LEIOMYOSARCOMA OF THE KIDNEY — A CASE REPORT WITH REVIEW OF THE LITERATURE

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Malignant smooth muscle tumours of the kidney are uncommon and to our knowledge no example of renal leiomyosarcoma in Singapore and Malaysia has been documented. Therefore, the following case report is presented. In addition, all the cases recorded in the literature are reviewed and some clinical features outstanding in these patients brought out.

CASE REPORT

The patient was a Chinese female, aged 40, who had noticed a swelling progressively increasing in size in the right side of the abdomen for the previous six months. Apart from losing about 20 pounds in weight during that period, there had been no associated symptoms. In particular there were no symptoms referable to the urinary tract. Recently she had experienced anorexia and felt generally unwell. At no time had she felt pain or was there haematuria.

On clinical examination she was seen to be ill and there was a very obvious swelling to be seen on the right side of the abdomen and which moved with respiration. The mass was easily ballotable from the loin, somewhat lobular in shape and firm in consistency. It was not tender to palpation. Her general appearance was consistent with her history of weight loss. There were no other physical signs of significance. Her blood pressure was 100/60.

On clinical grounds it was considered likely that the abdominal swelling was arising from the right kidney. An intravenous pyelograph showed on the control film areas of calcification in the site of the right kidney, which failed to concentrate the dye, whereas the function of the left kidney was good with a normal pyelographic picture (Fig. 1).

The radiological appearances were those of either a calcified renal carcinoma or calcification in a kidney destroyed by tuberculosis, a so-called auto-nephrectomy.

During the period that the patient was in hospital under investigation and being prepared for operation the swelling appreciably increased in size.

Investigations: Urine culture for myobacterium tuberculosis—negative. Urine culture—sterile. Blood haemoglobin—65%, white cell count—6,300/cu. mm. Blood urea—24 mgm.%. Urine microscopy—no red cells, occasional white cells, occasional epithelial cells, no casts, no crystals.

Exploration of the right kidney was performed on the 10th June 1966. A long right paramedian incision was made and a transperitoneal approach

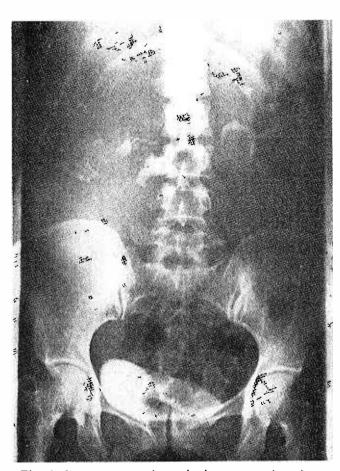


Fig. 1. Intravenous pyelograph shows normal pyelonephric appearances on the left side, no renal function on the right and calcification in the renal area on that side.

to the right kidney carried out. The renal origin of the abdominal mass was confirmed, and increased vascularity of the perinephric tissue was noted. Right radical nephrectomy was performed from which the patient made an uneventful recovery.

PATHOLOGY

On gross examination, the specimen consisted of a large firm, nodular mass, measuring 15 cm. × 12 cm. × 9 cm. the reniform shape still being roughly preserved (Fig. 2). It appeared to be enclosed by the remaining compressed renal tissue, from which the tumour mass could be readily separated. The cut-surface revealed firm, pinkish tumour tissue which was divided by fibrous strands into lobules of varying sizes (Fig. 3). Occasional areas of central necrosis and calcification were also noted. A length of normal looking ureter could be seen emerging from the mass, near its centre (Fig. 2).

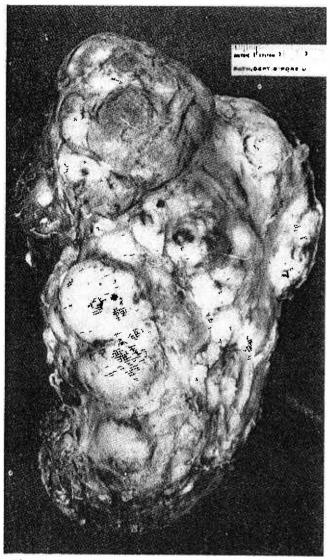


Fig. 2. External view of the specimen shows nodular appearance. A short stump of ureter is attached to the middle of the left margin.

