Historical Vignette

Saul Hertz, MD (1905-1950)

A Pioneer in the Use of Radioactive Iodine

Barbara E. Hertz, BS, MS, M.Ed & Kristin E. Schuller, BA, MAT

Greenwich, CT; Boston University School of Medicine

Correspondence to:

Barbara Hertz 6 Buckthorne Lane Greenwich, CT 06830 203-661-0777 htziev@aol.com

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Dr. Saul Hertz Poses a Question

Long before the first atomic bomb of World War II, Dr. Saul Hertz (1905-1950) (Fig. 1) took a profound step forward in the field of Nuclear Medicine. On November 12, 1936, Dr. Hertz attended a luncheon meeting at Harvard Medical School, with the President of the Massachusetts Institute of Technology (MIT), Dr. Karl Compton (1887-1954). Dr. Compton was discussing "What Physics can do for Biology and Medicine." Dr. Hertz, who was Director of the Thyroid Clinic at the Massachusetts General Hospital (MGH) (1931-1943), asked Dr. Compton, "Could iodine be made radioactive artificially?" (1). The question came quite spontaneously, as Dr. Hertz had been conducting studies on the effect of iodine on thyroid function. Dr. Compton responded by letter on December 15, 1936, describing the properties of radioactive iodine (1). A week later, Dr. Hertz wrote back that he hoped to do animal experiments and devise a useful therapy in patients with hyperthyroidism (1).

During the early months of 1937, the engineering skills of MIT and the medical expertise of MGH were brought together. Dr. Hertz was in charge of the biological and medical work with Dr. Arthur Roberts, a young physicist from MIT. Drs. Hertz and Roberts did their first series of experiments with ¹²⁸I on rabbits in late 1937. These early experiments involving 48 rabbits demonstrated that the normal thyroid gland concentrated ¹²⁸I, while the hyperplastic thyroid gland took up even more (2,3,4). In May 1938, the John and Mary Markle Foundation of New York City funded the building of a cyclotron at MIT with a \$30,000 donation. It was completed two years later in 1940. Experiments continued on rabbits during 1939 and 1940. Without a cyclotron, Hertz and Roberts were dependent on others for longer-lived radioactive isotopes such as ¹³¹I.

2

The First Patients

In late 1940, Dr. Hertz began using the cyclotron to produce ¹³⁰I and ¹³¹I, which he used in studies with subjects with Graves' hyperthyroidism (5). In early 1941, he administered ¹³⁰I to the first patients at MGH (Fig. 2). Gradually, a series of about thirty patients were treated and followed until Dr. Hertz joined the Navy during the war years (Figs. 3A,3B).

After the war, there was great interest in using atomic energy for peaceful purposes. In May 1946, JAMA published "Radioactive Iodine in the Study of Thyroid Physiology" (6), reflecting the success of Dr. Hertz's treatment with the first series of patients over a five year follow-up (Fig. 3). This firmly launched the use of radioactive iodine that has become the standard treatment for Graves' disease.

Dr. Saul Hertz uses Nuclear Fission in Cancer Treatment

Dr. Hertz established the Radioactive Isotope Research Institute in 1948; it's purpose was to apply fission products to the treatment of thyroid cancer, goiter and other malignant tumors. He extensively studied the use of radioactive iodine in the treatment of thyroid cancer (7), and the application of radioactive phosphorus and the influences of hormones on cancer as demonstrated by isotopic studies (8,9).

Remembering Saul Hertz, MD

Saul Hertz was a brilliant scholar and researcher who devoted his life to scientific work. He authored over thirty scientific publications on thyroid physiology, thyroid

disease and its treatment. Dr. Hertz pioneered tracer use of radioiodine in studying thyroid physiology, and was the first to administer therapeutic doses of radioactive iodine to treat thyroid disease. Mrs. George Bush, who was successfully treated for thyroid disease, wrote to Dr. Hertz's wife, Vitta Hertz, "It is comforting to know that so many people are well because of the scientific expertise of people like Dr. Hertz."

