# 1996 ANNUAL REPORT OF THE TUMOR REGISTRY



KING FAISAL SPECIALIST HOSPITAL & RESEARCH CENTRE RIYADH, KINGDOM OF SAUDI ARABIA

#### ACKNOWLEDGEMENTS:

The Cancer Program is a combined effort of many individuals. It is not possible to enumerate all the nurses, technicians, therapists, pharmacists, dentists, physicians, scientists, social workers and others whose work is primarily on behalf of the patient with cancer. In addition, nearly everyone associated with the hospital comes in contact with the cancer patient from time to time, frequently contributing significantly to their care. The staff of the Tumor Registry and members of the Tumor Committee recognize this hospital-wide involvement in the care of cancer patients. The information in this report is provided to assist all health care professionals to better understand the problems faced in treating patients with cancer.

The following Departments have assisted throughout the year and without their invaluable support this report would not be possible. The Tumor Registry staff acknowledges these Departments:

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#### 1996 ANNUAL REPORT OF THE TUMOR REGISTRY

#### TABLE OF CONTENTS

I.	KFSH&RC Cand	cer Program Activities	1
	Figures 1-A	to 1-D - Sample Cancer Registry Worksheet	3
II.	KFSH&RC Cano	er Patient Population	7
	Figure 2 -	Distribution of All Cases Accessioned By Year (1975-1996)	7
	Table 1 -	All Cases Seen at KFSH&RC (Male/Female and Pediatrics/Adults) By 5-Year Period (1975-1996)	8
	Figure 3 -	Distribution of All Cases By Nationality (1975-1996 and 1996)	8
	Figure 4 -	Distribution of All Cases By Geographic Region (1975-1996 and 1996)	9
	Trends in Re	elative Frequency of Cancer at KFSH&RC	10
	Figure 5 -	Distribution of 20 Most Common Malignancies (1975-1996)	10
	Table 2 -	Ten Most Common Malignancies By The Most Common Age Group at Diagnosis (1975-1996)	11
	Figure 6 -	Distribution of 10 Most Common Pediatric Malignancies (1975-1996)	12
	Figure 7 -	Distribution of 10 Most Common Pediatric Malignancies By Histology (1975-1996)	13
	Table 3 -	All Cases Seen at KFSH&RC By Site and Year (1975-1996)	14
	Table 4 -	All Cases Seen at KFSH&RC By Site & 5-Year Period (1975-1996)	15
	Figure 8 -	Distribution of All Cases By Age At Diagnosis (1975-1996 and 1996)	16
	Figure 9 -	Distribution of All Pediatric Cases By Age At Diagnosis (1975-1996)	17
	Table 5 -	All Cases Seen at KFSH&RC By Site, Sex, Class of Case and Summary Stage (1996)	18
	Table 6 -	Analytic Cases Seen at KFSH&RC By Site & Age (1996)	19
	Table 7 -	Analytic Male Cases Seen at KFSH&RC By Site & Age (1996)	20
	Table 8 -	Analytic Female Cases Seen at KFSH&RC By Site & Age (1996)	21

#### TABLE OF CONTENTS (cont'd)

	Figure 10 -	Distribution of 20 Most Common Malignancies (1996 Analytic Cases)	23
	Figure 11 -	Distribution of Pediatric Malignancies (1996 Analytic Cases)	24
	Figure 12 -	Distribution of 10 Most Common Pediatric Malignancies By Histology (1996 Analytic Cases)	24
	Table 9 -	Primary Site Table (1996)	25
	Table 10 -	Patients With Multiple Primaries (1996)	33
	Stage of Dis	sease at Diagnosis	35
	Figure 13 -	Distribution of Analytic Cases By Stage At Diagnosis (1996)	36
	Figure 14 -	Distribution of Analytic Cases By First Course of Treatment (1996)	36
	Table 11 -	AJCC TNM Group Stage of Analytic Cases of Major Sites By Year (1992-1996)	37
III.	Special Stud	lies	
	Extranodal N	Care For Patients With Advanced Cancer	3
		International Prognostic Index	4
IV.	Appendices		
	Appendix A -	- Requests for Tumor Registry Data	48
	Appendix B	Tumor Committee Members	50
	Appendix C -	- Summary of Cases Presented at the Tumor Board	51
	Appendix D -	- Summary of Oncology Grand Rounds Topics	52
v.	Glossary		53

#### I. KING FAISAL SPECIALIST HOSPITAL & RESEARCH CENTRE CANCER PROGRAM ACTIVITIES

#### TUMOR REGISTRY

The King Faisal Specialist Hospital and Research Centre (KFSH&RC) opened in June 1975 to provide specialized medical treatment to the people of Saudi Arabia and to promote the prevention of disease through research and education. It is a national and international tertiary care hospital for Oncology and the principal center for cancer therapy in Saudi Arabia.

The KFSH&RC Tumor Registry is a hospital-wide data system designed for the collection, management, and analysis of data on patients with the diagnosis of a malignant neoplasm (cancer). The Registry was established to meet one of the requirements for an Approved Cancer Program of the American College of Surgeons (ACoS) and is under the supervision of the Tumor Committee. The database now includes 32,980 malignant cases seen at KFSH&RC from June 1975 through December 31, 1996. More than 2,000 new cases are added annually.

The Registry is staffed with certified tumor registrars who support the database in case ascertainment, abstracting, follow up and statistical analyses. The basic source document is the patient's medical record from which pertinent information is abstracted for use in the Registry. The electronic data system used is the Cansur 3.0 designed by the ACOS in which the details of each diagnosed cancer case is entered and stored. (Please refer to Figures 1-A to 1-D for a sample data set.)

The data maintained in the Tumor Registry provides the statistics for the publication of the KFSH&RC annual report which summarizes the hospital's cancer experience. The data also supports a wide variety of reports at the request of physicians, researchers, and ancillary personnel. These reports support patient management and outcome, basic and clinical research investigations, educational publications and presentations, and resource utilization. In 1996, the Tumor Registry supported 38 data requests (see Appendix A for a listing of requests for Tumor Registry data). It also identified and reported to the National Cancer Registry 2,175 cases seen in 1996 that were diagnosed on or after 01 January 1994.

#### TUMOR COMMITTEE

The multidisciplinary Tumor Committee, which meets bimonthly, is the policy-making body of the Cancer Program at KFSH&RC (see Appendix B for membership listing). During 1996, the Committee provided professional and administrative guidance to the Tumor Registry and its most prominent achievements include the following:

Recruited an additional Assistant Tumor Registrar to complement the present staff of the Tumor Registry.

Coordinated with the Computer & Hospital Information Centre (CHIC) in adapting the computerized cancer staging forms. CHIC has finished entering 20 forms in the hospital computer system which could be printed from any terminal with access to the mainframe or the e-mail. These forms will be submitted to the Forms Committee for approval and implementation.

Helped in establishing multi-disciplinary combined clinics, the latest to be developed was the Combined Colorectal Clinic.

Published three brochures on three different cancers, i.e., breast cancer, ovarian cancer and lymphomas (including Hodgkin's Disease). A fourth one, on gastric and esophageal cancers, is ready for printing; and a fifth one, on colorectal cancer, is under final preparation.

#### TUMOR BOARD

This educational conference is held as frequently as twice a month for the benefit of the attending staff, house staff, allied health professionals and visiting attending staff from other hospitals. Cases of various types of malignant disease are selected for presentation on the basis of complexity, unusual manifestations of the disease, or interest. Each presentation includes an outline of the medical history, physical findings, clinical course, radiographic studies, and pathological interpretations. Following each presentation, there is an informal discussion of the case and a review of pertinent medical literature. Those attending are encouraged to share personal experience in the management of similar cases. Please refer to Appendix C for a summary of cases presented in 1996.

#### ONCOLOGY GRAND ROUNDS

This didactic conference is held every other week and is attended by the Medical staff and allied health professionals. Speakers are drawn from the KFSH&RC Medical and Research staff as well as from visiting guests. Please refer to Appendix D for a listing of the topics presented at the Oncology Grand Rounds in 1996.

## FIGURE 1-A

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# KING FAISAL SPECIALIST HOSPITAL AND RESEARCH CENTRE

CANCER REGISTRY WORKSHEET (CanSur 3.0)

			<u> </u>	
PF 10 TACS - ACCESSION	FILE MAINTENANCE		MARIITAL BIATUS AT DX:	I
ACCESSION HUMBER (ACSH) :	[8   7   0   1   2		1 - Single 3 - Soperated 5 - Widowod	
TUMOR SEQUENCE (SEQ):	<u>Lo</u>	<u>Lol</u>	Married 4 - Diveced 9 - Unknown	
Malignant/In situ tumors	Benign tumors	.	RELIGION:	
00 - One primary only 01 - First of two or more	XX - One primary only AA - First of two or more	.	(1) Murstim 03 - Hindu 08 - Other .	
98 - 98th or Interprimary	III - 6th or fater primary	1	02 - Christlen 04 - Buddhist 99 - Unknown	
99 - Unspecified sequence	II - Unspecified sequence	1.7.1	ALCOHOL USAGE:	1
THIS CANCER ACCESSION YEAR :	ľ. <u>o</u>	2.1.2.1	1 - Current alcohol usage . 3 - Never used alcohol	
MEDICAL RECORD NO. :	13.19.14.16.15	i.l.z.l	2 – Past history of alcohol usage 9 – Unknówn	
			FAMILY HISTORY OF CANCER:	
CASE STATUS :		[3]	Family history of cancer 9 - Utiknown	
0 - Suspense 1 - Incomplete			2 – No family history of cancer	
3 - Completed per Release 3		1	вмокіна/снешіна нівтолу (3	ł
PATIENT NAME			f - Current smoker clg. 5 - Shamma	
Last:			2 - Pest amokor 5 - Sixistra  3 Pettent never amoked 7 - Contbo	
First:			4 - Chat 8 - Other 9 - Unknown	
Second :			TOTAL PACK YEARS:	1
Third:				ı
ADDRESS AT DIAGNOSIS			INDUSTRY:	,
P.O. Box			OCCUPATION: Teacher	ļ
Riv	adh.	ļ	DATE ADMITTED: (mm/46/yyy)   0   1   /   2   0   /   1   9   8   7	7
\\ \frac{\pi_{\pi_{\pi_{\pi_{\pi_{\pi_{\pi_{\pi_	adh. Chy		DATEDISCHURGED:	7
RY zip Code:			(2222)	
1101.	<u> </u>		REPORTING SOURCE:	1
PF 11 TPAT - PATIEN	IT IDENTIFICATION	۱, ۱	(1) Inpatient 4 - Physician's office 7 - Death Cert.	•
; OH KOUAR	1 2 3 4 5		2 - Clinic/outpatient 5 - Nursing home 9 - Unknown 3 - Laboratory 6 - Autopsy	
DIRTHOATE:	[1]/[0]]/[1]9]	4 6	НО SPITAL REFERENCE FROM : 0 0 0 0 1 0 1	1
AGE AT DX :	Lol	4L1	Riyadh Central Hospital	- I
SEX:		L2		ı
1 - Mala 2 Femal	9 - Unknown		HOSPITAL REFEREND TO	. !
HATIOHALITY:	<u>(c</u>	<u>)                                    </u>		
01 - Amer, Can, Brit 05 - 0 02 - Egyptian 06 -	/ement 98 - Other Arab 99 - Other Ind, Pak African			
			A	

Form 310-13 (Rev. 3-10)

# FIGURE 1-B

PF 12 TIXT - MISCELLANEOUS TEXT	TCAN - Cancer Identification (Continue)
	GRADE:
PHYSICAL EXAM: 6-MO lix 2 cm mass rt breast	1 ~ Well differentiated (f) 5 · Y-cell
UOQ, mobile, no skin changes. 3x4 cm	2 - Mod well-differentiated (III) 6 - B cell  (3) Poorly differentiated (III) 7 that cell
	4 - Undifferentiated (IV) 9 - 1 Int stated, unknown
rt axillary LN. Lt breast NED.	
	LATERALITY: [1]
<del>-</del>	0 - Not paired organ 3 - Dt or ft unspecified
	Unight 4 Both, simultaneous
x mays/scans:01/20/87 Bilat Manmogram -	2 - Left 9 Unknown falcosity
	DX CONFIRMATION:
2x2.5x2.5 cm mass rt breast UOQ.	Positive histology
CXR, Bone Scan, U/S Abdomen - NED	2 - Cytology 7 - Badiography
The state of the s	4 - Pos. micro, confirm, NOS 8 - Clinical
	5 - Laboratory test/marker 9 - Unknown
	REGIONAL NODES EXAMINED:  1  9
COURTS (140, 01/25/87 FDX (4) DDX (4)	00 - No nodes examined 97 - 97 it makes examined
SCOPES/IAN: 01/25/87 ERA (+), PRA (+)	01 - One node examined 98 - Nodes examined, number unknown
	99 ~ Unknown if notes examined
}	REGIONAL NODES POSITIVE:
	00 - No nodes positive 97 - Positive nodes, number technolos
	01 - One node positive 96 - No nodes examined
	99 – Uoknown if any nodes +/-
	98 - 96 + nodes positive
OPERATIVES FINDINGS: 01/25/87 Rt Mod Rad Mas-	lo la
tectomy - no description of tumor.	TUMON SIZE (cm)  0  2    2
represent the department of the second of th	eg., 000 - No mass, -002 - 0.2 cm, -055 - 5.5 cm, -999 - Unknown
	DESIDUAL TUMON:
	(0)Hone 2 - Macroscopic 9 - Unknown
	i - Microscopic B - No respetion, MA
PAHIOLOGY/AUTOPSY: 87SP3286 01/25/87_Duct_	DISTANTMETS: 1:
Cell Ca, gr 3; 11/19 LN's. (tumor	0 Dono Mar. 4 -4 Liver 8 Lymph mode (distant) 2 :
	1 - Paritonoum 5 - Done 8 - Unknown/utter 3 :
size: 2.2x2x1.8 cm completely ex-	<b>,</b> ,
cised) Nipple & overlying skin NED.	<b>*</b>
	3-Piorra 7 - Skin
(largest LN 1.5 cm)	GEHERAL SUMMARY STAGE: 3
	0 - Institu 4 - Regional, both 2 & 3
	1 - Localized 5 - Regional, NOS
PF 13 TOAN - CANCER IDENTIFICATION	2 - Regional, direct extension 7 - Distant
DATEOFERITIAL DIAGROUS: (INNIVERSITY) 0 11 / 2 5 / 1 9 8 7	(3) Regional, nodes 9 - Unknown/unistageable
DATE OF #111/AL DIAGNOSIS (INVIDENTIAL SALE)	AICC STAGE:
CLASS OF CASE :	
	CLINICAL T 2 . N 1 . M 0 - STAGE GHOLIF 2 D
0: Dichere, ix elsewhere 4 - Dichere prior (1) Dicktration 5 - Dickt eutopsy	PATHOLOGICAL T 2   H 1 B M 0 STAGE GROUP 2 D
2 Axticro 9 - Unknown	
3 Fix allogations	OTHER*** TT
PHIRARY SHE - TEXT: Breast, Right UOQ	◆Tittl Codes ~ hise nipha codes as appropriate; eg. 12A, 2A, 12, 2,
해를 받아 것 같은 LE 1회 위기가 LE 도 레스( <u></u>	M18-18, M0-0, IS-In situ, X: Unknown)
CODE: [1   7 4   1	A A A ICO Stane Group - use plobe and a se suppositely
CODE : 12.1.212.12.1	**AUCC Stage Group - use alpha codes as appropriate;
	eg, 3A: Slago ⊞A, 1- Singo 1
	0 - Institu 2 - Stage II 4 - Stage IV
Historogy-icxi: Duct Cell Carcinoma, gr 3	
Histotogy-itxi: Duct Cell Carcinoma, gr 3	t - Stage III 9 - Unknown
	I "

