

# AMSER Rad Path Case of the Month:

## 44-Year-Old Man with Renal Incidentaloma

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# Patient Presentation

## Clinical history

- Asymptomatic 40M presents for evaluation of left kidney mass discovered on CT in 2018
- Interval changes noted on follow-up imaging this year

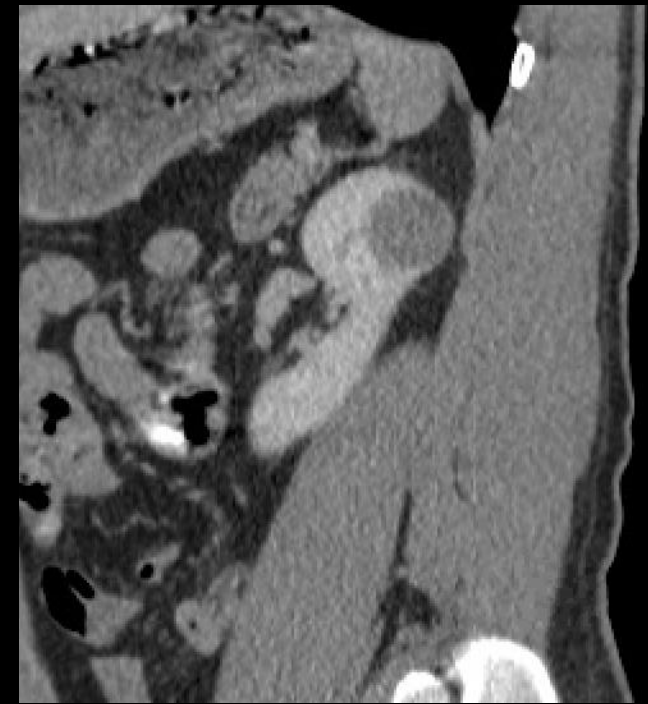
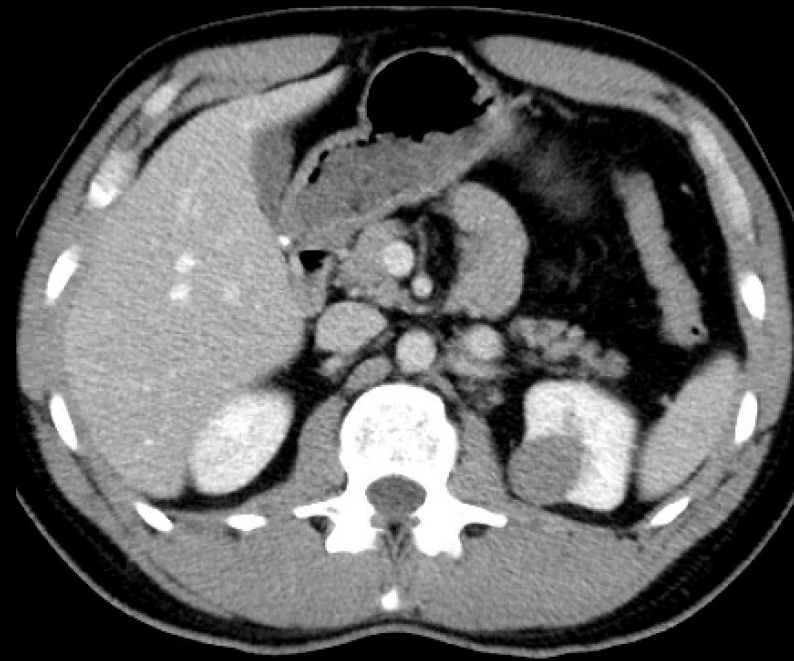
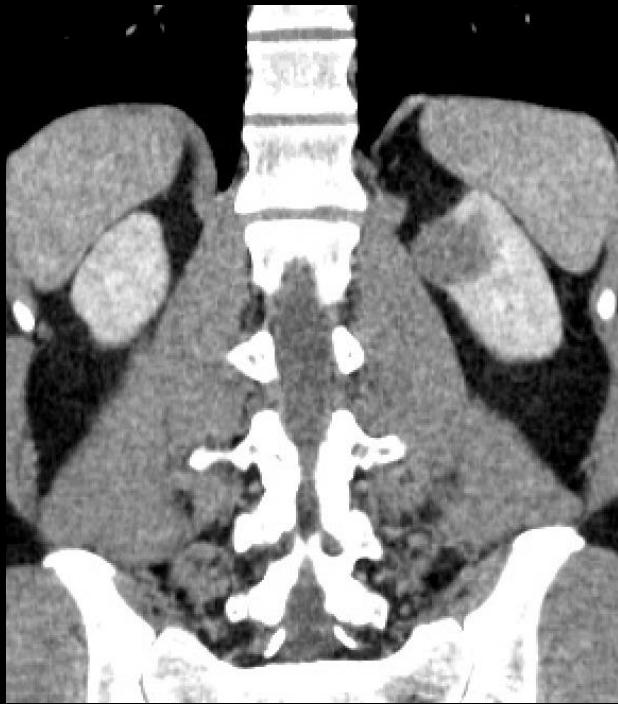
## Family history

