

Renal Tumors

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Renal Tumors

* Classification of Renal tumors:

I. Benign tumors:

- Cortical adenoma.
- Oncocytoma.

II. Malignant tumors:

- Renal cell carcinoma.
- Wilms tumor.

Renal adenoma : accidentally discovered & can not be differentiated from carcinoma and should be treated as cancer .

• **Renal Oncocytoma** : well encapsulated with central stellate scar

• grade 1

• grade 2

• grade 3 --- nuclear atypia

the diagnosis is pathological ;

it is premalignant & / or associated with other malignancies .

* *Angiomyolipoma*

(renal hamartoma) : rare benign tumour.

* U/S & CT scan:-are diagnostic due to high fat content that is shown as

*High intensity lesion on U/S

& negative density (- 20 to - 80 HF) on CT scan.

*The management is follow up if bleeds or size increase == surgical treatment by partial nephrectomy or embolization .

Renal cell carcinoma : RCC

(Adenocarcinoma of kidney ,hypernephroma ,clear cell carcinoma , alveolar carcinoma)

3 % of adult cancer

Aetiology :

-- cigarette smoking is the only proved risk factor .

-- structural changes of **chromos.**

3 in both sporadic & hereditary forms .

pathologically it originates from proximal renal tubules.

Grossly == yellow to orange due to lipid content .

Histopathology == shows mixed adenocarcinoma containing clear cells granular cells & occasionally sarcomatoid appearing cells .

Spread == direct spread through the renal capsule or direct extension into renal veins .

The most common site distant metastasis is the lung , liver , bone , lymph nodes and opposite kidney .

Tumor grades are 4 grades the highest the grade the worst prognosis .

Tumor staging :-

Stage I – confined within the kidney parenchyma .

Stage II - = = gerota fascia (including the adrenal) .

Stage IIIA- involves main renal vein or IVC .

Stage IIIB- = regional lymph nodes .

Stage IIIC = both local vessels & L.N .

Stage IV A - = adjacent organs (colon , pancreas) .

Stage IV B - distant metastasis .

Clinical findings :-

Mostly **asymptomatic** & discovered accidentally during imaging studies

A – symptoms & signs :+ gross hematuria

+ flank pain

+ palpable mass

+ metastatic symptoms as cough dyspnea

,convulsion , bone pain .

B – paraneoplastic syndrome :

+ erythrocytosis : due to increase erythropoitin or renal hypoxia .

+ hypercalcemia : = parathyroid hormone related peptide secretion .

+ hypertention : = increase rennin(refractory to drugs & respond after nephrectomy) .

+non metastatic hepatic dysfunction (stauffer syndrome) due to hepatotoxic product of tumour.

If nephrectomy fails to correct the syndrome === residual tumour .

C – laboratory findings

+ anemia = normochromic (of chronic disease)

+ GUE = hematuria .

+ ESR elevated

+ those of paraneoplastic syndrome.

