

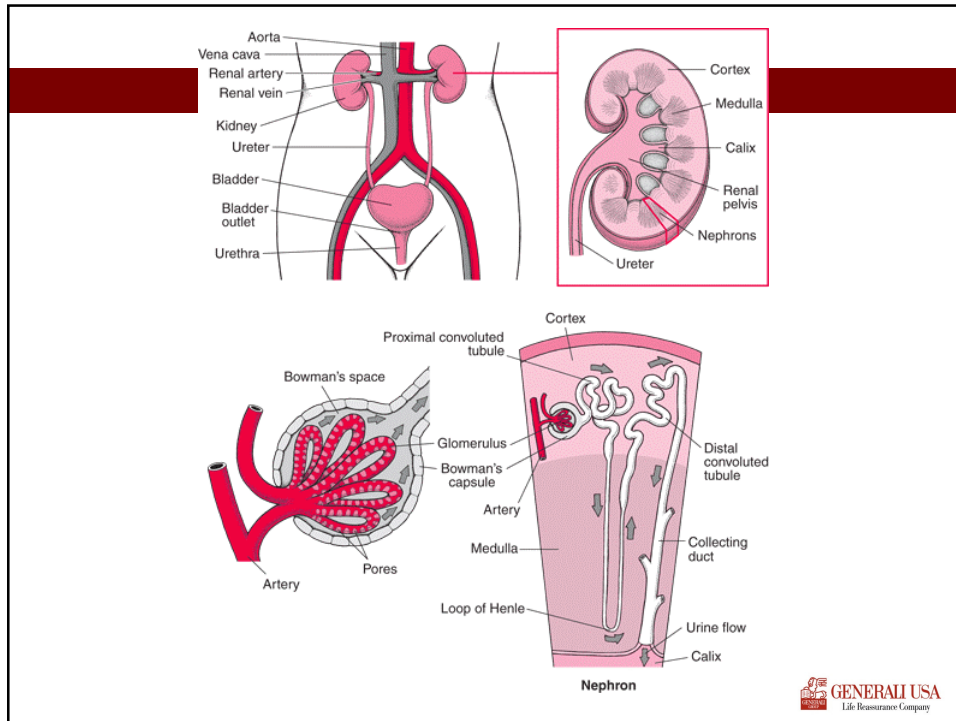
“Incidentalomas” of Kidney Imaging

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Imaging the Kidney

- **Review the Anatomy and Function**
- **Renal Mass**
 - Cysts
 - Solid Mass
- **Renal Parenchymal Disease**
- **Renovascular Disease**



Imaging the Kidney

- **Plain X-ray Kidneys, ureters, and bladder (KUB)**
 - Can detect calcifications, may detect renal outline if visible
 - Should be only the starting point
- **Intravenous Urography (IVU) a.k.a. Intravenous Pyelography (IVP)**
 - 25-40 mg of iodine (75-150 ml) injected IV as a bolus

60 seconds after contrast injection



Taal: Brenner and Rector's *The Kidney, 9th ed.*; Chapter 27 - Diagnostic Kidney Imaging

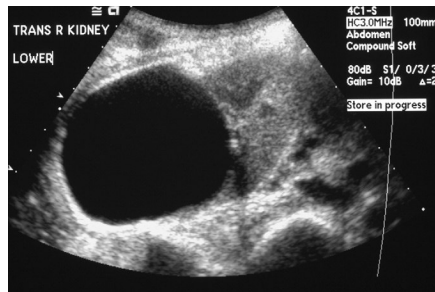
Imaging the Kidney

- **Renal Mass**
 - IVP is not a sensitive test for renal Mass
 - Ultrasound is a little better
 - Compared to Computed Tomography (CT)

Mass Size	IVP detects	Ultrasonography
< 1 cm	10%	26%
1-2 cm	21%	60%
2-3 cm	52%	82%
>3 cm	85%	85%