- No known family history of renal cancer or familial syndromes

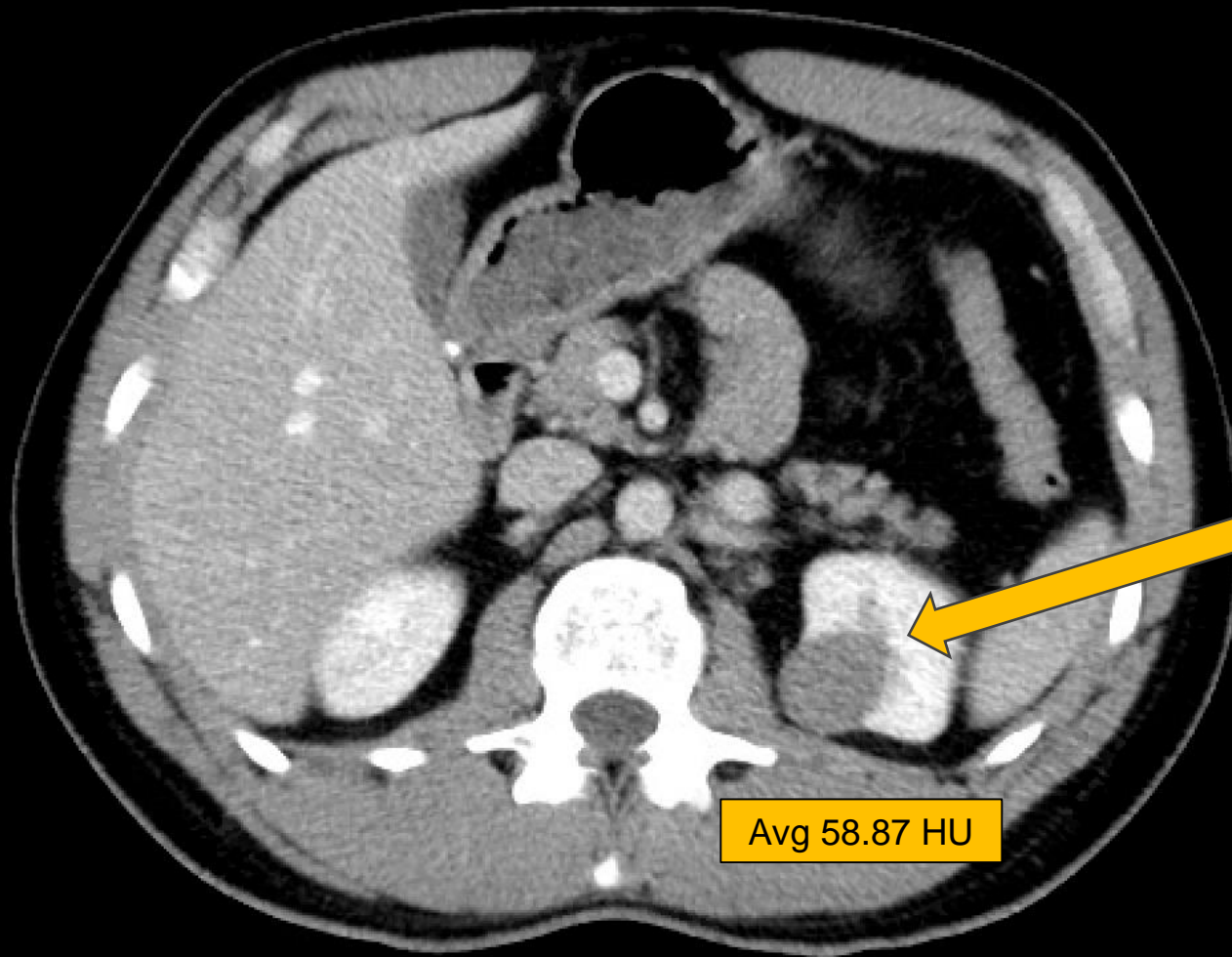
## Physical exam findings

- Unremarkable

# CT Abdomen with Contrast



# CT Abdomen with Contrast



Post-contrast