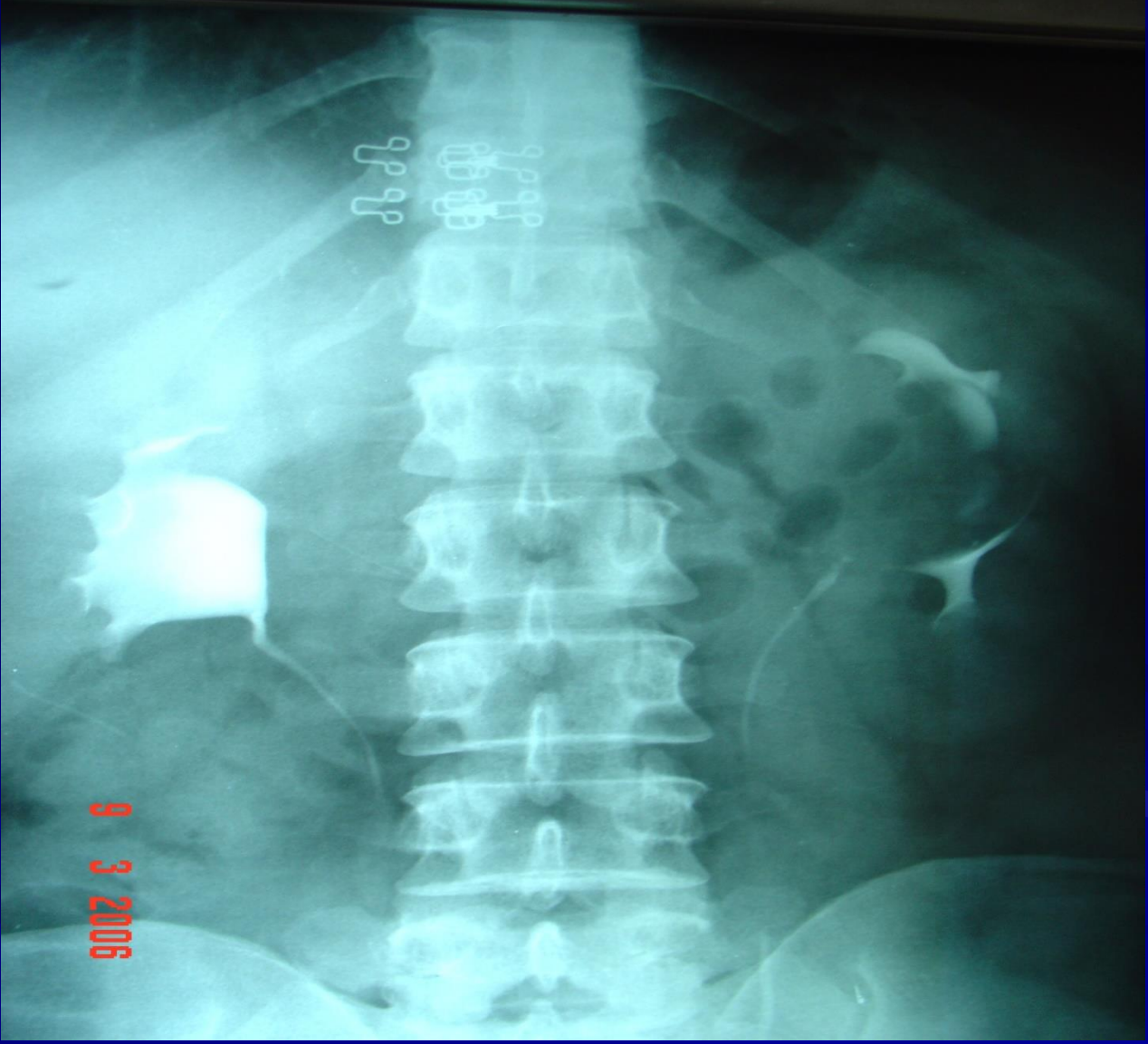
D – radiological findings:

U/S = can differentiate simple renal cyst from renal tumour.

IVP = show calcifications & function of kidney .

CT scan = method of choice , more sensitive and more accurate , tumour shown as :

- 1 - enhanced with contrast media .
- 2 – amputation of collecting system .
- 3 – calcification .
- 4 – ill defined border .
- 5 - lymph node enlargement.