Histologically, the tumour consisted of irregularly arranged spindle-shaped cells which showed whorled patterns in some areas. The nuclei were round, oval, elongated and of varying sizes; they were hyperchromatic and they exhibited numerous mitoses. Some multinucleated cells were also present. Myofibrils were demonstrated in the cells by means of special stains—Phosphotungstic acid haematoxylin, Mallory's Trichome (Figs. 4 and 5). Compressed and fibrotic renal structure was discernable at the periphery of the main tumour mass. Metastatic lesions were identified in a few lymphnodes removed together with the right kidney.

DISCUSSION

A survey of the literature shows that the case of renal leiomyosarcoma described by Pick (1893) in a dissertation quoting Ranke is probably the first such lesion reported. There is little doubt about its rarity (Mintz, 1937; Bruce and McNaught, 1954), as only 41 other cases, excluding the present one, have been documented (Table 1). Kerr's (1954) case was associated with a leiomyosarcoma of the stomach, and Leopold and Mogg (1964) described a combined carcinoma-leiomyosarcoma in an adult kidney.

Leiomyosarcoma of the kidney usually manifests itself between the ages of 40 years and 70 years (Bazaz-Malik and Gupta, 1966). The oldest patient was an 86-year-old woman (Tetelman and Lisa, 1945) and the youngest a 3-year-old boy (Pick, 1893). It occurs more frequently in females than in males, in the approximate ratio of 2:1. The right kidney is involved about twice as often as the left, and there is one instance where both kidneys are affected (Bhende, 1952).

Lumbar pain or tenderness and a lumbar mass on the side of the involved kidney are the commonest presenting symptoms. Haematuria occurs occasionally. Often only one of the mentioned symptoms presents itself, as in our patient whose complaint was that of a lumbar mass with no abdominal discomfort or haematuria. The duration of symptoms is usually from one month to one year, though histories as short as nine days (Briggs, 1955) and as long as ten years (Leopold and Mogg, 1964) have been recorded.

The period of survival after operation is usually not more than one year. Only one patient was reported to be well eighteen months after operation (Smith, 1935). Our own subject is apparently well in the sixth post-operative month.

The origin of this tumour is obscure. The commonly implicated sites are the renal capsule, smooth muscle rests in the renal parenchyma,

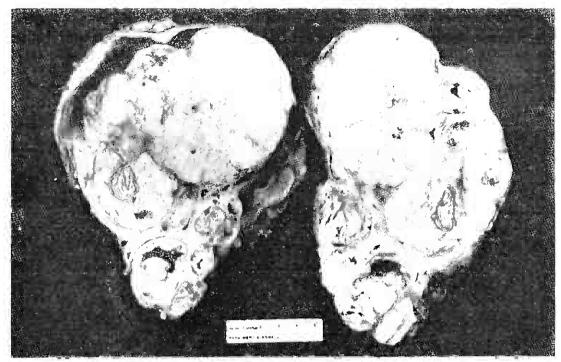


Fig. 3. Cut-surfaces show lobulated pattern with central areas of necrosis.

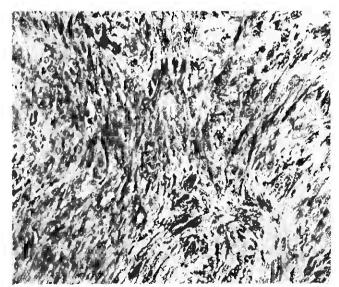


Fig. 4. Low-power photomicrograph shows irregularly disposed spindle cells. In some parts attempts at whorling are visible. (Phosphotungstic acid haematoxylin \times 150).

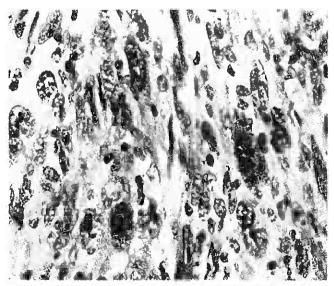


Fig. 5. High-power photomicrograph shows hyperchromatism and pleomorphism of round, oval and elongated nuclei. A few multinucleated cells are also present. (Haematoxylin and Eosin \times 500).