# FIGURE 1-C

PF 14 TRX = 1ST COURSE TREAT	MENT (SUNGERY, RADIATION)	PF 16 TNX3 - 1st COURS	
SURGLEY		CHEMOTHERAPY	
REASON:	lol	SUMMANY:	3
Can directed surg	6 - Reason unknown, no surg	AT THIS HOSPITAL:	l3 l
pedomed	7 - Patient/guardian refused	0 - No chamotherapy	7 - Patien/guardian refused
1 - Hot recommended	8 - Recommended, unk if done	1 - Chemotherapy, NOS	8 - Recommended, unk if done
2 · Contraindicated, other	9 – unknown	2 – Chemotherapy, single agent	9 - Unknown
SUMMANY : (Entire 1st course).	5,101	3 Chemothernpy, multi-agent c	
AT THIS HOSPITAL: •	5 lol	STARTED: (mar/dd/yyyy)	02 1/1131/1191817
• Refer to Appendix A In ConSur U		TEXT: 5-FU, Adria,	Ctx
STABLED: (mm/dd/yyyy)	0 1 4 5 4 1 9 8 7		
IEXT: Rt Mod Rad I	Mastectomy w/ Rt	HORMONE/STEROIDS	
Axillary Di		SUMMATIY :	<b>j</b> 1
MOHAHOAI		AT THIS HOSPITAL :	1
		0 - No hormonal the apy	7 - Patient/guardian refused
SUMMARY:	1.1	1) Hormonal therapy	B - Recommended, unk if done
AT THIS LIOSPITAL	[1]	2 – Endocrine sung/radiation	9 - Unknown
0 - No Badiatlor-therapy	5 - Radiation therapy, NOS	3 - Hormones + ender sing/ind	
(1) Ocaminadiation	7 - Pettent/guardian refused 8 - Recommended, unk if done	STARIED: (mm/dd/yyyy)	lo   2   1   9   1   9   8   7
2 - Radioactive Implants 3 - Radioisotopes	9 - Unknown	TEXT: Tamoxifen	
4 - Comir 1 + 2 or 3	• • • • • • • • • • • • • • • • • • • •		
. STARTED:(mm/dkf/yyyy)	[0] 8 / [2 9] / [1] 9 [8 ] 7 [	BIO-RESPONSE MODIFIER (BRM)	
1 .	lat	SUMMARY:	lo 1
TO BRAIT & CNS : (Lung & leukam 0 - None to CNS	8 - Recommended, unk if done	AT THIS HOSPITAL:	lo l
1 - Radiation therapy	(9) Unknown/not applicable	(0-)no BRIA	7 - Patlent/guardian refused
7 Patient/guardian refused	() s	1 - BRM	B - Recommended, unk if done
_	[3]	2 - Allo BMT	9 - Unknown
:D32 YR3DRU2\HOLLAIGAR oktabilqqa toU - 0	5 - Intraoperative radiation	3 - Auto BMT	
2 - Radiation before surgery	6 - Intrapperative plus 2, 3 or 4	STARTED: (mm/dd/yyyy)	
(3) Radiation after surgery	9 - Sequence unknown	1EXT:	
4 - Dolore & after surgery			
ıtxı: Chest Wall	6000		
· · · · · · · · · · · · · · · · · · ·		OTHER DX	l <sub>0</sub> l
- ,	· · · · · · · · · · · · · · · · · · ·	SUMMARY:	io i
		AT YILIS HOSPITAL:	6 - Unproved therapy
F   F   1   T   T   T   T   T   T   T   T   T	UO, THENAPY	1 - Other ca-directed ix	7 - Patient/guardian refused
Started Course	a Type Code Desc.	2 – Experimental carx	8 - Recommended, trik if done
men'del'yyyy		3 – Double-blied study	9 - Unknown
CHAININ D		STARTED: (minVd(Vyyyy)	1,1,171,1371,14.11
2 1 1/1 1/1 1 1	<u> </u>	TEXT:	
3   1   / 1   / 1   1			
	, , , , , , , , , , , , , , , , , , ,		
		<u> </u>	

# FIGURE 1-D

THE FORMAT AREA:  CAUSE OF DEATH ICD CODE:  CHIRILITY VITAL STATUS (1) Alive 2 Deat
CAUSE DE DUATH ROLOGOE:  CHINE IN VITAL STATUS (1) Alexe 2 Dead  CHINETH CAMBER STATUS (2) Alexe 2 Dead  CHINETH CAMBER STATUS (2) Alexe 2 Dead  1 Non-vidence of cancer 2 - Chinete concer 9 - Unknown  CHIALITY OF STRIVIVAL:  0 - Thermal 3 Ansh : 50% 6 - TIA, doad  1 - Synd Sindb 4 - Bodridden 9 Joknown  2 Ansh - 50%  1 - Synd Sindb 4 - Bodridden 9 Joknown  2 Ansh - 50%  STECIAL FIELDS (7 - Bypass edit, leave blank II edit not bypassed)  STECIAL FIELDS (1)  SECHO/SITEA IIST:  SECHO/SITEA IIST:  SECHO/SITEA IIST:  SECHO/SITEA IIST:  1 - 1 / 1: Hepatills
CHIRE IN VIAL STATUS (1) Alive 2 Dowl [1]  1 Hospidence of cancer (2-) Widence of cancer 9 ~ Unknown CKIALSTY OF STRIVIVAL: [9]  0 Hornid 3 And: 50% 6 - HA, dond 1 - Synt & and 2 - Hodridden 9 loknown 2 And: 50% 6 - HA, dond 5 - HA Condition 9 loknown 5 - HA Condition 9 loknown 1 - Synt & and 5 - SO% 50% 50% 50% 50% 50% 50% 50% 50% 50% 50
CHIDITH CARGER STATUS: [2]  1 Mossidence of cancer (2-) Vidence of cancer (9 - Unknown CRIACTLY OF STRIVIAL: [9]  0 - Hornal (1 - Symbol) (4 - Doddidlen (9) Joknown (2 - Anthology) (9) Joknown (9) J
CHIDITH CARGER STATUS: [2]  1 Mossidence of cancer (2-) Vidence of cancer (9 - Unknown CRIACTLY OF STRIVIAL: [9]  0 - Hornal (1 - Symbol) (4 - Doddidlen (9) Joknown (2 - Anthology) (9) Joknown (9) J
1 More idence of cancer 2 Vividence of cancer 9 - Unknown CKIALETY OF SHRIVIAL:  0 - More of Shrivian 3 Ando: 50% 8 - MA, doad 1 - Symbolid 4 - Dodridden 9 Joknown 2 - Ando - 50%  1ETHER FLAGS -  FAMILET (letter or asterisk, eg., *, A, B, I)  OVERINDE FIELDS (Y - Dynass edit, heave blank II edit not bypassed) SHEARST: AGE /SHEARST: SECHO/SITEARST: SECHO/SITEARST:  1 - 1 / 1 : Hepatills
CRIALITY OF STRIVIVAL:  0 - Hormal 3 Anabe: 50% 8 - HA, dond 1 - Symbanib 4 - Doddidlen 9 Joknown 2 Anabe: 50% 1 - Symbanib 4 - Doddidlen 9 Joknown SECHALITER IST: 2 Anabe: 50% SECHALITER IST:  SECHALITER: SECHALITER IST:  1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1 - 1
0 - Hormal 3 Anab : 50% 6 - HA, dond 1 - Synt Simb 4 - Bodridden 9 Joknown 2 Anab : 50%  1 ETHER FLAGS -  FAUCHAL Better or asterisk, eg., *, A, B, I)  STECIAL FIELDS:  1 1 - Hepatills
1 Synta and 4 Booldines (9 January SECHO/SITEA BST :  LETIFITIAGS - SPECIAL FIELDS :  PARENT Better or arterisk, eg., *, A, B, I) - / 1 : Hepatills
TETIFFITIAGS - SPECIAL FIELDS:  [Alterial Butter or asterisk, eg., *, A, B, I)  [1]  [1]  [2]  [3]  [4]  [5]  [6]  [7]  [7]  [7]  [8]
PATIENT: (letter or asterisk, eg., •, A, B, I)
4 1 1
(Chitye') fed. a mix change 1 . man changed 1.1
CUBINETHLY FOREIGN RESIDENT:
Y - Yes, toxelign resident, leave blank for all others)  # 4 : Consequences
CONTACTINEQUETICY:
(eq. (it) - One awath, 03 - 3 months, 12 - Annual (office-up) / 6 : Prognancy during dw/tx
MINICIPAL COMOTHOUS: 1.4 #7: Ronal Transplant
/ 8 : Immunodoficiency Disorder
PLACE OF DEATH (State of country - Georgide)
DECEMBER OF STREET
DATE: (mitted by yyy)   0,9   /(1   9) /(1   9)8   9
1975:  1 PF - 21 TADR - PATIENT NAME ADDRESS FILE  1 PF - 21 TADR - PATIENT NAME ADDRESS FILE
0 Herechicane 3 Distant resintence 1 tonalizations 4 - Hover fice MAILING NAME :
2 Regional recontence 9 Unknown SALUTATION:
DISTANTIMETS:
0 ft.t (1) tivis 0 1 yapshuooku(distant) 2: 51 ADDNESS 1: Riyadh
2 Ling 6 - CUS
3- Fleura 7 - Skin CITY:
(ULCTICE PHYSICIAL Riyadh
FILES CONE. HAME PROV.
The state of the s
2. OHER PHYSICIAL:
a DITER PHYSICIAN:       0  9 2 1 8 5  Surgeon PATIENT/GUANDIAN CODE: P - fratight G - Grandlan 1
4 DIER PHYSICIAL:
5. OHIER CHYSICIALE:
5 OHER PHYSICIAL:
AUDRESS 1:
INTERIOR FORTH: Riyadlı ADDRESS 2:
OFATH CERTIFICATE (ME NO:
PRIOV. [RY] ZEPCODE:
COMMENT:
DEFER HOSE, MINT: 89856

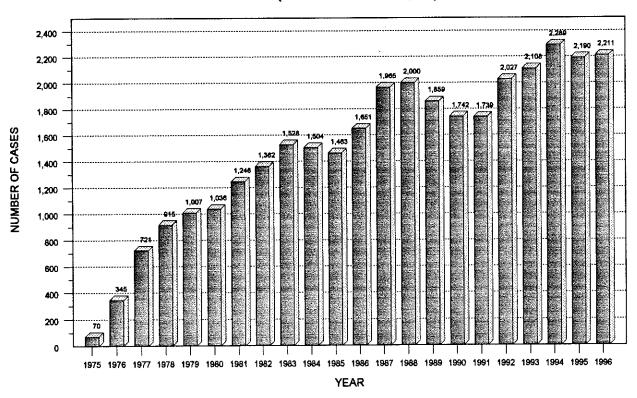
#### II. KFSHERC CANCER PATIENT POPULATION

A total of 2,211 cases were accessioned in 1996, with 1,098 males and 1,113 females or a male/female ratio of 1:1. This represents a 1% increase from 1995.

FIGURE 2

DISTRIBUTION OF ALL CASES ACCESSIONED BY YEAR

1975 - 1996 (TOTAL CASES = 32,980)



From the opening of the hospital (mid 1975) until December 1996, 32,980 cancer cases were registered (17,907 males and 15,073 females) with a male/female ratio of 1.2:1. There were 4,178 (12.7%) pediatric cases (0 to 14 years of age) and 28,802 (87.3%) adults (15 years old and above). A slight difference in the proportion was noted in 1996, 13.3% (295) for pediatrics and 86.7% (1,916) for adults.

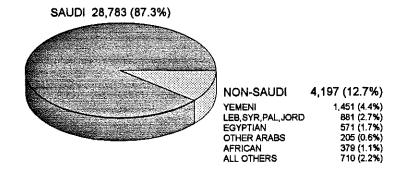
TABLE 1

ALL CASES SEEN AT KFSHERC (MALE/FEMALE & PEDIATRICS/ADULTS) BY 5-YEAR PERIOD 1975 - 1996

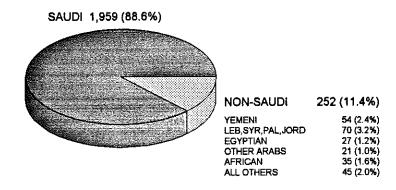
	1975-1	976*	1977-1	981	1982-1	986	1987-1	991	1992-199	6	TOTA	L
	No.	4	No.	*	No.	*	No.		No.	•	No.	8
MALE	280		2,981		4,150		4,969		5,527		17,907	
Female	135		1,945		3,358		4,336		5,299		15,073	
TOTAL	415		4,926		7,508		9,305		10,826		32,980	
M/F RATIO	2.1:1		1.5:1	-	1.2:1		1.2:1		1.0:1		1.2:1	
PEDIATRICS**	55	13.2	592	12.0	984	13.1	1,159	12.5	1,388	12.8	4,178	12.7
ADULTS	360	86.8	4,334	88.0	6,524	B6.9	8,146	87.5	9,438	87.2	28,802	87.3
TOTAL	415	- 100	4,926	100	7,508	100	9,305	100	10,826	100	32,980	100

- \* First two years of KFSH&RC partial operation.
- \*\* Pediatrics = 0 to 14 years of age; Adults = 15 years and above.

# FIGURE 3 DISTRIBUTION OF ALL CASES BY NATIONALITY 1975 - 1996 (TOTAL CASES = 32,980)



#### 1996 CASES (TOTAL = 2,211)



Saudi nationals totaled 1,959 (88.6%) in 1996 and the non-Saudi, 252 (11.4%). During the period 1975 to 1996, the former accounted for 87.3% (28,783) while the latter, 12.7% (4,197).

Geographically, the referral pattern in 1996 is mainly from the Riyadh Region with 31.4% of all cases, followed by the Eastern Province and the Makkah Region with 17.2% and 13.1%, respectively. These same regions had the most number of cases during the 22 years in review, i.e., 31.3% from Riyadh, 17.9% from Makkah and 14.0% from the Eastern Province.

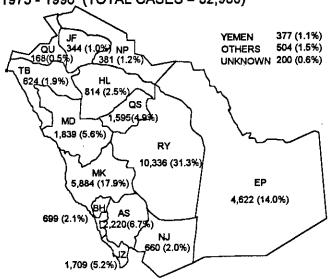
These percentages reflect KFSH&RC actual experience rather than adjusted to reflect the population of those regions.

#### FIGURE 4

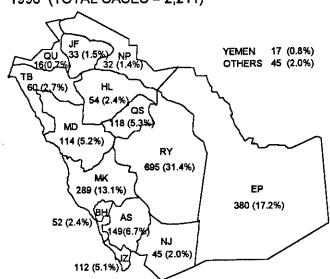
#### DISTRIBUTION OF ALL CASES BY GEOGRAPHIC REGION

(Based on Given Address at the Time of Diagnosis)





#### 1996 (TOTAL CASES = 2,211)



AS - ASIR BH - AL BAHA

AL JAWF

BH - AL BAHA
EP - EASTERN PROVINCE
HL - HAIL

MD - AL MADINAH
MK - MAKKAH
NJ - NAJRAN
NP - NORTHERN PROVINCE

JIZAN

JZ -

QS - AL QASSIM
QU - AL QURAYYAT
RY - RIYADH
TB - TABUK

#### TRENDS IN RELATIVE FREQUENCY OF CANCER AT KFSHARC

The crude relative frequency is the proportion of a given cancer in relation to all cases in a clinical or pathological series. Although such frequencies are subject to many biases, historically many elevated frequencies have been confirmed when complete cancer registration was introduced.

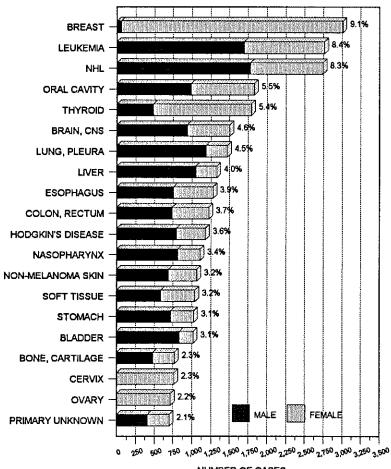
Biases that may have an effect on the relative frequencies of cancer cases at KFSH&RC include:

- possible nonusage of medical services by some of the population so that the hospital population may not reflect the disease state of the community
- resistance to examination by part of the female population
- absence of postmortem examinations/death certificates
- selective referral of certain malignancies because of a specialty service available
- eligibility criteria for admission to KFSH&RC
- age distribution of the population

Breast cancer led the list of total cancer cases seen from 1975 to 1996 with 9.1%, followed by Leukemia (8.4%), Non-Hodgkin's Lymphoma (8.3%), Oral Cavity (5.5%) and Thyroid (5.4%).

FIGURE 5

DISTRIBUTION OF 20 MOST COMMON MALIGNANCIES 1975 - 1996 (TOTAL CASES = 32,980)



NUMBER OF CASES

TABLE 2

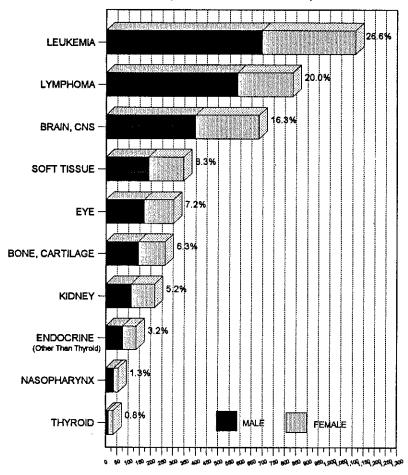
TEN MOST COMMON MALIGNANCIES BY THE MOST COMMON AGE GROUP AT DIAGNOSIS

1975 - 1996

SITE	AGE GROUP	No.	×	SITE	AGE GROUP	No.	×
Breast	40 - 49	924	30.7	Brain, CNS	00 - 09	504	33.5
	30 - 39	798	26.5		10 19	297	19.8
	50 - 59	614	20.4		20 - 29	168	11.2
	60 - 69	319	10.6	-	30 - 39	147	9.8
	20 - 29	215	7.2		50 - 59	145	9.6
SITE	AGE GROUP	No.	×	SIŢE	AGE GROUP	No.	x
Leukemia	00 - 09	865	31.3	Lung, Pleura	60 - 69	498	33.9
	10 19	509	18.4		50 - 59	371	25.2
	20 - 29	331	12.0		70 - 79	245	16.7
	30 - 39	328	11.9		40 - 49	203	13.8
	40 - 49	243	8.8		30 - 39	73	5.0
SITE	AGE GROUP	Nö.	×	SITE	AGE GROUP	No.	x
Non-Hodgkin's	60 - 69	454	16.6	Liver	60 - 69	420	31.6
Lymphoma	50 - 59	430	15.7		50 - 59	364	27.4
	40 - 49	341	12.4		70 - 79	204	15.4
	30 - 39	280	10.2		40 - 49	174	13.1
	70 - 79	278	10.1		30 - 39	62	4.7
SITE	AGE GROUP	No.	x	SITE	AGE GROUP	No.	x
Oral Cavity	60 - 69	461	25.2	Esophagus	60 - 69	406	31.8
	50 - 59	420	23.0	, -	50 - 59	283	22.1
	70 - 79	311	17.0		70 - 7 <del>9</del>	. 272	21.3
	40 - 49	236	12.9		40 - 49	131	10.2
	30 - 39	146	8.0		80 - 89	96	7.5
SITE	AGE GROUP	No.	×	SITE	AGE GROUP	No.	×
Thyroid	30 - 39	409	22.8	Colon, Rectum	50 - 59	288	23.6
,.	20 - 29	356	20.0	coron, needell	60 - 69	267	21.9
	40 - 49	316	17.6		40 - 49	201	16.5
	50 - 59	263	14.7		30 - 39	169	13.8
	60 - 69	200	11.2				
	OV - DY	200	11.2		70 - 7 <del>9</del>	150	12.3

Cancer among pediatrics (under the age of 15) accounted for 12.7% of all cases from 1975 to 1996. The five most common pediatric malignancies were Leukemia (26.6%), Lymphoma (20.0%) [NHL 11.8% and HD 8.2%], Brain/CNS (16.3%), Soft Tissue (8.3%) and Eye (7.2%).