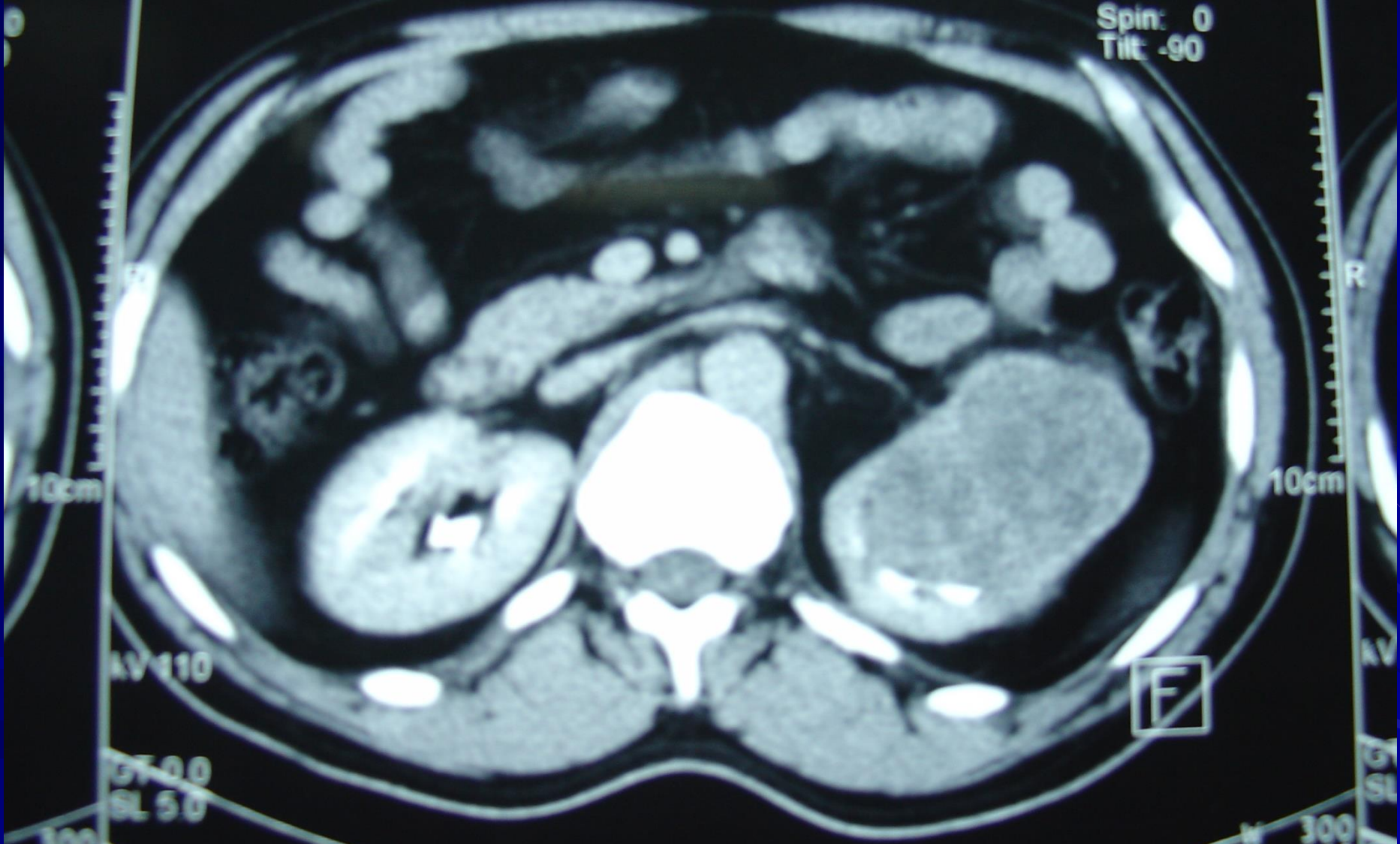


9 3 2006

502 IMA 3
MPR 2

502 IMA
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Spin: 0
Tilt: -90



CONTRAST

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02-MAY-2005

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SOMATOM PLUS 4
VC10C
H-SP

H

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D1	1	5.4
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D1	2	5.7
Ar	2	10

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01-JAN-1957
03-MAY-2005
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TP -535.7