CT demonstrates a mildly enhancing and heterogenous, exophytic mass in the upper pole of the left kidney

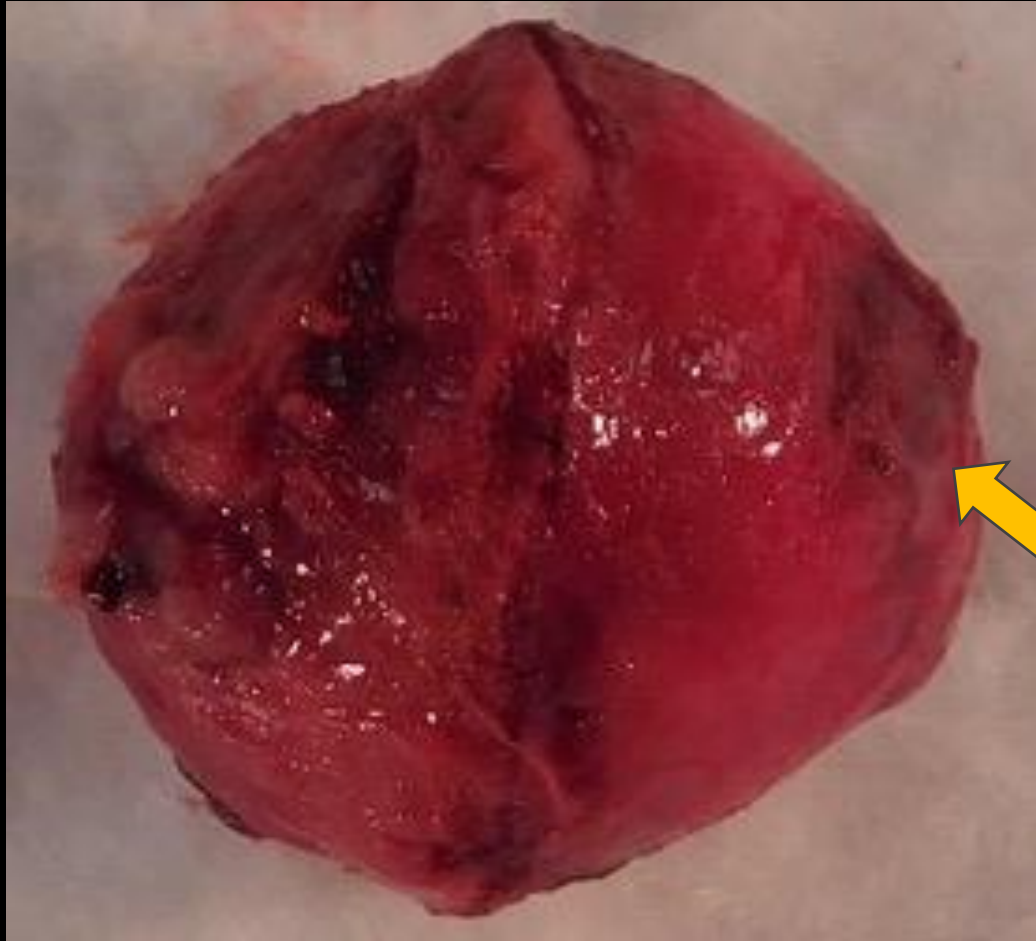
The lesion measures 3.2 x 2.3 cm which is increased from 0.8 x 0.5 cm in 2018