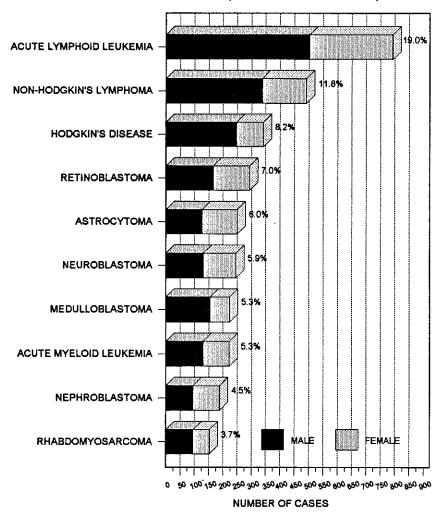
FIGURE 6
DISTRIBUTION OF 10 MOST COMMON PEDIATRIC MALIGNANCIES
1975 - 1996 (TOTAL CASES = 4,178)



NUMBER OF CASES

FIGURE 7

DISTRIBUTION OF 10 MOST COMMON PEDIATRIC MALIGNANCIES
BY HISTOLOGY, 1975-1996 (TOTAL CASES = 4,178)



32,980

2211

2190

2289

2108

2028

1739

1742

1859

2000

1965

1651

1463

1504

1528

1362

1246

1036

1008

915

2

345

TABLE 3

ALL CASES SEEN AT KFSH&RC BY SITE\* AND YEAR 1975 - 1996

1,828 1,105 1,105 1,105 1,224 1,328 1,324 1,304 1,304 1,304 1,503 1,039 1995 1994 1993 2222133213383238830002202383337288322238 1992 18 1990 506724868656867488445844684858688065 1989 2-4332-6384334434436-6356-6238992286486-63864-63 1988 1987 1986 1985 1984 1983 1981 334-653-7501245557 1980 1979 1978 1977 1976 1975 Hodgkin's Disease-LNs S Reticuloendothelium ¥on-Metanoma Skin ymphoid Leukemia Lymph Nodes Ayeloid Leukemia Extra-nodal Aultiple Myeloma - Extra-nodal Other Leukemias Bone, Cartilage Cidney, Urinary Other Endocrine Primary Unknown estis, Genital Geni tal Colon, Rectum Skin Melanoma .ung, Pieura Soft Tissue tasopharymx **Stal** Cavity Brain, CMS Other G.I. sophagus Prostate Pancreas rhyroid Jterus, stomach 31 adder Cervix -aryax 3reast Liver Vary 全

\* Includes Multiple Primary Neoplasms.

TABLE 4

ALL CASES SEEN AT KFSHRRC BY SITE\* AND 5-YEAR PERIOD

1975 - 1996

SITE	197	1975-1976**	1977-1981	1981	1982	1982-1986	1987	1987-1991	1992	1992-1996	1	TOTAL
	SE SE	ж	<b>₽</b>	×	S.	*	2	×	<b>₽</b>	×	N O	**
Oral Cavity	5	3.6%	310	6.3%	436	5.8%	534	Z.3	533	76.7	1,828	5.5%
Nasopharymx	7.	3.4%	192	3.9%	251	3.3%	331	3.6%	317	2.9%	1,105	3.4%
Esophagus	9	3.9%	304	6.2%	343	79.4	348	3.73	267	2.5%	1,278	3.9%
Stomach	17	4.1%	204	4.1%	787	3.8%	242	2.6%	275	2.5%	1,025	3.1%
Colon, Rectum	<u>*</u>	3.4%	166	3.4%	237	3.2%	357	3.8%	255	4.1%	1,221	3.7
Liver	23	5,3%	200	4.1%	312	4.2%	338	3.6%	456	4.2%	1,328	70.4
Pancreas	•	1.4%	29	1.4%	8	1.3%	88	0.9%	114	1.1%	374	1.1
Other G.I.	~	<u>۲</u>	25	1.2%	23	1.0%	101	1.1%	139	1.3%	377	1.1%
Larynx	9	1.4%	20	1.4%	\$	1.3%	138	1.5%	167	1.5%	780	1.5%
Lung, Pleura	7	3.4%	198	<b>70.7</b>	381	5.1%	177	¥.4	436	4.0%	1,470	4.5%
Multiple Myeloma	ī	1.2%	07	0.8%	09	0.8%	110	1.2%	123	1.1%	338	1.0%
Lymphoid Leukemia	<b>3</b>	4.3%	176	3.6%	326	4.3%	373	4.0%	414	3.8%	1,307	4.0%
Myeloid Leukemia	9	3.9%	215	47.4	287	3.8%	370	4.0%	416	3.8%	1,304	4.0%
Other Leukemias	-	0.2%	30	29.0	36	0.5%	45	0.5%	75	0.4%	154	0.5%
Reticuloendothelium	-	0.2%	4	0.1%	10	0.1%	2	0.1%	8	0.0%	22	0.1%
Bone, Cartilage	~	አ.	<b>103</b>	2.1%	172	2.3%	505	2.2%	280	2.6%	3	2.3%
Soft Tissue	5	3.6%	155	3,1%	210	2.8%	315	3.4%	344	3.2%	1,039	3.2%
Skin Melanoma	4	1.0%	33	۵.7	14	0.5%	7,	0.5%	£3	0.4%	165	0.5%
Non-Metanoma Skin Ca	16	3.9%	1%	4.0%	306	4.1%	255	2.7%	291	2.7%	1,064	3.2%
	27	6.5%	322	6.5%	633	8.4%	841	70.6	1184	10.9%	3,007	9.1%
Uterus, Genital	m	۲. ا	92	1.3%	124	۲. ۲.	177	 %	202	7.8	574	۲.
Cervix	<u>e</u>	2.4%	105	2.1%	185	2.5%	213	2.3%	250	2.3%	763	2.3%
Ovary	∞	<b>.</b> .	<b>8</b> 2	1.6%	151	2.0%	222	2.4%	524	2.3%	716	2.2%
Prostate	~	×.	32	<u>ک</u>	101	1.3%	116	1.2%	<b>1</b> %	1.8%	455	1.4%
Testis, Genital	4	1.0%	9	1.2%	71	0.9%	87	26.0	104	1.0%	326	1.0%
Bladder	=	۲.7	141	2.9%	197	2.6%	329	3.5%	338	3.1%	1,016	3.1%
Kidney, Urinary	Φ.	2.2%	87	1.8%	141	1.9%	195	2.1%	992	2.5%	869	2.1%
Eye	•	1.4%	ድ	1.9%	130	<u>ک</u>	142	1.5%	116	1.1x	687	1.5%
Brain, CNS	22	6.5%	155	3.1%	308	4.1%	443	4.8%	570	5.3%	1,503	79.7
Thyroid	10	2.4%	2	3.6%	333	4.4%	544	5.8%	724	<u>کر</u> ح	1,790	27.5
Other Endocrine	Ν.	0.5%	54	0.5%	54	0. 7X	36	<b>77.</b> 0	24	0.5%	173	0.5%
NHL - Lymph Nodes	23	5.5%	418	8.5%	488	6.5%	677	4.8%	3%	3.7%	1,774	2.4%
NHL - Extra-nodal	7	1.0%	ĸ	1.5%	<b>50</b> 8	2 %	303	3.3%	380	3.5%	896	2.9%
Hodgkin's Disease-LNs	32	۲.7	206	4.2%	238	3.2%	308	3.3%	383	3,5%	1,167	3.5%
MD - Extra-nodal	0	0.0%	0	0.0%	2	0.0%	-	20.0	^	0.1%	5	0.0%
Primary Unknown	14	3.4%	127	2.6%	138	1.8%	190	2.0%	231	2.1%	902	2.1%
All Other Sites	4	1.0%	35	۲.0	45	79.0	26	0.6%	9	79.0	201	79.0
TOTAL	415	100.0%	4,926	100.0%	7,508	100.0%	9,305	100.0%	10,826	100.0%	32,980	100.0%
											•	

\* Includes Multiple Primary Neoplasms.

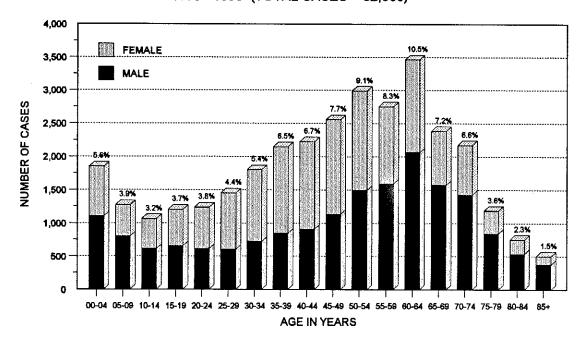
\*\* First Two Years of KFSH&RC Partial Operation.

The largest number of cases was noted in the 5th and 6th decades in males and in the 4th and 5th in females. In 1996, the mean age was 45.3, the median was 48.2 and the mode was 60. Pediatric malignancies are most common among children three years of age.

FIGURE 8

DISTRIBUTION OF ALL CASES BY AGE AT DIAGNOSIS

1975 - 1996 (TOTAL CASES = 32,980)



1996 (TOTAL CASES = 2,211)

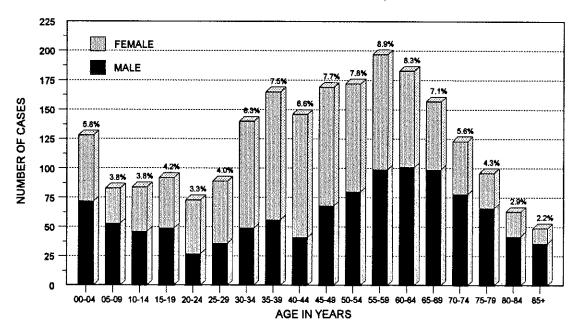
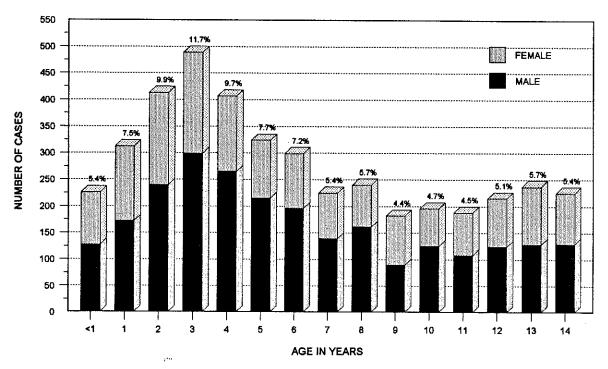


FIGURE 9

DISTRIBUTION OF ALL PEDIATRIC CASES BY AGE AT DIAGNOSIS

1975 - 1996 (TOTAL CASES = 4,178)



Of the 2,211 cases in 1996, 1,921 (86.9%) were **analytic** (defined as cases which were first diagnosed and/or received all or part of their first course of treatment at KFSH&RC. The remaining 290 cases (13.1%) were **non-analytic** (defined as cases diagnosed elsewhere and received all of their first course of treatment elsewhere). Out of the 1,921 analytic cases, pediatric cases totaled 257, with 149 males and 108 females.

See Table 5 for the distribution of cases by site, sex, class of case, and stage at diagnosis and Tables 6, 7 and 8 for the distributions of analytic cases by site, sex and age at diagnosis.

TABLE 5

ALL CASES SEEN AT KFSH&RC BY SITE\*, SEX, CLASS OF CASE AND SUMMARY STAGE

				•	)			ANAL	7 11 6 6	ASES	
SITE	TOTA	TAL	S	E X	CLASS OF	CASE**	9	RERAL	SUMMA	RYST	A G E
	Number	×	Male	Female	Analytic	Non-Anal	In Situ	Localized	Regional	Distant	Unstageable
Breast	277	12.5%	7	275	230	17	ľ	53	123	36	•-
Leukemia	165	7.5%	35	ይ	144	21	0	0	0	144	0
Thyroid	160	7.2%	94	114	135	52	0	58	63	12	~
Non-Hodgkin's Lymphoma	155	7.0%	86	25	134	21	0	38	52	7	0
Brain, CNS	127	5.7%	18	94	122	5	0	**	33	m	2
Oral Cavity	105	4.8%	20	55	26	80	0	20	63	14	0
Liver	95	4.3%	22	23	86	9	0	17	18	20	4
Colon, Rectum	76	4.3%	52	75	80	14	0	15	39	Ю	-
Lung, Pleura	88	3.9%	2	16	80	9	0	7	40	33	0
Nasopharymx	85	3.8%	26	92	ድ	9	0	2	67	%	2
Hodgkin's Disease	ĸ	3,4%	87	27	۲	4	0	٥	30	32	0
Non-Melanoma Skin Ca	92	2.9%	39	92	67	16	0	32	9	1	0
Soft Tissue	62	2.8%	33	82	87	77	0	27	16	ſΛ	0
Stomach	22	2.6%	37	20	51	9	0	7	30	13	-
Bone, Cartilage	25	2.6%	35	22	25	72	0	ΙΛ	33	14	0
Bladder	55	2.5%	47	80	94	ο.	0	22	16	_	_
Ovary	54	2.4%	0	24	36	5	0	12	2	54	-
Esophagus	55	2.4%	52	27	25	S.	0	16	18	13	0
Prostate	64	2.2%	67	0	33	5		5	7	72	_
Kidney, Urinary	84	2.2%	27	72	36	Φ.	0	15	٥	15	0
Cervix	25	2.1%	0	25	7,5	ıń	7	M	30	_	0
Primary Unknown	41	1.%	22	6	39	7	0	0	0	0	39
Larymx	36	1.6%	33	m	ጟ	7	0	13	17	4	0
Uterus, Genital	32	1.4%	0	32	92	9	0	17	9	M	0
Other G.I.	92	1.2%	=	5	8	œ	0	~	7	4	0
Eye	22	1.0%	12	우	16	•	-	4	10	-	0
Pancreas	20	0.9%	12	œ	16	4	0	_	0.	9	0
Multiple Myeloma	20	0.9%	12	∞	19	<del>-</del>	0	0	0	19	0
Testis, Genital	15	<u>ک</u>	15	0	13	7	0	Φ.	-	M	0
Other Endocrine	11	0.5%	٥	~	-	0	0	4	4	M	0
All Other Sites	9	0.5%	7	M	1		0	0	•	4	0
Skin Melanoma	<b>80</b>	27.0	m	2	•0	2	0	~:	2	2	0
TOTAL	2,211	100.0%	1,098	1,113	1,921	290	80	519	714	625	55
					ı						

\* Includes Multiple Primary Neoplasms.

<sup>\*\*</sup> Analytic Cases - cases which were first diagnosed and/or received all or part of their first course of treatment at KFSH&RC. Non-Analytic Cases - cases which were diagnosed elsewhere and received all of their first course of treatment elsewhere.

TABLE 6

ANALYTIC CASES SEEN AT KFSHRRC BY SITE\* AND AGE

Drail Cavity  Oral Cavity  Bosophayux  Esophagus  Stomach  Colon, Rectum  Colon, Colon, Colon  Colon, Colon  Colon, Colon  Colon, Colon  Colon  Colon, Colon	SITE	7-0	5-9	- 4	15- 19	50-	÷ &	ģ <u>%</u>	35- 39	-04	-67	50-5	55- 6	60 - 65	₽ ₹	κ. κ.	8 %	82 <del>+</del>	TOTAL
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iple Myeloma	g, Pleura	0	0	0	0											•	9	7	8
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0 0 2 6 13 19 18 20 10 10 6 0 4 0 0 1 0 0 0 0 0 0 0 1 0 0 0 0 0 0 0	in, CNS	54	27	<u>1</u> 3	2											-	0	0	122
6 0 4 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	roid	0	0	7	9											-	2	2	135
4 1 1 5 1 1 3 1 3 4  3 3 2 1 1 2 4 1 3 7  -LNs 2 10 10 15 6 4 7 4 0 5  0 0 0 0 0 0 0 0 0 0  0 1 0 0 0 0 1 1 5 5 2 3  0 0 1 1 1 1 5 0 0	er Endocrine	9	0	4	0											0	0	0	=
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	Other Sites	0	0	0	τ											-	0	0	2
TOTAL 114 72 71 81 64 80 110 140 125 149. 152	AL.	114	22	7	<b>8</b>						,	2 173	3 158	3 141	101	86	56	45	1.921

\* Includes Multiple Primary Neoplasms.

