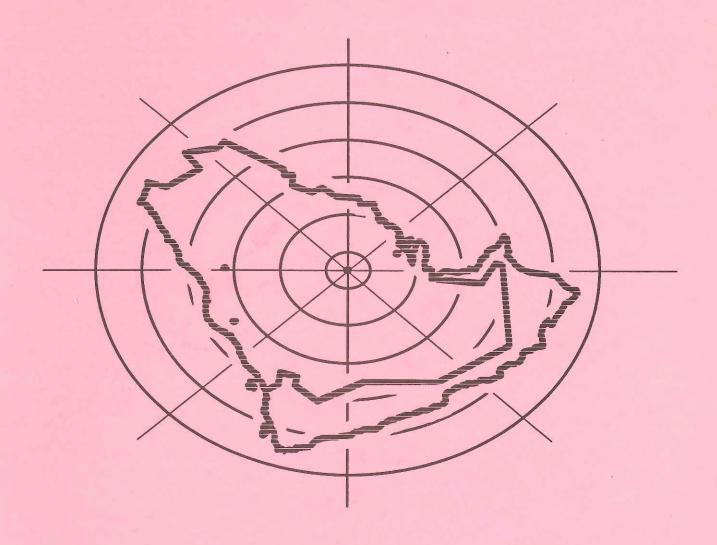
1993 ANNUAL REPORT OF THE TUMOR REGISTRY



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:

Department of Pathology & Laboratory Medicine Computer and Hospital Information Centre Medical Records Department Department of Oncology Home Health Care

SPECIAL THANKS TO:

Peter Ernst, M.D., Chairman, Tumor Committee (1993) Rajeh Sabbah, M.D., Acting Chairman, Oncology Department Mohd Maghazil, Computer & Hospital Information Centre Members of the Subcommittee on Annual Report:

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1993 ANNUAL REPORT OF THE TUMOR REGISTRY

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

TUMOR REGISTRY

<u>History</u>

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 hospital for Oncology and the principal center for cancer therapy in Saudi Arabia. There are about 500 inpatient beds.

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 more than 26,100 malignant cases seen at KFSH&RC from June 1975 through December 31, 1993. Approximately 2,000 new cases are added annually.

There are four (4) certified tumor registrars out of six (6) approved positions that 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 is the mechanism by 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.)

Data Use

Besides providing the statistics for the publication of the KFSH&RC annual report which summarizes the hospital's cancer experience, the data maintained in the Tumor Registry also support 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 1993, the Tumor Registry supported 52 special studies (see Appendix A for a listing of Special Studies requested in 1993).

Procedural and Administrative Changes During 1993

Oncology Clinic daily patient schedules are checked against the Master Index File in order to maximize new case identification on those patients without histopathology confirmation. These cases have the potential to be missed in the database if there is no inpatient admission since Medical Records Department does not code outpatient admissions.

A comprehensive review of the current automated system was conducted to include the data elements, entry, retrieval and edit functions. The recommendation was made to the Tumor Committee to obtain the updated Tumor Registry system from the American College of Surgeons as the current system is not meeting the needs of the clinicians or tumor registrars.

At the request of the Tumor Registry, the Computer and Hospital Information Centre (CHIC) developed a matching program of the Tumor Registry cases against the Admission/Discharge program. If there has been any patient activity since the <u>date of last contact</u> as noted on the Tumor Registry data set, the latest date the patient was seen is then electronically entered. This program is ran every two (2) months and has resulted in the follow up rate continuing to improve and it currently stands at 44% lost to follow up.

With the assistance of CHIC, an electronic productivity monitoring program directed at counting cases abstracted was developed. The tumor registrars rotate a manual quality control review of newly abstracted cases which supports the integrity of the data. Also, prior to microfiche, all medical records of deceased oncology patients 1975-1988 have been reviewed against the Tumor Registry data set to assure completeness and accuracy.

The Grade 8, Assistant Tumor Registrar and Grade 9, Associate Tumor Registrar job descriptions were revised in order to more clearly define job responsibilities and to facilitate recruitment. These were submitted to the Tumor Committee for approval.

In April, Tumor Registry personnel conducted a three-day training program in Tumor Registry Fundamentals for tumor registrars, data managers and data analysts from within the KFSH&RC and from outside hospital personnel.

Publications and Presentations

Michels D: Outcome Assessment - A Product of Comprehensive Followup. THE ABSTRACT, Vol 19:No 3, Journal of National Cancer Registrars Association, April 1993.

Michels D: <u>International Classification of Diseases-Oncology (ICD-O) Coding</u>. MODULE 7: FUNDAMENTAL TUMOR REGISTRY OPERATIONS PROGRAM. Edition 2, 1993, Cancer Department, American College of Surgeons.

Michels D: <u>Abstracting</u>. MODULE 11: FUNDAMENTAL TUMOR REGISTRY OPERATIONS PROGRAM. Edition 2, 1993, Cancer Department, American College of Surgeons.

Sackey K (presenter)....Anderson M, Atwood J, Becker N, Lange B, Michels D, Te O Neuroblastoma. RIYADH ONCOLOGY SYMPOSIUM, 27-28 April, 1993.

Michels D (Contributor, Oncology Care Indicator Development Task Force): <u>The Measurement Mandate</u>, On the road to performance improvement in health care. Joint Commission on Accreditation of Healthcare Organizations (JCAHO), 1993.

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 1993, the Committee provided professional and administrative guidance to the Tumor Registry and supported the following activities:

- A. Based on assessment of the existing electronic data system by Tumor Registry and Computer and Hospital Information Centre personnel, the Committee approved the recommendation that the updated software sysytem be purchased from the Americal College of Surgeons.
- B. A subcommittee was appointed to assess the hospital's need for patient educational brochures. The subcommittee is to review existing brochures and recommend to the departments if an update is needed. It is also to identify if patient brochures are needed on topics not available and recommend their development to the appropriate department.
- C. A subcommittee was appointed to develop guidelines for standardization of staging of cancer on newly diagnosed malignancies seen at KFSH&RC.

TUMOR BOARD

This educational conference is held as frequently as once weekly 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 1993.

ONCOLOGY GRAND ROUNDS

This didactic conference is held weekly 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 listing of the topics presented at the Oncology Grand Rounds in 1993.

FIGURE 1-A

PATIENT HAMEPIATE

KING FAISAL SPECIALIST HOSPITAL AND RESEARCH CENTRE

CANCER REGISTRY WORKSHEET (CanSur 3.0)

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Form 980-13 (Rev. 9-10)

FIGURE 1-B

PHYSICAL EXAM: 6-mo hx 2 cm mass rt breast UOQ, mobile, no skin changes. 3x4 cn rt axillary LN. Lt breast NED. x nays/scams:01/20/87 Bilat Mammogram - 2x2.5x2.5 cm mass rt breast UOQ. CXR, Bone Scan, U/S Abdomen - NED	TCAN - Cencer Identification (Continued) GRADE: 1 - Well differentiated (f) 5 - T-cell 2 - Mod well differentiated (fil) 6 - B-cell 3) Poorly differentiated (fil) 7 Null cell 4 - Undifferentiated (fil) 9 - Not stated, unknown LATERALITY: 0 - Not paked organ 3 - RI or it unspecified 4 - Brish, skeralization 7 - Inght 9 - Unknown laterality DX CONFIRMATION: 11 DX CONFIRMATION: 12 - Cytology 7 - Rachergraphy 4 - Post micro, confirm, NOS 5 - Clinical 9 - Unknown
SCOPES/IAB: 01/25/87 ERA (+), PRA (+)	REGIONAL NODES EXAMINED: 00 - No nodes examined 01 - One node examined 01 - One node examined 02 - 97 + noxitis examined 98 - Nodes examined, number unknown 99 - Unknown if notics examined 1 1 1 PEGIONAL NODES POSITIVE: 00 - No nodes positive 01 - One node positive 97 - Pusitive notics, number unknown 98 - No nodes examined 99 - Unknown if any nodus +/-
operatives findings: 01/25/87 Rt Mod_Rad_Mastectomy - no description of tumor.	96 - 98 + nodes positive TUMOR SIZE (cm) eg., 000 - No mass, 002 - 0.2 cm, 055 - 5.5 cm, 999 - Unknown RESIDUAL TUMOR: 1 - Microscopic
PATHOLOGY/AUTOPSY: 875P3286 01/25/87_Duct_Cell Ca, gr 3; 11/19 LN's. (tumor size: 2.2x2x1.8 cm completely excised) Nipple & overlying skin NED. (largest LN 1.5 cm)	DISTANT METS:
DATE OF INSTITUTE DESCRIPTION DATE OF INSTITUTE DIAGNOSIS: (man/dd/yyy) CLASS OF CASE: 0 - Dx here, rx elsowhere 1 Dx & rx here 2 - Rx here 3 Rx elsowhere 3 Rx elsowhere	ALCC STAGE: CLINICAL T 2 N 1 N 0 STAGE GROUP 2 B OTHERASS T 1 N N N N STAGE GROUP 2 B OTHERASS T 1 N N N STAGE GROUP 1 N STAGE GROUP 1 N N N N STAGE GROUP 1 N N N N N STAGE GROUP 1 N N N N N N N N N N N N N N N N N N
CODE: Breast, Right UOQ	eg, 3A- Stage IIIA, 1-Stage 1
HISTOLOGY - TEXT: Duct Cell Carcinoma, gr CODE: [8 5 0 0],[3	1 - 5(agu 1

