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### **Abstract**

Computer processed adrenal imaging using dual-radioisotopes, 6 beta-iodomethyl-19-nor-cholest-5(10)-en-3 beta-ol-131I and 99mTc-phytate was performed in 12 patients with primary aldosteronism and 4 with Cushing's syndrome due to adrenocortical tumor. Adreno-photoscanning and hepato-photoscanning were performed in the same position 2-4 days following intravenous administration of radiocholesterol. The scintigraphic information was stored on cassettes and scan subtraction and a digital-computer method for data smoothing were performed on an oscilloscope. The tumor site could be determined in all cases until day 4 by this computer processed image.

**KEYWORDS:** new adrenal imaging technique, dual radioisotopes, adrenocortical tumor.

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# A NEW ADRENAL COMPUTER IMAGING TECHNIQUE USING DUAL-RADIOISOTOPES

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Abstract. Computer processed adrenal imaging using dual-radioisotopes, 6ß-iodomethyl-19-nor-cholest-5(10)-en-3ß-ol-<sup>131</sup>I and <sup>99m</sup>Tc-phytate was performed in 12 patients with primary aldosteronism and 4 with Cushing's syndrome due to adrenocortical tumor. Adreno-photoscanning and hepatophotoscanning were performed in the same position 2-4 days following intravenous administration of radiocholesterol. The scintigraphic information was stored on cassetes and scan subtraction and a digital-computer method for data smoothing were performed on an oscilloscope. The tumor site could be determined in all cases until day 4 by this computer processed image.

Key words: new adrenal imaging technique, dual radioisotopes, adrenocortical tumor.

The use of scintigraphy and venography has greatly improved the diagnosis of adrenal tumors. Usual scintigraphy is more sensitive than retroperitoneal pneumography and has fewer complications than venography, but diagnosis takes longer and small adrenocortical tumors are difficult to visualize. We developed a new technique for early diagnosis of small adrenocortical tumors employing the dual-radioisotope techniques and digital image-subtraction methods used in pancreas visualization (1,2).

#### MATERIALS AND METHODS

Patients with primary aldosteronism. Ten female and two male patients between 29 and 56 years of age were examined (Table 1-a). Hypertension (160-230 mmHg systolic), muscle weakness, headaches and paresthesia were observed in most patients. Serum potassium levels were low, between 2.1-3.2 mEq/1. Plasma renin activity levels were low at 0.05-0.94 ng/ml/h at rest, while plasma aldosterone concentrations were high at 28.0-205 ng/dl.

Patients with Cushing's syndrome due to adrenocortical tumor. The patients were four females between 25 and 44 years of age with symptoms such as moon face, obesity, amenorrhea and hypertension (160-198 mmHg systolic) (Table 1-b). The plasma cortisol levels at 9 a.m. were increased to 29.5-34.5  $\mu$ g/dl, while the plasma ACTH levels were suppressed to undetectable levels (under 25 pg/ml) in all cases. Urinary 17-OHCS excretion was neither suppressed with 16mg of dexamethasone nor increased with 3g of

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## metyrapone.

Methods of adrenal imaging and adrenal angiography. Five drops of Lugol's iodine solution was given b.i.d. prior to injection of the radioactive tracer and continued for six days to block thyroid accumulation of <sup>131</sup>I. Then, a tracer dose of 0.3-0.4 mCi. (0.6 μCi/kg/b.w.) of 6β-iodomethyl-19-nor-cholest-5(10)-en-3β-ol-<sup>131</sup>I (NCL-6-<sup>131</sup>I, Daiichi Radioisotope Laboratories, Tokyo) was injected intravenously (Fig. 1). On day 2 to 4 after tracer administration, adreno-photoscanning was conducted from the anterior side using a gamma camera (Nuclear-Chicago Corp.) (Fig. 2-A). Hepato-photoscanning was also conducted in the same position after an intravenous injection of 3.0 mCi of <sup>99m</sup>Tc-phytate (Daiichi Radioisotope Laboratories, Tokyo) (Fig. 2-B). These <sup>131</sup>I and <sup>99m</sup>Tc images were