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1.5

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TOTAL

TABLE 7

ANALYTIC MALE CASES SEEN AT KFSH&RC BY SITE\* AND AGE 1996

TOTAL

SITE	7-0	5-9	<del>,</del> 5	- <del>5</del> 1	50-	- <del>5</del> 2	30-	35-	<b>\$ \$</b>	45-	50-	55-	\$ <b>\$</b>	69	2 %	ķ &	8 - - -	85+
Oral Cavity Nasopharynx	00	00	0 2	o -	o -	۰ -	2 4	2 2	<b>⊖</b> ∞	4 0	<b>~ ™</b>	2 <del>L</del>	s 4	ın M	٥ ٥	۰ د	~ 0	0 0
Esophagus	0	0	0	0	0	· •	0	8	0		~	. 0	~	ı K	· <del>-</del>	. 40	, <del>, .</del>	~
Stomach	0	0	0	0	0	0	_	0	-	0	7	æ	7	4	4	м	m	_
Colon, Rectum	0	0	0	~	0	4	_	_	7	4	8	9	9	7		m	0	
Liver	0	<del></del>	0	0	0	0	0	m	_	m	7	12	=	10	=	ထ	m	m
Pancreas	0	0	0	0	0	0	0	_	_	2	2	_	0	0	_	-	0	
Other G.I.	0	0	0	0	0	0	0	0	0	_	m	<b>,</b>	0	_		0	_	0
Larynx	0	0	0	0	0	0	0	0	_	2	7	'n	-	~	ī	-	m	2
Lung, Pleura	0	0	0	0	0	0	0	7	м	4	œ	11	٥	1	m	4	9	4
Multiple Myeloma	0	0	0	0	-	0	0	~	0	0	~	o	~	•	0	7	0	-
Lymphoid Leukemia	19	ĸ	7	9	M	_	_	-	0	0	0	0	_	0	0	<b>-</b> -	0	0
Myeloid Leukemia	4	4	0	7	M	4	M	4	2	0	_	-	0	0	0	0	-	0
Other Leukemias	-	_	0	<b>-</b>	0	0	0	0	0	_	0	0	0	0	0	0	0	0
Bone, Cartilage	<b>f</b> orm	4	9	∞	7	~	_	~	0	0	-	0	7	0	0	0	-	0
Soft Tissue	М	-	2	~	-	7	m	7	0	50	<b>-</b>	_	0	<b>,</b>	_	0	0	0
Skin Melanoma	0	0	0	0	0	0	-	<b>,</b>	0	0	0	0	0	0	0	0	0	0
Non-Melanoma Skin Ca	0	0	-	-	0	-	~	m	<b>-</b>	2	0	2	5	2	2	_	~	m
	0	0	0	0	0	0	0	0	0	-	0		0	0	0	0	0	0
Uterus, Genital	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Cervîx	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Ovary	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Prostate	0	0	0	0	0	0	0	0	0	•	~	-	M	'n	7	7	7	9
Testis, Genital	-	0	0	0	m	M	Α.	-	2	0	0	0	0	0	0	-	0	0
Bladder	-	0	0	0	0	_	0	2	_	7	9	m	~	4	ī	ī	m	Ŋ
Kidney, Urinary	•	0	0	0	0	0	o	0	_	0	0	m	4	'n	~	~	0	0
Eye	œ	0	0	0	0	0	0	0	0	0	<b>-</b>	0	0	0	0	0	0	0
Brain, CNS	ŧ.	16	7	M	N	2	4	м	м	m	ΙΛ	7	4	4	0		0	0
Thyroid	0	0	-	7	7	2	m	7	4	2	2	<b>,</b>	_	4	_	0	0	2
Other Endocrine	4	0	4	0	0	_	0	0	0	0	0	0	0	0	0	0	0	0
NHL - Lymph Nodes	M	<b>-</b> -	0	-	0	-	C4	_	_	M	5	m	0	2	~	м	M	0
WHL - Extra-nodal	-	М	7	-	_	2	m	_	M	7	7	~	_	īV	2	2	0	<b></b>
Hodgkin's Disease-LNs	_	æ	9	٥	-	m	4	7	0	7	_	_	4	0	0	0	0	<b>~</b> ~
HD - Extra-nodal	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Primary Unknown	0	-	0	0	0	-	0	2	<del></del>	7	~	~	~1	m	-	~	~	0
All Other Sites	0	0	0	-	_	0	0	0	0	0	_	_	0		_	<b>-</b>	0	0

\* Includes Multiple Primary Neoplasms.

TABLE 8

ANALYTIC FEMALE CASES SEEN AT KFSHRRC BY SITE\* AND AGE

un of the control of	SITE Oral Cavity	7-0	2-6	÷ 4 - 0	<u>2</u> 5 2 2 2	26-	52 23	3 34	39 39	04 40	45- 2 2	0.74 PZ 17	55 59 6	09 % E	% % .v	5 % E	ķε 40	8 % %		82+
Rectum  Bectum  Bectum	Nasopharynx Esophagus	~ 0	00	00	v 0	0 0	o <del>-</del>	<b>4</b> –		0 7	m N	ın N	0 4	- 4	~ -	0 2	0 ~	0 M		- 0
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ass 6.1.  Pleura 6.1  Pleura 6.1	Colon, Rectum	<b>5</b> 6	<b>-</b>	<b>5</b> (	<b>-</b> (	v	<b>v</b> (	- (	<b>~</b> •	۰ ،	4 •	Λ (		Λ t	vn •	٧,	<b>V</b> 1	4 (		- 4
Feura 1e Myeloma 15 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Pancreas	<b>-</b> -	- 0	<b>&gt;</b>	<b>.</b>	<b>-</b> -	<b>.</b>	<b>-</b>	- c	c		<b>~</b> -	<b>4</b> C	<b>~</b> ←		<b>4</b> -	າ 🗢	<b>-</b>		<b>-</b>
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Interview   Color	Lung, Pleura	0	0	0	0	0	0	0	0	0	7	7	м	-	7	M	~	0		
Interchanta   15	Multiple Myeloma	0	0	0	0	0	0	<del>-</del>	-	0	_	0	M	-	0	•	0	0	_	_
Checkenia   3   3   1   3   3   1   4   6   4   0   1   0   2   0   0   0   0   0   0   0   0	Lymphoid Leukemia	15	4	9	2	-	-	0	-	0	0	0	0	0	0	0	-	0		_
Leukemias 0 0 0 0 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0	Myeloid Leukemia	M	м	-	M	m	-	4	9	4	0	-	0	7	0	0	0	0	_	_
Centilege 1 1 1 8 5 3 0 1 0 0 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0	Other Leukemias	0	0	0	0	0	-	0	0	0	0	0	0	0	0	0	0	0	_	_
Tissue 5 0 2 1 2 0 1 3 1 4 2 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Bone, Cartilage		<b>-</b> -	∞	S	M	0	-	0	0	0	0	-	0	0	0	0	0	_	_
Helenoma Skin Ca	Soft Tissue	'n	0	7	-	~	0	_	M	<b>+-</b>	4	7	2	0	0	0	0	0	_	_
Welenoma Skin Ca         1         0         1         0         1         0         1         0         1         1         2         42         3         3         2         1         1         3         2         1         1         3         2         0         0         0         0         1         1         1         2         4         3         3         4         1         2         1         7         2         1         7         2         1         2         0	Skin Metanoma	0	0	0	0	0	-	_	_	<b>(</b>	0	0	0	0	0	0	0	0	_	_
ist  us, Genital  us, Commission  us, Genital  us, Commission  us, Genital  us, Commission  us, Commissi	Non-Melanoma Skin Ca	0	φo	0	-	0	-	<b>-</b> -	2	м	M	7	-	-	M	2	0	0	0	_
us, Genital         0         1         0         1         2         2         0         1         7         2         3         3         1         1         1         1         1         1         1         1         1         1         1         1         1         2         3         3         4         1         2         0 <th< td=""><td></td><td>0</td><td>0</td><td>0</td><td>-</td><td><b>(</b>-</td><td>Ξ</td><td>22</td><td>45</td><td>37</td><td>32</td><td>77</td><td>, K)</td><td>17</td><td>٥</td><td>M</td><td>-</td><td>7</td><td>N</td><td></td></th<>		0	0	0	-	<b>(</b> -	Ξ	22	45	37	32	77	, K)	17	٥	M	-	7	N	
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its, Genital 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Ovary	0	-	2	M	4	m	7	M	4	м	M	4	-	m	7	-	0	_	_
Hermore III	Prostate	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	٥	_
New Contract         O <t< td=""><td>Testis, Genital</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>0</td><td>3</td><td>_</td></t<>	Testis, Genital	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	3	_
ley, Urinary 5 2 0 0 0 1 0 0 3 2 1 2 1 0 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Bladder	0	0	0	0	0	0	0	0	•	0	-	-	0	7	0	-	0	_	_
n, CNS  oid  oid  oid  oid  - Lymph Nodes  1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Kidney, Urinary	'n	2	0	0	0	-	0	0	м	7	-	7	•-	0		0	0	0	_
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id 0 0 1 4 11 17 15 13 6 8 5 10 5 2 1 1 2 Endocrine 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Brain, CNS	•	=	•	2	~	7	7	-	0	m	7	-	~	m	0	0	0	0	
Extra-nodal 2 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Thyroid	0	0	-	4	-	17	15	13	9	∞	Ŋ	욘	'n	7	_	-	7	0	_
Lymph Nodes 1 0 1 4 1 0 1 0 2 1 4 1 5 0 1 3 1 Extra-nodal 2 0 0 0 0 0 1 0 0 3 2 4 2 5 1 2 1 ints Disease-LNs 1 2 4 6 5 1 3 2 0 1 0 0 0 0 1 0 0 1 0 0 0 0 0 0 0 0 0	Other Endocrine	7	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	-	_
Extra-nodal 2 0 0 0 0 0 1 0 0 3 2 4 2 5 1 2 1 1 1 1 1 1 1 1 1 1 1 1 1 2 1	NHL - Lymph Nodes		0	-	4	<b>~</b> ~	0	_	0	7	-	4	-	'n	0	_	m	-		_
ints Disease-LNs	MHL - Extra-nodal	7	0	0	0	0	0	_	0	0	M	2	4	~	10		7	•	_	_
Extra-nodal 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	Hodgkin's Disease-LNs		7	7	9	ın	_	м	7	0	-	0	0	0	0	0	-	0	0	_
ry Unknown 0 0 0 0 0 0 1 3 1 1 2 1 3 2 3 0 0 ther Sites 0 0 0 0 1 0 0 0 0 0 0 0 there sites 0 0 0 0 0 1 0 0 0 0 0 0 0 0 0 0 0 0 0	HD - Extra-nodal	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	_	_
ther Sites 0 0 0 0 0 1 0 0 0 0 1 0 0 0 0 0 0 0 0	Primary Unknown	0	0	0	0	0	0	-	м	<b>e</b>	*	2	-	M	7	M	0	0	•	
48 27 33 39 41 48 73 93 89 80 85 71 53 37 28 18	All Other Sites	0	0	0	0	0	-	0	0	0	0	-	0	0	-	0	0	0	-	_
	TOTAL	87	23	33	ŝ	14	84	ĸ	8	8	8	8	85	F	53	37	82	∞	12	•

\* Includes Multiple Primary Neoplasms.

#### TRENDS IN RELATIVE FREQUENCY OF CANCER AT KFSH&RC (cont'd)

The relative frequencies of primary cancers seen at KFSH&RC are very different from the Western world. Common tumors of the West (lung, colon, and prostate) are much less frequent here while leukemia and thyroid cancer, among others, are more common. The following 1996 analytic cases exhibit significant differences in trends from those of the West when compared to the data published in Cancer Facts & Figures - 1996, by the American Cancer Society:

Breast - The most common malignancy seen at KFSH&RC is breast cancer, comprising 12.0% of all cases, as compared to 13.7% of all neoplasms diagnosed in the U.S.A. It affects mostly women less than the age of 50, while in the U.S.A., those more than 50 years of age are mostly affected. As in the Western countries, it is the number one cancer among women.

Leukemia - Leukemia constitutes 7.5% of all cases seen at KFSH&RC, as compared to about 2% of all neoplasms diagnosed in the U.S.A. The male/female ratio is 1.2:1. It is the second most common type of malignancy seen in males and the third most common in females. It is also the most common malignancy among pediatrics.

Thyroid - 3.6% of all male malignancies in KFSH&RC are thyroid tumors. However, they represent 10.5% of female malignant neoplasms, second to breast cancer. The male/female ratio is 0.3:1. Thyroid cancer accounts for only 1% of all cases in the U.S.A. and 2% of female malignancies.

Non-Hodgkin's Lymphoma - The most striking feature is the unusually high crude relative frequency of non-Hodgkin's lymphoma, accounting for 7.0% of all cases. The male/female ratio is 1.7:1. In the U.S.A., NHL accounts for only about 4% of all cancer.

Brain/CNS - Primary malignant neoplasm of the brain and CNS accounts for 6.4% of all malignancies and ranks second among the most common pediatric malignancies. The male/female ratio is 1.8:1. This is comparatively higher than in the West with only 1.3% of all cases.

Oral Cavity - A high crude relative frequency rate was also noted in cancer of the oral cavity. In Western countries, oral cancer accounts for no more than 3% of all cancers, whereas at KFSH&RC it represents 5.1% of the cases. The male/female ratio is 0.9:1, and 2.1:1 in the West.

Liver - The relative frequency of liver cancer at KFSH&RC (4.6%) is more than three times higher than that of the West (1.5%). The male/female ratio (3.2:1) is also significantly higher than in the West (1.2:1).

Lung - Frequency of lung cancer is much lower than in Western countries, most likely reflecting the much lower levels of smoking and industrial pollution. In the U.S.A., primary lung cancer represents 13.0% of all cancer cases (12.9% in males, and 13.1% in females). At KFSH&RC, 4.2% of all diagnoses are lung cancer; in males it is the fifth most common tumor, constituting 6.8% of male malignancies and 1.6% in females. The male/female ratio is 4.3:1, in the West, 1.3:1.

Colo-Rectal - Markedly less common than in the West, this disease represents only 4.2% of all tumors. In the U.S.A. it constitutes 9.8% of newly diagnosed cancer cases. Dietary factors, particularly lower animal fat intake, may play a role. The male/female ratio at KFSH&RC is 1:1.

Hodgkin's Disease - The incidence of Hodgkin's lymphoma is comparatively more frequent at KFSH&RC than in Western countries. In the U.S.A. it constitutes 0.6% of all cancers, compared to 3.7% at KFSH&RC. The male/female ratio is 1.7:1, in the West, 1.1:1.

Bone - A higher crude relative frequency rate is seen in bone cancer. It constitutes 0.2% of the all cancers in most centers in the West, but is 2.7% of the cases at KFSH&RC. The male/female ratio at KFSH&RC is 1.6:1.

Soft Tissue - KFSH&RC cases show a higher rate of soft tissue malignancies than the U.S.A., with 2.5% against the latter's 0.5% of all cases. The male/female ratio is 1.1:1, in the West, 1.2:1.

Prostate - The observed rate of prostatic cancer in men is much lower than in the West, where it is one of the most common male cancers (constituting 36% of the male malignancies). This is in contrast to the KFSH&RC experience, where prostatic cancer makes up only 4.1% of the male cancer. This is probably due to the population age difference. Prostate cancer is a disease chiefly of old men and the population of Saudi Arabia is, in general, very young.