IBN ALBITAR C.S.C.
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H-SP-CR

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An 1 171
Di 2 4.0
An 2 63

SPI 10

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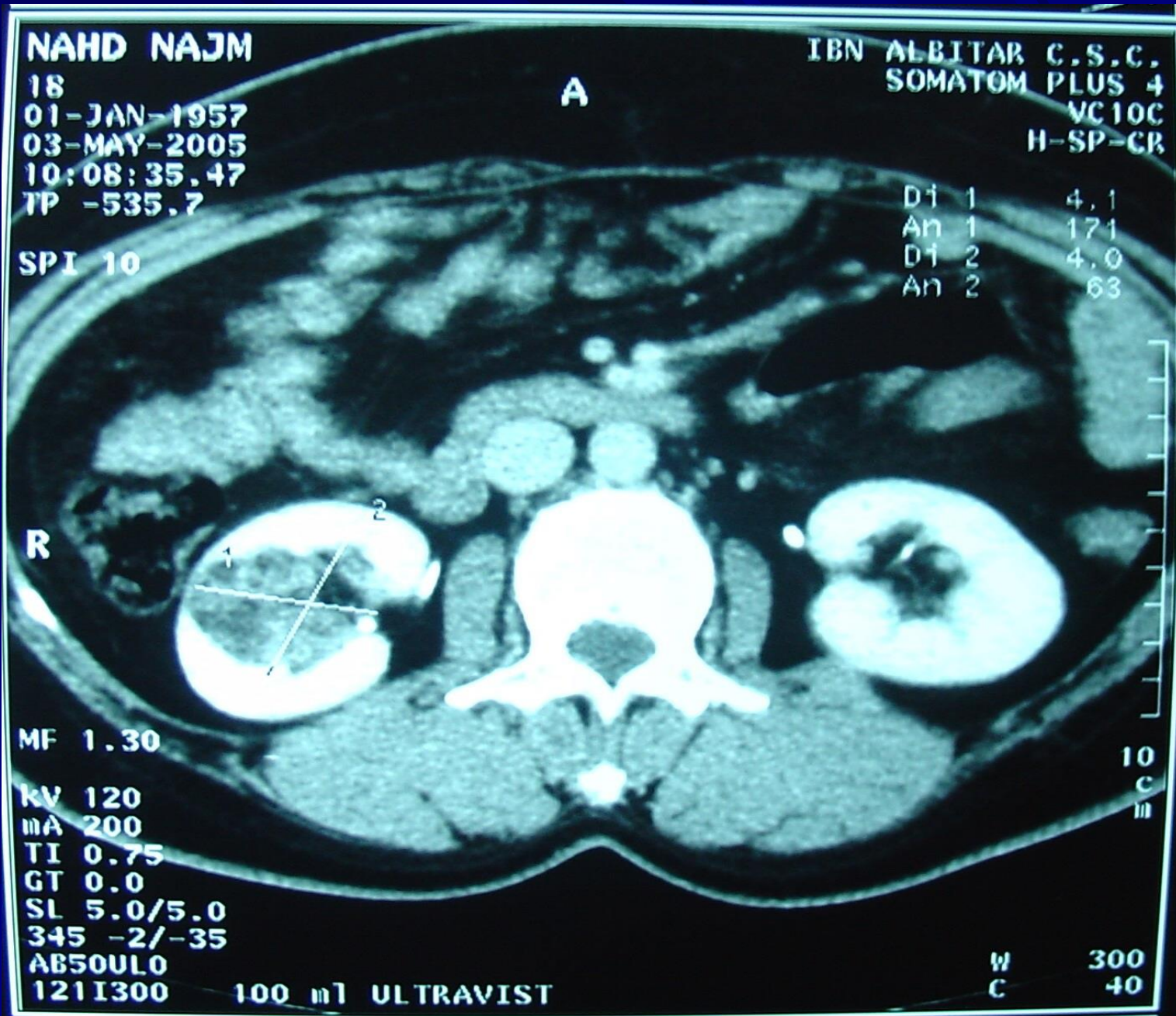
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chest CT or brain CT may be required for metastasis .

Chest X-ray = to exclude pulmonary metastasis .

Renal angiography : of limited role .

Radionuclide imaging = if patient allergic to contrast media

= for bone metastasis .

MRI = superior to CT in assessing inferior vena cava involvement.

FNA = is of limited role, to differentiate secondaries to kidney & abscess.

Differential diagnosis

- 1 – simple renal cyst = U/S can diagnose it (98% accuracy).
- 2 – renal abscess = fever , pain , leukocytosis . (FNA can differentiate)
- 3 – angiomyolipoma =hypotense on CT & hyperechoic on U/S.
- 4 – benign renal mass .
- 5 – transitional cell carcinoma of renal pelvis .
- 6 – adrenal cancer = CT +/- MRI .
- 7 – metastasis to kidney == FNA can be helpful .

Treatment:-

1 – localized disease (I , II , IIIA):

- * **radical nephrectomy** is the primary treatment
==>removal of the kidney & gerota , the adrenal , proximal half of ureter& lymph nodes up to transection of renal vessels (regional LN removal still controversial).
- ***embolization** (angioinfarction) for
=large tumour when it is difficult to reach the renal vessels early during nephrectomy .
= palliative for non resectable tumour .
- * **radiotherapy** (radioresistant) controversial as adjuvant or neoadjuvant

- * laparoscopic radical nephrectomy.
- * partial nephrectomy for small tumour < 4 cm & in single kidney .
- * watchfull waiting : in highly selected patient especially old age < 3 cm

2 – disseminated disease :

- * palliative radical nephrectomy for severe hemorrhage , unremitting pain , paraneoplastic syndrome .
- * combined nephrectomy & removal of metastatic foci .
- * radiotherapy as palliative because it is radioresistant .
- * hormonal therapy = not so usefull .
- * chemotherapy = not so usefull .
- * radioimmunotherapy = promising .
 - biological response modifier = promising .