FIGURE 1-C

PF 14 TRX1 ~ 1ST COURSE TREATMENT (SURGERY, RADIATION)	PF 16 TRX3 - 1st COURSE TREATMENT (CHEMO, HORMONES, BRM, OTHER)
SUNGERY	СНЕМОТНЕПАРУ
REASON:	SUMMARY: 31
Can directed surg 6 - Reason unknown, no surg - performed	AT THIS HOSPITAL:
7 - Patient/guardian refused 1 - Not recommended	0 - No chemotherapy 7 - Patient/guardian refused
8 - Recommended, unk if done 2 - Contraindicated, other	1 - Chemotherapy, NOS B - Recommended, unk if rione
9 - unknown	2 - Chemotherapy, single agent 9 - Unknown
SIAMMARY : (Entire 1st course).	3 Chemotherapy, multi-agent combination
AT THIS HOSPITAL: • 5.10	STARTED: (mm/dd/yyy) 02 /13 /19 8 7
Refer to Appendix A In CanSur User Manual for site specific codes.	TEXT: 5-FU, Adria, Ctx
STARTED: (nim/dd/yyyy) 011/25/1987	
TEXT: Rt Mod Rad Mastectomy w/ Rt	HORMONE/STEROIDS
Axillary Dissection	SUMMARY: 1
	AT THIS HOSPITAL :
RADIATICIN	0 - No hormonal therapy 7 - Patient/guardian refused
SUMMARY:	(1) Hormonal therapy 8 - Recommended, unk if done
AT THIS HOSPITAL	2 - Endocrine surg/radiation 9 - Unknown
0 · No Radiation therapy 5 - Radiation therapy, NOS [1]	3 - Hormones + endor surg/rad
1 Resultinadiation 7 - Patient/guardian refused	STARTED: (mm/dd/yyyy) 0 2 / 0 9 / 1 9 8 7
2 Redioactive implants 8 - Recommended, unk if done	TEXT: Tamoxifen
3 Radiolsotopes 9 - Urknown	
4 - Comb 1 + 2 or 3	
STARTED: (mm/vdd/yyyy) 0 8 / 29 / 1 98 7	BIO-RESPONSE MODIFIER (BRM)
TO BRAIN & CNS : (Lung & leukemia cases only)	SUMMANY:
None to CNS B - Recommended, unk if done	AT THIS HOSPITAL:
t Badiation therapy 9 Unknown/not applicable	0 No BRM 7 - Patien//guardian refused
7 - Patient/guardian refused	1 - BRM 8 - Recommended, unk if done
RADIATION/SURGERY SEQ:	2 - Allo BMT 9 - Unknown
0 - Not applicable 5 - Intreoperative radiation	3 - Auto BMT
2 - Radiation before surgery 6 - Intraoperative plus 2, 3 or 4	STARTED:(mm/dd/yyyy)
Radiation after surgery Sequence unknown Foliore & after surgery	1EXT :
TEXT: Chest Wall 6000	OTHER RX
	SUMMARY:
	AT THIS HOSPITAL: 0
FF 18 TFU2 - \$UB, THERAPY	(0) to other callification 6 - Unproven therapy
Stated Course Type Code Desc.	1 - Other ca-directed rx 7 - Patient/guardian refused
mit/dd/yyyy	2 - Experimontal carx 8 - Recommended, unk if done
	3 - Double-blind study 9 - Unknown
	STATIED: (minVdd/yyyy)
2] /] /	TEXT:
3 / /	
TO THE CONTRACT OF CALL	`

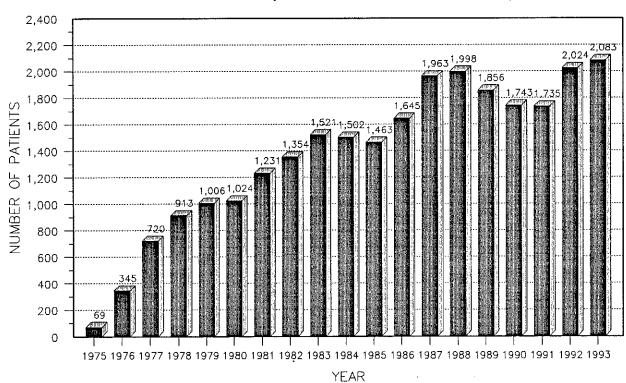
FIGURE 1-D

PF 17 TFU1 - FOLLOW-UP INFORMATION	PF 20 TREM - REMARKS/SPECIAL DATA ITEMS
LAST CONTACT/DEATH: (ppm/dx//yyy) [1] 0 / [1 7] / [1 9 8 9]	FREE FORMAT AREA:
CAUSE OF DEATH ICD CODE:	NEMARKS: Mother died of breast cancer.
, , , , , , , , , , , , , , , , , , , ,	
CURRENT VITAL STATUS (1) Alive 2 Dead 11	•
CURRENT CANCER STATUS:	The second secon
1 No synderice of cancer (2-) Cvidence of cancer 9 - Unknown	
CHIALITY OF SURVIVAL:	OVERRIDE FIELDS (Y - Bypass edit, leave black if edit not bypassed)
0 - NA doad 3 - Amb 50% 8 - NA doad	SITEARIST:
1 Syru∧) 4 Bodridrien (9 Inknown	AGE /SITE/IIST:
2 Amh 50%	
PAUFNA (letter or asterisk, eq. a. A. B. U)	SPECIAL FIELDS :
t 1	/ 1: Hepatitls
CONTACT (eg 0 first contact, 3 - flord contact) 1 .	#3: Burn Scar
Y - Yes, fixeign resident, leave blank for all others)	# 4 : Consanguinity
CONTACT PREQUENCY:	5: Predisposing Factors
(eg. 01 - One month, 03 - 3 months, 12 - Armial follow-up)	# 6: Pregnancy during dx/tx
UNUSUAL CONDITIONS 1 +	to the state of th
	# 8 : Immunodeficiency Disorder
PLACE OF DEATH (State of country - Geocode)	/9:
RECURRENCE INFOLIMATION	/ 10:
DATE (min/dd/yyyy) [0,9] /[1+0] /[1+9,8+9]	
TYPE: 9 Therecuriers,e 3 Distant recurrence	PF - 21 TADR - PATIENT NAME ADDRESS FILE
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2 - Выдиналиськоенно 9 - Unknown	SÁLUTATIÓN:
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2 OTHER PHYSICIAN: 0 7 1 6 7 8 Rad, Onc	COMMENT:
3 OHER PHYSICIAN: 10 9 2 1 8 5 Surgeon	PATIENT/GUARDIAN CODE: P - Patient G - Guardian 1 1
4 OTHER PHYSICIAN: 111411111	PF - 22 TCON - CONTACT NAME/ADDRESS FILE MAINTENANCE
S. OHIER PHYSICIAN:	CONTACT NUMBER: (0 - First contact, 1 - Second, 9 - Tenth) MAILING NAME:
6 OTHER PHYSICIAN: 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Riyadh Central Hospital
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HEXT HOSP FOR FID:	Riyadh
DEATH CERTIFICATE FILE INO: 1. 1. 1. 1. 1. 1.	gity:
	Riyadh
	PROV. RY) ZIP CODE:
	IELEPHONE: (111) [111] EXT. [11]
	COMMENT:
	REFER HOSP, MRN: 89856

II. KFSH&RC CANCER PATIENT POPULATION

A total of 2,083 cases were accessioned in 1993, with 1,048 males and 1,035 females or a male/female ratio of 1.01:1. This represents a 2.9% increase from 1992.

FIGURE 2
DISTRIBUTION OF ALL CASES ACCESSIONED BY YEAR
1975 - 1993 (TOTAL CASES = 26,195)



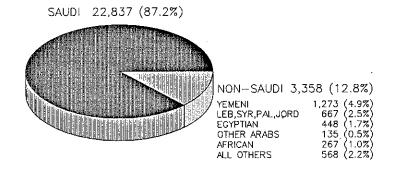
From the opening of the hospital (mid 1975) until December 1993, 26,195 cancer cases were registered (14,466 males and 11,729 females) with a male/female ratio of 1.2:1. There were 3,301 (12.6%) pediatric cases (0 to 14 years of age) and 22,894 (87.4%) adults (15 years old and above). Only a slight difference in the proportion was noted in 1993, 13.4% (279) for pediatrics and 86.6% (1,804) for adults.

TABLE 1 ALL CASES SEEN AT KFSH&RC (MALE/FEMALE & CHILDREN/ADULTS) BY 5-YEAR PERIOD 1975 - 1993

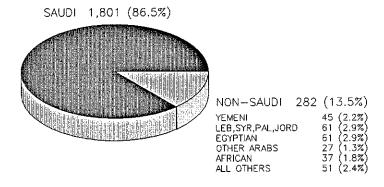
	1975-1976*	1977-1981	1982-1986	1987-1991	1992-1993	TOTAL
	No. %					
MALE	280	2,969	4,136	4,961	2,120	14,466
FEMALE	134	1,925	3,349	4,334	1,987	11,729
TOTAL	414	4,894	7,485	9,295	4,107	26,195
M/F RATIO	2.1:1	1.5:1	1.2:1	1.1:1	1.1:1	1.2:1
CHILDREN** ADULTS	55 13.3	587 12.0	985 13.2	1,157 12.4	517 12.6	3,301 12.6
	359 86.7	4,307 88.0	6,500 86.8	8,138 87.6	3,590 87.4	22,894 87.4
TOTAL	414 100	4,894 100	7,485 100	9,295 100	4,107 100	26,195 100

FIGURE 3 DISTRIBUTION OF ALL CASES BY NATIONALITY

1975 - 1993 (TOTAL CASES = 26,195)



1993 CASES (TOTAL = 2,083)



Saudi nationals totalled 1,801 (86.5%) in 1993 and the non-Saudi, 282 (13.5%). During the period 1975 to 1993, the former accounted for 87.2% (22,837) while the latter, 12.8% (3,358).

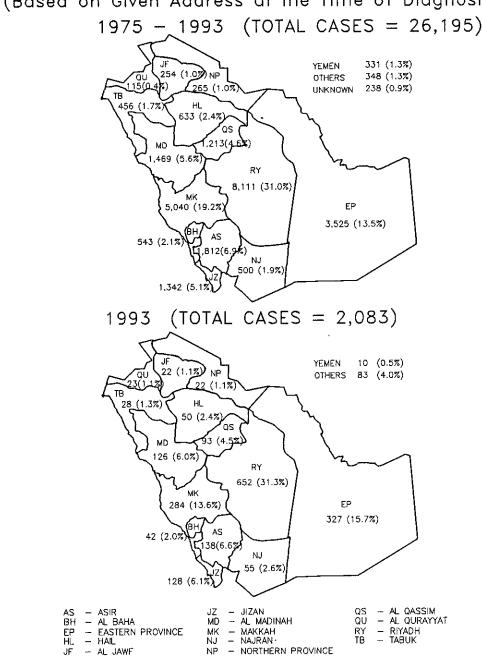
^{*} First two years of KFSH&RC partial operation.
** Children = 0 to 14 years of age; Adults = 15 years and above.

Geographically, the referral pattern is mainly from the Riyadh Region with 31.3% of all cases, followed by the Eastern Province and the Makkah Region with 15.7% and 13.6%, respectively, in 1993. The same regions had the most number of cases during the 19 years in review, i.e., 31.0% from Riyadh, 19.2% from Makkah and 13.5% 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)



TRENDS IN RELATIVE FREQUENCY OF CANCER AT KFSH&RC

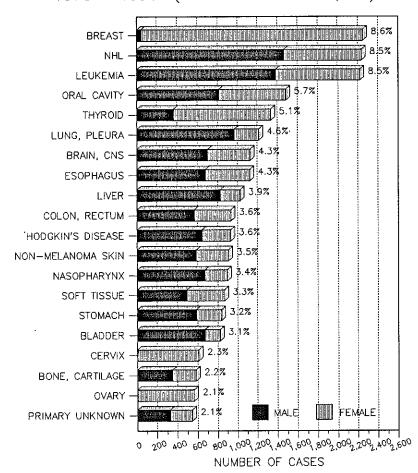
The crude relative frequency is the proportion of a given cancer in relation to all cases in a clinical or pathological series. Although such frequencies are subject to many biases, historically many elevated frequencies have been confirmed when complete cancer registration was introduced.