Imaging the Kidney – Renal Cysts

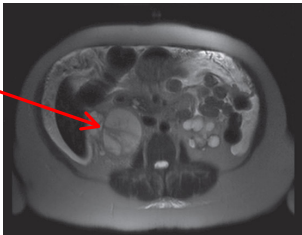
- **Renal cysts – found incidentally**
 - Quite common – Most persons age 60+ will have one or more, found in 50% of people over age 50
 - Rare in those under age 25
 - Usually cortical in location
 - If seen on IVP, Ultrasound (US) is a good follow-up test to characterize
 - If a cyst has septa or internal echoes, a CT or MRI is warranted to evaluate



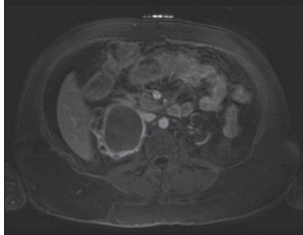
Imaging the Kidney – Renal Cysts		
Classification	Description	Risk of malignancy
Bosniak I	Round or oval, thin walled, no septations, 0-20 Hounsfield CT Units, no enhancement with IV contrast	Benign
Bosniak II	As above with a few septations and/or a few calcifications	Very low risk, f/u 6-12 months
Bosniak III	Thick wall, calcifications, Hounsfield density 0-20, no enhancement or nodules	60% chance
Bosniak IV	Thick wall, thick septations, coarse calcifications, Hounsfield Units >20, enhance with contrast	Consider malignant, proceed with work-up

Imaging the Kidney – Renal Cysts

- **Complex cysts contain blood or proteinaceous material, may have density of 50-80, still have thin walls and no nodules**
- **Magnetic Resonance Imaging (MRI) may be superior to CT, especially with subtraction techniques, in differentiating complex cysts from cystic neoplasm**




Complex structure with Septations
T1 weighted



Gadolinium, does not enhance
Needle aspirated – hemorrhagic cyst

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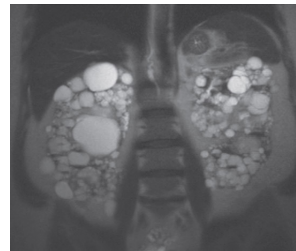


Imaging the Kidney

- **Autosomal dominant polycystic kidney disease (APCKD)**
 - Accounts for 6-10% of patients on dialysis in the US
 - Begins in the 3rd or 4th decade
 - Pain in the flank, back, or abdomen is the most common complaint
 - If no change in the kidneys or early cysts by age of 19 – unlikely affected
 - If no findings by age 40 – extremely unlikely to be affected



Sonography



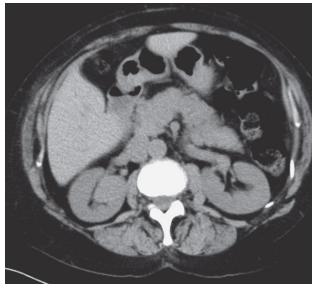
CT T2 weighted

Imaging the Kidney

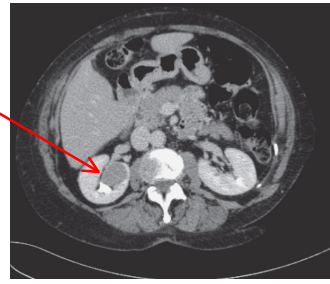
- **Benign Solid Renal Mass**
 - Renal adenoma is the most common benign neoplasm
 - Corticomedullary in location
 - Almost always < 2-3 cm in size
 - Demonstrate uniform enhancement on CT
 - Hamartomas (angiomyolipomas)
 - Solitary unilateral in women aged 30-50, often painful
 - Multiple, bilateral in those with tuberous sclerosis
 - The presence of Fat in the lesion ensures the diagnosis
 - < 4cm can be monitored, with surgery for symptoms or hemorrhage
 - Oncocytoma
 - Uncommon, benign tumor originating from the epithelium in the proximal collecting tubule
 - Oncocytic renal cell carcinomas occur, and surgery is usually needed to make the diagnosis
 - Fibromas, Myomas, Lipomas, and Hemangiomas are other uncommon, benign tumors of the kidney