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Figure legend

Figure 1. Saul Hertz, MD (circa 1945)

Figure 2. Saul Hertz, MD uses a multicounter to analyze the distribution of radioactive iodine in a patient.

Fig. 3. (A.) Saul Hertz's original, handwritten table detailing the first series of MGH subjects whose hyperthyroidism was not cured by administration of ¹³¹I. (B.) Original, handwritten table detailing the first series of MGH subjects whose hyperthyroidism was successfully treated with ¹³¹I.







	TABLE I AN ANALYSIS JF CASES "NOT CURED" BY RA-I+KI (TO MARCH-'46)								% OF RO-I (URINE) EXCRETED - 72HRS			
SERIES NO.	CASE-HOSP.NO.	BMR PRIOR TO I 130	DOSAGE OF 1130 and DATES OF ADMINISTRATION	BMR PRIOR TO SUB-TOTAL	POST-OF BMR			TOTAL THE		ESTIMATED THYROID WT. BEFORE 1'30	FOLLOWING TH ADMINISTRAT OF 1130	
1	ELIZABETH D. MGH-173954	+30	2.1 mC 3-31-41 3.4 1.3 mC 4-16-41 mC	(-5)(-7)	(-29)	34	INVOLUTION	470 220	660 240	35	20 28	
5	LILLIAN R. MGH-308552	+35	5.7mC 7-16- 41	PLANNED EXPERIMENT	(-20)	31	HYPERPLASIA NO INVOL-	1000	1150	40	27	
10	GLADYS B. MGH-121922	+55	0.7mC 2-2- '42	(+3)	(-26)	26 30}56	HYDERPLASIA MOD. INVOL.	120	80	60	38	
14	WILFRED B. MGH-363179	+50	15mC 7-15-42	(-15)	(-24)	55	HYPERPLASIA + INVOLUTION	650	- 1	60	71	
16	CARMELLA D. MGH-255820	+25	10mc 8-11-42	(-8)	(-24)	28	INVOLUTION	1800		45	6	
19	РЕГЕК С. МС-H-369233	+65	15 mC 8-25-42 8 mC 3-8-43 5 mC 3-9-43 mC	(+8) (+13)	(+36) 70 (-18)	35	SL. HYPERPLASE TINVOLUTION	2000	_	60	9 15 7	
2	MARGARET B. MGH-300230	+35	1.4mC 5-10-'41 0.9mC '41 5.6 2.4mC '42 mC 0.8mC '42	41 5.6 PERSISTENT THYROTOXICOSIS				160 110 120 100	140 100 120 100	40	54* 48 78	
4	CAMILLE SCH* MGH-309302	+30	3.6m(7-14-41) 5.8 2.2m(7-31-41) mC + EYES BETTER, NO GOITER BMR (+2) OFF MED 4 YRS				270	300 180	60	55 56		
3	RUTH M. +5		3.4mC 6-6-41 REMISSION FOR IVR - THEN					430	\$10	45	45	
3	MGH-304558		20mC 1-9-46 (RECENTLY FOR TRUE RECURRENCE)					4300	-	30 (RECURRENT)	35	