Biases that may have an affect 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 speciality 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 1993 with 8.6%, followed by Non-Hodgkin's Lymphoma (8.5%), Leukemia (8.5%), Oral Cavity (5.7%) and Thyroid (5.1%).

FIGURE 5
DISTRIBUTION OF 20 MOST COMMON MALIGNANCIES
1975 - 1993 (TOTAL CASES = 26,195)



Cancer among children under the age of 15 accounted for 12.6% of all cases from 1975 to 1993. The five most common childhood malignancies were Leukemia (26.2%), Lymphoma (21.3%)[NHL 13.0% and HD 8.3%], Brain/CNS (15.3%), Eye (8.4%) and Soft Tissue (7.8%).

FIGURE 6
DISTRIBUTION OF 10 MOST COMMON CHILDHOOD MALIGNANCIES
1975 - 1993 (TOTAL CASES = 3,301)

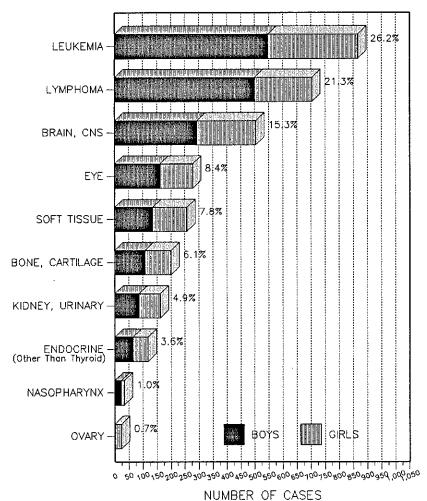


Table 2 shows the number of all malignant cases seen at KFSH&RC from 1975 to 1993 by site and year and Table 3, the 5-year summaries.

TABLE 2

ALL CASES SEEN AT KFSH&RC BY SITE* AND YEAR 1975 - 1993

TOTAL	1,485	968	124	837	933	027	302	562	374	214	263	056	070	123	20	580	998	146	910	220	430	610	563	319	257	822	541	410	128	335	148	595	640	930	545	180	!	26 105
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1991	101	6	99	35	8	99	12	<u>5</u>	34	83	54	22	7	12	0	37	83	٥	24	168	30	32	37	16	15	77	35	9	8	11	∞	7	25	26	40	Ţ		1736
1990	103	62	23	51	\$	54	7	2	56	2	13	26	2	•	-	36	Z	'n	67	168	34	7,	94	23	19	58	34	28	81	35	9	26	22	26	0,4	7	:	174.2
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SITE	Oral Cavity	Nasopharvox	Esophagus	Stonach	Colon. Rectum		Pancreas	Other G.f.	Larvnx	Lung, Pleura	Multiple Myeloma	Lymphoid Leukemia	Myeloid Leukemia	Other Leukemias	Reticuloendothelium	Bone, Cartilage	Soft Tissue	Skin Melanoma	Non-Melanoma Skin Ca	Breast	Uterus, Genital		Ovarv	Prostate	Testis, Genital	Bladder	Kidney, Urinary	Eve	Brain, CNS	Thyroid	Other Endocrine	NHL - Lymph Nodes	NHL - Extra-nodal		Primary Unknown	All Other Sites	און סוופן פונפפ	LATOR

* Includes Multiple Primary Neoplasms.

TABLE 3

ALL CASES SEEN AT KFSH&RC BY SITE* AND 5-YEAR PERIOD 1975 - 1993

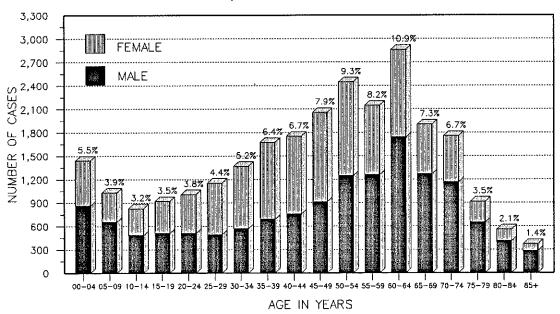
	107	1975-1976**	1977-1981	1081	1982.	1982-1986	1987	1987-1991	1992-1993	1993	TOTAL	AL.
<u>.</u>	O.N.	¥	2	ж	N _O	×	2	*	9	×	9	×
Oral Cavity	5	3.6%	301	6.2%	432	5.8%	528	5.7	509	5.1%	1,485	5.7%
Nasopharynx	14	3.4%	194	70.7	248	3,3%	328	3.5%	112	2.7%	968	3.4%
Esophagus	16	3.9%	306	6.3%	338	4.5%	349	3.8%	115	2.8%	1,124	4.3%
Stomach	17	4.1%	201	4.1%	585	3.8%	243	2.6%	8	2.2%	837	3.2%
Colon, Rectum	14	3.4%	159	3.2%	235	3.1%	357	3.8%	168	4.1%	933	3.6%
Liver	22	5.3%	203	4.1%	311	4.2%	335	3.6%	156	3.8%	1,027	3.9%
Pancreas	9	1.4%	99	1.3%	88	1.3%	88	0.9%	95	1.1%	302	1.2%
Other 6.1.	~	1.7%	26	1,1%	22	1.0%	104	1.1%	9	1.5%	586	1.1%
Larynx	9	1.4%	23	1.5%	26	1.3%	138	1.5%	58	1.4%	374	1.4%
Lung, Pleura	14	3.4%	195	4.0%	391	5.2%	777	4.8%	172	4.2%	1,214	79.4
Multiple Myeloma	١n	1.2%	41	0.8%	9	0.8%	111	1.2%	95	1,1%	263	1.0%
Lymphoid Leukemia	18	4.3%	175	3.6%	325	4.3%	375	7. 0%	163	70.7	1,056	4.0%
Myeloid Leukemia	16	3.9%	216	77.7	287	3.8%	370	70.7	151	3.7%	1,040	70.4
Other Leukemias	-	0.2%	53	29.0	35	0.5%	43	0.5%	13	24.0	123	0.5%
Reticuloendothelium	-	0.2%	4	0.1%	10	0.1%	'n	0.1%	0	0.0%	20	0.1%
Bone, Cartilage	7	1.7%	66	2.0%	173	2.3%	199	2.1%	102	2.5%	580	2.2%
Soft Tissue	15	3.6%	169	3.5%	230	3.1%	335	3.6%	117	2.8%	866	3.3%
Skin Melanoma	7	1.0%	33	0.7%	77	0.6%	43	0.5%	22	0.5%	146	0.6%
Non-Melanoma Skin Ca	17	4. 1%	197	70.4	313	4.2%	267	2.9%	116	2.8%	910	3.5%
Breast	27	6.5%	320	6.5%	627	8.4%	841	80.6	435	10.6%	2,250	8,6%
Uterus, Genital	2	0.5%	62	1.3%	119	1.6%	170	1.8%	11	1.9%	430	1.6%
Cervix	10	2.4%	100	2.0%	185	2.5%	213	2.3%	102	2.5%	610	2.3%
Ovary	80	1.9%	28	1.6%	153	2.0%	227	2.4%	26	2.4%	293	2.1%
Prostate	7	1.7%	36	0.7%	86	1.3%	115	1.2%	63	1.5%	319	1.2%
Testis, Genital	4	1.0%	26	1.1%	8	0.9%	83	26.0	87	1.2%	257	1.0%
Bladder	11	2.7%	136	2.8%	197	2.6%	328	3.5%	150	۲. ۲.	822	3.1%
Kidney, Urinary	٥	2.2%	89	1.8%	145	1.9%	194	2.1%	104	2.5%	541	2.1%
Eye	9	1.4%	89	1.8%	128	1.7%	134	1.4%	53	1.3%	410	1.6%
Brain, CNS	27	6.5%	152	3.1%	307	4.1%	443	4.8%	199	78.7	1,128	4.3%
Thyroid	10	2.4%	179	3.7%	330	77.7	244	5.9%	272	29.9	1,335	5.1%
Other Endocrine	7	0.5%	22	25.0	29	0.8%	41	0.4%	54	29.0	148	
NHL - Lymph Nodes	23	2.6%	777	80.6	515	%6.9	677	4.8%	166	70.7	1,595	6.1%
NHL - Extra-nodal	ĸ	<u>ک</u>	48	1.0%	165	2.2%	262	3.1%	132	3.2%	640	2.4%
Hodgkin's Disease	32	7.7%	203	4.1%	539	3.2%	308	3.3%	148	3.6%	930	3.6%
Primary Unknown	14	3.4%	119	2.4%	127	1.7%	190	2.0%	35	2.2%	245	2.1%
All Other Sites	4	1.0%	5 ,5	0.9%	07	0.5%	65	22.0	27	o.72	180	۳. 0
TOTAL	414	100.0%	768'7	100.0%	7,485	100.0%	9,295	100.0%	4,107	100.0%	26, 195	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 1993, the mean age was 52.4, the median is 50.3 and the mode is 60.0. Childhood malignancies are most common among children three years of age.

FIGURE 7
DISTRIBUTION OF ALL CASES BY AGE AT DIAGNOSIS
1975 - 1993 (TOTAL CASES = 26,195)



1993 (TOTAL CASES = 2,083)

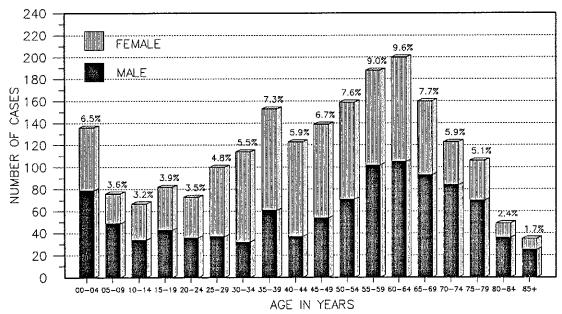
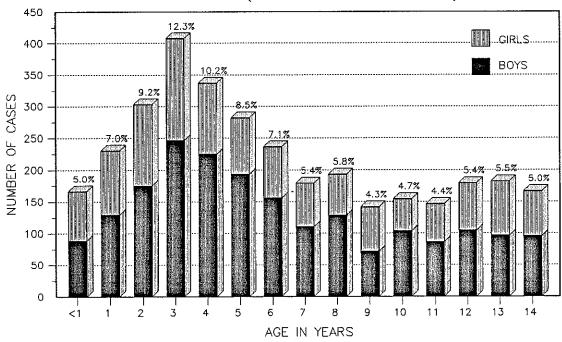


FIGURE 8

DISTRIBUTION OF ALL PEDIATRIC CASES BY AGE AT DIAGNOSIS

1975 - 1993 (TOTAL CASES = 3,301)



Of the 2,083 cases in 1993, 1,570 (75.4%) 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 513 cases (24.6%) were non-analytic (defined as cases diagnosed elsewhere and receiving all of their first course of treatment elsewhere). Out of the 1,570 analytic cases, pediatric cases totalled 211, with 123 boys and 88 girls.