Table 1-a. Clinical symptoms and laboratory data in 12 patients with primary aldosteronism

Cas no	e Patients	Age	Sex	Clinical symptom	Blood pressure	Serum K	PRA	PA
1.	M.K.	29	$\mathbf{F}$	Headache, paralysis	210/130	2.9	0.10	74.0
2.	C.T.	29	F	Paralysis, muscle weakness	200/102	2.8	0.36	45.8
3.	M.S.	39	F	Headache, paralysis, visual disturbance	170/100	3.2	0.05	28.0
4.	M.M.	40	F	Paralysis, muscle weakness	200/108	2.4	0.94	76.5
5.	E.I.	32	F	Headache, paralysis, edema	172/120	2.5	0.1	36.0
6.	K.U.	53	F	Headache, polyuria	220/110	2.8	0.24	134
7.	T.N.	49	F	Paralysis, edema	200/100	2.1	0.80	42.0
8.	T.O.	48	$\dot{\mathbf{M}}$	Palpitation	220/130	3.0	0.30	40.0
9.	Y.A.	37	F	Headache, paralysis, muscle weakness	160/110	2.4	0.30	42.0
10.	K.T.	37	$\mathbf{F}$	Paralysis, muscle weakness	170/94	2.6	0.25	205
11.	T.I.	54	M	Headache, paralysis, muscle weakness	230/130	2.6	0.40	40.1
12.	Y.Y.	56	F	Muscle weakness, edema	175/90	2.5	0.50	32.0

PRA: Plasma renin activity, PA: Plasma aldosterone

Table 1-b, Clinical symptoms and laboratory data in 4 patients with Cushing's syndrome due to adrenocortical tumor

Case no	Patients	Age	Sex	Clinical symptom	Blood pressure	P-F	Ú-17 OHCS	P- ACTH
13.	H.M.	30	F	Moon face, obesity, buffalo hump	186/100	29.5	17.84	n.d.
14.	T.M.	39	F	Moon face, obesity, buffalo hump, amenorrhea, muscle weakness	198/110	31.9	15.11	n.d.
15.	E.H.	25	F	Moon face, obesity, amenorrhea	180/130	34.5	20.0	n.d.
16.	T.O.	44	F	Moon face, obesity, buffalo hump, amenorrhea	160/102	32.2	17.3	n.d.

P-F: Plasma cortisol, U-17 OHCS: Urinary 17-hydroxy corticosteroid,

P-ACTH: Plasma adrenocorticotropic hormone

displayed as a 64 × 64 matrix on an oscilloscope, photographed and stored digitally on cassete. The region of interest specific to the liver was defined by profile display of the liver image with <sup>99m</sup>Tc (Fig. 2-C) and average <sup>131</sup>I/<sup>99m</sup>Tc ratios for the matrix subtraction were determined after <sup>131</sup>I and <sup>99m</sup>Tc data were transferred from cassete to TOSBAC 40 computer (Toshiba, Tokyo). Thereafter a computer-processed adrenal image was obtained on an oscilloscope by display with a three-level gray scale (Fig. 2-D).

Adrenal venography was performed on eleven patients using the method of Reuter et al. (3) and adrenal arteriography was conducted on one patient with primary aldosteronism (case 9) using the method of Kahn and Nichrosz (4).

Fig. 1. Structural formulae for CL-19-131 and NCL-6-131 I

#### RESULTS

Table 2 presents the results of retroperitoneal pneumography, adrenal angiography, adrenal imaging, tumor site and tumor size.

In primary aldosteronism patients, retroperitoneal pneumography did not show a clear tumor shadow. A suspicious shadow was evident in only 3 patients (Cases 4,5 and 7). The tumor site was detected definitely by adrenal venography in four of nine cases. A clear tumor angiogram was not obtained in Case 2 because the tumor was in the upper posterior region of the adrenal vein and in Case 3 because the tumor was too small. Venous catheterization for measurement of aldosterone levels located the tumor site in six of nine cases. In Case 9 the tumor site was detected by selective adrenal arteriography. However, our adrenal computer imaging located the tumor site at day 2 after radioactive tracer injection in Cases 3, 5, 8, 9, 10 and 12, and at day 4 in Cases 1, 2, 4, 5, 7 and 11 (Figs. 2-4). In all primary aldosteronism cases, unilateral adrenalectomy or extirpation of the tumor was successful. Tumor size ranged from  $0.9 \times 0.9 \times 0.6$  to  $3.0 \times 3.0 \times 1.5$  cm (Table 2) and adenoma was comfirmed histologically.

