SHORT COMMUNICATION

Cancer in Pregnancy

Hassan S O Abduljabbar, MD, FRCSC*
Hisham Ramadani, MD, FRCSC**

Cancer incidence in pregnancy occurs in approximately 2 in 1,000 pregnancies at King Abdulaziz University Hospital, Jeddah, Saudi Arabia. Twelve cases of cancer in pregnancy are presented, and some of the effects of cancer therapy in pregnancy are discussed.

Cancer in pregnancy is an unsusual event. It occurs in approximately 1 in 1,000 pregnancies¹. The obstetrician and gynaecological oncologist is faced with a therapeutic dilemma involving surgical, perinatal, obstetric, physiological and moral issues. It is a very devastating experience to the patient and her treating physician; and it is a very frustrating situation to the physician because of the lack of information available about malignant disease in pregnancy, and that there are two individuals: the mother and the fetus.

Twelve cases of cancer in pregnancy were seen at King Abdulaziz University Hospital over two years period and the effects of cancer therapy in pregnancy are discussed.

CASES AND RESULTS

Twelve cases of cancer in pregnancy were referred to King Abdulaziz University Hospital over two years

period. The total number of deliveries were 5,617. Therefore, the rate of cancer in pregnancy at our institution was 2.1 in 1,000 pregnancies.

Cancer distribution according to site was as follows: 3 cancer of the cervix, 3 lymphoma, 2 gastro-intestinal cancer, 2 cancer of the breast, 1 ovarian cancer and a thyroid cancer.

Table 1 is a summary of the 12 cases.

DISCUSSION

In our institution cancer occured in 2.1/1000 pregnancies. It had been estimated that cancer may complicate 1 in every 1000 pregnancies. Table 2 shows the incidence of cancer in pregnancy^{1,4-8}.

Anatomical and physiological changes that occur during pregnancy might obscure early neoplasm and make the diagnosis very difficult on clinical grounds. The best example of this is carcinoma of the ovary. Execerbation of some cancer can occur during pregnancy such as breast cancer. Increase in the vascularity and lymphatic drainage might have an effect on the dissemination of cancer⁸.

^{*} Diplomat American Board of Obstetrics & Gynaecology Consultant and Assistant Professor

^{**}Department of Obstetrics & Gynaecology King Abdulaziz University Hospital Jeddah Saudi Arabia

Table 1

Summary of 12 cases of cancer in pregnancy

Case No.	Diagnosis	Age	Gravida	GA	Stage	RX	Outcome
1	Ca. Cervix	31		28	IB	SUR	C/S 980 gm EARLY NND
	Ca. Cervix			+2	0.10 0.10	RAD	SVD 300 gm BOY, L & W
	Ca. Cervix	32	gang 4	40	II I	RAD	SVD 3000 gm BOY, L & W
4	*Lymphoma ³		midi 7	25	IV	CHEM	SVD 1970 gm BOY, L & W
5	*Lymphoma ³	25	2	20	IV	СНЕМ	ABORTION 475 gm
6	Lymphoma ^{2,3}	38	7 for	23	IA	SUR	C/S 2400 GM BOY,L & W
7	Ca. Breast		3	38	II - d chemother ancer in pre-	SUR RAD CHEM	INDUCED VAG. DEL. 3100 gm GIRL, L & W
	Ca. Breast		4	37	II ed to the seco	SUR RAD CHEM	INDUCED VAG. DEL 3000 gm BOY, L & W
9	Ca. Stomach	35	4	35	IV	SUR	C/S 2460 gm BOY, L & W
10	C-R Cancer	29	3	23	III	RAD	C/S APH SUR 650 gm EARLY NND
11 11 11 11 11 11 11 11 11 11 11 11 11	Ca Ovary	25			ub ii IB q s	SUR CHEM	C/S 2500 gm BOY, L & W
12	Ca. Thyroid	28	2 180	35	nferioryena crion of the gastr in the gastr	SUR	SVD 2980 gm GIRL, L&W
SUR — Surg RAD — Radi SVD — Spor	iotherapy ntaneous Vaginal Delivery o-rectal	Creary RK, Re Practice, 2nd ed. Gusberg SB, Shi York, Edinburg Livingstone: 199 Muller BJ, And	C L C	IND — CHEM — A & W — C/S — Cx —		erapy d well a Section t	ADJOTHERA Prognant pai

Table 2 Incidence of cancer in pregnancy

Site/Type	Estimated Incidence / 1000 pregnancy				
Cervix Uteri		28 IB			
Non-invas	ive	1.3			
Invasive		1.0			
Breast		0.33			
Melanoma		0.14			
Ovary		0.10			
Thyroid		Unknown			
Leukaemia		0.01			
Lymphoma		0.01			
Colo-rectal		0.02			

Patient with cancer in pregnancy, poses therapeutic dilemma, because the patient is usually young and a second life is involved. Termination of pregnancy might be required. In our Islamic society, termination of such cases is permitted.