See Table 4 for the distribution of cases by site, sex, class of case, and stage at diagnosis and Tables 5, 6 and 7 for the distributions of analytic cases by site, sex and age at diagnosis.

TABLE 4

ALL CASES SEEN AT KFSH&RC BY SITE*, SEX, CLASS OF CASE AND SUMMARY STAGE 1993

* Includes Multiple Primary Neoplasms.

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 5

ANALYTIC CASES SEEN AT KFSH&RC BY SITE* AND AGE 1993

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Oral Cavity Masopharynx Esophagus Stomach Colon, Rectum Liver Pancreas Other G.I. Larynx Lung, Pleura Myeloid Leukemia Hyeloid Leukemia Other Sisase H.D Extra-nodal Hodgkin's Disease H.D Extra-nodal Primary Unknown All Other Sites	7-0	000000000000000000000000000000000000000	ያ
	SITE	Oral Cavity Nasopharynx Esophagus Stomach Colon, Rectum Liver Pancreas Other G.I. Larynx Lung, Pleura Multiple Myeloma Lymphoid Leukemia Myeloid Leukemia Myeloid Leukemia Other Sites Other Sites	TOTAL

* Includes Multiple Primary Neoplasms.

TABLE 6

ANALYTIC "MALE" CASES SEEN AT KFSH&RC BY SITE* AND AGE 1993

SITE	7-0	6-5	10-	15.	20-	\$ \$	ጵጵ	35-	-07	45-	2, 20	55-	\$ 3	69	2	ŔR	8 %	85+	TOTAL
Oral Cavity Nasopharynx Esophagus Stomach Colon, Rectum Liver Pancreas Other G.I. Larynx Lung, Pleura Multiple Myeloma Lymphoid Leukemia Other Leukemia Other Cartilage Soft Tissue Skin Melanoma Non-Melanoma Non-Melanoma Ron-Melanoma Non-Melanoma Non-Me	000000000000000000000000000000000000000	00000000000000000000000000000000000000	, , , , , , , , , , , , , , , , , , ,	0 W 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0-00-000000000000000000000000000000000	0+004000000+-00000000000000000000000000	04-0000000000000004-00-	MW0-4004>-00-2000000000000000000000000	MUOO-0000M000000000000000000000000000000						404-044-045M0-0	00000000000000000000000000000000000000	, иоомомоом-ооооомоооооии-о	NONUT-000++00+00000000000000000000000000000	25 1 1 0 0 0 0 1 3 4 5 5 6 5 6 5 6 5 6 5 6 5 6 5 6 5 6 5 6
Thyroid Other Endocrine NHL - Lymph Nodes NHL - Extra-nodal Hodgkin's Disease H.D Extra-nodal Primary Unknown All Other Sites	00000000	00-00000	00	-000-00-	40-4400 m	-0048-00	-0800-	WO4000	-00M000- %	00072000		-0%00000 (-04-0040	-ommooro (-004000- [-0-4400- H	00-000-0 K	000-00-0	7 9 23 33 54 75 6 75 6 75 6 75 6 75 6 75 6 75 6 7
TOTAL	29	36	88	33	27	82	55	3	92	38	52	69	2	29	<u></u>	አ	Q	<u>^</u>	000

* Includes Multiple Primary Neoplasms.

TABLE 7

ANALYTIC "FEMALE" CASES SEEN AT KFSH&RC BY SITE* AND AGE 1993

TOTAL

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* 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 soft tissue sarcoma, among others, is more common. The following 1993 analytic cases exhibit significant differences in trends from those of the West:

Breast - The most common malignancy seen at KFSH&RC is breast cancer, comprising 13.0% of all cases, as compared to about 16% 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 8.5% 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 1.7:1.It is the most common type of malignancy seen in males and the third most common in females. It is also the most common malignancy in children under the age of 15.

Non-Hodgkin's Lymphoma - The most striking feature is the unusually high crude relative frequency of non-Hodgkin's lymphoma, accounting for 7.8% of all cases. The male/female ratio is 1.6:1. In the U.S.A., NHL accounts for only about 4% of all cancer.

Thyroid - 2.2% of all male malignancies in KFSH&RC are thyroid tumors. However, they represent 11.1% of female malignant neoplasms, second to breast cancer. The male/female ratio is 0.2:1. Thyroid cancer accounts for only 1.1% of all cases in the U.S.A. and 1.6% of female malignancies.

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 5.2% of the cases. The male/female ratio is 0.7:1.

Brain/CNS - Primary malignant neoplasm of the brain and CNS accounts for 5.0% of all malignancies and ranks second among the most common childhood malignancies. The male/female ratio is 1.2:1. This is comparatively higher than in the West with only 1.5% of all cases.

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 about 15% of all cancer cases (17% in males, and 12% in females).

At KFSH&RC, 4.4% of the diagnoses are lung cancer, although in males it is the third most common tumor, constituting 7.8% of male malignancies and 1.2% in females. The male/female ratio is 5.9:1.

Colo-Rectal - Markedly less common than in the West, for which dietary factors (particularly lower animal fat intake) may play a role, this disease represents only 3.3% of all tumors. In the U.S.A. it constitutes 13% of newly diagnosed cancer cases. The male/female ratio at KFSH&RC is 1.0:1.

Esophagus - The incidence of esophageal carcinoma is comparatively more frequent at KFSH&RC than in Western countries. In the U.S.A. it constitutes 1% of all cancers, compared to 2.7% at KFSH&RC. The male/female ratio is 1.3:1.

Liver - Although the relative frequency of liver cancer at the KFSH&RC (3.3%) is almost the same as that of the West, the male/female ratio appears to be significant and may be an area for future research investigations. KFSH&RC has 5.5:1 and the West, 1.2:1.

Nasopharynx - A higher crude relative frequency rate is seen in nasopharyngeal cancer. It constitutes less than 1% of the pathologically diagnosed cancers in most centers in the West, but is 3.4% of the cases at KFSH&RC. The male/female ratio is 3.5:1.

Soft Tissue - KFSH&RC cases show a higher rate of soft tissue malignancies than the U.S.A., with 2.2% against the latter's 0.5% of all cases. The male/female ratio is 1.1:1.

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 14% of the malignancies). This is in contrast to the KFSH&RC experience, where prostatic cancer makes up only 0.7% 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 9 DISTRIBUTION OF 20 MOST COMMON MALIGNANCIES 1993 ANALYTIC CASES (TOTAL CASES = 1,570)

MALE FEMALE LEUKEMIA 84 (11.1%) BREAST 204 (25.1%) NHL 75 (9.9%) THYROID 90 (11.1%) LUNG, PLEURA 59 (7.8%) LEUKEMIA 49 (6.0%) BLADDER 46 (6.1%) NHL 48 (5.9%) LIVER 44 (5.8%) ORAL CAVITY 47 (5.8%) NASOPHARYNX 42 (5.6%) CERVIX 44 (5.4%) OVARY 36 (4.4%) BRAIN, CNS 42 (5.6%) HODGKIN'S DISEASE 36 (4.8%) BRAIN, CNS 36 (4.4%) ORAL CAVITY 35 (4.6%) UTERUS, GENITAL 30 (3.7%) COLON, RECTUM 26 (3.4%) COLON, RECTUM 26 (3.2%) LARYNX 26 (3.4%) BONE, CARTILAGE 21 (2.6%) BONE, CARTILAGE 25 (3.3%) KIDNEY, URINARY 21 (2.6%) ESOPHAGUS 24 (3.2%) HODGKIN'S DISEASE 21 (2.6%) NON-MELANOMA SKIN 23 (3.0%) ESOPHAGUS 18 (2.2%) TESTIS, GENITAL 21 (2.8%) SOFT TISSUE 17 (2.1%) STOMACH 19 (2.5%) PRIMARY UNKNOWN 16 (2.0%) KIDNEY, URINARY 19 (2.5%) BLADDER 13 (1.6%) SOFT TISSUE 18 (2.4%) NASOPHARYNX 12 (1.5%) OTHER G.I. 11 (1.4%) THYROID 17 (2.2%) PRIMARY UNKNOWN 12 (1.6%) LUNG, PLEURA 10 (1.2%)

FIGURE 10

DISTRIBUTION OF CHILDHOOD MALIGNANCIES 1993 ANALYTIC CASES (TOTAL CASES = 211)

MALE LEUKEMIA 38 (30.9%) LYMPHOMA 26 (21.1%) BRAIN, CNS 21 (17.1%) EYE 9 (7.3%) BONE, CARTILAGE 7 (5.7%) SOFT TISSUE 7 (5.7%) ENDOCRINE 6 (4.9%) (Other Than Thyroid) NASOPHARYNX 2 (1.6%) NON-MELANOMA SKIN 2 (1.6%)

KIDNEY 2 (1.6%)

TESTIS, GENITAL 1 (0.8%)

MEDIASTINUM 1 (0.8%) PRIMARY UNKNOWN 1 (0.8%)

FEMALE

LEUKEMIA 26 (29.5%) BRAIN, CNS 16 (18.2%) BONE, CARTILAGE 11 (12.5%) LYMPHOMA 11 (12.5%) SOFT TISSUE 8 (9.1%) KIDNEY 5 (5.7%) EYE 4 (4.5%) OVARY 3 (3.4%) ENDOCRINE 2 (2.3%) (Other Than Thyroid)

OTHER G.I. 1 (1.1%)

NON-MELANOMA SKIN 1 (1.1%)

TABLE 8

PRIMARY SITE TABLE
(INCLUDES MULTIPLE PRIMARIES)
1 9 9 3

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
(NOS - Not Otherwise Specified)	2,083	1,048	1,035
LIP	4	1	3
Squamous Cell Carinoma			
TONGUE	27	14 12	13 13
Squamous Cell Carinoma Verrucous Carcinoma	25 2	2	0
MAJOR SALIVARY GLANDS	7	3	4
Mucoepidermoid Carcinoma	· 3	1	2
Carcinoma, NOS	2	2	0
Adenoid Cystic Carcinoma Acinar Cell Carcinoma	1 1	0 0	1
GUM	17	9	8
Squamous Cell Carcinoma	14	6	8
Non-Hodgkin's Lymphoma	2	2	0
Verrucous Carcinoma	1	1	0
OTHER PARTS OF MOUTH	17	8	9
Squamous Cell Carcinoma	12	5 0	7 2
Adenoid Cystic Carcinoma Mucoepidermoid Carcinoma	2 1	1	Õ
Non-Hodgkin's Lymphoma	ī	1	0
Malignant Neoplasm	1	1	0
OROPHARYNX	9	4	5
Non-Hodgkin's Lymphoma	, 6	4	2
Squamous Cell Carcinoma	2	0	2 1
Carcinoma, NOS	_	_	_
NASOPHARYNX	61 35	48 28	13 7
Squamous Cell Carcinoma Undifferentiated Carcinoma	13	11	2
Carcinoma, NOS	10	8	2
Lymphoepithelial Carcinoma	1	0	1
Non-Hodgkin's Lymphoma	1	1 0	0 1
Hodgkin's Disease		_	
HYPOPHARYNX Squamous Cell Carcinoma	25	10	15
PHARYNX, NOS Squamous Cell Carcinoma	1	1	0
ESOPHAGUS	47	25	22
Squamous Cell Carcinoma	42	22	20
Adenocarcinoma, NOS	2	2	0
Carcinoma, NOS	2 1	0 1	2 0
Malignant Neoplasm	1	+	U

