Peer Reviewed **Article of Interest**

A PICTORIAL PRESENTATION OF CT KIDNEY LESIONS IN ADULTS, PARTICULARLY KIDNEY CANCER: PART 5

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ABSTRACT
There are many pathologies that affect the kidneys. The focus of this paper is computed tomography of kidney cancer. Other imaging modalities for kidney pathologies are briefly discussed. A few examples of benign tumours and masses are presented. The bulk of the paper is a pictorial presentation of kidney cancer and its spread to other organs and bones. A range of unenhanced and enhanced CT images are included for self-assessment.

Keywords: renal cell carcinoma, lymphadenopathy, oncocytoma, imaging modalities

LAY ABSTRACT
Imaging of the kidneys is done to find out whether a mass in a kidney may be a benign tumour, cyst or cancer. If it is a cancer then it is important to check whether there is spread to other organs. Examples of computed tomography (CT) are used to describe the different patterns of benign and cancer lesions.

12. KIDNEY CANCER TREATMENT, FUSION IMAGING AND DUAL ENERGY CT

There are several methods used for kidney cancer treatment. Surgery is used for radical or partial nephrectomy.[43] Minimally invasive surgery includes laparoscopic and robotic surgery.[43] Non-surgical tumour treatment includes radiofrequency ablation, cryoablation, and therapies using medication.[43] Radiation therapy may be used. Drugs (e.g., serofenib, axitinib, pazopanib), which are given by mouth or via the bloodstream, can be used to treat kidney cancer. These drugs are called systemic therapies for use in targeted therapy, immunotherapy, and chemotherapy.[43]

In terms of imaging modalities literature reports the role of US/CT fusion imaging to guide radiofrequency ablation of residual tumour.[44] Dual energy CT (DECT) may be used for grading of clear cell renal cell carcinoma[45] and for imaging of renal masses.[16, 46]

13. KEY POINTS

- There are no available routine tests to detect kidney cancer
- Imaging (e.g., ultrasound, CT, MRI, PET/CT) plays an important role in the visualisation of renal pathology
- There are benign renal tumours (e.g., oncocytoma and angiomyolipoma) and malignant tumours (e.g., renal cell carcinoma)
- Renal pathology may be in one kidney or both
- Multiphase CT of the abdomen is used to visualise lesions in the kidney and spread to adjacent organs
- Mean attenuation values seem to be reliable in terms of determining which renal masses on unenhanced CT need further evaluation
- Enhancement values in a ROI that are more than 10-12 HU above that of pre-contrast measurements are considered significant
- RCC shows significant enhancement above pre-contrast measurements: typically over 100 HU in the corticomedullary phase (also called angionephrographic or late arterial phase) and 60 HU in the delayed phase
- A routine should be followed for CT scans to assess spread of kidney cancer to other organs, lymph nodes and bones
- Important to examine the venous system (e.g., renal veins and IVC) for spread
- Important to examine the adrenals for spread
- The position of both the spleen and the pancreatic tail usually changes to lie in the left renal bed in left post-nephrectomy patients
- The bowel usually occupies the right renal bed in right post-nephrectomy patients

14. CONCLUSION

Early detection of kidney cancer is important in terms of treatment and management of patients. There are no available routine tests to detect kidney cancer. Radiology imaging thus plays an important role to visualise renal pathology. Radiographers, as members of the imaging team, should be able to perform pattern recognition of kidney anatomy in terms of size, shape and anatomical position.

This paper focuses on CT images of

kidney cancer and its spread to other organs, lymph nodes, venous system and bones. A range of examples are presented to highlight the different patterns of benign and malignant masses seen on unenhanced and enhanced CT images.

15. SELF-ASSESSMENT

When assessing images it is important to remember that an image's pattern may be normal including normal variants and congenital abnormalities or abnormal indicating pathology. Figures 15a to 15w are a range of examples covered in the text and the figures in the different sections.

Kindly email sorsaoffice@gmail.com to request a copy of the self-assessment answers (for members only).

16. ACKNOWLEDGEMENTS

Professor PJ Pickhardt from Wisconsin University is thanked for providing the oncocytoma CT images. Professor D Kim from Wisconsin University is thanked for providing the lymphoma CT images.

17. CONFLICT OF INTEREST

The authors declare no conflict of interest.

