains cytologically malignant cells



3.UROTHELIAL

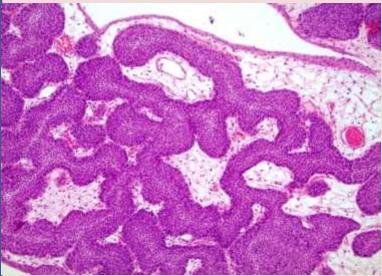
PAPILLOMA

Delicate fibrovascular cores with exophytic pattern lined by normal appearing urothelium with out atypia



4.INVERTED PAPILLOMA

Benign urothelial tumor characterized by inverted growth pattern with no to minimal cytologic atypia
May invaginate extensively into lamina propria but not into muscular wall

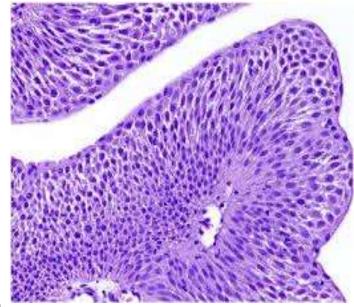


5.Papillary urothelial neoplasm of low malignant potential

- Papillae are lined by *multilayered epithelium* thicker than papilloma
- Cell density appears to be increased
- Polarity is preserved
- Minimal or absent cytological atypia

Differential diagnosis:

- Urothelial papilloma
- Non invasive low grade papillary urothelial carcinoma



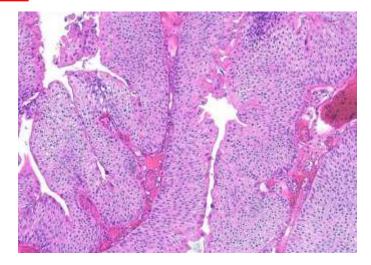
6. Non invasive papillary urothelial carcinoma

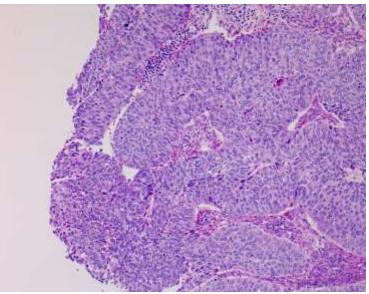
Low grade:

- Papillae lined by multilayered epithelium with frequent branching
- Mild anisocytosis and nuclear atypia

High grade:

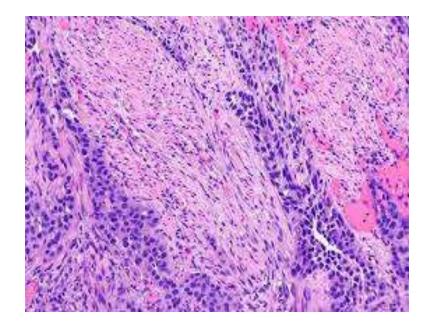
- Moderate to marked cytologic pleomorphism
- Mitotic figures are frequent





Invasive urothelial carcinoma

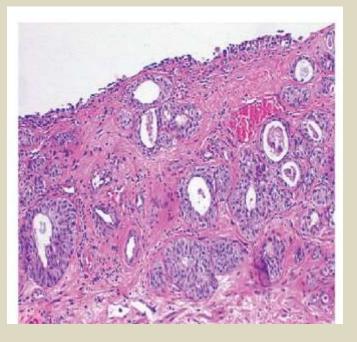
- Defined as urothelial tumor that invades beyond the basement membrane
- Irregular infiltration to level of muscularis propria is diagnosis of malignancy



NESTED VARIANT

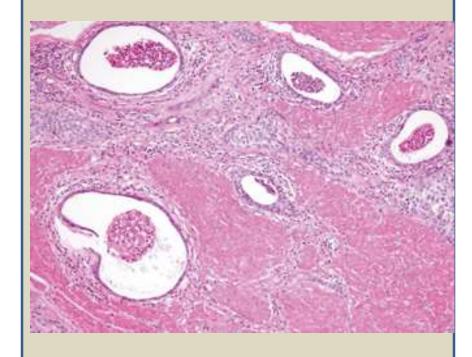
•Deceptively bland •Irregular distribution of bland urothelial cells in suburothelial tissues