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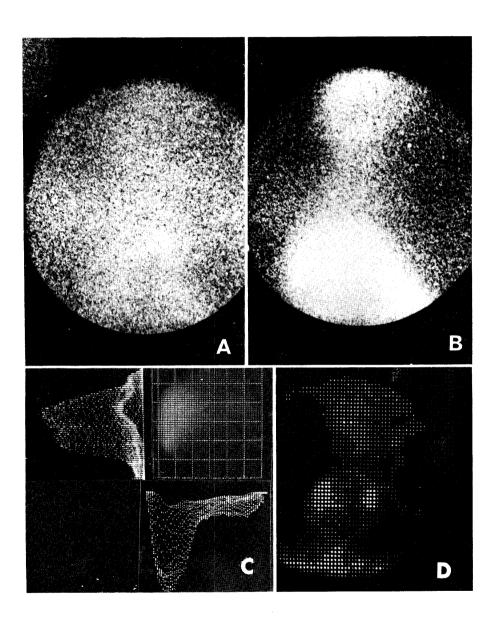


Fig. 2. Computer processed adrenal image of a primary aldosteronism patient (Case 3) A: Routine adreno-photoscan (anterior view) 2 days after intravenous injection of 0.3 mCi of NCL-6-131I. B: Routine hepato-photoscan in the same position. C: Profile display of the liver image with 99mTc. D: Computer processed adrenal image after matrix subtraction and display with three-level gray scale.

## Adremal Computer Imaging

TABLE 2. SUMMARY OF RESULSTS

Case	Patients	PRP	Adrenal venography	aldostarono	Adrenal imaging	Tumor site	Tumor size
1.	M.K.	Negative	Right	Right	Right	Right	1.8×1.7×1.3 cm
2.	C.T.	Negative	Susp. of right	Susp. of right	Right	Right	1.8×1.7×1.3 cm
3.	M.S.	Negative	Susp. of right	Right	Right	Right	0.9×0.9×0.6 cm
4.	M.M.	Susp. of left	Left	Left	Left	Left	1.1×1.1×1.1 cm
5.	E.I.	Susp. of left	Left	Susp. of left	Left	Left	1.4×1.4×1.2 cm
6.	K.U.	Negative	R: unsucc. L: negative	Left	Left	Left	1.2×0.8×0.6 cm
7.	T.N.	Susp. of ritht	Right	Right	Right	Right	1.5×1.5×0.5 cm
8.	T.O.	Negative	R: unsucc. L: negative	Left	Right	Right	2.0×1.5×1.5 cm
9.	Y.A.	Negative	Left*	Not performed	Left	Left	1.8×1.8×1.4 cm
10.	K.T.	Negative	Not performed	Not performed	Right	Right	3.0×3.0×1.5 cm
11.	T.I.	Negative	R: unsucc. L: negative	Right	Right	Right	1.0×0.9×0.8 cm
12.	Y.Y.	Negative	Not performed	Not performed	Right	Right	1.4×1.4×1.1 cm
13.	H.M.	Left	Susp. of left		Left	Left	$2.5 \times 2.5 \times 2.0$ cm
14.	T.M.	Right	Right		Right	Right	6.0×5.0×4.0 cm
15.	E.H.	Negative	Not performed		Left	Left	2.5×2.2×1.6 cm
16.	T.O.	Left	Not performed		Left	Left	2.2×2.0×0.6 cm

\* Selective adrenal arteriography was performed PRP. Pneumo retroperitoneum

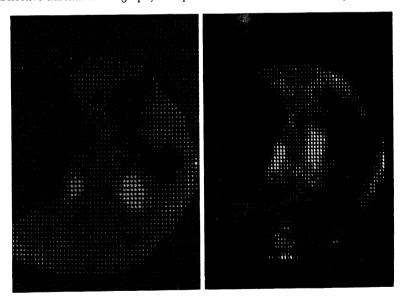


Fig. 3. (left) Computer processed adrenal image (anterior view) of a primary aldosteronism patient (Case 4) on day 4.