#### FIGURE 10

#### DISTRIBUTION OF 20 MOST COMMON MALIGNANCIES 1996 ANALYTIC CASES (TOTAL CASES = 1,921)

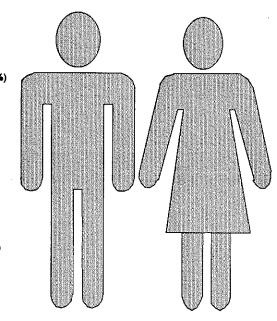
MALE **FEMALE** NHL 84 (8.8%) BREAST 229 (23.8%) **LEUKEMIA 80 (8.4%)** THYROID 101 (10.5%) BRAIN, CNS 79 (8.3%) LEUKEMIA 64 (6.6%) LIVER 68 (7.1%) NHL 50 (5.2%) LUNG, PLEURA 65 (6.8%) ORAL CAVITY 50 (5.2%) NASOPHARYNX 53 (5.5%) BRAIN, CNS 43 (4.5%) ORAL CAVITY 47 (4.9%) CERVIX 42 (4.4%) HODGKIN'S DISEASE 45 (4.7%) COLON, RECTUM 41 (4.3%) BLADDER 40 (4.2%) OVARY 39 (4.1%) COLON, RECTUM 39 (4.1%) NASOPHARYNX 26 (2.7%) PROSTATE 39 (4.1%) HODGKIN'S DISEASE 26 (2.7%) THYROID 34 (3.6%) UTERUS, GENITAL 26 (2.7%) STOMACH 34 (3.6%) **ESOPHAGUS 25 (2.6%)** BONE, CARTILAGE 32 (3.3%) **SOFT TISSUE 23 (2.4%)** LARYNX 31 (3.2%) NON-MELANOMA SKIN 21 (2.2%) NON-MELANOMA SKIN 28 (2.9%) LIVER 21 (2.2%) SOFT TISSUE 25 (2.6%) BONE, CARTILAGE 20 (2.1%) ESOPHAGUS 22 (2.3%) KIDNEY, URINARY 18 (1.9%) KIDNEY, URINARY 21 (2.2%) PRIMARY UNKNOWN 18 (1.9%) PRIMARY UNKNOWN 21 (2.2%) STOMACH 17 (1.8%)

#### FIGURE 11

# DISTRIBUTION OF PEDIATRIC MALIGNANCIES 1996 ANALYTIC CASES (TOTAL CASES = 257)

#### MALE

LEUKEMIA 41 (27.5%) BRAIN, CNS 38 (25.5%) HODGKIN'S DISEASE 15 (10.1%) BONE, CARTILAGE 11 (7.4%) NHL 10 (6.7%) EYE 8 (5.4%) **ENDOCRINE 8 (5.4%)** (Other Than Thyroid) **SOFT TISSUE 6 (4.0%)** KIDNEY, URINARY 4 (2.7%) NASOPHARYNX 2 (1.3%) LIVER 1 (0.7%) NON-MELANOMA SKIN 1 (0.7%) TESTIS, GENITAL 1 (0.7%) **BLADDER 1 (0.7%)** THYROID 1 (0.7%) PRIMARY UNKNOWN 1 (0.7%)



#### FEMALE

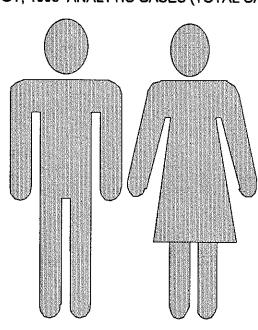
LEUKEMIA 32 (29.6%) BRAIN, CNS 23 (21.3%) BONE, CARTILAGE 10 (9.3%) **SOFT TISSUE 7 (6.5%)** KIDNEY, URINARY 7 (6.5%) HODGKIN'S DISEASE 7 (6.5%) EYE 6 (5.6%) NHL 4 (3.7%) OVARY 3 (2.8%) NASOPHARYNX 2 (1.9%) ENDOCRINE 2 (1.9%) (Other Than Thyroid) UTERUS, GENITAL 1 (0.9%) THYROID 1 (0.9%) ORAL CAVITY 1 (0.9%) NON-MELANOMA SKIN 1 (0.9%) LIVER 1 (0.9%)

FIGURE 12

DISTRIBUTION OF 10 MOST COMMON PEDIATRIC MALIGNANCIES BY HISTOLOGY, 1996 ANALYTIC CASES (TOTAL CASES = 257)

#### MALE

ALL 31 (20.8%)
HODGKIN'S DISEASE 15 (10.1%)
ASTROCYTOMA 13 (8.7%)
MEDULLOBLASTOMA 11 (7.4%)
NHL 10 (6.7%)
EWING'S SARCOMA 8 (5.4%)
RETINOBLASTOMA 7 (4.7%)
GLIOMA, NOS 8 (4.0%)
NEUROBLASTOMA 5 (3.4%)
RHABDOMYOSARCOMA 5 (3.4%)



#### FEMALE

ALL 25 (23.1%)
HODGKIN'S DISEASE 7 (6.5%)
ASTROCYTOMA 7 (6.5%)
MEDULLOBLASTOMA 6 (5.6%)
RETINOBLASTOMA 6 (5.6%)
NEPHROBLASTOMA 6 (5.6%)
EWING'S SARCOMA 5 (4.6%)
GLIOMA, NOS 5 (4.6%)
AML 5 (4.6%)
OSTEOSARCOMA 5 (4.6%)

TABLE 9

PRIMARY SITE TABLE
(INCLUDES MULTIPLE PRIMARIES)
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Primary Site Table (cont'd)

SITE HISTOLOGY (NOS-Not Otherwise Specified)	ALL CASES	MALE	OULTS FEMALE	PED: MALE	IATRICS FEMALE
PHARYNX	2	1	1	0	0
Non-Hodgkin's Lymphoma	_	_	_	•	•
ESOPHAGUS	52	25	27	0	0
Squamous Cell Carcinoma	49	22	27	Ŏ	Ŏ
Adenocarcinoma, NOS	2	2	Ö	Ō	ō
Carcinoma, NOS	ī	1	0	0	0
STOMACH	71	44	27	0	0
Adenocarcinoma, NOS	31	20	11	0	0
Non-Hodgkin's Lymphoma	14	7	7	0	Ō
Signet Ring Cell Carcinoma	8	6	2	0	0
Adenocarcinoma, Intestinal Type	5	4	1	0	0
Carcinoma, NOS	4 .	3	1	0	0
Squamous Cell Carcinoma	2	1	1	0	0
Leiomyosarcoma	2	1	1	0	0
Malignant Neoplasm, NOS	2	0	2	0	0
Linitis Plastica	1	0	1	0	0
Mucinous Producing Adenocarcino	ma 1	1	0	0	0
Carcinoma, Diffuse Type	1	1	0	0	0
SMALL INTESTINE	13	6	5	1	1
Non-Hodgkin's Lymphoma	7	3	2	1	1
Adenocarcinoma, NOS	3	2	1	0	0
Mucinous Adenocarcinoma	1	1	0	0	0
Carcinoid Tumor	1	0	1	0	0
Neuroendocrine Carcinoma	1	0	1	0	0
COLON	37	19	17	1	0
Adenocarcinoma, NOS	23	11	12	0	0
Carcinoma, NOS	4	2	2	0	0
Non-Hodgkin's Lymphoma	3	2	0	1	0
Mucinous Adenocarcinoma	2	1	1	0	0
Adenoca in Tubulovillous Adenoma	a 2	1	1	0	0
Carcinoid Tumor	1	0	1	0	0
Signet Ring Cell Carcinoma	1	1	0	0	0
Adenocarcinoma in Adenomatous Po	olyp 1	1	0	0	0
RECTUM/RECTOSIGMOID JUNCTION/ANUS	60	35	25	0	0
Adenocarcinoma, NOS	43	25	18	0	0
Mucinous Producing Adenocarcino	na 8	3	5	0	0
Signet Ring Cell Carcinoma	3	2	1	0	0
Squamous Cell Carcinoma	3	2	1	0	0
Carcinoma, NOS	2	2	0	0	0
Cloacogenic Carcinoma	1	1	0	0	0
LIVER/INTRAHEPATIC BILE DUCTS	95	71	22	1	1
Hepatocellular Carcinoma	88	68	20	0	0
Cholangiocarcinoma	3	1	2	0	0
Mucinous Adenocarcinoma	1	1	0	0	0
Carcinoma, NOS	1	1	0	0	0
Rhabdomyosarcoma	1	0	0	1	0
Malignant Neoplasm, NOS	1	0	0	0	1

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### Primary Site Table (cont'd)

SITE HISTOLOGY	ALL (	CASES	AI	ULTS	PED	IATRICS
(NOS-Not Otherwise Spec	ified)		MALE	FEMALE	MALE	FEMALE
GALLBLADDER/EXTRAHEPATIC BILE	DUCTS	20	8	12	0	0
Adenocarcinoma, NOS		14	5	9	Ō	Ō
Carcinoma, NOS		3	1	2	Ŏ	Ö
Mucinous Adenocarcinoma		1	1	Ö	ō	Ŏ
Signet Ring Cell Carcino	ma	1	ī	Ō	Ö	Ŏ
Papillary Adenocarcinoma		1	ō	1	ō	ŏ
PANCREAS		20	12	8	0	0
Adenocarcinoma, NOS		12	8	4	Ō	Ō
Carcinoma, NOS		6	3	3	ō	Ö
Carcinoid Tumor		1	1	Ō	ō	Ö
Malignant Neoplasm, NOS		1	0	1	ō	ō
OTHER G.I. SITES		2 .	1	0	1	0
Non-Hodgkin's Lymphoma						
NASAL CAVITIES/ACCESSORY SINU	SES	12	8	3	1	0
Squamous Cell Carcinoma		3	3	, O	0	0
Esthesioneuroblastoma		3	1	2	0	0
Non-Hodgkin's Lymphoma		3	2	0	1	0
Carcinoma, NOS		2	1	1	0	Ō
Adenoid Cystic Carcinoma		1	1	0	0	Ō
LARYNX		37	33	4	0	0
Squamous Cell Carcinoma		32	29	3	0	0
Verrucous Carcinoma		1	1	0	0	0
Adenosquamous Carcinoma		1	1	0	0	Ō
Papillary Squamous Cell (	Carcinoma	1	1	0	0	Ō
Neuroendocrine Carcinoma		1	1	0	Ō	Ō
Malignant Neoplasm, NOS		1	0	1	Ö	Ō
BRONCHUS/LUNG		85	69	16	0	0
Squamous Cell Carcinoma		36	33	3	ō	ō
Adenocarcinoma, NOS		22	13	9	ŏ	Ö
Carcinoma, NOS		8	7	í	Ö	Ö
Small Cell Carcinoma		4	3	ī	Ö	Õ
Large Cell Carcinoma		3	2	ī	Ö	0
Clear Cell Carcinoma		3	3	ō	0	0
Bronchiolo-Alveolar Adend	carcinoma	2	2	Õ	Ö	0
Adenosquamous Carcinoma	Julionia	2	2	0	Ö	0
Malignant Neoplasm, NOS		2	2	0	0	0
Lymphoepithelioma-Like Ca	ercinoma	1	1	0	Ö	0
Carcinoid Tumor	II CITIONIA	1	Ŏ	1	Ö	
Non-Hodgkin's Lymphoma		1	1	Ō	0	0
PLEURA		2	2	o	0	•
Mesothelioma		•	-	U	U	0
THYMUS/MEDIASTINUM		7	2	3	0	2
Malignant Thymoma		2	1			
Neuroblastoma		1	0	1 0	0	0
Leiomyosarcoma		1	0		0	1
Ganglioneuroblastoma		_		1	0	0
Limphospitholises Tile C		1	0	0	0	1
Lymphoepithelioma-Like Ca	rcinoma	1	1	0	0	0
Carcinoma, NOS		1	0	1	0	0

Primary Site Table (cont'd)

AULTIPLE MYELOMA  SONE MARROW  Acute Lymphoid Leukemia  Acute Myeloid Leukemia  Chronic Myeloid Leukemia  Chronic Lymphoid Leukemia  Acute Promyelocytic Leukemia  Chronic Myelomonocytic Leukemia	20 165 84 34 25 6	12 46 14	8	0	0
Acute Lymphoid Leukemia Acute Myeloid Leukemia Chronic Myeloid Leukemia Chronic Lymphoid Leukemia Acute Promyelocytic Leukemia Chronic Myelomonocytic Leukemia	84 34 25	14	34		
Acute Myeloid Leukemia Chronic Myeloid Leukemia Chronic Lymphoid Leukemia Acute Promyelocytic Leukemia Chronic Myelomonocytic Leukemia	34 25			46	39
Chronic Myeloid Leukemia Chronic Lymphoid Leukemia Acute Promyelocytic Leukemia Chronic Myelomonocytic Leukemia	25		5	34	31
Chronic Lymphoid Leukemia Acute Promyelocytic Leukemia Chronic Myelomonocytic Leukemia		12	10	6	6
Acute Promyelocytic Leukemia Chronic Myelomonocytic Leukemia	6	11	12	1	1
Chronic Myelomonocytic Leukemia	•	4	2	0	0
	6	3	2	0	1
Nauta Mualamanasitis Taribanis	3	0	1	2	0
Acute Myelomonocytic Leukemia	2	0	1	1	0
Acute Monocytic Leukemia	2	1	. 0	1	0
Megakaryocytic Leukemia	1	0	0	1	0
Hairy Cell Leukemia	1	1	0	0	0
Malig Lymphoproliferative Disease	1 .	0	1	0	0
PLEEN Non-Hodgkin/g Tymphoma	2	1	0	0	1
Non-Hodgkin's Lymphoma					
BONE & CARTILAGE	60	24	13	13	10
Ewing's Sarcoma	25	7	5	8	5
Osteosarcoma, NOS	16	7	1	3	5
Chondroblastic Osteosarcoma	4	1	2	ī	ō
Chondrosarcoma, NOS	$\bar{4}$	3	1	Ö	Ō
Non-Hodgkin's Lymphoma	3	1	1	ì	Ö
Malig Adamantinoma of Long Bone	2	ī	ī	ō	Ö
Juxtacortical Osteosarcoma	ī	ī	ō	ŏ	Ö
Small Cell Osteosarcoma	ī	ī	Õ	ő	Ö
Malig Giant Cell Tumor of Bone	ī	ō	í	Ö	ő
Plasmacytoma	ī	í	Ō	ŏ	Ö
Chordoma	ī	ī	Ŏ	0	0
Peripheral Neuroectodermal Tumor	1	ō	ĭ	ŏ	ő
CONNECTIVE/SUBCUTANEOUS/SOFT TISSUE	62	26	21	8	7
Malignant Fibrous Histiocytoma	8	3	4	1	Ö
Leiomyosarcoma	7	2	5	ō	Õ
Non-Hodgkin's Lymphoma	6	2	4	Õ	Ö
Synovial Sarcoma	5	3	1	Ö	1
Alveolar Rhabdomyosarcoma	4	1	Ō	1	2
Myxoid Liposarcoma	4	2	1	ī	0
Spindle Cell Sarcoma	3	2	1	ō	Ö
Rhabdomyosarcoma, NOS	. 3	ō	ō	ĭ	2
Sarcoma, NOS	3	2	Ŏ	ō	1
Peripheral Neuroectodermal Tumor	3	1	1	1	ō
Clear Cell Sarcoma of Tendon	2	ō	2	ō	ŏ
Liposarcoma	2	2	ō	ŏ	ő
Malignant Neurilemmoma	2	2	Ö	ő	0
Embryonal Rhabdomyosarcoma	ī	0	0	1	0
Fibrosarcoma, NOS	1	0	0	1	0
Malignant Neuroepithelioma	1	0	1	0	0
Ganglioneuroblastoma	1	0	0	0	1
		1	0		
Malignant Mesenchymoma	1	0		0	0
Malignant Hemangioendothelioma	1	_	1	0	0
Kaposi's Sarcoma	1	1	0	0	0
Neuroblastoma	1	0	0	. 1	0
Extraskeletal Chondrosarcoma Extraskeletal Ewing's Sarcoma	1 1	1	0 0	0	0

Primary Site Table (cont'd)

SITE (NOS-Not	HISTOLOGY  therwise Specified)	ALL CASES	ai Male	OULTS FEMALE	PED: MALE	IATRICS FEMALE
SKIN (MELANOMA)		8	3	5	0	0
SKIN (NON-MELANC	DMA)	69	40	26	2	1
Basal Cell	Carcinoma	27	14	12	0	1
Squamous Ce	ll Carcinoma	20	10	8	2	0
Kaposi's Sa	rcoma	10	10	0	0	0
Dermatofibr		7	3	4	0	0
Mycosis Fun	goides	4	3	1	0	0
	denocarcinoma	1	0	1	0	0
BREAST, FEMALE		277	0	277	0	0
Duct Cell C	arcinoma	218	0	218	0	0
Lobular Car	cinoma	20	0	20	0	0
Paget's Dis	ease & Duct Cell Ca	10	0	10	0	0
Carcinoma,		7	0	7	0	0
Medullary C		5	Õ	5	ō	Ō
Comedocarci		4	ŏ	4	Ö	Ö
	lenocarcinoma	3	ō	3	Ŏ	Ö
Paget's Dis		2	0	2	Ŏ	Ŏ
Adenocarcin		2	ŏ	2	Ŏ	Õ
	's Lymphoma	2	ŏ	2	ő	Ö
	y Carcinoma	1	ő	1	ő	ő
	bular Carcinoma	1	Ö	1	ŏ	ő
		_	Ö	1	0	Ö
Hemangiosar	: Carcinoma, Non-Invasi coma	1	0	1	ŏ	ŏ
BREAST, MALE		2	2	0	0	0
Duct Cell C	arcinoma	1	1	0	0	0
Papillary C		1	1	0	0	0
CERVIX UTERI		47	0	47	0	0
Squamous Ce	ell Carcinoma	41	0	41	0	0
Adenocarcin	oma, NOS	3	0	3	0	0
Clear Cell	Carcinoma	1	0	1	0	0
Adenosquamo	us Carcinoma	1	0	1	0	0
Carcinoma,	NOS	1	0	1	0	0
PLACENTA		9	0	9	0	0
Choriocarci	noma	8	0	8	0	0
Trophoblast	ic Tumor	1	0	1	- 0	0
CORPUS UTERI		19	0	18	0	1
Adenocarcin		9	0	9	0	0
Endometrial	Stromal Sarcoma	4	0	3	0	1
	d Carcinoma	2	0	2	0	0
Papillary S	erous Carcinoma	1	0	1	0	0
	lenocarcinoma	1	0	1	0	0
Mullerian M		1	0	1	0	0
Leiomyosaro	oma	1	0	1	0	0
OVARY		54	0	51	0	3
	Serous Cystadenocarcino		0	7	0	0
Adenocarcin		7	0	7	0	0
Carcinoma,		6	0	6	0	0 0
	denocarcinoma	5	0	5	0	0
	d Carcinoma	4	0	4	0	0