Primary Site Table con't

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
STOMACH Adenocarcinoma, NOS Non-Hodgkin's Lymphoma Signet Ring Cell Carcinoma Squamous Cell Carcinoma Carcinoma, NOS Adenosquamous Carcinoma Malignant Neoplasm	57 32 15 3 2 1	35 23 6 2 2 1 0	22 9 9 1 1 1 0
SMALL INTESTINE Non-Hodgkin's Lymphoma Adenocarcinoma, NOS Neuroendocrine Carcinoma Carcinoid Tumor	15 11 2 1	9 6 1 1	6 5 1 0
COLON Adenocarcinoma, NOS Mucinous Adenocarcinoma Non-Hodgkin's Lymphoma Adenocarcinoma in Adenomatous Polyp Hodgkin's Disease Carcinoma, NOS Malignant Neoplasm	30 13 6 6 2 1 1	18 6 6 4 0 0 1	12 7 0 2 2 1 0
RECTUM/RECTOSIGMOID JUNCTION/ANUS Adenocarcinoma, NOS Mucinous Adenocarcinoma Squamous Cell Carcinoma Adenocarcinoma in Adenomatous Polyp Adenocarcinoma in Villous Adenoma Mucin-Producing Adenocarcinoma Signet Ring Cell Carcinoma Small Cell Carcinoma Basaloid Carcinoma Carcinoma, NOS Malignant Neoplasm	57 35 8 5 2 1 1 1 1 1	31 20 5 3 0 0 0 1 0	26 15 3 2 2 1 1 1 0 1
LIVER/INTRAHEPATIC BILE DUCTS Hepatocellular Carcinoma Cholangiocarcinoma Malignant Neoplasm Combined Hepatocellular & Cholangioca Carcinoma, NOS	81 74 3 2 1 1	68 63 2 2 0 1	13 11 1 0 1
GALLBLADDER/EXTRAHEPATIC BILE DUCTS Adenocarcinoma, NOS Carcinoma, NOS	21 20 1	6 6 0	15 14 1
PANCREAS Adenocarcinoma, NOS Malignant Neoplasm Non-Hodgkin's Lymphoma Carcinoma, NOS	22 15 4 2	16 11 3 2 0	6 4 1 0
RETROPERITONEUM/PERITONEUM Endodermal Sinus Tumor Non-Hodgkin's Lymphoma	2 1 1	o 0 0	2 1 1

Primary Site Table con't

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
OTHER G.I. SITES Adenocarcinoma Carcinoid Tumor Non-Hodgkin's Lymphoma Signet Ring Cell Carcinoma Malignant Neoplasm	8 2 2 2 1 1	5 2 1 0 1	3 0 1 2 0
NASAL CAVITIES/ACCESSORY SINUSES Squamous Cell Carcinoma Non-Hodgkin's Lymphoma Carcinoma, NOS Adenoid Cystic Carcinoma Solitary Plasmacytoma Adenocarcinoma, NOS	11 4 2 2 1 1	7 2 1 1 1 1	4 2 1 1 0 0
LARYNX Squamous Cell Carcinoma Carcinoma, NOS Verrucous Carcinoma	31 28 2 1	29 27 1 1	2 1 1 0
BRONCHUS/LUNG Adenocarcinoma Squamous Cell Carcinoma Carcinoma, NOS Small Cell Carcinoma Undifferentiated Carcinoma Carcinoid Tumor Bronchio-Alveolar Adenocarcinoma Large Cell Carcinoma Giant Cell Carcinoma Malignant Neoplasm Mixed Small & Large Cell Carcinoma Solid Large Cell Carcinoma Adenosquamous Carcinoma Signet Ring Cell Carcinoma	85 32 22 6 4 4 4 3 2 2 2 2 1 1 1	75 26 21 6 4 3 2 2 2 2 1 1 1	10 6 1 0 0 1 2 0 0 0 0 0
PLEURA Mesothelioma Adenocarcinoma	4 3 1	2 1 1	2 2 0
THYMUS/MEDIASTINUM Malignant Thymoma Neuroblastoma Hodgkin's Disease Malignant Neoplasm Malignant Neurilemmoma Non-Hodgkin's Lymphoma Endodermal Sinus Tumor Squamous Cell Carcinoma Ganglioneuroblastoma Ewing's Sarcoma	14 2 2 2 2 2 1 1 1 1 1	10 1 2 1 1 1 0 1 1 1	4 1 0 1 1 0 1 0 0 0 0
MULTIPLE MYELOMA	22	15	7

Primary Site Table con't

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
BONE MARROW	186	121	65
Acute Lymphoid Leukemia	76	47	29
Acute Myeloid Leukemia	38	23	15
Chronic Myeloid Leukemia	38	23	15
Chronic Lymphoid Leukemia	10	10	Ō
Acute Myelomonocytic Leukemia	8	6	2
Acute Promyelocytic Leukemia	6	4	2
Acute Leukemia, NOS	3	3 2	0
Acute Monocytic Leukemia	3 1	0	1 1
Chronic Myelomonocytic Leukemia	1	1	Ō
Megakaryocytic Leukemia Erythroleukemia	1	i	ŏ
Leukemia, NOS	ī	ī	Ö
	_	20	22
BONE & CARTILAGE	57 24	30 11	27 13
Osteosarcoma, NOS	15	8	7
Ewing's Sarcoma Chondrosarcoma, NOS	7	5	2
Non-Hodgkin's Lymphoma	5	ĭ	4
Chondroblastic Osteosarcoma	2	2	Õ
Juxtacortical Osteosarcoma	<u>1</u>	ī	0
Solitary Plasmacytoma	1	1	0
Sarcoma, NOS	1	1	0
Malignant Neoplasm	1	0	1
CONNECTIVE/SUBCUTANEOUS/SOFT TISSUE	53	27	26
Neuroblastoma	7	3	4
Leiomyosarcoma	6	3	3
Embryonal Rhabdomyosarcoma	4	2	2
Malignant Fibrous Histiocytoma	4	3	1
Rhabdomyosarcoma, NOS	3	2	1
Synovial Sarcoma	· 3	2	1
Sarcoma, NOS	3	2	1
Malignant Neoplasm	3	1 1	2 1
Spindle Cell Sarcoma Alveolar Rhabdomyosarcoma	2 2 2	ō	2
Fibrosarcoma, NOS	2	1	1
Peripheral Neuroectodermal Tumor	2	ī	ī
Chordoma	2	Ž	ō
Non-Hodgkin's Lymphoma	2	2	0
Myxoid Liposarcoma	1	0	1
Liposarcoma	1	0	1
Clear Cell Sarcoma	1	0	1
Hemangiosarcoma	1	0	1
Malignant Hemangiopericytoma	1	0	1
Malignant Neurilemmoma	1	0	1
Malignant Triton Tumor	1	1 1	0
Ewing's Sarcoma	1	_	_
SKIN (MELANOMA)	8	3	5

Primary Site Table con't

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
SKIN (NON-MELANOMA)	55	37	18
Squamous Cell Carcinoma	25	19	6
Basal Cell Carcinoma	15	10	5
Basosquamous Carcinoma	7	4	3
Mycosis Fungoides	2	1	1
Kaposi's Sarcoma	1	1	0
Dermatofibrosarcoma Protuberans	1	1	0
Sweat Gland Adenocarcinoma	1	0	1
Sebaceous Adenocarcinoma	1 1	0	1 1
Adenoid Cystic Adenocarcinoma	1	1	Ô
Adenocarcinoma, NOS	_	_	_
BREAST, FEMALE	248	0	248
Duct Cell Carcinoma	204	0	204
Carcinoma, NOS	11	0	11
Paget's Disease & Duct Cell Carcinoma	5	0	5
Adenocarcinoma, NOS	5	0	5
Medullary Carcinoma	4	0 0	4 4
Intraductal Carcinoma	4 4	0	4
Lobular Carcinoma Comedocarcinoma	3	0	3
	3	0	3
Malignant Neoplasm Cystosarcoma Phyllodes	1	ŏ	1
Tubular Adenocarcinoma	1	Ö	ī
Inflammatory Carcinoma	ī	ō	ī
(Mucinous) Colloid Carcinoma	ī	ō	ī
Non-Hodgkin's Lymphoma	ī	Ō	1
	1	1	0
BREAST, MALE Paget's Disease & Duct Cell Carcinoma	ī	1	ŭ
CERVIX UTERI	50	0	50
Squamous Cell Carcinoma	34	0	34
Adenocarcinoma, NOS	· 8	0	8
Adenosquamous Carcinoma	2	0	2
Carcinoma, NOS	2	0	2
Mucin-Producing Adenocarcinoma	1	0	1
Papillary Adenocarcinoma	1	0	1
Clear Cell Adenocarcinoma	1	0	1
Undifferentiated Small Cell Carcinoma	1	0	1
PLACENTA	12	0	12
Choriocarcinoma	10	0	10
Trophoblastic Tumor	2	0	2
CORPUS UTERI	20	0	20
Adenocarcinoma, NOS	25 15	Ö	15
Endometrial Stromal Sarcoma	2	ŏ	2
Serous Papillary Carcinoma	1	ŏ	1
Adenosarcoma	i	ŏ	ī
Carcinoma, NOS	ī	Õ	ī

Primary Site Table con't

SITE HISTOLOGY	ALL CASES	MALES	FEMALES
OVARY	52	o	52
Adenocarcinoma, NOS	9	0	9
Dysgerminoma	7	Ō	7
Mucinous Cystadenocarcinoma	6	0	6
Carcinoma, NOS	4	0	4
Serous Cystadenocarcinoma	3	0	3 3
Papillary Adenocarcinoma	3	0	3
Papillary Serous Cystadenocarcinoma	3	0	3
Endodermal Sinus Tumor	3	0	3
Mucinous Adenocarcinoma	2	0	2 2
Malignant Teratoma	2	0	2
Papillary Serous, Borderline Malignancy	2 2	0	2
Serous Tumor, Bordeline Malignancy	1	Ö	1
Serous Papillary Carcinoma Endometrioid Carcinoma	1	Ö	ī
Papillary Cystadenocarcinoma	i	ő	i
Cystadenocarcinoma, NOS	1	ŏ	ī
Choriocarcinoma, NOS	i	Ö	ī
Squamous Cell Carcinoma	ī	ō	ī
OTHER FEMALE GENITAL ORGANS	4	0	4
Squamous Cell Carcinoma	3	0	3
Melanoma	1	0	1
PROSTATE	26	26	0
Adenocarcinoma, NOS	23	23	0
Mucinous Adenocarcinoma	1	1	0
Carcinoma, NOS	2	2	0
TESTIS	27	27	0
Seminoma, NOS	12	12	0
Mixed Germ Cell Tumor	11	11	0
Endodermal Sinus Tumor	2	2	0
Non-Hodgkin's Lymphoma	2	2	0
OTHER MALE GENITAL ORGANS	2	2	0
Kaposi's Sarcoma	1	1	0
Squamous Cell Carcinoma	1	1	0
URINARY BLADDER	87	69	18
Papillary Transitional Carcinoma	36	27	9
Transitional Cell Carcinoma	33	28	5
Squamous Cell Carcinoma	13	10	3
Carcinoma, NOS	3	2	1
Solid Carcinoma	1	1	0
Malignant Neoplasm	1	1	0
KIDNEY/URETER	52	27	25
Renal Cell Carcinoma	36	22	14
Nephroblastoma	8	2	6
Transitional Cell Carcinoma	2	2	0
Carcinoma, NOS	2	0	2
Renal Cell Carcinoma, Granular Cell	ī	0	2 1
	ī	0	1
Adenocarcinoma, NOS Non-Hodgkin's Lymphoma		0 1	1 0