Avg 58.87 HU

# DDX (based on imaging)

- Renal Cell Carcinoma
- Oncocytoma
- Minimal Fat Angiomyolipoma (AML)
- Urothelial Mass of Pelvis and Collecting System
- Lymphoma
- Metastasis

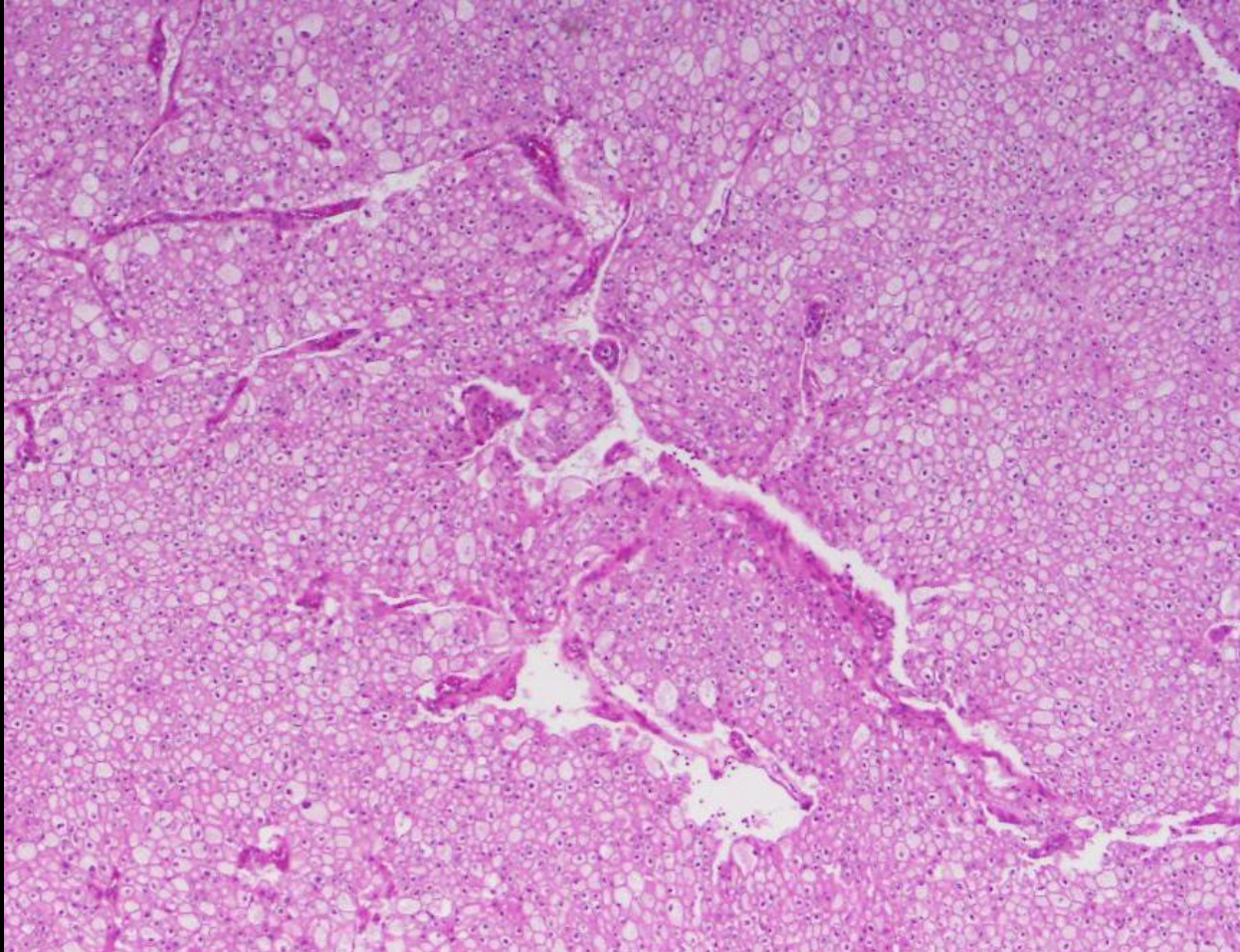
# Gross Pathology



3.1 x 2.9 x 2.9 cm left nephrectomy