#### Primary Site Table (cont'd)

SITE (NOS-No	HISTOLOGY ALI t Otherwise Specified)	L CASES		OULTS FEMALE		LATRICS FEMALE
	· · · · · · · · · · · · · · · · · · ·		<del>.</del>			4.5
OVARY (cont'd)		4	•		_	•
	/stadenocarcinoma Sinus Tumor	4	0	4	0	0
	– –	3	0	1	0	2
Dysgerminon		3	0	3	0	0
	Serous, Borderline Malig	3	0	3	0	0
	Cystadenocarcinoma	2	0	2	0	0
	adenoma, Borderline Mali		0	2	0	0
	ystadenoma, Border Malig	2	0	2	0	0
	Neoplasm, NOS	2	0	2	0	0
Serous Cyst	tadenocarcinoma	1	0	1	0	0
Mixed Germ	Cell Tumor	1	0	0	0	1
Teratoma		1	0	1	0	0
Malignant 0	Granulosa Cell Tumor	1	0	1	0	0
OTHER FEMALE GE	NITAL ORGANS	4	o	4	0	0
Squamous Ce	ell Carcinoma	2	0	2	0	0
Basal Cell		ī	· ō	1	Õ.	-
	denocarcinoma	1	ŏ	ī	o .	Ö
PROSTATE		49	49	0	0	0
Adenocarcin	noma	44	44	Ö	. 0	Ö
Carcinoma,		5	33	0	ŏ	0 '
TESTIS		14	12	0	2	0
Seminoma		6	-6	0	ō	ŏ
Mixed Germ	Call Tumon	3	3	0	0	0
				-	-	-
	Sinus Tumor	3	1	0	2	0
Teratocarci	nabdomyosarcoma Inoma	1 1	1 1	0	0 0	0 0
	TAT ODGANG		•	•	•	
OTHER MALE GENI: Squamous Ce	ell Carcinoma	1	1	0	0	0
URINARY BLADDER		56	47	8	1	0
Transitiona	al Cell Carcinoma	22	18	4	ō	Ö
	Pransitional Cell Ca	19	18	1	ŏ	Ö
	ell Carcinoma	8	5	3	Ö	0
		_				-
Adenocarcin		2	2	0	0	0
Carcinoma,		2	_	0	0	0
	Rhabdomyosarcoma	1	0	0	1	0
Carcinosard		1	1	0	0	. 0
Non-Hodgkir	n's Lymphoma	1	1	0	0	0
KIDNEY/URETER		48	21	14	6	7
Renal Cell	Carcinoma	24	17	7	0	0
Nephroblast	coma	11	0	0	5	6
	Adenocarcinoma	5	2	3	0	0
Adenocarcin		2	ō	2	ō	ō
	al Cell Carcinoma	ī	ŏ	1	ŏ	ŏ
Chromophobe		i	1	0	Ö	0
	ell Carcinoma	1	0	1		
					0	0
	Sarcoma of Kidney	1	0	0	1	0
Carcinoma,		1	1	0	0	0
Malignant N	Weoplasm, NOS	1	0	0	0	1

Primary Site Table (cont'd)

SITE	HISTOLOGY (NOS-Not Otherwise Specified)	ALL CASES	MALE	OULTS FEMALE	PED: MALE	IATRICS FEMALE
EYE		24	4	1	10	9
R	etinoblastoma	16	0	0	8	8
M	[elanoma	3	2	0	0	1
N	on-Hodgkin's Lymphoma	2	1	0	1	0
	quamous Cell Carcinoma	1	1	0	0	0
	mall Cell Carcinoma	1	0	1	0	0
M	alignant Neoplasm, NOS	1	0	0	1	0
BRAIN		126	43	22	36	25
G	lioblastoma	27	15	8	2	2
A	strocytoma, NOS	23	12	6	4	1
M	Medulloblastoma	19	1	1	11	6
M	Malignant Glioma, NOS	15	5	1	4	5
P	ilocystic Astrocytoma	14	0	0	7	7
A	naplastic Astrocytoma	5	3	1	0	1
0	ligodendroglioma	5	3	2	0	0
P	rimitive Neuroectodermal Tumor	4	0	1	2	1
N	on-Hodgkin's Lymphoma	4	2	2	0	0
P	leomorphic Xanthoastrocytoma	3	1	0	1	1
E	pendymoma	3	0	0	2	1
F	ibrillary Astrocytoma	1	0	0	1	0
_	Germinoma	1	1	0	0	0
C	arcinoma, NOS	1	0	0	1	0
M	Malignant Neoplasm, NOS	1	0	0	1	0
OTHER	NERVOUS SYSTEM	6	1	1	4	0
M	Malignant Glioma, NOS	2	0	0	2	0
M	Malignant Meningioma	2	1	1	0	0
P	ilocytic Astrocytoma	1	0	0	1	0
N	on-Hodgkin's Lymphoma	1	0	0	1	0
THYRO	ID.	163	46	113	2	2
P	apillary Carcinoma	· 115	28	85	1	1
	apillary & Follicular Adenoca	19	6	13	0	0
	Medullary Carcinoma	8	5	2	0	1
	ollicular Adenocarcinoma	7	2	5	0	0
A	naplastic Carcinoma	6	2	4	0	0
C	arcinoma, NOS	3	1	1	1	0
	on-Hodgkin's Lymphoma	3	2	1	0	0
M	Malignant Neurilemmoma	1	0	1	0	0
T	rabecular Adenocarcinoma	1	0	1	0	0
OTHER	ENDOCRINE GLANDS	11	1	0	8	2
	Meuroblastoma	5	ō	ō	4	_ 1
	Germinoma	4	Ō	0	4	0
	ineoblastoma	1	0	0	0	1
	fixed Germ Cell Tumor	<u>-</u>	1	Ō	Ö	ō
	NODES, NON-HODGKIN'S LYMPHOMA	81	44	27	8	2
	Excluding Extra-Nodal Lymphoma)				_	_
	arge Cell, Diffuse	26	16	10	0	0
	Jon-Hodgkin's Lymphoma	11	6	3	2	0
	mmunoblastic	7	5	2	0	0
	arge Cell, Follicular	7	4	3	0	0
	ymphoblastic	4	0	1	2	1
K	(i−1	4	2	1	1	0

Primary Site Table (cont'd)

SITE	HISTOLOGY	ALL	CASES	AI	OULTS	PEDIATRICS		
	(NOS-Not Otherwise Specified)			MALE	FEMALE	MALE	FEMALE	
LYMPH N	ODES, NON-HODGKIN'S LYMPHOMA	(cont	′d)					
	cluding Extra-Nodal Lymphoma							
Bu	rkitt's		4	0	0	3	1	
Sma	all Lymphocytic		4	2	2	0	0	
Sma	all Cleaved Cell, Follicular		3	3	0	0	0	
	Cell Rich B-Cell		2	2	0	0	0	
Moi	nocytoid B-Cell		2	1	1	0	0	
Mi:	xed Small & Large Cell, Diffu	se	2	1	1	0	0	
	llicular, NOS		2	2	0	0	0	
Lyn	mphoplasmacytic		1	0	1	0	0	
	xed Small Cleaved & Lge Cell,	Foll	1	0	1	0	0	
Sm	all Cleaved Cell, Diffuse		1	0	1	0	0	
LYMPH N	ODES, HODGKIN'S DISEASE		75 .	33	19	15	8	
No	dular Sclerosis		61	24	18	12	7	
Ly	mphocytic Predominance		7	5	1	0	1	
	xed Cellularity		5	2	0	3	0	
	dgkin's Disease, NOS		2	2	0	0	0	
PRIMARY	UNKNOWN		41	21	19	1	0	
Ad	enocarcinoma		15	8	7	0	0	
Ca	rcinoma, NOS		13	4	9	0	0	
Ma	lignant Neoplasm, NOS		3	1	2	0	0	
C1	ear Cell Adenocarcinoma		2	2	0	0	0	
Ne	uroendocrine Carcinoma		2	1	1	0	0	
Sq	uamous Cell Carcinoma		1	1	0	0	0	
si	gnet Ring Cell Carcinoma		1	1	0	0	0	
	pillary Carcinoma		1	1	0	0	0	
	enosquamous Carcinoma		1	1	0	0	0	
	rcinoid Tumor		1	1	0	0	0	
Ma	lignant Gastrinoma		1	0	0	1	0	

TABLE 10

PATIENTS WITH MULTIPLE PRIMARIES
1 9 9 6

PRIMARY SITE HISTOLOGY 1996 (NOS-Not Otherwise Specif	OTHER PRIMARIES ied) (PREVIOUS OR CONCURRENT)	ALL	MALE	FEMALE
		65	32	33
ORAL CAVITY		4	3	1
Sq Cell Ca-Buccal Mucosa	Bladder - Sq Cell Ca	1	1	0
Sq Cell Ca-Buccal Mucosa	Tongue - Sq Cell Ca	1	1	0
Verrucous Ca - Lower Gum	Prostate - Adenoca	1	1	0
NHL - Tonsil	Breast - Duct Cell Ca	1	0	1
NASOPHARYNX		2	2	0
Undiff Carcinoma	Larynx - Sq Cell Ca	1	1	0
Undiff Carcinoma	Ethmoid Sinus - Sq Cell Ca	1	1	0
ESOPHAGUS		1	1	0
Squamous Cell Carcinoma	Larynx - Sq Cell Ca	1	1	0
STOMACH		2	1	1
Adenocarcinoma	Larynx - Sq Cell Ca	1	1	Ö
Leiomyosarcoma	Uterus - Adenocarcinoma	ī	ō	1
COLON		2	2	0
Carcinoma, NOS	Prostate - Adenocarcinoma	1	1	0
Adenocarcinoma	Ileum - NHL	1	1	0
RECTUM		3	2	1
Adenocarcinoma	Hodgkin's Disease	1	0	1
Adenocarcinoma	Colon - Adenocarcinoma	1	1	0
Mucinous Adenocarcinoma	Colon - Adenoca in Adenoma- tous Polyp	1	1	0
LIVER		1	1	0
Hepatocellular Ca	Tongue - Sq Cell Ca			
LARYNX		1	0	1
Squamous Cell Carcinoma	Breast - Duct Cell Ca			
LUNG		3	3	0
Carcinoma, NOS	Bladder - Trans Cell Ca	1	1	0
Bronchiolo-Alveolar Ca	Larynx - Sq Cell Ca	1	1	0
Adenosquamous Ca	Kidney - Renal Cell Ca	1	1	0
BONE MARROW		1	0	1
Acute Myeloid Leukemia	Breast - Duct Cell Ca	1	0	1
SOFT TISSUE		1	. 1	0
Malig Fibrous Histiocytoma	Skin - Dermatofibrosarcoma			
SKIN		1	1	0
Basal Cell Carcinoma*	Kidney - Renal Cell Ca LNs - NHL	-	_	

PRIMARY SITE HISTOLOGY 1996 (NOS-Not Otherwise Spec		ALL ASES	MALE	FEMALE
BREAST		22	0	22
Duct Cell Carcinoma	Contralateral Breast	18	0	18
Duct Cell Carcinoma	Ovary - Malig Teratoma	1	0	1
Medullary Carcinoma*	Contra Breast-Duct Cell Ca Same Breast - Duct Cell Ca	1	0	1
Lobular Carcinoma	Contralateral Breast	1	0	1
Carcinoma, NOS	Hodgkin's Disease	1	0	1
OVARY		3	0	3
Endometrioid Ca	Breast - Duct Cell Ca	ĭ	0	1
Mucinous Cystadenoca	Skin - Basosquamous Ca	ī	0	1
		1	0	1
Serous Cystic Tumor, Borderline Malig	Contra Ovary - Pap Serous Cystadenoma, Border Malig	Ŧ	U	1
PROSTATE		2	2	0
Adenocarcinoma	Kidney - Renal Cell Ca	1	1	0
Adenocarcinoma*	Kidney - Renal Cell Ca Bladder - Pap Trans Cell Ca	1	1	0
URINARY BLADDER		3	3	0
Papillary Trans Cell Ca	Kidney - Renal Cell Ca	1	1	0
Carcinosarcoma		1	1	0
Carcinoma, NOS	Larynx - Sq Cell Ca Thyroid - Pap & Follicular Ca	_	1	0
KIDNEY		3	3	0
Renal Cell Carcinoma	Skin - Sq Cell Ca	1	1	0
Renal Cell Carcinoma	Bladder - Trans Cell Ca	1	1	0
Renal Cell Carcinoma	Bladder - Pap Trans Cell Ca	1	1	0
CONJUNCTIVA		1	1	0
Squamous Cell Carcinoma	LNs - NHL			
BRAIN		1	1	0
Glioblastoma	Acute Lymphoid Leukemia			
THYROID		3	2	1
Follicular Adenoca	Breast - Duct Cell Ca	1	0	1
Follicular Adenoca	Small Intestine - NHL	1	1	0
Papillary Carcinoma*	Jejunum - Adenocarcinoma LNs - NHL	1	1	0
PINEAL GLAND		1	0	1
Pineoblastoma	Retinoblastoma			
LYMPH NODES		3	3	0
Non-Hodgkin's Lymphoma	Kidney - Renal Cell Ca	1	1	0
Non-Hodgkin's Lymphoma	Skin - Kaposi's Sarcoma	1	1	0
Non-Hodgkin's Lymphoma	Jejunum - Adenocarcinoma	1	1	0
UNKNOWN PRIMARY		1	0	1
Adenocarcinoma	Gallbladder - Adenoca In-Situ			

<sup>\*</sup> Patient has three primary malignancies.

#### STAGE OF DISEASE AT DIAGNOSIS

Stage in any malignant process may be defined as the particular step, phase, or extent in a tumor's development, which is one of the predictors for outcome and treatment selection assigned at the time of initial diagnosis. The microscopic appearance, extent, and biological behavior of a tumor, as well as host factors, play a part in prognosis and are therefore important in staging.

The SEER (Surveillance, Epidemiology, and End Results) Summary Staging Guide was utilized for all stageable cases. This system summarizes the disease categories into four general staging groups (i.e. in situ, localized, regional, and distant). Stage categories are based on a combination of clinical observations and operative-pathological evaluation.

Summary Staging Definitions:

IN SITU: Intraepithelial, noninvasive, noninfiltrating

LOCALIZED: Within organ

a. Invasive cancer confined to the organ of origin

b. Intraluminal extension where specified

REGIONAL: Beyond the organ of origin

a. By direct extension to adjacent organs/tissues

b. To regional lymph nodes

c. Both (a) and (b)

DISTANT: Direct extension or metastasis

a. Direct continuity to organs other than above

b. Discontinuous metastasis

c. To distant lymph nodes

Systemic diseases, i.e., leukemia and multiple myeloma and cases of unknown primary were disregarded in graphically illustrating the stages for all analytic cases seen at KFSH&RC in 1996 (Figure 13). The 16 cases unstageable at diagnosis were those patients who refused further diagnostic workup or further workup was not possible due to the patients' state of health; e.g. terminal cases or those with co-morbid conditions. Please refer also to Table 5 for the distribution of the 1996 analytic cases by site and stage at diagnosis.

In addition to the SEER Summary Staging, the cases were also staged according to the American Joint Committee on Cancer (AJCC) TNM system. This scheme is based on the premise that cancers of similar histology or site of origin share similar patterns of growth and extension. This system is based on the assessment of three components:

T: Extent of the primary tumor

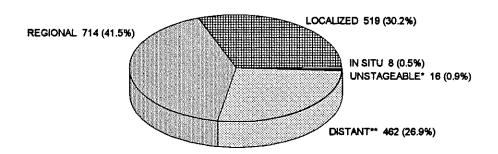
N: Absence or presence and extent of regional lymph node involvement

M: Absence or presence of distant metastasis

Analytic cases of four major sites, i.e., breast, lung, nasopharynx and Hodgkin's Disease are presented in Table 11 with their clinical group stage and yearly comparative figures from 1992 to 1996. The pathologic group stages of stomach and colorectum are also presented in the same table.