Primary Site Table con't

SITE	HISTOLOGY	ALL CASES	MALES	FEMALES
EYE			39	2217
Retinoblast	oma	33	18	15
Squamous Ce	ell Carcinoma	3	2	1
Melanoma		2	1	1
Sarcoma		1	1	0
BRAIN		85	51	34
Glioblaston	·· ·	26	14	12
Astrocytoma		19	11 9	8 4
Medulloblas	·	13 6	4	2
Malignant G		5	4	1
Malignant N	n's Lymphoma	4	4	ô
Ependymoma	(eopius	3	i	2
	oglioma, NOS	3		2
	c Medulloblastoma	1	1	0
	Neuroectodermal Tumor	1	0	1
Anaplastic	Astrocytoma	1	1	0
	Astrocytoma	1	0	1
	Sinus Tumor	1	o o	1
Germinoma		1	1	0
OTHER NERVOUS	SYSTEM	9	6	3
Astrocytoma	a, NOS	4	2	2
Non-Hodgkir	n's Lymphoma	2	2	0
Malignant (Glioma	1	0	1
Ependymoma		1	1	0 0
Ewing's Sar	coma	1	1	U
THYROID		136	29	107
Papillary (Carcinoma, NOS	96	20	76
	Follicular Adenocarcinoma	22	0	22
	n's Lymphoma	, 5	4	1 3
Medullary (5 3	2 0	3
· · ·	Adenocarcinoma Adenocarcinoma	2	ĭ	1
	Anaplastic Type	1	ō	ī
	Adenocarcinoma	ī	ĭ	ō
Carcinoma,		ī	ī	Ö
		10	7	3
OTHER ENDOCRING		8	6	2
Germinoma	Sille.	1	ĭ	Õ
Malignant I	Neoplasm	ī	ō	1
-	-	78	46	32
	ON-HODGKIN'S LYMPHOMA Extra-Nodal Lymphomas)	70	40	32
Large Cell		33	18	15
	Lymphoma, NOS	8	3	5
Small Cell		7	6	1
	tic Lymphoma	6	4	2
Lymphoblast	tic Lymphoma	5	3	2
Non-Hodgki	n's Lymphoma, NOS	4	3	1
Mixed Smal	l Cleaved & Large Cell, Foll:		2	1
	l & Large Cell, Diffuse	3	2 2	1 0
Burkitt's 1		2	*7	7.1

Primary Site Table con't

SITE	HISTOLOGY	ALL CASES	MALES	FEMALES
	NON-HODGKIN'S LYMPHOMA	(Cont'd)		
	ng Extra-Nodal Lymphomas)		^	0
	mphocytic Lymphoma	2	0	0
	Lymphoma	2	0	2
Lympnocy	tic Lymphoma	± 1	0	1 1
	ithelioid Lymphoma	1	1	Ó
T-Cell L	ymphoma	-	-	Ū
LYMPH NODES,	HODGKIN'S DISEASE	68	46	22
Nodular	Sclerosis	36	24	12
	llularity	15	8	7
	s Disease, NOS	13	11	2 1
Lymphocy	tic Predominance	4	3	1
PRIMARY UNKN	IOWN	42	21	21
	cinoma, NOS	16	5	11
Carcinom		11	5	6
	Cell Carcinoma	5	1	4
	t Neoplasm	3	3	0
	locrine Carcinoma	1	1	0
Undiffer	entiated Carcinoma	1	1	0
Malignan	t Teratoma	1	1	0
Transiti	onal Cell Carcinoma	1	1	0
Melanoma	L Company	1	1	0
Mucinous	Adenocarcinoma	1	1	0
Signet F	ling Cell Carcinoma	1	1	0

TABLE 9

PATIENTS WITH MULTIPLE PRIMARIES
1 9 9 3

PRIMARY SITE HISTOLOGY 1993	OTHER PRIMARIES (PREVIOUS OR CONCURRENT)	ALL CASES	MALES	FEMALES
		50	24	26
ORAL CAVITY		2	1	1
Lymphoepithelial Ca-NP Sq Cell Ca-Hypophapharynx	Thyroid Tongue	1 1	0 1	1 0
ESOPHAGUS		2	2	0
Adenocarcinoma Malig Neoplasm	Chr Lymphocytic Leukemia Tonque	1	1 1	0 0
STOMACH	-	1	0	1
Non-Hodgkin's Lymphoma	Thyroid			
SMALL INTESTINE Non-Hodgkin's Lymphoma*	Descending Colon Cecum	1	0	1
COLON		1	1	0
Mucinous Adenoca-Sigmoid	Descending Colon			
RECTUM Adenocarcinoma	Thyroid	1	1	0
LIVER		4	2	2
Hepatocellular Ca Hepatocellular Ca	Tongue Bucal Mucosa	1 1	0 1	1 0
Hepatocellular Ca	Skin	ī	Õ	1
Hepatocellular Ca	Thyroid	1	1	0
GALLBLADDER Adenocarcinoma	Stomach - NHL	2 1	0 0	2 1
Carcinoma In Situ	Ovary	ī	ŏ	1
PANCREAS		2	2	0
Adenocarcinoma Adenocarcinoma	Vocal Cord Liver - HCC	1 1	1 1	0 0
LUNG	22,02	1	1	0
Carcinoid Tumor	Thyroid	-	-	•
BONE MARROW		2	1	1 .
Acute Myeloid Leukemia Acute Myeloid Leukemia	Non-Hodgkin's Lymphoma Non-Hodgkin's Lymphoma	1 1	0 1	1 0
BONE	Non-nougkin a Dymphoma	1	1	0
Ewing's Sarcoma	Non-Hodgkin's Lymphoma	•	•	•
CONNECTIVE TISSUE		1	1	0
Leiomyosarcoma	Rectum			_
SKIN Pagal Coll Carginoma	Nagonharuny	6 1	4 0	2 1
Basal Cell Carcinoma Basal Cell Carcinoma	Nasopharynx ST - Malig Histiocytoma	i	0	1
Basal Cell Carcinoma	Skin - Sq Cell Ca	1	1	0
Basal Cell Carcinoma	Skin - Sq Cell Ca	1	1	0
Melanoma	Skin - Basal Cell Ca Tonque	1 1	1 1	0 0
Basosquamous Carcinoma	toudae		1	U

Multiple Primaries con't

PRIMARY SITE HISTOLOGY	OTHER PRIMARIES (PREVIOUS OR CONCURRENT)	ALL CASES	MALES	FEMALES
BREAST		9	0	9
Duct Cell Carcinoma	Contra Breast	1	Ō	1
Duct Cell Carcinoma	Contra Breast	1	Ō	1
Duct Cell Carcinoma	Contra Breast	1	0	1
Duct Cell Carcinoma	Hodgkin's Disease	1	0	1
Duct Cell Carcinoma	Non-Hodgkin's Lymphoma	ī	0	1
Inflammatory Carcinoma	Contra Breast-Duct Cell	Ca 1	0	1
Paget's Disease & Duct Ca	•••••	1	Ó	1
Tubular Adenocarcinoma	Lobular Ca In Situ	ī	Õ	ī
Carcinoma, NOS	Thyroid	ĩ	ō	1
Carcinoma, Nos	Injioid		-	_
CERVIX		1	0	1
Squamous Cell Carcinoma	Breasț			
PROSTATE		2	2	0
Adenocarcinoma	Thyroid	<u></u>	ī	0
Adenocarcinoma	Thyroid	ī	ī	Ō
#F 1 DDDD	-	1	1	0
BLADDER Transitional Cell Ca	Kidney	•	-	J
	2	2	•	
KIDNEY	_	2	1	1
Renal Cell Carcinoma	Larynx	1	1	0
Renal Cell Carcinoma	ST - Liposarcoma	1	0	1
BRAIN & CNS		5	2	3
Glioblastoma	Acute Lymphoid Leukemia	1	0	1
Glioblastoma	Non-Hodgkin's Lymphoma	1	Ō	1
Glioblastoma	Bone	ī	ŏ	1
Malignant Neoplasm	Thyroid	ĩ	ī	ō
Malignant Neoplasm	Non-Hodgkin's Lymphoma	ī	ī	ŏ
Marrane Mechiasm	Hon Hondyrin a nimbrious	_		
THYROID		2	1	1
Papillary Carcinoma	Lip ,	1	1	0
Papillary Carcinoma	Colon	1	0	1
LYMPH NODES		1	0	1
Non-Hodgkin's Lymphoma	Hodgkin's Disease	ī	ŏ	ī
MOH-HORDER B TAMBHOME	"ordur" o proceso	-	•	-

^{*} 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

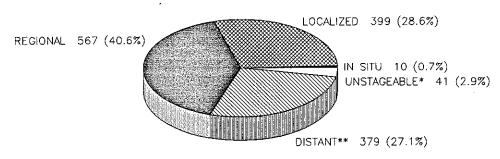
b. Discontinuous metastasis

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 1993. The 41 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 4, page 17, for the distribution of the 1993 analytic cases by site and stage at diagnosis.