•Worse prognosis



MICROCYSTIC VARIANT

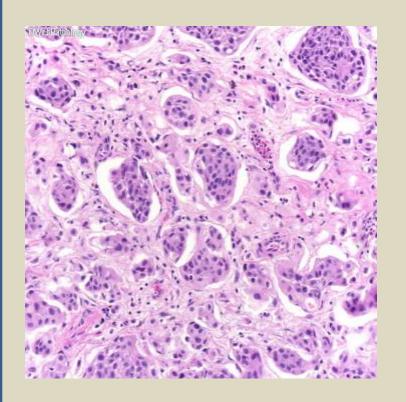
•UC with striking cystic pattern in which the cysts contain necrotic material or pale pink secretions •Favourable prognosis



MICROPAPILLARY VARIANT

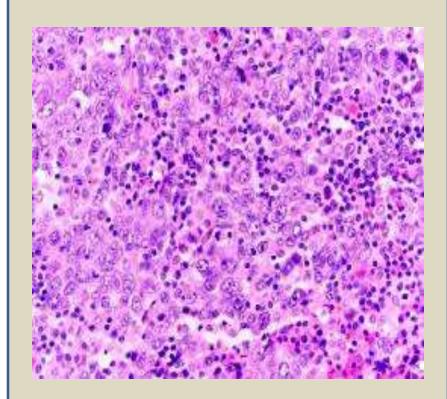
•Characterized by back to back retraction spaces and multiple epithelial aggregates in a single retraction space

•Favourable prognosis



LIKE VARIANT

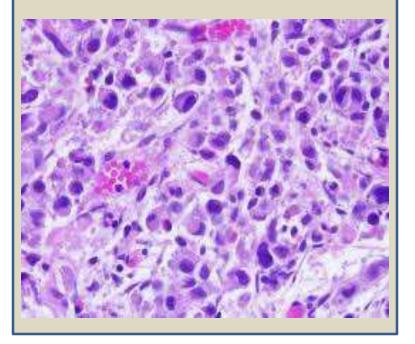
- Appearance that of a non keratinising carcinoma associated with heavy inflammatory infiltrate
 - Favourable prognosis



PLASMACYTOID

VARIANT

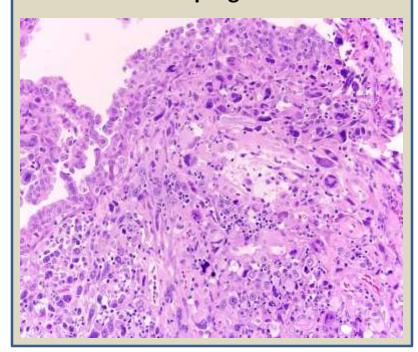
Monomorphic round neoplastic cells are discohesive, grow singly or in clusters
Poor prognosis



SARCOMATOID

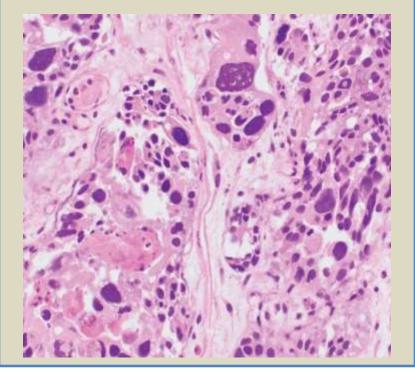
VARIANT

 High grade neoplasm of the bladder in which a malignant epithelial component coexist with areas having spindled sarcoma like appearance
 Poor prognosis