MODALITIES OF TREATMENT OF CANCER IN PREGNANCY

Surgery, radiotherapy and chemotherapy are the usual management options for cancer in pregnancy⁸⁻¹⁰.

SURGERY

Surgery should be postponed to the second trimester if possible, for evaluation or treatment of cancer particularly intra-abdominal cancer. The incidence of spontaneous abortion is reduced.

Removal of ovarian cysts or complete ovarian resection may be safely accomplished in the second trimester when the placental production of progesterone replaces corpus luteum cyst⁹.

The supine position of the patient during surgery might produce hypotention and hypoxia to the fetus and therefore, 15 degree wedge under the right hip will produce left uterine displacement of the inferior vena cava. Progesterone produces marked relaxation of the gastrointestinal smooth muscle leading to delay in the gastric emptying time and therefore may result in fastric dilatation¹⁰.

RADIOTHERAPY

Pregnant patients receiving radiotherapy directed to the pelvis for pelvic malignancies will suffer a fetal demise and usually spontaneously abort. Patients receiving supra-diaphragmatic irradiation will receive only minor exposure, due primarily to internal radiation scatter, and may safely carry an early pregnancy. However supra-diaphragmatic irradiation in late pregnancy may expose the growing fetus to excessive radiation that will produce an acceptable fetal injury^{10,11}.

CHEMOTHERAPY

Information on the effect of chemotherapy on the fetus remains incomplete⁸, but avoidance of cytotoxic chemotherapy should be the rule, not only in the organogenesis phase but in whole period of the first trimester of pregnancy⁹. Even a single chemotherapeutic agent, in the first trimester will lead to either spontaneous abortion or congenital anomalies of the offspring¹⁰. In the second and third trimester, a single chemotherapeutic agent rarely causes any congenital anomalies. In recent years, multiple reports confirm that even multiple agents chemotherapy in the second and third trimester rarely results in congenital anomalies.

Long-term sequence due to intra-abdominal exposure to cytotoxic chemotherapy remains unknown. It is possible that deleterious effect in the offspring of women exposed in utero to cytotoxic chemotherapy will occur. These patients require long-term monitoring.

REFERENCES

- Allen HH, Nisker JA. Cancer in Pregnancy: Therapeutic Guidelines. Mount Kisco, New York: Futura Publishing Co, 1986.
- Abduljabbar H, Ghazzawi B, El-Hosseiny M. Primary ovarian lymphoma in pregnancy: A case report. Ann Saudi Med 1990; 10:453-6.
- Ramadani HM, Abduljabbar H, Ghazzawi B. Non-Hodgkin's lymphoma in pregnancy. Saudi Med J 1991;12:32-6.
- Pollack ES. The epidemiology of cancer and the delivery of medical care services. Pub Health Rept 1984;99:476-85.
- Cutler SJ, Young JL Jr. Third National Cancer Survey; Incidence Data. Natl Cancer Inst Monogr 1975;41:20-32.
- Chung A, Birnbaum SJ. Ovarian cancer associated with pregnancy. Obstet Gynecol 1973;41:211-9.
- Breese MW. Cancer of thyroid gland in women of childbearing age. Am J Obstet Gynecol 1963;86:616-9.
- Disaia PJ, Creasman WT. Clinical Gynecologiconcology. 3rd ed. St Louis, Washington DC, Toronto: The CV Mosby Co, 1989:511-2.
- Creasy RK, Resnik R. Maternal Fetal Medicine: Principle and Practice. 2nd ed. Philadelphia, WB Saunders Co: 1989.
- Gusberg SB, Shingleton HM, Deppe G. Female Genital Cancer New York, Edinburg, London, Melbourne. New York, Churchill Livingstone: 1986.
- 11. Muller HJ. Artificial transmutation of the gene. Science 1927;66:84.