#### FIGURE 13

# DISTRIBUTION OF ANALYTIC CASES BY STAGE (SEER) AT DIAGNOSIS - 1996 (TOTAL CASES = 1,719)



\*Excludes Unknown Primaries (39 cases)
\*\*Excludes Leukemia and Multiple Myeloma (163 cases)

#### FIGURE 14

# DISTRIBUTION OF ANALYTIC CASES BY FIRST COURSE OF TREATMENT (SINGLY OR IN COMBINATION) 1996 (TOTAL CASES = 1,921)

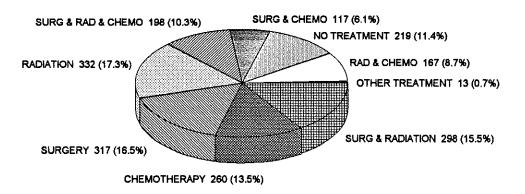


TABLE 11

AJCC CLINICAL THM GROUP STAGE OF ANALYTIC CASES OF MAJOR SITES\* BY YEAR

1992 - 1996

					8	REAST						
Stage	1	992	1 9	993	1.9	994	1 9	995	1 '	996	Т О	TAL
-	No	×	No	×	No	×	No	×	No	×	No	*
0	0	0.0	2	0.9	0	0.0	2	1.0	4	1.7	8	0.8
1	4	2.8	19	9.0	9	4.6	11	5.6	22	9.6	65	6.7
2A	22	15.4	29	13.7	44	22.6	39	19.7	52	22.6	186	19.0
2B	33	23.0	46	21.7	41	21.0	37	18.7	37	16.1	194	19.8
3A	21	14.7	26	12.3	14	7.2	19	9.6	19	8.3	99	10.1
3B	32	22.4	37	17.4	31	15.9	29	14.6	22	9.6	151	15.4
4	23	16.1	41	19.3	27	13.8	32	16.2	36	15.6	159	16.3
Unstageable	8	5.6	12	5.7	29	14.9	29	14.6	38	16.5	116	11.9
Total	143	100.0	212	100.0	195	100.0	198	100.0	230	100.0	978	100.0
					Ĺ	UNG						
Stage	1	992	1 9	993	1 1	994	1 1	995	1 1	996	ΤO	TAL
_	No	x	No	×	No	x	No	x	No	×	No	×
1	5	7.3	3	4.5	13	16.9	8	10.3	2	2.6	31	8.4
2	1	1.5	2	3.0	4	5.2	2	2.6	4	5.1	13	3.6
3A	13	19.1	6	9.0	4	5.2	9	11.5	6	7.7	38	10.3
3B	11	16.2	23	34.3	24	31.1	22	28.2	26	33.3	106	28.8
4	35	51.5	24	35.8	17	22.1	28	35.9	27	34.6	131	35.6
Unstageable	3	4.4	9	13.4	15	19.5	9	11.5	13	16.7	49	13.3
Total	68	100.0	67	100.0	77 -	100.0	78	100.0	78	100.0	368	100.0
					NASOP	HARYNX						
Stage	1	992	1 (	993	1 (	994	1 (	9 5	1 (	996	тο	TAL
o tage	No	×	No	, , <u>,</u> ,	No	×	No	x	No	×	No	* *
0	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0
1	Ö	0.0	1	1.9	1	1.9	1	1.8	ō	0.0	3	1.0
ż	ō	0.0	Ċ	0.0	1	1.9	3	5.3	2	2.5	6	2.1
3	3	6.5	4	7.4	4	7.5	7	12.5	7	8.9	25	8.7
4	42	91.3	47	87.0	47	88.7	44	78.6	67	84.8	247	85.8
Unstageable	1	2.2	2	3.7	0	0.0	1	1.8	3	3.8	7	2.4
Total	46	100.0	54	100.0	53	100.0	56	100.0	79	100.0	288	100.0

<sup>\*</sup> Excludes Lymphoma Cases

Table 11 (contid)

# AJCC CLINICAL GROUP STAGE OF ANALYTIC CASES OF MAJOR SITE BY YEAR 1992 - 1996

					HODGKII	4'S DISEAS	Έ					
Stage		992	1	993	1	994	1	995	1	996	то	TAL
	No	x	No	X	No	×	No	*	No	x	No	×
1A	8	12.7	12	21.1	12	17.9	16	20.5	8	11.3	56	16.6
18	3	4.8	3	5.3	2	3.0	1	1.3	1	1.4	10	3.0
2A	15	23.8	11	19.3	19	28.4	19	24.4	20	28.2	84	25.0
2B	8	12.7	6	10.5	3	4.5	10	12.8	10	14.1	37	11.0
3A	11	17.4	6	10.5	9	13.4	10	12.8	6	8.4	42	12.5
3B	7	11.1	8	14.0	12	17.9	11	14.1	9	12.7	47	14.0
4A	2	3.2	1	1.8	1	1.5	2	2.6	ź	4.2	9	2.7
4B	9	14.3	10	17.5	9	13.4	9	11.5	14	19.7	<b>5</b> 1	15.2
Total	63	100.0	57	100.0	67	100.0	78	100.0	71	100.0	336	100.0

# AJCC PATHOLOGIC TNM GROUP STAGE OF ANALYTIC CASES OF MAJOR SITES\* BY YEAR 1992 - 1996

					STO	<b>MACH</b>						
Stage	1	992	1	993	1	994	1	995	1	996	τo	TAL
	No	×	No	×	No	*	No	×	No	*	No	, K L
0	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0	0	0.0
1A	2	6.5	0	0.0	1	2.1	1	2.1	ž	3.9	6	3.0
18	1	3.2	0	0.0	3	6.4	2	4.3	3	5.9	9	4,4
2	4	12.9	6	21.4	7	14.9	4	8.5	0	0.0	21	10.3
3A	3	9.7	4	14.3	6	12.8	5	10.6	13	25.5	31	15.2
38	1	3.2	3	10.7	3	6.4	6	12.8	4	7.8	17	8.3
4	5	16.1	0	0.0	1	2.1	2	4.3	1	2.0	9	
Unstageable	15	48.4	15	53.6	26	55.3	27	57.4	28	54.9	111	4.4 54.4
Total	31	100.0	28	100.0	47	100.0	47	100.0	51	100.0	204	100.0
					COLON	, RECTUM						
Stage	1.9	992	1 9	9 3	1 9	94	1.9	9 9 5	1 (	96	т о	TAL
	No	*	No	×	No	×	No	x	No	×	No	* X
0	0	0.0	0	0.0	1	1.6	0	0.0	0	0.0	1	0.3
1	5	7.3	2	3.8	3	4.7	4	5.7	3	3.8	17	5.1
2	11	16.2	8	15.4	8	12.5	16	22.9	17	21.2	60	18.0
3	21	30.9	12	23.1	19	29.7	14	20.0	10	12.5	76	22.8
4	3	4.4	8	15.4	5	7.8	5	7.1	6	7.5	27	8.1
Unstageable	28	41.2	22	42.3	28	43.7	31	44.3	44	55.0	153	45.7
Total	68	100.0	52	100.0	64	100.0	70	100.0	80	100.0	334	100.0

<sup>\*</sup> Excludes Lymphoma Cases

# PALLIATIVE CARE FOR PATIENTS WITH ADVANCED CANCER Alan Gray, MB ChB, FRACP, FRACR Adnan Ezzat, MB, BCh, FRCPC, FACP King Faisal Specialist Hospital & Research Centre

#### Introduction

The true incidence of cancer in the Kingdom of Saudi Arabia (KSA) is just now being measured. Hospital statistics reflect patterns of referral, rather than the real incidence of the various types of cancer. In 1993 a National Cancer Registry (NCR) was established, collecting data from all regions, and publishing its first report in 1996. Over the next few years, the NCR will establish a baseline for the incidence of cancer, and provide a means of monitoring the success of its treatment for the whole country.

Much about cancer in KSA can be implied from the experience of other countries. Cancer is particularly a disease of old age when it is one among many causes of morbidity and mortality. Although cancer becomes more common as we age, it differs from other diseases of the elderly, in that cancer can occur at any age. With its large population of young people, cancer in children is also a significant problem in KSA. 12.6% of the cancer patients seen at King Faisal Specialist Hospital and Research Centre (KFSH&RC) are children. This is a much higher proportion than are seen in similar referral centers in the West. (1) The average life expectancy for men and women in the Middle East is estimated to be 62 years. By comparison, in the West life expectancy is 75 years or more. (2) With the rapid improvements in nutrition, sanitation, housing, and the highly effective vaccination programs that have been developed, the Saudi populations life expectancy will rapidly increase. The national census conducted in 1992, shows that the average life expectancy here is already 70 years. We can reasonably predict that this increase in life expectancy will be accompanied by an alteration in the patterns of disease similar to that in Western countries; where infectious diseases leading to premature death have been replaced by more chronic illnesses of old age, such as heart disease, cancer, and cerebro-vascular disease. In the U.S.A., cancer accounts for 22% of all deaths, second only to heart disease. In general, in Western countries, one in every three people will develop cancer during their lifetime, and one in every four will die of it.

#### Prevention

McDonald of Canada has provided a useful description of four phases of cancer "prevention" covering all aspects of the disease:

- 1. Prevention of the disease (public education and policy).
- Prevention of advanced disease (early diagnosis programs).
- Prevention of death (anti-cancer treatment).
- 4. Prevention of suffering (palliative care) (4)

Cancer is one of the most treatable of all chronic illnesses. This aspect of cancer is often overlooked, because the disease arouses such fear. It is the most feared of all illnesses, and its diagnosis is usually associated in the patient's mind with premature and unpleasant death. In fact, however, in the West 40-50% of cancer patients will be cured; that is, they will have a normal life expectancy for their age and sex. (2) In children the cure rates are higher, over 60%. (5) However, these figures rely upon the fact that 70% of cancer patients in the West present with "early" cancer, stage 1 and 2. At KFSH&RC only 30% of patients present with localized stage 1 or 2 disease. Nearly 70% have stage 3 or 4 advanced disease with a consequently much worse prognosis. (1) As these patients are accepted at this major tertiary referral center on the basis of their likelihood of responding to treatment, other centers are likely to see even more advanced disease. It is likely that 80% or more of cancer patients in this country present with advanced disease, when cure rates will be low. Even so after surgery, chemotherapy, or radiotherapy, or by the use of one or all three modalities, dramatic remissions can be obtained which may last from months to

years before the patient's condition deteriorates again. At present however, probably 70% or more of cancer patients in Saudi Arabia will eventually die of their disease.

While there is constant incremental progress in the treatment of cancer, with improvements continually being introduced into the Kingdom, major progress could be made if patients could be encouraged to present early. This involves firstly, improving the public's knowledge about their personal health problems and the benefit of presenting early with their symptoms, and secondly, to ensure that patients with symptoms do not have a delay in diagnosis and treatment.

#### The Public's Perception of Cancer

Bedikian and Saleh interviewed one hundred Saudi patients with cancer. They reported 92% had an adverse reaction to the diagnosis, and the median duration of symptoms was from 3-5 months. None had received professional assistance for these problems, and the authors recommended psycho-social support be available to patients as part of their total management. (6) Bedikian in a further study interviewed two hundred and fifty healthy Saudis on their attitudes and knowledge of cancer. This study revealed a considerable degree of fear and anxiety about this disease. (7) Similarly, Ibrahim et. al. interviewed six hundred adults on their attitudes to cancer and confirmed the high level of fear and misperceptions about the disease. (8) Both groups of authors called for more comprehensive health education and awareness of cancer as a treatable disease.

#### Palliative Care

Dying is as natural an event as being born. This is recognized by Islam and its acceptance of death as an expression of God's will. In Western countries where the population has a long life span and cancer is common, dying of cancer has a special dimension that has made it the most feared of all illnesses. This is because cancer deaths are rarely easy. The distress it causes is legend. Eighty percent of dying cancer patients will suffer pain, and in 60% it will be severe and require strong analgesics of the morphine type. (9) Pain is only one of the distressing symptoms caused by advanced cancer. Anorexia, weight loss, tiredness, malaise, shortness of breath, and confusion are some of the other many symptoms that cancer can cause as it spreads to vital organs. Added to the physical distress of advancing cancer is the emotional distress of impending death.

The explosion of medical technology and the opportunity to finally cure major disease began at the end of the nineteenth century. This new era may have obscured the need to provide a supportive and caring environment for patients with chronic illnesses. Cancer was once considered by some to be an untreatable disease, and eminent physicians were known to discharge cancer patients once they were diagnosed, on the grounds that nothing further could be done for them. Presently, an enormous amount of knowledge has evolved on the nature and treatment of cancer. Much of this research for newer and better methods of cure has been concentrated on changes at a cellular level. However, in the last twenty years, there has also evolved a body of knowledge on the treatment of the symptoms of advanced cancer, which has failed anti-cancer treatments; this has now become a specialized area of interest for doctors and nurses called palliative medicine. While its practitioners are few, its attitudes are important to medical systems in general, for the following reasons:

#### 1. Providing Continuity of Care

In most Western countries patient care is based upon the primary care physician (family practitioner). This doctor will have a formal or informal "contract" with an individual or family to undertake their medical care and act on their behalf at all times. A well organized family practice or polyclinic will have a computerized "template" of routine examinations, screening investigations, and a complete medical record of the patient and their family environment. When a patient has a terminal illness, they have a doctor they know and can trust, and in turn the doctor knows both the patient and their family. This continuity of care is important, particularly for chronic illnesses and for a terminal disease. It means

that "no cure" is not the same as saying "no care". Unfortunately at present in Saudi Arabia, patients with either a terminal or chronic illness have their continuing care, even as outpatients, provided mainly by a hospital service. (11) This type of care should become an essential part of the primary health care system of the Kingdom.

#### 2. Cure Or Palliation?

The emphasis for cancer patients in Saudi Arabia is on curative treatment. There is a strong tendency here to deny the patient information on their illness. This is a problem shared by other countries, but for the last twenty years in Western countries, doctors have treated the individual patient as an autonomous agent, capable of receiving information and acting upon it. This allows the individual patient to give true informed consent by understanding the advantages and disadvantages of the proposed treatment. This also allows the patient to more fully participate and cooperate with the treatment, and by such compliance improve the chance of success. (12) In Saudi Arabia, the predominant principle is "beneficence", where the patient is viewed as one member of the larger family and the family is responsible for the patient. The consent for the patient's treatment is usually a substitute consent by the family, whose purpose is to avoid disturbing the patient emotionally. By this means, the family consider they are protecting the patient from harm, and that if the patient is told the truth, then they will not be able to adapt to the situation and may lose hope. This attitude seems to particularly apply to female patients. However, Abu Aisha wrote that, it is recognized in Saudi Arabian law that a woman is legally considered to be a responsible citizen, and it follows therefore that an adult female patient of sound mind has the right to give her own consent after being adequately informed of the nature of her illness. (13) There is now ample evidence that patients cope better with a serious illness if they are informed. (12) Without such vital information on their illness Saudi patients will continue to travel overseas at considerable cost, with the unrealistic expectation that they can find a cure somewhere. Unfortunately, the development of market medicine elsewhere in the world feeds this expectation, by giving treatment which may be neither necessary or appropriate. If the only purpose of medicine is the saving of people from death, then obviously, medicine cannot win. The art of good medicine, is to decide when life sustainment is no longer possible, and therefore when to allow death to occur without further impediment. (14) Physicians should also bear some responsibility for the quality of their patient's death. (15)

#### 3. The Relief of Pain

Chronic pain, particularly as it occurs in cancer, has several distressing features:

- It gets progressively worse.
- It has no meaning.
- It creates a feeling of hopelessness.
  - It dominates the patient's life.
- It can destroy their will to live.

One part of pain is its perception, the other the emotional response to it. This is why people experience different degrees of pain. Pain is precisely what the patient says it is, and hurts as much as they say it hurts. Pre-conceived ideas of how much pain patients will or should have are best avoided. There is a general lack of knowledge of pain relief in the Kingdom, and in many hospitals adequate analgesics of the morphine type are simply not available. There is an unreasonable fear of morphine addiction amongst patients and their families, but studies have convincingly shown that addiction is never a problem in a terminal illness. (16)

The important point for doctors to know is that no patient needs to endure pain so severe that their work, sleep, and quality of life are ruined. It should be rare for dying cancer patients to have uncontrolled pain. The World Health

Organization recommends a three step process in pain control moving from simple analgesics such as tylenol on the first step, to tylenol and codeine based compounds ± NSAID's on the second step, and morphine or similar compounds for patients with severe or step 3 pain. (19) Each of these regimens require that medications be given regularly around the clock, not PRN. This requires careful monitoring by the physician of the patient's pain level, and their response to medications. Regular analgesics and continuity of care is the secret to good pain control.

#### Cultural Aspects

The family unit is the structural foundation of Saudi society. The impression is that patients here cope better with a terminal illness in their home, than happens elsewhere in the West. This is probably because of the close family bonds and their strong Islamic faith with its obligation to provide for parents or children in case of need, and to help make their lives as comfortable as possible. The corollary of this is, that a small input of medical and nursing care, results in a magnified response by the extended family in the care of a patient. This can be very gratifying to the doctor and nurse.

There is an impression that palliative care in the Kingdom is confused with euthanasia, which is totally forbidden by Islam. (17) Palliative care is the moral and ethical alternative to euthanasia. Providing comfort, relieving distress, controlling pain, and offering a service that is available 24 hours a day, sustains the patient's hope. It is not based on the false hope of providing inappropriate and ineffective treatments. This society quite rightly sets great store by hope, and palliative care increases that hope, that each day may be more comfortable than the last.

# The King Faisal Specialist Hospital & Research Centre Palliative Care Program

In 1991 a Home Health Care Program was developed at KFSH&RC. This was in response to concerns by staff regarding the distress suffered by patients with terminal cancer, and the loss of the benefits of good quality medical care once the patient returned home. (18) Two years of planning and research went into developing the program. What began as essentially a nursing service for patients with a terminal illness, developed over the years into a nursing and medical service. These efforts resulted in the development of a Section of Palliative Care Medicine in the Department of Oncology in 1996; this appears to be the first such service in the Arab world.

At present, a multi-disciplinary team includes: three palliative care consultants, six nurses, five translator/drivers, a palliative care nurse clinician, and social workers. This team provides total care for the patient and support for the family, as a unit. The program consults with other health disciplines to provide the most appropriate care. The service runs 24 hours a day, seven days a week. To date, over 1350 patients living in Riyadh have been referred to the Home Health Care program. An increasing number of other patients who live outside Riyadh are being followed up in the Palliative Care outpatient clinics. While the Home Health Care Program in Riyadh includes cases of chronic illness, palliative care for a terminal illness makes up 75-80% of the workload. At present the program has between 40-50 terminally ill patients at any one time being cared for at home in Riyadh. Six in-patient beds are provided in the hospital. Many patients require at least one admission for control of symptoms that cannot be managed at home, and for stabilization of symptoms before they are transferred either to home or to a hospital elsewhere in the Kingdom. The average duration of stay has been 6-7 days.