specimen. There is a clear delineation between the renal parenchyma on the left and mass on the right

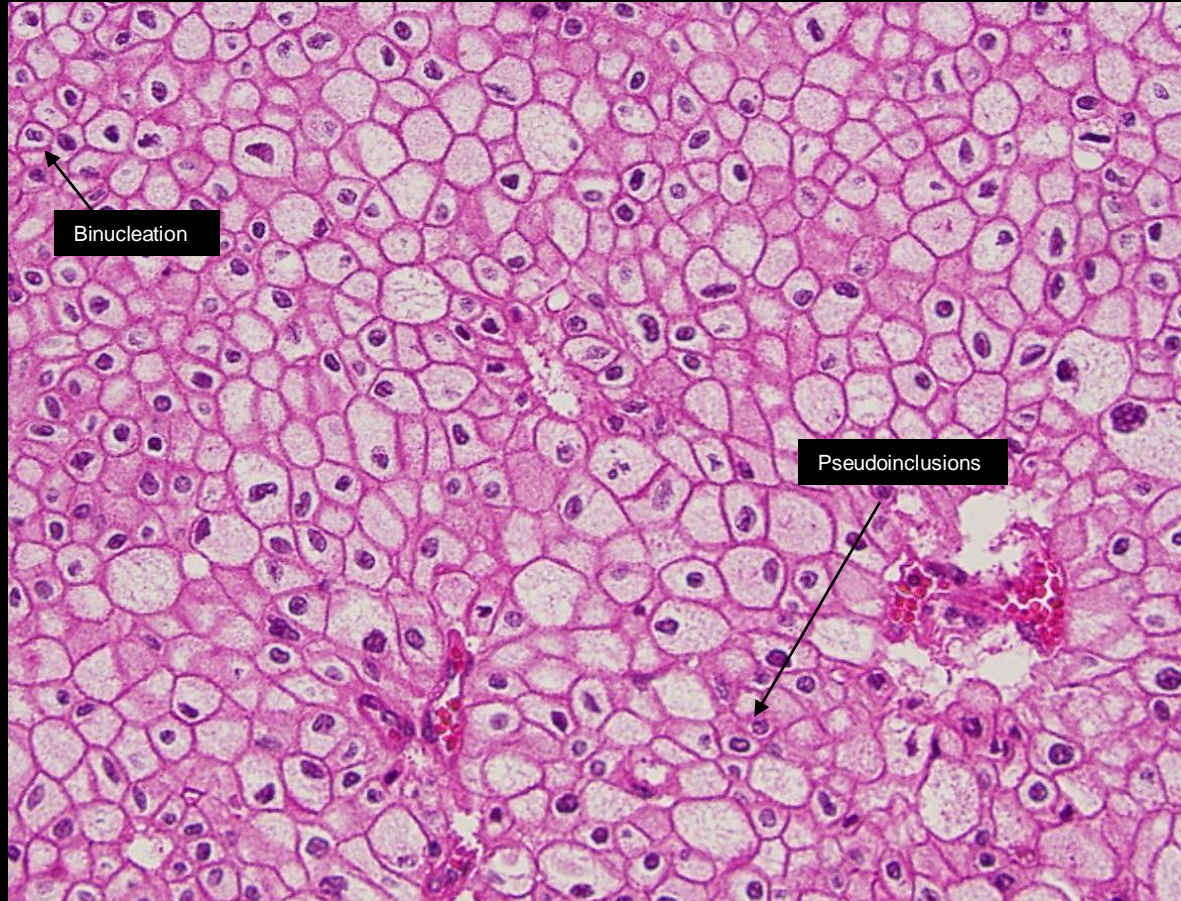
The mass is well-circumscribed and completely encapsulated apart from a 0.8 x 0.4 cm brown and friable defect, which may represent necrosis