TABLE I
REPORTED CASES OF RENAL LEIOMYOSARCOMA

Author	hor	Year	Age	Sex	Symptoms	Duration	Side	Physical and Gross	Gross Pathological	Matagrasic	
		, [Smorth		anic	Findings	Findings	Metastasis	Survival
Pick quoting Ranke 1893		~	33	Σ	Symptoms of hydro-	9 months	T	N.S.	Z.S.	N.S.	N.S.
1919	1919	_	72	Σ	Right Lumbar	3 months	ĸ	Z.S.	Tumour upper pole	Z.S.	v: Z
Brandt 1919	1916	_	89	Σ	N.S.	Z.S.	J	Z.S.	$15 \times 10 \times 7$ cm. Tumour upper pole	l ungs, liver	S. Z
Mackenzie and 1929	192	6	Z S	Z.S.	s. Z	Z.S.	Z.S.	Z.S.	$13 \times 11 \times 10$ cm. N.S.	, Z	
1929	192	0	26	Ľ	Right Lumbar pain	14 months	24	Z.S.	Large mass lower pole	 1ocally infil-	Z.
Crosbie and 1932	1932	Λ.	35	Щ.	Right Lumbar pain	6 months	ĸ	Z.S.	Large mass upper pole	rating N.S.	υ: Ζ
Cooke 1933	193	~	09	(I.,	Right Lumbar mass	5 years	ĸ	Š.	Large mass whole	Z.S.	Died ! vear
1933	193	~~	71	Σ	Haematuria	Z.S.	Z.S.	S. Z.	kidney N.S.	\$\frac{\sigma}{Z}	r oper vear
Smith 1935	1935		Z.S.	Z.S.	Z.S.	z.s.	si Z	ý. Ž	Z.S.	Locally infil-	i operation Well 14 vears
Mintz 1937	193.		45	LL,	Left Lumbar pain	6 months		Deformity of	8 cm, in diameter	ಯ	r opei
Mintz 1937	1937	~~~	4	<u>{1.,</u>	Right Lumbar pain	i year	ĸ	calyces Hydronephrosis	628 gm.	S.Z	S Z
1937	1937		37	Σ	Abdominal mass	5 months	_	of ureters Left abdominal	Inoperable bilateral	o, Z	Died 5 days
1937	1937		46 55	ω Σ	Haematuria Right Lumbar pain Right Lumbar pain	5 weeks 2 weeks	X X	mass Filling defect Defect of	tumours Mass lower pole Tumour whole kidnev	N.S. Mesenteric	after operation N.S.
Weisel, Dockerty 1943	1943		Z.S.	S.S.	Haematuria N.S.	Z.S.	Z.S.	calyces N.S.	13 × 9 × 5 cm. N.S.	lymph nodes	5 V. Z.
weisel, Dockerty 1943	1943		s. Z	s; S	S.Z.	Z.S.	Z.S.	Z.S.	N.S.	ý. Z	si Z
Weisel, Dockerty 1943	1943		S Z	Z.S.	Z.S.	Z.S.	Z.S.	N.S.	Z.S.	S.Z.	si. Z
Crabtree Tetelman and Lisa 1945	1944 1945		43 86	ᄕᇄᅜ	Left Lumbar mass Right Lumbar mass	S.S.	コペ	si si Z Z	1000 gm. 1000 gm.	N.S. Mesenteric	s si ZZ
Tetelman and Lisa 1945 Bagolan 1950	1945 1950		47	<u>г</u> .	Autopsy material Right Lumbar pain	z z z	기 &	S.S. S.S.	2×2 cm. Masses both kidneys	lymph nodes N.S. N.S.	N.S. Died I day
Blum and Frehling 1951	1951		25	ĹĽ,	Left Lumbar lump	2 years	<u> </u>	Deformity of	Large mass lower pole	S.Z.	after operation
Blum and Frehling 1951	1951		65	(L,	Right Lumbar lump	4 years	<u>~</u>	Displaced ca-	Large mass upper pole	Ś.Z.	after operation N.S.
Petkovic 1951	1951		48	Σ	Left Lumbar pain	Z.S.		beformed lower	Lower pole	S. S.	Died 6 months
Petkovic 1951	1951		52	江	naematuna Left renal colic. Kaematuria	z.s.	٦	calyces Deformed upper calyces	Upper pole	œ.z	5 - 5

TABLE 1 (Continued)