#### Conclusion

What are the aims of a good health service? The American President's Commission for the study of ethical problems in medicine provided the following definition: "to provide treatment that will restore patients to as near normal or usual a

quality of life as is possible under the circumstances" or, put more simply, "to maximize the patient's well-being". $^{(19)}$ 

The patient's perception of this is that the health services will:

- Restore them to good health.
- Improve their function. 2.
- Relieve their suffering (particularly if the first two are not possible). 3.

That we fail to cure all our patients, does not mean that we should fail to care for them and relieve their suffering. The purpose of a palliative care service at the KFSH&RC is "the care and study of patients with active, progressive, far advanced disease; for whom cure is impossible, the prognosis predictably short, and the focus of care is the quality of life". Palliative Care has broadened the spectrum of health services available to cancer patients in the Kingdom. It has proven that it is possible to provide palliative care of a standard similar to that in a Western country. Cultural aspects of life in the Middle East, in particular the strong family bonds, the acceptance of matters of life and death, and the emphasis on hospitality, actually enhance and promote this form of care, which empowers Saudi families to care for dying patients until the end of life.

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#### EXTRANODAL NON-HODGKIN'S LYMPHOMA:

#### THE PREDICTIVE VALUE OF THE INTERNATIONAL PROGNOSTIC INDEX

A. Ezzat, A. Jamshed, F. Zwaan, Y. Khafaga, M. Raja, S. Bazarbashi, A. Rostom and M. Ranginwala

Non-Hodgkin's lymphoma (NHL) is common in Saudi Arabia, accounting for 7.8% of all newly diagnosed malignancies. It ranks second in males and fourth in females. Only 20% of the patients are diagnosed at an early stage, according to the 1994 National Cancer Registry Report. At King Faisal Specialist Hospital and Research Centre (KFSH&RC) the majority (93%) of NHL are of aggressive histology, with extranodal involvement occurring in 34%.

It is increasingly recognized that a number of prognostic factors may influence the outcome in aggressive NHL. In recent attempts to categorize NHL patients uniformly, members of the international non-Hodgkin's lymphoma prognostic factor project, formulated a predictive model using recognized prognostic factors such as: age, stage, performance status (PS), number of extranodal sites and serum level of lactic dehydrogenase (LDH). This International Prognostic Index (IPI) classified patients into low, low-intermediate, intermediate-high and high risk categories; depending on the number of risk factors where complete remission (CR), relapse free survival (RFS) and overall survival (OS) correlate with the prognostic subgrouping. This index, however, included all patient groups.

A study was performed at KFSH&RC to test the predictive value of IPI in patients with early stage extranodal NHL. This included 106 consecutive patients referred to KFSH&RC between 1985 and 1994, who met the study criteria of age >15 years, stage IE and stage IIE extranodal NHL of intermediate and high grade histologies (excluding lymphoblastic lymphoma), as per the International Working Formulation (IWF). The patient characteristics are shown in Table 1 and the extranodal sites are shown in Table 2.

The prognostic index for each patient was calculated by scoring the absence or presence of a risk factor as 0 or 1, respectively. These were: age (<60 vs >60 years), performance status (ECOG 0, 1 vs >2), extranodal site (0, 1 vs >1) and serum LDH (normal vs elevated). Patients were grouped into three risk categories: low risk 80 (76%), low-intermediate 13 (12%) and intermediate-high 13 (12%). None was in the high risk group.

The management of these patients utilizing either radiation and/or chemotherapy is shown in Table 3. Anthracyclin based combination chemotherapy was used in the majority (85%). The WHO criteria for response and the statistical program SPSS were used to analyze the data. Clinical complete remission (CR) was achieved in 97 (91%), partial remission (PR) in 4 (4%), progressive disease (PD) in 2 (2%), while 3 (3%) are not assessable for response because they did not complete the treatment. Twenty patients have relapsed (including 4 who died of other diseases) and 9 were lost to follow up. At the time of analysis, 73 patients remain alive and disease free, 8 are alive with active disease and 25 had died (including 8 deaths due to other causes). CR rates and 5 years relapse free and overall survivals in relation to the index categories are shown in Table 4 and Figures 1 and 2.

It could be concluded, therefore, that the International Prognostic Index is a good predictor model in extranodal, aggressive NHL in the KFSH&RC patient population. While no substantial change in therapeutic intervention is required for patients at low and low-intermediate risk categories, patients who are at intermediate-high risk should be considered for more aggressive or experimental approaches.

Table 1
PATIENT CHARACTERISTICS

		Number	Percentage
Sex	Male	66	62
	Female	40	38
Age	<60	65	61
	>60	41	39
B Symptoms	Yes	47	44
<b>2 1</b> 1 mp • • · · · · ·	No	59	56
Performance Status	0-1	71	67
WHO	≥2	35	33
Bulk of Disease	<10cm	56	53
_	>10cm	25	23.5
	Unknown	25	23.5
Serum LDH	Normal	74	70
	Elevated	32	30
Histology	Intermediate	91	86
(IWF)	High (excl lymphoblastic)	15	14

Table 2
EXTRANODAL SITES

Site	Number	Percentage
Gastrointestinal	34	32
Thyroid	16	15.1
Soft Tissue	6	5.7
Head and Neck	44	41.5
Others	. 5	4.8
Bone	1	0.9
TOTAL	106	100

Table 3
PRESCRIBED TREATMENTS

Treatment	Number	Percentage
Radiation	20	19
Chemotherapy	74	70
Radiation/Chemotherapy	12	11
TOTAL	106	100

Table 4
COMPLETE REMISSION AND SURVIVAL RATES BY INDEX CATEGORY

IPI Category	Number	CR (%)	5 yr RFS(%)	5 yr OS(%)
Low	80	94	72	83
Low-Intermediate	13	100	53	65
Intermediate-High	13	70	32	41

Figure 1
OVERALL SURVIVAL BY INDEX
PRIMARY EXTRANODAL NHL

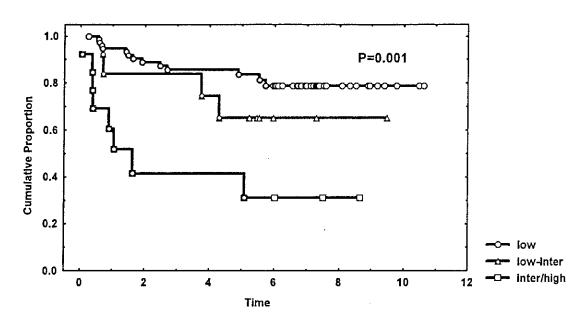
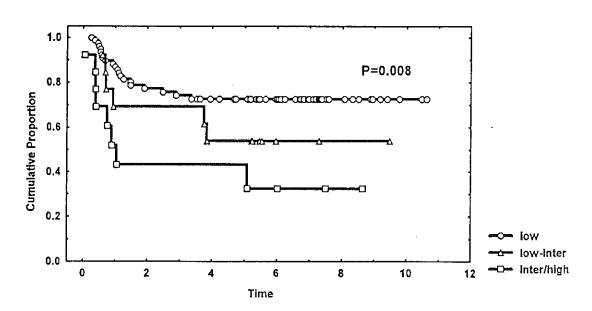


Figure 2
RELAPSE FREE SURVIVAL BY INDEX
PRIMARY EXTRANODAL NHL



#### APPENDIX A

1996 REQUESTS FOR TUMOR REGISTRY DATA
\*Publication \*\*KFSH&RC Presentation \*\*\*Outside KFSH&RC Presentation

January Adult Extraskeletal/Soft Tissue Osteosarcoma w/ Site, Stage, Treatment, Vital Status as of	Dr. M. Raja
Last Date of Contact (MR Numbers) (1975-1995) Breast Non-Hodgkin's Lymphoma (MR Numbers) (1975-present)	Dr. A. Ezzat
Pediatric Medulloblastoma Cases (MR Numbers) (Nov 1994-present)	Dr. M.Mustafa
Breast Cancer Cases w/ Mastectomy as Primary Treatment (MR Numbers) (1994-1995)	Dr. A. Ezzat
Breast Cancer Cases w/ Neoadjuvant FAC (MR Numbers) (1994-1995)	Dr. A. Ezzat
February	
Pediatric Acute Lymphoblastic Leukemia (MR Numbers) (1990-1995)	Dr. A. Al Nasser
Extranodal Head & Neck Non-Hodgkin's Lymphoma, Stages I & II, w/ Age, Sex, Site, B/A	Dr. A. Ezzat
Symptoms, Treatment (MR Numbers) (1980-1994) Pediatric Neuroblastoma Cases (MR Numbers) (1995-present)	Dr. M.Mustafa
Adult AML Cases w/ Treatment & Vital Status as of Last Contact Date (MR Numbers)(1981-1995)	Dr. E. Sahovic
1994 Malignant Cases by Site, Sex, Pediatrics Vs All Cases	Ministry of Health
April Acute Myeloid Leukemia Cases (MR Numbers) (1994-1995)	Ms. R. Gumaer
Мау	
Metastatic Breast Cancer Cases w/ Date of Dx and Site/s of Metastasis (MR Numbers) (1975-1991)	Dr. A. Ezzat
Breast Cancer Cases with Disease Relapse, w/ Date of Relapse and Site/s of Relapse	Dr. A. Ezzat
(MR Numbers) (1975-1991) Adult Head & Neck Sarcoma Cases (MR Numbers) (1980-1990)*	Dr. A. Ezzat
100 Non-Hodgkin's Lymphoma Cases w/ At Least 5 Yrs Survival, w/ Site, Histology, Stage, Age at Dx and Treatment (MR Numbers)	Dr. A. Kandil
Pediatric Cases with Second Primaries, w/ Sites, Histologies, Age, Dates of Dx and Treatment (MR Numbers) (1993-1995)**	Dr. M. Mahr
Lung Cancer Cases by Age, Sex, Region, Histology, Smoking Hx & Shamma/Shisha Usage (1985-1994)	Ministry of Petroleum
Breast Medullary Cancer Cases (MR Numbers) (1975-1995)	Dr. A. Ezzat
June Childhood Cancer Cases by Histology and Sex	D. M. Mustafa
(1975-1994) Adult Hodgkin's Disease Cases (MR Numbers) (1992-present)*	Dr. S. Bazarbashi

# Appendix A (cont'd)

(1992-present)* Urinary Bladder Cancer Cases by Treatment and by Dr Histology (1975-1994)**	r. S. Bazarbashi r. S. Bazarbashi
	r. S. Bazarbashi
	r. A. Ezzat
Adult CNS Leukemia Cases, Pre-Treatment and Post- Dr BMT (Last 10 years)	r. J. Chacko
July	
(MR Numbers) (1975-1995)	r. A. Ezzat
Esophageal Squamous Cell Carcinoma Cases by Dr the Clinical TNM Stage (1992-1995)	r. S. Bazarbashi
August	
Pediatric B-Cell Acute Lymphoblastic Leukemia Dr (MR Numbers) (1991-present)	r. M.Mustafa
	r. M. Mustafa
	c. A. Al Amro
	. M. Mustafa
	r. A. Gray
	r. W. Mourad
September	
Pediatric Acute Myeloid Leukemia (MR Numbers) Dr (1981-1996)	. H. El Solh
1995 Malignant Cases by Site, Sex, Pediatrics vs Min All Cases	inistry of Health
October	
Breast Cancer Cases vs Total Cancer Cases (1975- Dr August 1996)	. M. Vora
Pediatric Germ Cell Tumor Cases w/ Age, Sex, Site, Dr Histology, Date of Dx, Stage and Treatment (MR Numbers) (1975-1996)	. I. Fawaz
November	
	. H. Schultz

#### APPENDIX B

#### 1996 Tumor Committee Members

William Allard, D.M.D. Hamad Al Daig Saud Al Dossary Rita Anderson Shouki Bazarbashi, M.D.\* Edward De Vol, Ph.D. Poul Eriksen, M.D. Adnan Ezzat, M.D. Mohd Hannan, Ph.D. Stig Ingemansson, M.D. Justin Martin, M.D.\*\* Peter McArthur, M.D. Dolores K. Michels-Harper, C.T.R. Tumor Registry Lamia NouNou Assem Rostom, M.D. Rajeh Sabbah, M.D. \*\*\* Jens O. Sieck, M.D. Ofelia B. Te, C.T.R. Beth Ann Tomasek\*\*\* Edward Wiebe, M.D. Ferdinand Zwaan, M.D.

Dentistry CHIC Media Affairs Nursing Medical Oncology BS&SC Research Centre Obstetrics and Gynecology Medical Oncology B&MR Research Centre Surgery Pathology Surgery Social Services Radiation Oncology Chairman, Oncology Medicine Tumor Registry Quality Assurance Radiology Hematology/Oncology

- \* Chairman
- \*\* Deputy Chairman \*\*\* Ad hoc Members

## APPENDIX C

## SUMMARY OF CASES PRESENTED TUMOR BOARD - 1996

SITE	NO.
Lymphatic System Non-Hodgkin's Lymphoma Hodgkin's Disease	1 2
Brain	3
Breast	2
Liver	1
Ovary	1
Testis	1
Orbit	. 1
Unknown Primary w/ Metastasis	1
Metastatic Tumor to Palate	1
Neck Tumor	1
Mediastinal Lymphangioma	1
Spinal Cord Compression	1

Tumor Board Coordinator: Dr. Shouki Bazarbashi

#### APPENDIX D

## 1996 SUMMARY OF ONCOLOGY GRAND ROUNDS TOPICS

09	Jan	The Danish Breast Cancer Co-operative Group (DBCG)	Dr.	н.	Schultz
16	Jan	BMT in Children: KFSH&RC Experience and Future Directions	Dr.	н.	El Solh
12	Mar	BMT in Patients with Chronic Myeloid Leukemia	Dr.	N.	Chaudhri
26	Mar	Dose Intensive High Dose Chemotherapy with Autologous Bone Marrow and Stem Cell Transplantation im Breast Cancer Patients	Dr.	н.	Upadhyaya
09	Apr	Histiocytic Disorders of Children	Dr.	ĸ.	McClain
14	May	Psychoneuro Immunology: The Bone Marrow Transplant Model	Dr.	A.	Sullivan
28	May	Clinical Pathways	Ms.	J.	Al Dihan
11	June	Fanconi's Anemia "From Care To Cure"	Dr.	M.	Ayas
25	June	Intravesical Therapy: Does It Affect The Natural History of Superficial Bladder Cancer?	Dr.	s.	Bazarbashi
	July July	Stereotactic Radiosurgery The Measurement of Prostatic Specific Membrane Antigen - A New Prognostic Marker			Al Amro Murphy
24	Sept	Aplastic Anemia: Results of Immuno- suppressive Treatment & Bone Marrow Transplant	Dr.	F.	Zwaan
08	Oct	Molecular Monitoring of Bone Marrow Engraftment in the Allogeneic Setting	Dr.	М.	Gyger
12	Nov	Unproven Cancer Treatments - A North American Experience	Dr.	М.	Brigden
	Dec Dec	Drug and Radiation Interaction Secondary Malignancy in Childhood			Khafaga Al Mahr

#### V. GLOSSARY OF TERMS

Accessioned: Patients are entered into the Tumor Registry by the year in which they were first seen at KFSH&RC for each primary cancer.

Age of Patient: Recorded in completed years at the time of diagnosis.

Analytic Cases: Cases which were first diagnosed and/or received all or part of their first course of treatment at KFSH&RC.

Non-Analytic Cases: Cases diagnosed elsewhere and received all of their first course of treatment elsewhere.

Case: A diagnosis or finished abstract. A patient who has more than one primary is reported as multiple cases.

**Crude Relative Frequency:** The proportion of a given cancer in relation to all cases in a clinical or pathological series.

First Course of Treatment: The initial tumor-directed treatment or series of treatments, usually initiated within four months after diagnosis.

Stage of Disease: Determined at the time of the first course of treatment.

#### SEER (Surveillance, Epidemiology and End Results) Summary Staging:

In Situ: Tumor meets all microscopic criteria for malignancy except invasion.

Local: Tumor is confined to organ of origin.

Regional: Tumor has spread by direct extension to immediately adjacent organs and/or lymph nodes and appears to have spread no further.

**Distant:** Tumor has spread beyond immediately adjacent organs or tissues by direct extension and/or has either developed secondary or metastatic tumors, metastasized to distant lymph nodes or has been determined to be systemic in origin.

AJCC (American Joint Committee on Cancer) TNM Staging: A classification scheme based on the premise that cancers of similar histology or site or origin share similar patterns of growth and extension.

#### T+N+M = Stage

T: Extent of primary tumor

N: Extent of regional lymph node involvement

M: Distant Metastasis

Clinical Stage: Classification based on the evidence acquired before treatment. Such evidence arises from physical examination, imaging, endoscopy, biopsy, surgical exploration and other relevant findings.

Pathologic Stage: Classification based on the evidence acquired before treatment, supplemented or modified by the additional evidence acquired from surgery and from pathologic examination of the resected specimen.

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