		TABLE II-A ON B.		OF EXAM						
series No.	CASE-HOSP. NO	2005 0 5 1/20	BMR	BMR LEVEL	TIME OFF	THYROID SIZE'46	ESTIMATED THYROID WT. (gm.)	7 OF Ra-I EXCRETED 72-HOURS	ESTIMATED IRRIDIATI 12 HOUR	
6	MICHAEL K. MGH-227382	2.3mC 7-24-41 4.0 1.7mC 7-30-41 mC	+45	DEC'42 (-9) MAY'43(-16) JAN'46 (-7)	4. YR 5.+	N	45	35 22	320 280	390 300
7	ALLISOND. (A. MGH=319927	ET 9.) 1.4 mC 9-19-4/ 2.9 1.5 mC 9-21-41 mC	* + 65	1-8-46 (-6)	4 YRS.	N	45	9 20(?)	260 260 ^(?)	230
8	NAOM 1 K.(A MEH-32/155	CONTRACTOR OF THE OWNER	+30	7-17-45 (-3) 3-27-46 (+4)	7 MOS	FIRM 2XN	40	15	300	250
9	MILDRED G. MGH-322935	4,9m(11-26-'41	+30	5-8-'45(-10)	4 YRS.	N	60	17	650	420
11	FRANCES H MGH-198910	5.8mc 4-9-42	+37	7-9-142 (-12) 2-24-144(+9) 2-3-146(-13)	3.5YRS.	~	60	17	750	380
12	FERDINAND L. MGH-354330	7.5mC 5-15-'42	+55	2-3-46(-13)	3 YRS.	HARD 1.5 X N	60-75	26	950	500
13	DOROTHY Р. МGH-585541	12mc 6-9-142	+30	3- '43 (+6) 2-3- '46 (-10)	3YRS.	\sim	40	71	750	
15	MARY M. MGH-362811	6m(8-11-42) 10 4m(8-11-42) mC	+35	4-145(-6) 2-3-46(+2)	IOMOS.	N	40	10	2000	
17	GEORGE T. BCH-1076956	13mc 8-13-342	+50	6-10 ·44 (-15) 1-6-'46 (-9)	3YRS,+	N	60	14	1300	
18	JEANETTE G. MGH-367094	10.5mC 8-15-42	+35	8-22-'44(+11) 2-16-'46(+1)	3 YRS. +	\sim	40	15	2000	
20	ANNE D. MGH-233271	10mc 11-14 42	+50	4-3- 45(-1) 2-16-46(-5)	2 YR5.+	N	45	20	1600	
21	RICHARD T. BIH-67686	14mC 11-20-'42	+45	1-8-46 (-13)	3YRS.+	N .	50	15(?)	2000	
22	ESTHER R. MGH 38504	13m(3-9-'43	+20	6-30-'43 (-8)	2 YR5.+	"?N"(LMD.)	55	33	2200	
23	MARGARET D. MGH-385741	8m (3-15-43) 18 10m (3-16-43) mC	+55	6-9-43(-11) 2-16-46(-3)	2 YRS.+	FIRM 1.5XN	75	76 67	500	
24	VANE ANNE F. MEH-397402	10.5mC 3-26-43 15 4.5mC 3-27-43 mC	+40	12-'45(-5)	2 YRS.+	N (DR.J.C (ZILHAR	(.) DT) 50	57? 31	1000	
25	SOPHIE R. MGH-397951	16mC 4-2-'43	+44	9-28-44 (-7) 4-27-45 (+9) 3-20-46(+14)	2 YRS.+	N(DRJ.C (AUB)	3/1	20.6 63.0	750	
26	BESSIE W. METAB,#23843	12mC 4-6-'43	+39	1-16-46 (+2)	2 YR5.+	N	45	85	350	11
27	WINIFRED K. MGH-398698	13 0 4-12-43	+40	7-17-45(-16) 2-15-46(-10)	2 YRS.+	N	50	33	1600	6-7
28	MARGARET H. p.p. Dr Hertz	10.5mC 4-13- 43 21 11.0mC 4-13- 43 mC	+55	12-45 (-15) 2-3-46 (+6)	2 YRSt	N	75		2000	
29	JULIAZAF, RY MCH-395852	8mC 5-29-43)12 4mC 3-30-43 mc	+30	246 (+4)	2 YR5+	N	55	10 53(?)	1200 250	
		* 8 DAY ISOTOPE DURING DECAY NOT VEAGURED	, IMF	Y ARE I HEK	EFINI	SS OF I EXCESS	ODINE FROM	THYPOL	0	