REPORTED CASES OF RENAL LEIOMYOSARCOMA

ż	Author	Year	Age	Sex	Symptoms	Duration	Side	Physical and Pyelogram Findings	Gross Pathological Findings	Metastasis	Survival
26	Kretschmer	1952	62	Z	Right Lumbar pain	3 months	R	Filling defect	Adherent large mass	Z.S.	Well 9 months
27	Bhende Bhende	1952	30	ΣΣ	Autopsy material Autopsy material	s; s; ZZ	78	N.S. N.S. N.S. N.S. N.S. N.S. N.S. N.S.	6 cm. in diameter Kidney	Si Si	
30	Bhende Bruce and Mc- Naught	1952	10 59	다.다.	Autopsy material Loss of weight Right Lumbar lump	s; s; ZZ	Both	N.S. No function at all	24 × 14 × 8 cm. Multiple nodules Large mass lower pole	N.S. Inguinal and a xillary	N.S. Recurrence after 9 months
31	Higbee and Atkin	1954	63	<u></u>	Right Lumbar pain	6 weeks	~	Double right kidney with	Mass upper pole	N.S.	S. Z.
32	Lazarus and Fried-	1954	39	Σ	Paintess haematuria	6 months	~	mass Defective filling	20 × 15 × 15 cm.	Lung, liver	Died 71 weeks
33	man Kerr	1954	44	Σ	S.Z.	Z.S.	~	upper pole	150,	s.Z.	alter operation N.S.
۶. م.	Briggs	1955	35	Ц	Profuse painless	9 days	≃.	Gross deforma-	N.S. 1 .: 5 CM.	Z.S.	S.Z.
35	Briggs	1955	75	Ľ	Right Lumbar lump	3 weeks	~	Large mass in	Growth at lower pole	Locally infil-	Well 10 months
36	Briggs	1955	08	<u>т</u>	Left abdominal pain	I month	⊣	right flank Non - filling of left upper	24 cm. in diameter	nating N.S.	atter operation Well 3 months after operation
37	Briggs	1955		ш.	Increasing tiredness and weight loss	l year	_	Mass in left loin Non - function-	30 cm. in diameter	χ. .χ.	Died 7 months after operation
398	Frehling and Lev Gupta, Nagrath	1956 1963	51	Σı	Right Lumbar pain Right Lumbar pain	N.S. 5 months	~ ~	N.S.	N.S. Mass 10 > 7 \ 3 cm.	S. S.	50 50 27.
04	and bhagwal Leopold and Mogg	1964	<u>*</u>	Σ	naematuria Left renal pain Haematuria later	10 years	1	Non-functioning kidney Distorted mid- dle and upper calyces	Upper part kidney 14 × 12 × 10 cm.	S.Z.	S: Z
4 4	Lamarina Bazaz ₁ Malik and Gupra	1964 1966	63 4 8 8	ŭ ∑	Left Lumbar paín Haematuria	2 vears	_	Tender mass in left lumbar region	15 × 13 · 7 cm.	Lung	Alive after 1 year
£4	Lee and Tinckler (present case)	9961	40	Ц.	Right Abdominal lump Weight loss	6 months	ĸ	calyces Mass in right lein No function on right side Calcification	Mass 15 × 12 · 9 cm. replacing kidney	Lymph nodes in right renal	Well 6 months after operation

walls of renal vessels and the renal pelvis (Zuckerman et al, 1947; Lazarus and Friedman, 1954; Crosbie and Pinkerton, 1932; Bhende, 1952). In our case, it had probably arisen from the region of the renal pelvis and had enlarged to compress the renal parenchyma.

The tumour is locally infiltrating (Berry, 1929; Smith, 1935) and is known to metastasise to lungs, liver (Brandt, 1919; Lazarus and Friedman, 1954), mesenteric lymphnodes (Patch, 1937; Tetelman and Lisa, 1945), and inguinal and axillary lymphnodes (Bruce and McNaught, 1954). Our patient showed secondary deposits in the lymphnodes in the right renal hilus, but no lymphnode involvement elsewhere.

SUMMARY

A case of renal leiomyosarcoma in a 40-year-old Chinese female is described. This brings the number of such cases reported in the literature to 43. The clinical features, prognosis and commonly believed sites of origin of the growth are briefly reviewed.

ACKNOWLEDGEMENTS

Our thanks are due to Mr. T. T. Tan for preparing the photographs, and Miss Y. H. Chan for typing the script.

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