Imaging the Kidney

- **Renal Malignancy**
 - **Renal Cell Carcinoma**
 - 85% of primary renal malignancies
 - Usually occurs in the 6th decade, male: female ~ 2:1
 - Flank pain, hematuria and flank mass seen in ~ 10%
 - Persistent, painless hematuria with negative cystoscopy and IVP needs to have a CT or MRI of the kidneys



Non-Contrast CT



Nephrographic Phase

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Imaging the Kidney

DIFFERENTIAL DIAGNOSIS OF CHRONIC RENAL PARENCHYMAL DISEASE

•No papillary/calyceal abnormality

•Diffuse parenchymal loss

•A. Bilateral

- Chronic glomerulonephritis
- Diffuse small-vessel disease
- Hereditary nephropathies

•B. Unilateral

- Renal artery stenosis
- Post irradiation
- Rare:
 - Hypoplastic kidney
 - Postobstructive atrophy

Focal parenchymal loss

- Infarct
- Previous trauma

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Renal Parenchymal Disease, including Renal Failure, Renovascular Disease and Transplantation



Imaging the Kidney

- **Papillary/calyceal abnormality**
- *Diffuse parenchymal loss*
 - Obstructive nephropathy
 - Generalized reflux nephropathy
- *No Parenchymal Loss*
 - Papillary necrosis
 - Tuberculosis
 - Medullary sponge kidney
 - Megacalyces
 - Pelvicalyceal cyst
- *Focal Parenchymal Loss*
 - Focal reflux nephropathy (chronic atrophic pyelonephritis)
 - Tuberculosis
 - Calculus disease

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Disease and Transplantation



Imaging the Kidney



B

Chronic pyelonephritis, dilated calyces, thinned cortex, calcifications,
And surrounding inflammation

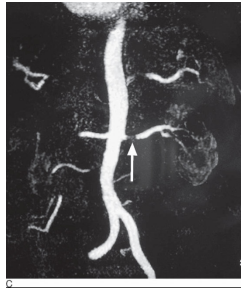
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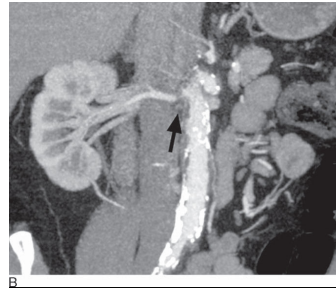
Imaging the Kidney – Vascular disease

- The most common cause of renal artery stenosis is atherosclerosis (70–80%)
- It occurs most often in men over the age of 50 with risk factors for vascular disease
- It usually involves the origin (ostial) or the proximal third of the renal artery
- Plaques within the aorta often cause ostial lesions
- Eccentrically placed atheromatous plaques in the proximal renal artery may also be seen
- If the kidney is small or there are distal plaques, there may be little benefit to intervention

MRA



CTA



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Imaging the Kidney

- Fibromuscular dysplasia (FMD) is the 2nd most common cause of renal artery stenosis (15-20%)
- FMD typically occurs in young women, and should be investigated in a young person with hypertension
- It is a heterogeneous group of conditions, with medial fibroplasia the commonest form
- There are multiple short stenoses and is seen as a 'string of beads' in the distal main artery and major branches, and may be bilateral
- Medial fibroplasia dilates easily at lower pressures with excellent long-term results, while the other types, characterized by smooth stenoses, do not always dilate satisfactorily.



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Imaging the Kidney

- **Renal artery aneurysms are rare**
 - They can be congenital, mycotic, post-traumatic, atherosclerotic, vasculitic or associated with FMD
 - Treat if causing hypertension or if there is a risk of rupture
 - Risk of rupture
 - aneurysm size (> 2.5 cm)
 - the absence of calcification
 - an association with pregnancy.
 - If the aneurysm is at risk of rupture, selective renal arteriography will be necessary to plan the appropriate treatment
- Renal infarction is usually due to thromboembolic occlusion of a renal artery
 - Principal source is atrial fibrillation
 - Others: aortic aneurysm, atheroma, vasculitis or trauma.
 - Clinical presentation: with pain and hematuria
 - Over time, the infarcted area decreases in size, with cortical scar formation
 - In vasculitis there are multiple small infarcts with patchy or wedge-shaped areas of altered perfusion