
Pattern of urological malignancies in 2010 – an audit from a tertiary referral centre

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Cancer is a major public health problem worldwide. Among various types of malignancies, genitourinary malignancies, except testicular cancer are notorious for causing high mortality and morbidity. Currently, 1 in 4 deaths in the United States is due to cancer. In USA, prostate cancer is the commonest cancer in men and accounts for the second commonest cause for the cancer death in men (1). Patterns of these cancers are variable according to time, region, and ethnic groups. We conducted an audit on the incidence of urological malignancies in a single urological unit at the National Hospital of Sri Lanka.

Method

Newly diagnosed cases of all histopathologically confirmed urological malignancies in the year 2010 were

collected from the clinic notes and operation register retrospectively and analysed.

Results

There were 76 urological malignancies diagnosed in the year 2010 and Table 1 shows their age and sex distribution.

In this study, bladder cancer was the commonest urological malignancy accounting for 42.10% of all the cases. Their median age was 67.5 years (range 50 - 90) and the male to female ratio was 3.7:1 (Table 1). Three fourth of them had painless or painful macroscopic haematuria at presentation (Table 2).

Table 1. Age and sex distribution

Site of malignancy	Number of patients	% of each malignancy	Male	Female	Average age years
Bladder	32	42.10	25	7	67.51
Upper urinary tract	1	01.31	1		42
Prostate	31	40.78	31	-	70.7
Kidney (RCC)	9	11.84	8	1	57.44
Penis	2	02.63	2	-	53.5
Testis	1	01.31	1	-	20
Total	76	100.00	68	8	66.7

Table 2. Clinical presentation of bladder cancer and number of patients

Presentation	No of patients
Haematuria (painless/painful)	21
Haematuria + LUTS	4
LUTS	6
Liver secondaries	1
Lower abdominal pain	1

Histological type, grade and pathological stage of bladder cancers are shown in Table 3. Out of 23 cases of nonmuscle invasive transitional cell carcinoma (TCC), 14 were low grade. All muscle invasive cancer in this study were high grade. There were two pure squamous cell carcinoma, one TCC with squamous differentiation and one sarcomatoid tumour in the study.

Prostate carcinoma was the second commonest malignancy. The median age at presentation was 70.7 years (54-84). While the commonest presentation was lower urinary tract symptoms (18 cases) acute urine retention was the second commonest (5 cases). All patients with prostate carcinoma presented with locally advance disease except three patients who presented with symptoms of distant metastases. Histologically all were small acinar type adenocarcinoma except one case of papillary TCC. The Gleason sum score was 8 or more in 20 patients (64.5%). More than half of the patients had their serum PSA >60 ng/l.

All nine cases of renal cell carcinoma were clear cell variety except one being chromophobe. Their mean age at presentation was 57.4 years (53-72) and male to female ratio was 9 to 1. Six patients had macroscopic haematuria at presentation while two cases were detected incidentally. According to the TNM classification, there were 3,4 and 2 of pT1, pT2 and pT3a respectively.

Table 3. Histological type, grade and pathological stage

Type	Stage	Grade	Number		% of patients
TCC	pTa	Low grade	9	10	30.30
		High grade	1		
	pT1	Low grade	5	13	39.39
		High grade	8		
	pT2	Low grade	0	7	21.21
		High grade	7		
SCC	pT1	MD*	1	2	06.06
	pT2	MD*	1		
Sarcomatoid carcinoma			1	1	03.03

*Moderately differentiated

Table 4. Clinicopathological presentation of prostate carcinoma

No.	Age	Presentation	DRE	Serum PSA	Gleason sum score
1	79	LUTS	M	43.4	10(5+5)
2	65	LUTS	M	>100	9(5+4)
3	72	LUTS	E	250	9(5+4)
4	82	LUTS	M	>2000	6(3+3)
5	64	LUTS	M	>400	7(4+3)
6	54	LUTS	M	271	8(4+4)
7	83	ARU	M	33.3	9(5+4)
8	69	LUTS	M	23.7	8(4+4)
9	67	LUTS	M	>100	8(4+4)
10	71	LUTS	M	957	7(3+4)
11	68	LUTS	E	306	9 (5+4)
12	68	LUTS	E	4330	9 (4+5)
13	69	LUTS	M	6.6	10(5+5)
14	67	LUTS	M	142	8(3+5)
15	65	Vertebral metastasis	M	23	8(3+5)
16	71	Lower limb lymphoedema	M	>100	9 (5+4)
17	66	LUTS	M	734	9 (4+5)
18	73	Cerebral metastasis	M	30.6	7(4+3)
19	67	ARU	M	67	9 (5+4)
20	71	LUTS	E	45.8	7(4+3)
21	74	ARU	M	>400	9 (4+5)
22	67	LUTS	E	18.8	6 (3+3)
23	84	ARU	M	12.1	9 (4+5)
24	68	ARU	M	>100	7(3+4)
25	76	LUTS	M	6.75	9 (5+4)
26	81	Haematuria	B	15.7	7(4+3)
27	74	HPCR	M	16.7	6(3+3)
28	65	Haematuria	E	6.8	9 (4+5)
29	64	LUTS	M	13.2	9 (5+4)
30	74	Paraplegia	M	3920	7(3+4)
31	74	Lower limb lymphoedema	M	4.1	papillaryTCC

LUTS = Lower Urinary Tract Symptoms, ARU = Acute Urinary Retention, HPCR = High Pressure Chronic Retention.