FIGURE 11

DISTRIBUTION OF ANALYTIC CASES BY STAGE AT DIAGNOSIS - 1993 (TOTAL CASES = 1,396)

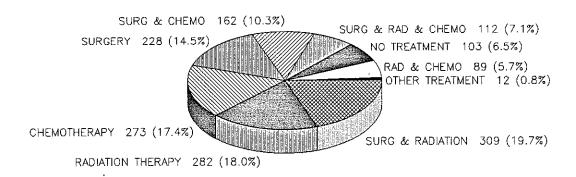


*Excludes Unknown Primaries (28 cases)

**Excludes Leukemia and Multiple Myeloma (146 cases)

FIGURE 12

DISTRIBUTION OF ANALYTIC CASES BY FIRST COURSE OF TREATMENT (SINGLY OR IN COMBINATION) 1993 (TOTAL CASES = 1,570)



GASTROINTESTINAL NON-HODGKIN'S LYMPHOMA IN ADULTS AT KING FAISAL SPECIALIST HOSPITAL AND RESEARCH CENTRE (1985-1992)

Adnan Ezzat, MB Bch ABIM FRCPC, M.A. Abdulkareem, MD Madras Raja, MD, Ofelia B. Te, CTR, Mary Anderson, CTR & Dolores K. Michels, CTR

The incidence of any malignant disease in the Kingdom of Saudi Arabia is not known, but hopefully the newly created National Cancer Registry will be able to shed some light on this issue in the near future. Since the opening of the King Faisal Specialist Hospital and Research Centre (KFSH&RC) in 1975 and up to December 1992, a total of 24,050 cases of various malignant disease were registered at our hospital-based Tumor Registry. Non-Hodgkin's lymphoma (NHL) is the most common malignant disease referred to KFSH&RC, representing 8.6% of all malignant cases.

In this report we will give an overview on extranodal NHL referred to our institution between 1985 and 1992 with special emphasis on the gastrointestinal NHL (GI NHL).

A total of 1,117 NHL cases were referred to KFSH&RC during the study period of which 438 (39.2%) were extranodal in origin. We will concentrate here on 321 extranodal analytic cases involving adult patients. Adult is defined as age of 15 years or above and analytic cases are patients who received either part or the whole treatment at our hospital.

The anatomic distribution of analytic extranodal NHL is shown in Figure 1. Waldeyer's ring lymphomas in this report are included in the head and neck lymphomas, not with the gastrointestinal. The median age for all cases was 54 years (range 15-91). The majority of the patients were Saudi nationals (90%) and the male:female ratio was 1.4:1. The histopathology according to International Working Formulation (IWF) revealed: low grade (2.5%), intermediate (79.1%), high grade (15%), and not classified, 3.4%.

Using the Ann Arbor Staging System, we found Stage I in 24.3%, Stage II in 57.6%, Stage III in 5.6%, Stage IV in 12.2% and unstaged 0.3%. Constitutional B-symptoms were found in 39% of all cases. Frequently, multimodality treatment (surgery, radiation and/or chemotherapy) were used in managing these patients, depending on but not limited to such factors as site and stage. A total of 145 cases had surgical resection, radiation therapy as adjunctive therapy or for palliation was used in 111 cases while chemotherapy was given to 244 cases. The crude survival rate for all extranodal lymphomas in this series was 59.4% at 73.7 months (Figure 2).

For the 122 (38%) primary GI NHL in this report, the median age was 51.5 years (range 15-91), the majority were Saudi (92%) and the male: female ratio was 1.6:1. The age distribution as compared to all patients with extranodal NHL is shown in Figure 3. In patients with GI NHL, the stomach was the most common site (70%), followed by small intestinal involvement (22%) and the colon, 8%. Applying the IWF for the GI NHL, we found low grade lymphoma in 2.5%, intermediate 80.3%, high grade 14.8% while 2.4% were unclassified. Over the study period, we did not encounter a single case of Mediterranean lymphoma, otherwise known as immuno-proliferative small intestinal disease (IPSID). Ann Arbor Staging revealed: 23% Stage I, 59.8% Stage II, 8.2% Stage III, 8.2% Stage IV, while 0.8% could not be staged. Constitutional B-symptoms were found in 61% of GI NHL.

For gastric lymphoma, 53/85 underwent surgery at presentation; partial gastrectomy in 36, total gastrectomy in 16 and local excision in one patient. Surgery rendered 17 patients with residual disease, 29 without, while in 7 patients this could not be ascertained. In small intestinal NHL, 20/27 underwent simple removal of the tumor or resection ± anastomosis while one patient had radical surgery. In colonic NHL, 7/10 underwent hemicolectomy and one patient had total colectomy.

Various chemotherapy regimens were used in managing these patients ranging from first to third generation anti-lymphoma therapy depending on the available protocol at the time. A total of 100 patients with gastrointestinal NHL received chemotherapy. The complete response (CR) was 49%, partial response (PR) 13%, progressive disease (PD) 8%, while 30% were not evaluable (as they received the chemotherapy as adjuvant therapy or for microscopic disease at resection margins). Radiation therapy was given to 17 patients; CR and PR were seen in 8 and 2 patients, respectively. The other 7 patients were not evaluable (NE) as for the above reasons.

When all modalities were combined, the CR was 63.9%, PR 8.2%, PD 10.7% and NE in 17.2%. The overall survival for all GI NHL at 73 months was 58.1% which was similar to all cases of extranodal NHL (Figure 2).

Our study, as compared with recent reports from Italy¹, the Netherlands² and Britain³, is summarized in Table 1. The number of cases in our report in relation to the study period is higher than the others with the exception of the Netherlands since their cases were reported from a regional registry the population of which is not stated. Our patients are younger than in other reports. However, this should be interpreted with caution since we cannot account with any certainty for the referral bias (young patients seem to be referred more than older ones because of the belief that they stand a better chance of survival). An additional fact is that the birth date as used in official documents has only recently been introduced in Saudi Arabia.

The anatomical distribution within the gastrointestinal tract varies from one country to country. Ours was similar to the Milan series. There is no perfect staging system for NHL in general and GI NHL in particular, though the Ann Arbor Staging is widely applied and showed marked variation among the four series. Again there are several histological classification that can be used in lymphomas but perhaps the International Working Formulation is the most common. Both staging and histology variations make comparisons difficult among different series. This coupled with the fact that great emphasis is to ascertain the origin of the GI NHL and whether it is of MALT (mucosa-associated lymphoma tissues) origin or not. Only the Netherlands series could be compared with ours since they used IWF and both revealed the majority to be of intermediate or high grade origin.

There are no standard approaches in the management of GI NHL, but frequently surgery, radiation, chemotherapy and/or any combination of these modalities are used in the treatment. We did not attempt to analyze the outcome of patients in these series according to the treatment modalities used.

The overall survival (OS) for GI NHL in both Milan and the British series reported 42% and 44%, respectively at 10 years. The OS from the Netherlands series and ours revealed 50% and 62% at 4 and 5 years, respectively.

In conclusion, our series is similar to other reports in many aspects and all of them underline the need for better histological classification, more precise staging system and standardization of treatment according to site, stage and histology.

REFERENCES:

- Tondini C, Giardini R, Bozzetti F, Valagussa P, Santoro A, Bertulli R, Balzarotti M, Rocca A, Lombardi F, Ferreri AJM and Bonadonna G. Combined modality treatment for primary gastrointestinal non-Hodgkin's lymphoma. The Milan Cancer Institute Experience. Annals of Oncology 4; 831-837, 1993.
- 2. Otter P, Bieger R, Kluin PM, Hermans J and Willemze R on behalf of the Study Group. Primary gastrointestinal non-Hodgkin's lymphoma in a population-based registry. Br J Cancer 60: 745-750, 1989.

3. Morton JE, Leyland MJ, Vaughan Hudson G, Vaughan Hudson B, Anderson L, Bennett MH and MacLennan KA. Primary gastrointestinal non-Hodgkin's lymphoma: A review of 175 British national lymphoma investigation cases. Br J Cancer 67; 776-782, 1993.

Table 1
SUMMARY OF GI NHL

	Milan ¹	Netherlands ²	Britain ³	KFSH&RC
Number of cases	135	96	175	122
Study period (yrs)	20	5	14	8
Age (median) ≥ 50 y < 60 y	NA NA 75%	65 NA NA	NA 73% NA	51.5 54% 69%
Male:female ratio	1.3:1	1.3:1	1.9:1	1. 6:1
Site of disease Stomach Small Intestine Large Bowel > 1 site Mesenteric	73% 15% 9% 4% -	53% 14% 16% 3% 14%	45% 33% 15% 7%	70% 22% 8% - -
Stage I II III IV Unknown	56% {28% { 16% 0	34% · 39% 3% 24% O	19% 54% 1% 26% 0	23.0% 59.8% 8.2% 8.2% 0.8%
B-symptoms	NS	NS	33%	61%
Histology Low grade Intermediate High grade Unclassified	30% {70% { 0	16% 48% 33% 3%	NA NA NA	2.5% 80.3% 14.8% 2.4%
Overall Survival (time)	42% 10 yrs	50% 4 yrs	44% 10 yrs	62% 5 yrs

NA - Not Applicable NS - Not Stated

Figure 1
DISTRIBUTION OF ADULT ANALYTIC EXTRANODAL NHL BY SITE

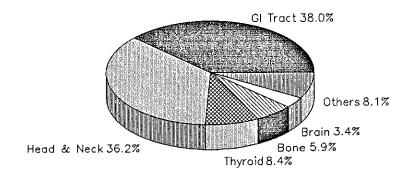


Figure 2
ADULT ANALYTIC EXTRANODAL NHL OVERALL SURVIVAL

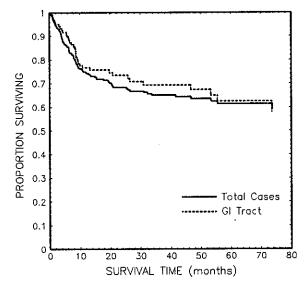
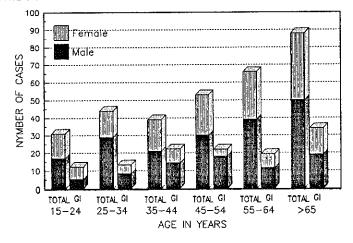