18. CONTRIBUTIONS OF THE AUTHORS

JHB (LSG), RvdV (NMU) and LM (Dbn) contributed to the draft manuscript. LM edited the manuscript. Annotation of the images (figures) was done by LM. JHB wrote the legends for all the figures.

Figure 15a

Figure 15b

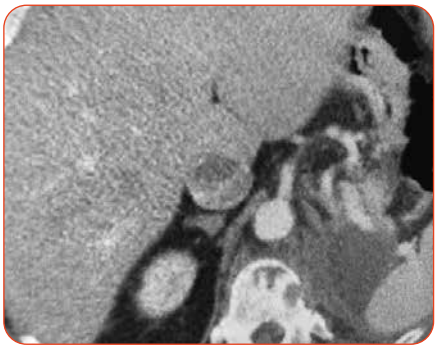

Figure 15c

Figure 15d

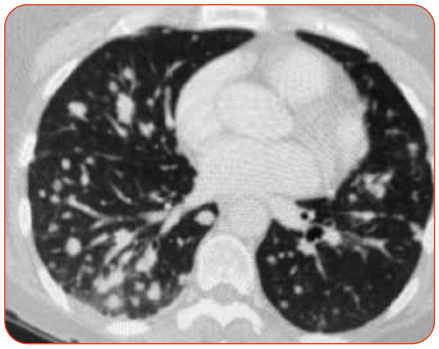

Figure 15e

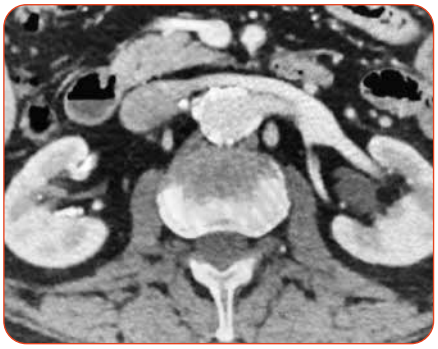

Figure 15f

Figure 15g

Figure 15h

Figure 15i

Figure 15j

Figure 15k

Figure 15l

Figure 15m

Figure 15n

Figure 15o

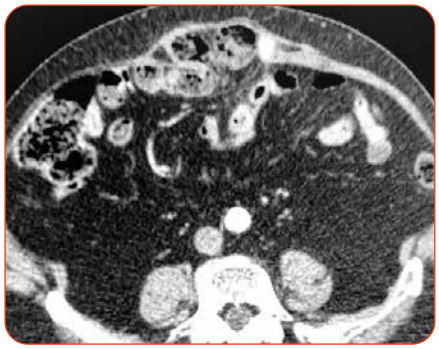

Figure 15p

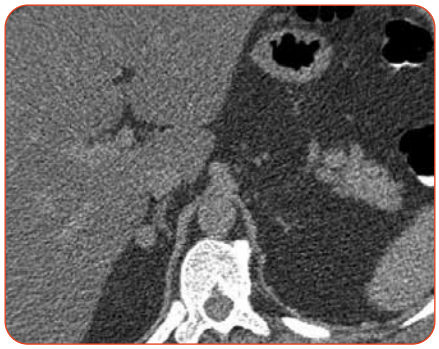

Figure 15q

Figure 15r

Figure 15s

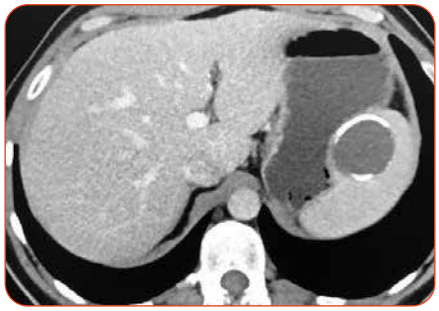

Figure 15t

Figure 15u

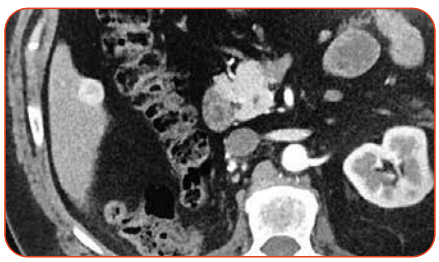

Figure 15v

Figure 15w

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