# Micro Path



Low power view demonstrates sheets of large polygonal cells oriented along hyalinized stroma

# Micro Path

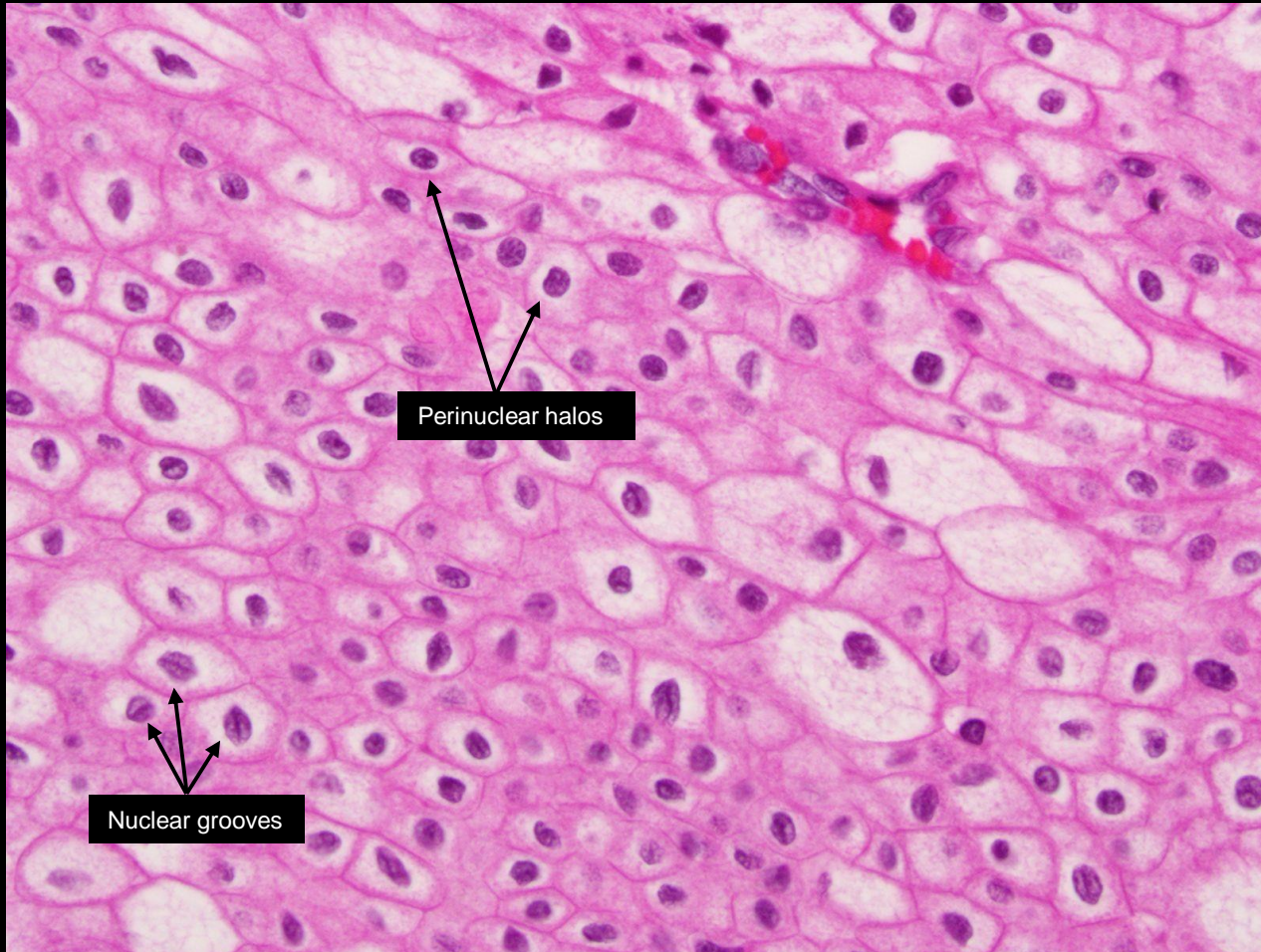


Cells demonstrate perinuclear halos within a flocculent cytoplasm that condenses around the edges, giving the appearance of thick prominent “vegetable-cell” borders

Cells are frequently binucleate and some have nuclear pseudoinclusions



# Micro Path



Nuclei have a koilocytic “wrinkled raisinoid” appearance

Perinuclear halos and nuclear grooves are visualized throughout the view

Note again the flocculent or finely reticulated cytoplasm, better seen at higher power

Final Dx:

Chromophobe Renal Cell Carcinoma

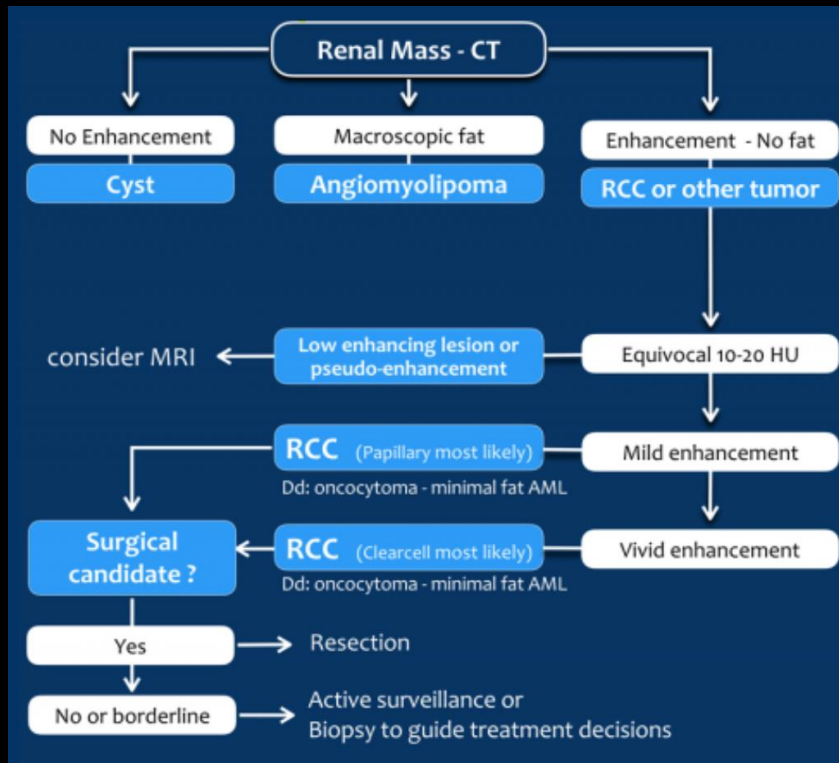
# Case Discussion

## Renal Incidentaloma Work-up

- Indeterminate masses require additional imaging
  - Too small to conclusively interpret as benign or malignant
  - Improper imaging protocol to sufficiently assess all relevant features
- MRI and multiphase renal protocol CT, with and without contrast
  - Similar diagnostic accuracy, though CT is preferred when masses have cystic components
- Increasing tumor size, male gender, and contrast enhancement are strong predictors for malignancy and higher tumor grade
  - Should be considered when deciding on continued surveillance vs biopsy/excision

# Case Discussion

## CT Characteristics To Describe Masses:



Adapted from "Solid Masses," by Reingard, van der Zon-Conijn, and Smithuis. TheRadiologyAssistant.