Figure 3
DISTRIBUTION OF ADULT ANALYTIC EXTRANODAL NHL BY AGE



APPENDIX A

1993 SPECIAL STUDY REQUESTS FOR TUMOR REGISTRY DATA
*Publication **KFSH&RC Presentation ***Outside KFSH&RC Presentation

T	
January Neuroblastoma Cases with Age & Site (MR Numbers) (1983-1991)***	CRU for Dr. Sackey
Cerebellar Astrocytoma Cases with Age & Sex (MR Numbers) (1975-1993)*	Dr. M. Hassounah
Ependymoma, Ependymoblastoma & Subependymoma Cases with Site, Histology, Age & Sex (MR Numbers) (1975-1993)*	Dr. M. Hassounah
B-Cell A.L.L. Pediatric Patients (MR Numbers) (1981-1993)	CRU for Dr.Martins
Acute Leukemia (ALL & AML) (MR Numbers)(1991-1993) Brain Stem Glioma/Astrocytoma Cases (MR Numbers) (1975-1993)*	Dr. Al Fiar Fahad Dr. M. Hassounah
Breast Cancer Cases which Had Radiation as Part of Initial Management (MR Numbers) (1985-1990) Marjolin's Ulcer (Squamous Cell Ca of Skin) Cases	CRU for Radiation Oncology Dr. S. El Akkad
(MR Numbers) (1975-1993) Total Number of Nasopharyngeal Cancer and All	Dr. Sheth
Malignant Cases in the Registry (1975-1993) Pituitary/Hypothalamic Tumor Cases (MR Numbers) (1975-1993)	Dr. M. Ahmed
February	
Breast Cancer Cases by Year, Age & Nationality (Saudi/Non-Saudi) (1989-1992)	Ministry of Health
Orbital Rhabdomyosarcoma Cases (MR Numbers) (1975-1993)	Dr. S. El Akkad
Retinoblastoma Cases (MR Numbers) (1975-1993)	Dr. S. El Akkad
Optic Glioma Cases (MR Numbers) (1975-1993)	Dr. S. El Akkad
Pediatric Benign Cases (MR Numbers) (1975-1993)	Dr. A. Ali
March	
Extra-Nodal NHL Adult Cases, as much information as can be extracted from the Registry (1985-1992)*	Dr. A. Ezzat
Leukemia, Lymphoma and Solid Tumor Cases by Year and Age Group (1975-1992)	Dr. S. Taher
Pediatric Cancer Cases by Histology, Site, Age Group, Sex, Sex Ratio, Stage at Dx (1982-1991)***	Dr. K. Sackey
Esophageal Cancer Cases by Stage at Dx and Treatment (1990-1991)	Dr. S. Bazarbashi
Osteogenic Sarcoma Cases, Total & Pediatrics, with Site, Sex, Age at Dx, Vital Status (MR Numbers) (1975-1992)*	Dr. S. Lindahl
Pediatric Neuroblastoma Cases (MR Numbers) (1989-1993)	Dr. A. Kofide
April	
Total Number of Osteogenic Sarcoma Cases (1981-1993)*	Dr. B. Sanjay

May				
	Pediatric A.M.L. Cases (MR Numbers) (1988-1992) Cervical Cancer Cases (MR Numbers) (1975-1993)* Ovarian Cancer Cases (MR Numbers) (1975-1993)* Cancer of the Vulva Cases (MR Numbers) (1975-1993)* Female Malignant Cases, Age 15-45, (MR Numbers) (1975-1993)***	Dr. Dr. Dr.	Y. Y. Y.	Giri Bakri Bakri Bakri Rejjal
	Breast Cancer Cases (MR Numbers) (1985-1991) Pediatric Malignant Brain & CNS Tumors with Histo- logy, Sex & Age at Dx (MR Numbers) (1975-1993)**			search Unit rtins
	Hypopharyngeal, Post-Cricoid & Pyriform Fossa Cancer Cases (MR Numbers) (1985-1993)	Dr.	z.	Mahasin
June	Glomus Jugulare and Carotid Body Tumor Cases (MR Numbers) (1975-1993)	Dr.	к.	Taibah
July	•			
-	Cancer Cases Treated By Radiation (as 1st course) by Year and Site. Ratio of Radiation Cases to All Malignant Cases (1988-1992)	Dr.	A.	Flores
Augu	st			
•	Bladder Cancer Cases (Analytic) by Year, Sex, Age, Saudi/Non-Saudi, Region, Histology, Stage at Dx, Treatment Modality (MR Number Listing, too) (1983-1992)*	Dr.	М.	Manzi
	Teratoma, Germinoma, Endodermal Sinus Tumor, Embryo- nal Ca, Choriocarcinoma of the Brain & CNS (MR Numbers) (1975-1992)*	Dr.	E.	Al-Shail
	Signet Ring Cell Carcinoma of the Colon Cases (MR Numbers) (1975-1993)	Dr.	F.	Kahlifa
	Glioma and Astrocytoma (grades 3 & 4) Treated by Radiation (as 1st course) by Site, Sex, Age Group and Stage (1988-1992)***	Dr.	Α.	Gray
	Adult Kidney Cancer Cases (histologically proven) by Year, Sex, Age, Treatment Modality & Stage (1983-1992)***	Dr.	A.	H. Kardar
	Stomach Cancer Cases with Age, Sex, Region and Histology (MR Numbers)	Dr.	s.	Shebib
Sept	ember			
-	Malignant and Benign Thyroid Cases by Year (1990-1993)	Dr.	М.	Ahmed
Octo	ber			
	Thyroid Cancer Cases with Sex, Age, Region, Class of Case, Histology and Stage (MR Numbers) (1989-1993)	Dr.	N.	Farid
	Osteogenic Sarcoma Cases (MR Numbers) (1991-1993)**			Wierzbicki
	Slides of 2 Graphs in the 1991 Annual Report of the Tumor Registry***	Dr.	A.	Al-Nasser
	Head & Neck Cancer Cases by Site, Histology, Sex, Age, Region, Saudi/Non-Saudi, Class of Case, Stage at Dx & Tx Modality (1985-1992)***	Dr.	ĸ.	Taibah
	Pediatric A.L.L. Cases by Year and Age Group (1975-1992)***	Dr.	A.	Al-Nasser
	Colon and Rectal Cancer Cases by Site and Histology (1987-1992)***	Dr.	М.	Manji

November

Slides of 6 Graphs in the 1991 Annual Report of the

Tumor Registry***
Oropharynx (Tonsils, Base of Tongue, Pharyngeal Wall),

Vallecula, Soft Palate & Uvula Cancer Cases with Site, Histology, Stage & Tx Modality (1985-1992)

Thyroid Cancer Cases, downloading of some info into a diskette for patient follow-up (1975-1993)

Sarcoma of the Uterus and Endometrial Carcinoma with Age, Histology, Histology Grade and Stage (1975-1992)***

Dr. A. Al-Nasser

Dr. A. Kandil

Mr. K. Abdulkareem

Dr. Y. Bakri

December

Squamous Cell Ca, Adenoca and Sarcoma of the Larynx, Mouth, Pharynx, Nose, Paranasal Sinuses, Salivary Gland Cases w/ Age, Sex, Histology, Stage, Date of Last Follow-up and Vital Status (1986-1992)

Pediatric Malignant Cases with Second Primary (MR Numbers) (1975-1992)**

Hodgkin's Disease Cases with Nasopharyngeal
Involvement (MR Numbers) (1975-1993)

Involvement (MR Numbers) (1975-1993)
Male Breast Cancer Cases (MR Numbers) (1989-1993)

Ovarian Cancer Cases with Histology, Grade and Stage (MR Numbers) (1975-1993)*

Dr. A. Flores

Dr. M. Mahr

Dr. D. Pradhan

Dr. A. Ezzat

Dr. Y. Bakri

APPENDIX B

1993 Tumor Committee Members

A. M. Abdulkareem, M.D.
M. Ashraf Ali, M.D.**
William Allard, D.M.D.
Hamad Al Daig
Peter Ernst, M.D.*
Adnan Ezzat, M.D.
Mohd Hannan, Ph.D.
Peter McArthur, M.D.
Dolores K. Michels, C.T.R.
Lamia NouNou
Robin Pavillard, M.D.
Assem Rostom, M.D.
Rajeh Sabbah, M.D.***
Sultan Al Sedairy, Ph.D.
Jens O. Sieck, M.D.
Jamal Al Subhi, M.D.
Beth Ann Tomasek***

Surgery Pathology Dentistry CHIC Medical Hematology Medical Oncology B&MR Research Centre Surgery Tumor Registry Social Services Quality Assurance Radiation Oncology Chairman, Oncology B&MR Research Centre Medicine Obstetrics/Gynecology Quality Assurance

^{*} Tumor Committee Chairman

^{**} Deputy Chairman

^{***} Ad hoc Members

APPENDIX C

SUMMARY OF CASES PRESENTED KFSH&RC TUMOR BOARD - 1993

SITE	NO.
Parotid Gland	1
Liver	1
Leukemia	2
Cutaneous T-Cell Lymphoma	1
Urinary Bladder	1
Brain	2
Thyroid	1
Lymphatic System Hodgkin's Disease Non-Hodgkin's Lymphoma	4 3
Aggressive Fibromatosis	1

Tumor Board Coordinator: Dr. Shouki Bazarbashi

APPENDIX D

1993 SUMMARY OF ONCOLOGY GRAND ROUNDS TOPICS

05	Jan	Interstitial Re-Irradiation for Recurrent or New Primary Gynecologic Malignancies	Dr. Randall
	Jan Jan	Total Body Irradiation Esophagus Cancer	Dr. Wynne Dr. A. Flores
02	Feb	Bone Marrow Transplantation in Children with Inborn Errors of Metabolism	Dr. P. Van Dijken
	Feb Feb	Taxol: A Review Recent Trends in Pediatric Radiation Oncology	Dr. Belanger Dr. Theriault
30	Mar	New Zealand Cancer Statistics	Dr. A. S. Abdelaal
06	Apr	Is More Better? Anthracyclines in Metastatic Breast Cancer	Dr. M. Dalmark
13	Apr	Minimal Residual Disease in Certain Hematological Disorders	Dr. M. Jackson Dr. B. Meyer Dr. R. Nounou
20	Apr	Hypercalcemia of Malignancy	Dr. A. Al-Nasser
04	May	Head and Neck Cancer	Drs Wierzbicki, Raja, Abdallah, Taibah, Osoba
11	May	Quality of Life for Cancer Patients	Dr. Osoba
	June June	Chemotherapy for Bladder Cancer Prostate Cancer - Methods of Early Detection	Dr. F. Freiha Dr. F. Freiha
	July July	Interferons Immunophenotyping of Acute Leukemia in KFSH in 1992	Dr. J. Berry Dr. S. Khalil
	Aug Aug	Brachytherapy Chronic Myeloid Leukemia: Pathogenesis and Therapy	Dr. A. Flores Dr. P. Stryckmans
24	Aug	Studies on Chemotherapy by Oral Route in Metastatic Breast Cancer	Dr. M. Dalmark
31	Aug	Palliative Care: Who Needs It?	Dr. A. Gray
05	Oct	Multiple Drug Resistance: Is It Reversible?	Dr. H. Solh
12	Oct	Hemorrhagic Cystitis Following Allo- geneic & Autologous Bone Marrow Transplantation	Dr. A. Martins
19	Oct	Third Generation Anti-NHL Combination	Dr. M. A. Raja Dr. A. Ezzat
26	Oct	Chemotherapy at KFSH&RC Hyperthermia Enhances Chemotherapy	Dr. A. Kandil
02	Nov	A Diagnostic Dilemma	Dr. K. Rao Dr. A. Al-Nasser
09	Nov	GM-CSF Modulation of Macrophage/ Killer Cell Function	Dr. P. Ernst

23 Nov	C-MYC and P53 Mutations in Burkitt's Lymphoma and their Prognostic Significance	Dr. Kishor G. Bhatia
30 Nov	Management of Medulloblastoma	Dr. A. Jamshed
14 Dec 21 Dec	Good, Bad and the Ugly: Free Radicals?! Geographical Epidemiology w/ Emphasis on Childhood Malignancies in the Middle East	
28 Dec	Male Fertility Following Cancer Therapy	Dr. A. Rostom

Oncology Grand Rounds Coordinator: Dr. Kwesi Sackey

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 Summary Staging Guide:

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.