Fig. 4. (right) Computer processed adrenal image (anterior view) of a primary aldosteronism patient (Case 5) on day 2.

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In Cushing's syndrome due to adrenocortical tumor (Cases 13-16), the tumors were large and easily located by retroperitoneal pneumography in three of four cases. Adrenal computer imaging showed marked radiocholesterol uptake at the tumor site in all cases (Fig. 5) including case 13 in whom the adrenal venography was not typical. A later unilateral adrenalectomy showed the presence of adrenal tumors, ranged from  $2.2 \times 2.0 \times 0.6$  to  $6.0 \times 5.0 \times 4.0$  cm (Table 2), and comfirmed adenoma histologically.

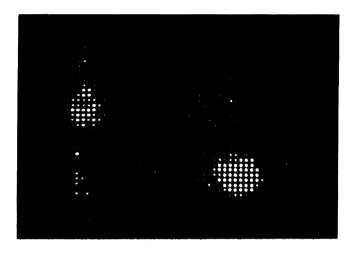


Fig. 5. Computer processed adrenal image (anterior view) on day 4 of a patient with Cushing's syndrome due to lt. adrenal tumor (Case 16).

#### DISCUSSION

Significant progress has been made in locating adrenocortical tumors by adrenal venography and adrenal scintigraphy. Adrenal venography and venous catheterization for measurement of aldosterone levels are good for diagnosing adreno-cortical disease especially for primary aldosteronism (5). However, they are technically difficult and can be complicated by extravasation (6-8).

In adrenal imaging, Nagai and associates (9) in 1968 reported visualization of human adrenal glands by radioiodinated stigmastrol, but only a large adrenocortical tumor could be detected. In 1971 Blair and associates (10) synthesized 19-iodocholesterol (CL-19-<sup>131</sup>I) (Fig. 1) as a possible human adrenal scanning agent by iodinating the C-19 position of cholesterol. They reported that the isotope had a biological half-life of 3.15 days and an effective half-life of 2.26 days in man. Lieberman *et al.* (11) and Conn *et al.* (12) reported the clinical usefulness of this agent. However, some authors (13,14) reported difficulty in detecting small adrenocortical tumors. In regard to this, Moses *et al.* (15) reported computer display with background subtraction and Conn *et al.* (12,16) described a

dexamethasone-modified adrenal scintiscan. But these methods require further study of the time needed for diagnosis. Recently, NCL-6-131 was successfully synthesized by Kojima et al. (17), Maeda et al. (18) and Sarker et al. (19). Tissue distribution studies with NCL-6-131 in male rats revealed that the ratio of adrenal gland-to-liver concentration was 113 on day 1, 330 on day 3 and 851 on day 7 after radiocholesterol administration (17). These ratios were very high compared with the 157 (20) on day 7 with CL-19-131 I. NCL-6-131 is a superior adrenal scanning agent to CL-19-131 I. Some authors (21-23) reported the clinical usefulness of NCL-6-131I, but it was difficult to obtain satisfactory images because of liver images overlapping with the right adrenal gland 2-4 days. We performed adreno-photoscanning for two to four days following administration of NCL-6-131 When a substantial amount of 131 had accumulated in the adrenal glands, then performed hepato-photoscanning using 99mTc-phytate in the same position. The overlapping liver images of the right adrenal gland were eliminated by matrix subtraction and clear images of both adrenal glands were obtained on an oscilloscope. The tumor site could be predicted preoperatively in all cases by this method. Thus, our methods of adrenal computer imaging should be more suitable for detecting adrenocortical tumors in outpatient department because of its sensitivity and simplicity.

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