- Presence of fat

- Density less than water or soft tissue
- Suggestive of angiomyolipoma

- Enhancement

- Increased signal > 20 Hounsfield units after contrast
- Absence is likely benign (cyst)
- Presence is concerning for malignancy

- Cystic components

- Findings are characterized by the Bosniak criteria
  - Septa and/or calcifications – benign
  - Wall irregularity and/or enhancement – likely malignant

# Case Discussion

## Background

- Chromophobe variant is the third most common subtype of renal cell carcinoma (RCC)
  - Arises from intercalated cells of the collecting system
  - Accounts for 5-7% of all RCCs
- Mean incidence in the 6<sup>th</sup> decade with no gender predilection
- Generally presents at an earlier stage with a better prognosis than other RCC variants
  - 5- and 10-year cancer specific survival (CSS) are 93% and 88.9% respectively

# Case Discussion

## Radiologic Features

- Solid and sharply demarcated from renal cortex, may be lobulated
- Contrast enhancement is often homogenous on CT and MRI
- Most are hypovascular to renal cortex with moderate contrast uptake (80-100 HU) on CT
- Distinguishing from other mimics:
  - Tends to be more homogenous and less intense than clear cell variant
  - Indistinguishable from oncocytomas on imaging
  - Lacks low signal on fat-suppression MRI seen in AML or clear cell variant

# Case Discussion

## Histopathologic Features

- Arranged in sheet or alveolar pattern around thick hyalinized vessels
- Two cytomorphologic variants of cells:
  - Classic: Large polygonal cells with pale reticulated cytoplasm and koilocytic nuclei
  - Eosinophilic: Round cells with dense, pink, granular cytoplasm and round nuclei
- Central nuclei with perinuclear clearing results in prominent cell borders with "plant-cell appearance"
- Nuclei are often binucleate or grooved and pseudoinclusions can sometimes be present

# References:

- ACR Appropriateness Criteria® Indeterminate Renal Mass. *American College of Radiology*. Retrieved October 15, 2020, from <https://acsearch.acr.org/docs/69367/Narrative/>.
- Pierorazio, P., Campbell, S. (2020). Diagnostic approach, differential diagnosis, and management of a small renal mass. In S. Shah (Ed.), *UpToDate*. Retrieved October 15, 2020, from <https://www.uptodate.com/contents/diagnostic-approach-differential-diagnosis-and-management-of-a-small-renal-mass>
- Reinhard, R., van der Zon-Conijn, M., Smithuis, R. (2016). Kidney: Solid masses. *The Radiology Assistant*. Retrieved October 12, 2020, from <https://radiologyassistant.nl/abdomen/kidney/solid-masses>.
- Atkins, M., Choueiri, T. (2020). Epidemiology, pathology, and pathogenesis of renal cell carcinoma. In S. Shah (Ed.), *UpToDate*. Retrieved October 17, 2020, from <https://www.uptodate.com/contents/epidemiology-pathology-and-pathogenesis-of-renal-cell-carcinoma>
- Muglia, V., Prando, A. Renal cell carcinoma: histological classification and correlation with imaging findings. *Radiol Bras*. 2015;48(3):166-174.
- Anderson, D., Tretiakova, M. (2017) Chromophobe. *PathologyOutlines.com*. Retrieved October 7, 2020, from <http://www.pathologyoutlines.com/topic/kidneytumormalignantccchromo.html>
- Granter, S., Renshaw, A. Fine-needle aspiration of chromophobe renal cell carcinoma. *Cancer*. 1997;81(2):122-128.