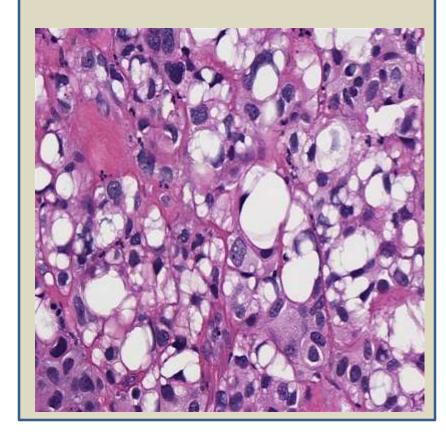
GIANT CELL VARIANT

 Biphasic appearance of the tumor, consisting of sheets and nodules of mononuclear cells and scattered osteoclast like giant cells,poor prognosis



LIPID CELL VARIANT

It consists of infiltrating nests of epitheloid cells with abundant vacuolated cytoplasm
Presents at higher stage
Poor prognosis

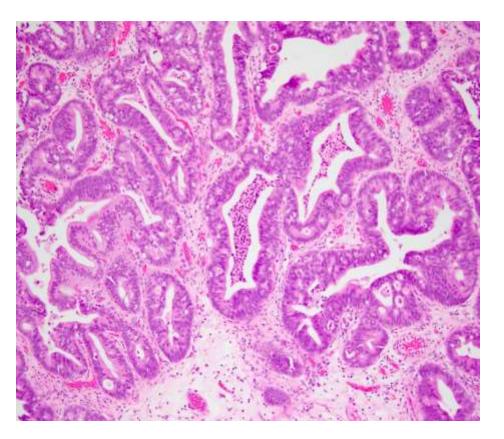


CASE NO:4

Clinical history:

- A 30 year old male patient came with the complaints of pain abdomen, hematuria since 5 days
- Associated with increased frequency of micturition
- H/o of similar complaints in his grand father
 INVESTIGATIONS:
- **CECT** shows a large 3.4x3.7x3.4cm sized polypoidal intraluminal lesion noted in right anterior fundal region of bladder





Neoplastic glands are lined by pleomorphic mucin producing pseudostratified columnar epithelium with central necrosis



ADENOCARCINOMA OF BLADDER (ENTERIC TYPE)

Adenocarcinoma of bladder

- Adenocarcinoma is a malignant neoplasm derived from the urothelium with histologically pure glandular phenotype
- Bladder adenocarcinomas constitute 2% of the malignant tumors of the bladder

Etiology:

- Long standing intestinal metaplasia particularly in patients with bladder exostrophy
- Chronic irritation and obstruction
- Non functioning bladder
- schistosomiasis

- Hematuria is most common symptom and associated with irritative voiding symptoms
- Mucusuria occurs in some patients
- Grossly cases appear as fungating masses that ulcerate the mucosa and invade the bladder wall

Adenocarcinoma variants:

- 1. Enteric
- 2. Mucinous
- 3. Mixed

1.Enteric adenocarcinoma:

• Glands are lined by

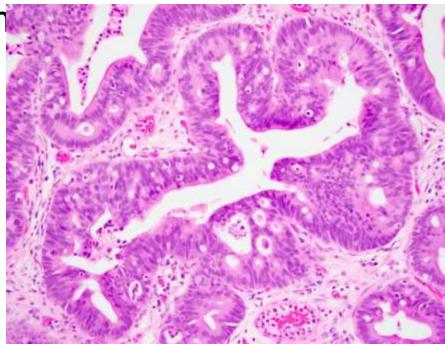
Pseudostratified

mucin secreting epitheliun

with various degrres of

pleomorphism and with

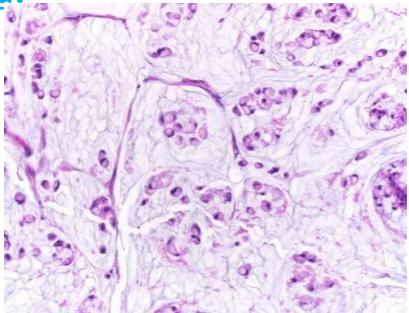
central necrosis



2. Mucinous adenocarcinoma:

The tumor cells form nests

floating in abundant extracellular mucin



3. <u>Mixed adenocarcinoma:</u>

Tumors with mixture of enteric and mucinous patterns

Prognosis:

 Overall prognosis for primary adenocarcinoma of the urinary tract has been *poor*



CLINICAL HISTORY:

- A 34 years male complaints of lower abdominal pain since 5 days, radiating to back
- H/O alcohol intake present

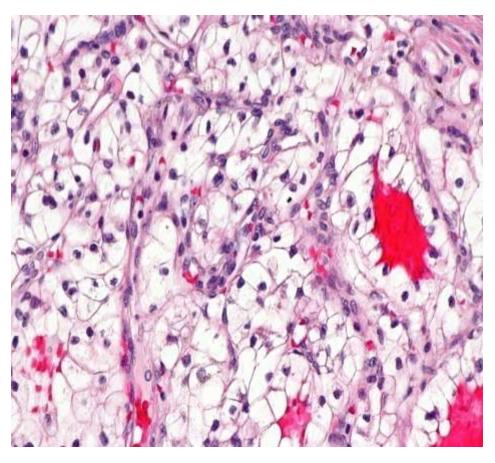
Investigations:

CECT abdomen:

 A well defined heterogenous lesion of approximately 2.1x2.1cms in upper pole of left kidney







MICROSCOPY



RENAL CELL CARCINOMA – CLEAR CELL VARIANT

RENAL CELL CARCINOMA

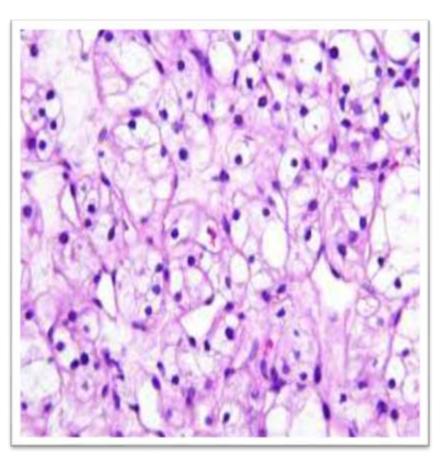
RISK FACTORS:

- 1. Tobacco smoking
- 2. Obesity
- 3. Hypertension
- 4. Acquired cystic kidney disease
- 5. Occupational exposure
- 6. Genetic susceptibility

CLASSIFICATION OF RENAL CELL

CARCINOMA

- 1. Clear cell Rcc
- 2. Papillary Rcc
- 3. Clear cell papillary Rcc
- 4. Chromophobe Rcc
- 5. Collecting duct carcinoma
- 6. Renal medullary carcinoma
- 7. Spindle cell carcinoma
- 8. Succinase dehydrogenase deficient Rcc
- 9. Acquired cystic disease associated Rcc



Clear cell carcinoma

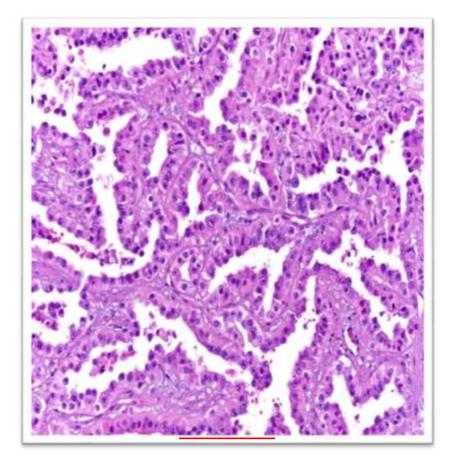
POOR PROGNOSIS

Differential diagnosis:

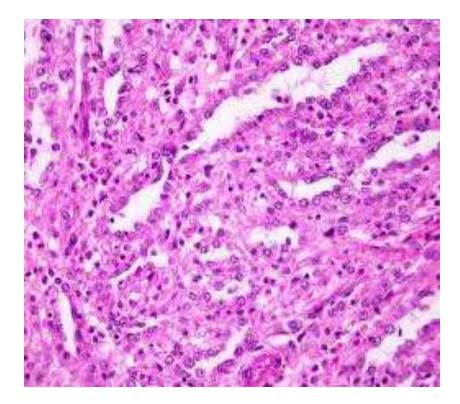
1.Chromophobe Rcc

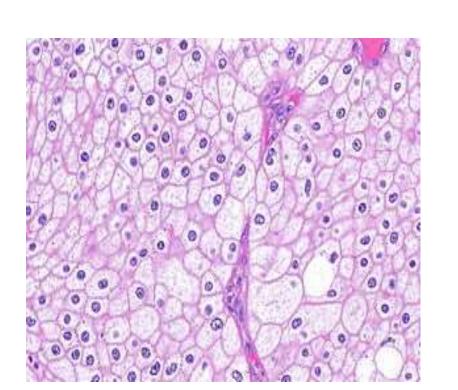
2.Papillary Rcc

3.Clear cell papillary Rcc



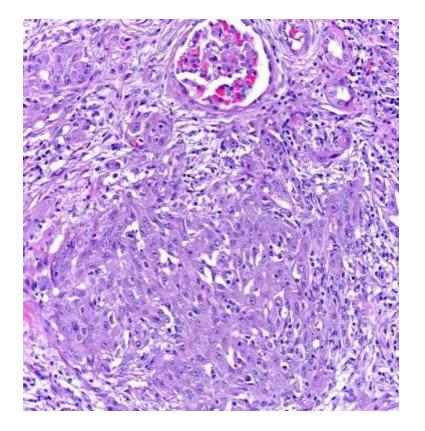
Papillary renal cell carcinoma BETTER PROGNOSIS Differential diagnosis: Papillary adenoma

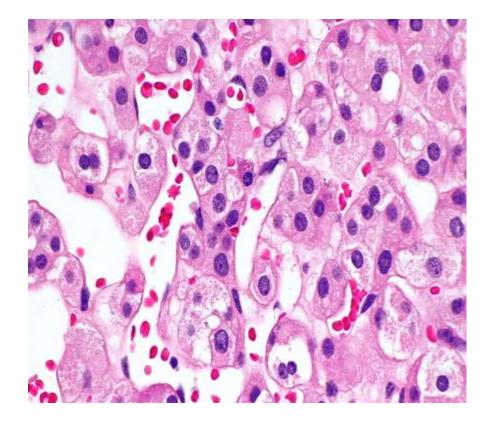




COLLECTING DUCT CARCINOMA

POOR PROGNOSIS Differential diagnosis: 1.Papillary rcc 2.Renal medullary carcinoma CHROMOPHOBE CARCINOMA BETTER PROGNOSIS THAN CLEAR CELL CARCINOMA Differential diagnosis •Clear cell renal cell carcinoma





RENAL MEDULLARY CARCINOMA

<u>Differential diagnosis</u> •Collecting duct carcinoma

SUCCINATE DEHYDROGENASE DEFICIENT RCC

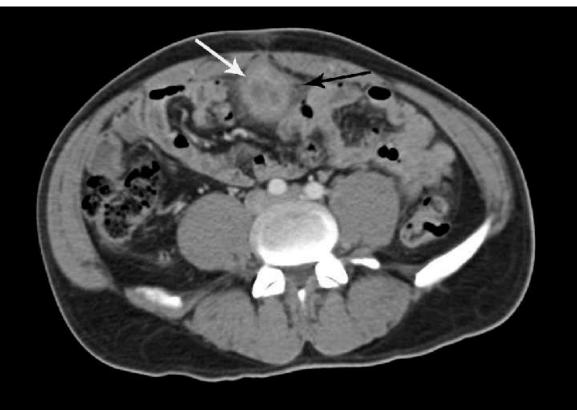
FAVOURABLE PROGNOSIS



CASE HISTORY:

- A 11 years female complaints of lower abdominal pain, swelling below the umbilicus on and off since 11 months
- On examination a ill defined soft palpable lump noted infraumbilically

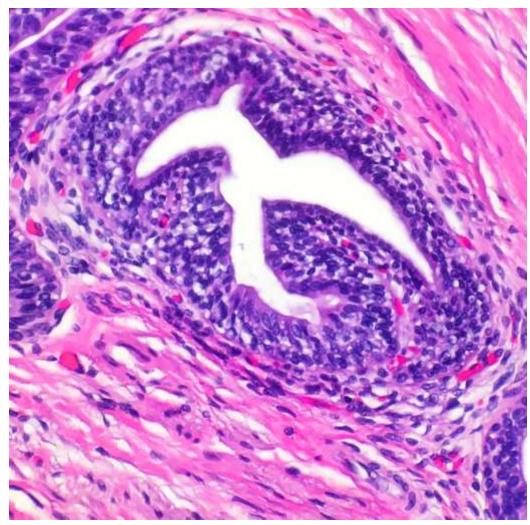
 CT abdomen shows a tubular structure measuring 5.4cm in length extending from superior aspect of urinary bladder to umbilicus –PATENT URACHUS



GROSS AND MICROSCOPY





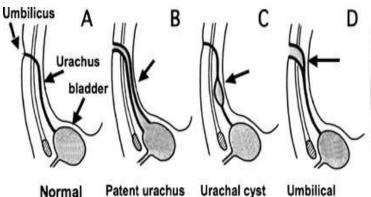


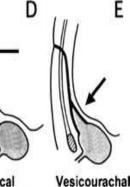


URACHAL CYST

URACHAL CYST

- Urachal cyst is a sinus remaining from the allantois during embryogenesis
- It is a cyst occurs in the remnants between the umbilicus and bladder
- Clinically important when infected, dilated or neoplastic





diverticulum

urachal cyst

and sinus

 Ford in the units

 Ford in the units

 For or burning with wenders

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- Lined by flattened atrophic urothelial epithelium
- These cysts are surrounded by a thin layer of fibro muscular tissue

DIFFERENTIAL DIAGNOSIS:

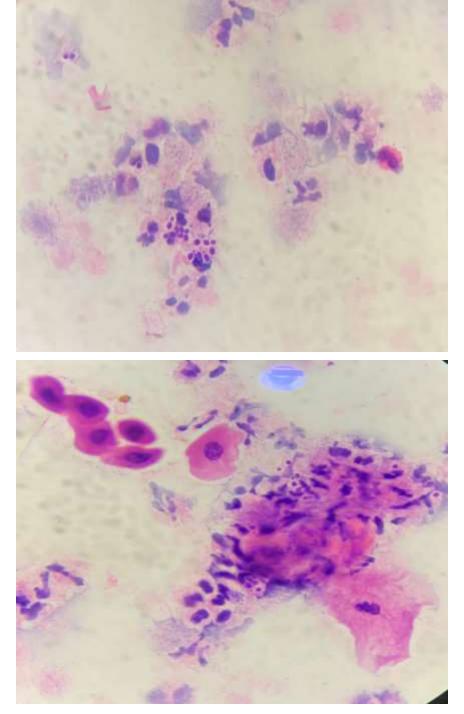
1.Omphalitis

2. Patent omphalomesenteric duct

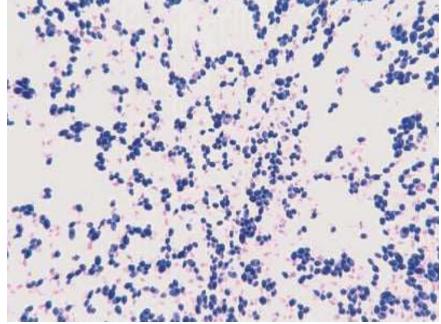


Case history:

- A 80 year old male patient complaints of burning micturition since 1 month
- k/c/o diabetic since many years
- O/E the urine is turbid









CANDIDURIA