Table 5. Clinical presentation of renal malignancy and number of patients

Presentation	No of patients
Haematuria	4
Haematuria and loin pain	1
Haematuria and LOW	1
Anaemia	1
Incidental	2

There was one testicular cancer in a 20-year old man which presented as a testicular mass and it was classic seminoma stage 1.

Out of two penile cancers, one presented as phimosis while the other was detected during follow up for benign penile lesion. Both were histologically well differentiated and pathological stages were pT2 and pT1a.

Discussion

According to the 2010 cancer statistics, the commonest noncutaneous malignancy in USA men is carcinoma of the prostate accounting for 25% of all cancers and is the second commonest cause of cancer death i.e. 11%

of all cancer death. Urinary bladder cancer, kidney and renal pelvis cancers are the 4th and 7th leading cancers in men respectively. Kidney and renal pelvis malignancy is the commonest urological malignancy in female and is the 8th leading cancer (1). According to the cancer statistic data 2005, prostate cancer, being 8th leading cancer (5% of all cancers), is the only urological malignancy included among the first ten leading cancers in Sri Lanka (2).

In our study, number of Pca was slightly less than that of bladder cancer. As the PSA based screen detected Pca is not included, actual number of Pca would be much greater.

In USA, Pca incidence in 2005 was 170 per 100000 population (3,4). In Sri Lanka it was 3.1 in the same year (2). Worldwide, it is the fourth commonest male malignancy with great variation between countries and ethnic groups, Asia having the lowest incidence rate (1.9 cases per 100000 population per year in Tianjin, China) while African-American having the highest (172 cases per 100000 population per year) (5). Reason for such diversity would be due to access to and quality of health, accuracy of cancer registries and penetrance of PSA screening, environmental and genetic predisposition. Over the five year commencing from 2001 to 2005, Pca incidence has fluctuated in Sri Lanka (250,297,259,273,303 respectively) (2).

Table 6. Comparison of age distribution of Pca at presentation in US, in Sri Lanka and in our study (2,3)

Age	Sri Lanka		Study group		USA
	Number	Percentage	Number	Percentage	Percentage
<34	0		0	0	0
35-44	3	0.9	0	0	0.6
45-54	19	6.27	1	3.22	8.9
55-64	63	20.79	2	6.22	29.9
65- 74	126	41.58	22	70.96	35.3
75-84	92	30.36	6	19.35	20.7
>85			0	0	4.6

Table 7. Stage distribution and 5-year relative survival by stage at diagnosis for 1999-2006, all races, males (3)

Stage at diagnosis	Stage distribution%	5-year relative survival (%)
Localized (confined to primary site)	80	100
Regional (spread to regional lymphnodes)	12	100
Distant (cancer has metastasized)	4	30
(unstaged)	3	75

Table 8. Cancer incidence by the reporting year and site 2001-2005 in Sri Lanka (2)

Site of malignancy	2001	2002	2003	2004	2005	Total
Prostate	250	297	259	273	303	1382
Bladder	136	142	99	120	156	653
Kidney	80	90	85	103	100	458
Penis	75	63	69	44	73	324
Testis	44	33	33	35	42	187
Total	585	625	545	575	674	3004

Above comparison shows that peak age group for Pca is 65-74 years in both study group and Sri Lankan study, in keeping with USA Pca incidence while median age for Pca in our study is higher (70.7 years) than that of USA (67 years).

In our study, all cases were advanced stage at diagnosis. Three patients presented with distant metastases (one with hemiparesis following brain metastases and two with vertebral metastases). Two patients presented with unilateral lower limb lymphoedema indicating regional lymph node involvement.

The number of bladder cancers is slightly higher than that of prostate cancers in our study (32 vs 31). Similar study performed in the same unit for two year period from January 1994 to December 1995 shows higher incidence of bladder cancer (88 vs 62) (6). According to the Sri Lanka Cancer Registry, incidence of bladder cancer is much lower than that of prostate cancer (2) (Table 7). This difference is due to the fact that bladder

cancer is almost exclusively managed by the urological surgeons at the start of the treatment while other cancers such as cancer of the prostate and kidney are predominantly managed by the general surgeons in Sri Lanka (6).

At presentation, approximately 75-85% of all patients with bladder cancer have non-muscle invasive disease, an almost similar finding (75%) in our study also (7). However, the largest bladder cancer study performed in Sri Lanka shows that muscle invasive disease at presentation is seen in nearly half of the patients (8).

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