

Title	Wegener's granulomatosis manifested by 67Ga- citrate scanning
Author(s)	大島,統男;安河内,浩;町田,喜久雄
Citation	日本医学放射線学会雑誌. 1975, 35(12), p. 1092- 1097
Version Type	VoR
URL	https://hdl.handle.net/11094/16857
rights	
Note	

The University of Osaka Institutional Knowledge Archive : OUKA

https://ir.library.osaka-u.ac.jp/

The University of Osaka

Wegener's granulomatosis manifested by ⁶⁷Ga-citrate scanning

Motoo Ohshima*, Hiroshi Yasukochi and Kikuo Machida Department of Radiology, University Branch Hospital, University of Tokyo Faculty of Medicine, Tokyo, Japan

Research Code No.: 722

Key Words: Wegener's granulomatosis, 67 Ga-citrate, Lung, Large

nodular densities, Focal glomerulonephritis

67Ga-citrate スキャンにて陽性像を呈した Wegener の肉芽腫

東京大学医学部附属病院分院放射線科 大島 統男* 安河内 浩 町田喜久雄

(昭和50年4月24日受付) (昭和50年7月24日最終原稿受付)

我々は、 両肺 に 腫瘤状陰影 が 出没し、 ⁶⁷Ga-citrate によるシンチグラムで 著明な 陽性像を呈し、更に巣状糸球体炎を伴つた一剖検例を経験したので報告する.

症例は44歳の男性で発熱及び腰痛を主訴に来院し、胸部×線写真で左肺尖部に比較的柔らかな腫瘤状陰影及び左胸水を認めた(図1,2).化学療法を行ない、肺腫瘤状陰影は一時消腿した(図3).しかし2年半後の胸部×線写真では両肺に多発性腫瘤状陰影が出現した(図4,5).同時期に

『Ga-citrate シンチグラムを施行したところ,胸 部X線写真に相当して陽性像を認めた(図 6).再 び化学療法を施行したところ,今回も肺病変には 効果があつた(図 7).しかし患者は全身状態が悪 化し死亡した.剖検所見は i)両肺に granulomatous fibrosis の瘢痕(図 8) ii) focal glomerulonephritis(図 9) iii) necrotizing colitis(図 10) 等の所見があり,病理学的診断は Wegener's granulomatosis であつた.

Abstract

This paper reports an autopsy case with Wegener's granulomatosis complicated with severe focal glomerulonephritis. In this case large nodular densities appeared intermittently in the bilateral lung fields and showed positive scan by ⁶⁷Ga-citrate.

^{*}Present Address: Department of Radiology, Nagoya University School of Medicine, Nagoya, Japan. 名古屋大学医学部放射線医学教室

Introduction

Since Wegener reported the syndrome of destructive rhinitis, pulmonary lesions and glomerulonephritis as Wegener's granulomatosis in 1936¹²⁾, there has been much work published in the literature on Wegener's granulomatosis⁽³⁾⁴⁾⁵⁾⁽⁸⁾. Carrington and Liebow reported thereafter the limited forms, only involving the lung without evidence of glomerulitis and rhinitis⁽¹⁾. It is reported herewith an autopsy case of Wegener's granulomatosis where intermittent appearance of nodular shadows demonstrated in the lungs, with positive uptakes by ⁶⁷Ga-citrate scanning, and with severe focal glomerulonephritis.

Case Report

A 44 year old male visited Tokyo University Branch Hospital with the chief complaints of fever and lumbago in February 1970. He had suffered from mild diabetes mellitus. Intravenous pyelogram was within normal limits, but urine culture showed multiple Gram positive rods. Chest radiogram revealed large nodular densities in the apex of the left lung and pleural effusion as shown Fig. 1 and 2. The patient was, therefore, suspected pulmonary carcinoma and was admitted to our hospital. Laboratory findings on admission, as shown in Table 1, indicated that the erythrocyte sedimentation rate was accelerated, multiple Gram positive rods were found by urine culture and the cytology of sputum was in class II to III. As illustrated in Fig. 1 and 2, an abnormal massive shadow having a rather sharply marginated contour but no notching nor calcification was found in the chest radiogram. Differential diagnosis of neoplasm and granuloma was clinically discussed between radiologists and surgeons, but the conclusion was not obtained on this case. Surgeons suspected the possibility of pulmonary carcinoma and we insisted that the chemotherapy is the first choice. Therefore, chemotherapy with 10 g of fluorouracil (5-FU), 4 g of Endoxan, 40 mg of Mitomycin C and 10 mg of Toyomycin (FAMT) was carried out.

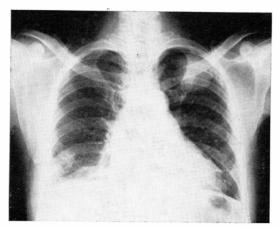


Fig. 1. Chest X-ray film in February 1970, 10 days after the admission showing the large nodular densities with 5cm in diameter in the apex of the left lung and slight pleural effusion.

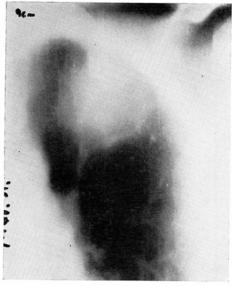


Fig. 2. Tomography at the same period as Fig. 1 (9cm).

Table 1

1970- 2	NORMAL BLOOD CHEMISTRY
	ESR 98/1 h., 105/2 hrs.
	WBC 6700
	URINAL CULTURE: G (+) ROD MULTIPLE
	Wa R (-), Tb (-)
	CYTOLOGY OF SPUTUM: CLASS II-III
1972-10	NORMAL BLOOD CHEMISTRY
	URINALYSES: Protein (+), Glucose (-)
	RBC $2-3/f$, WBC $1-2/f$.
	ESR 45/h., 76/2 hrs.
	WBC 9200
	SPUTUM: Histiocytes, Monocytes
	CRP(++)
1973- 1	BLOOD CHEMISTRY: UN 80, GOT 204, GPT 186,
	LDH 750, Al-P 14.
	ESR 44/1 h., 87/2 hrs.
	WBC 13,900-26,200
	URINALYSIS: Protein $(++)$, Glucose $(-)$,
	UROBILINOGEN $(++)$
	RBC multiple, WBC $(-)$
	CYTOLOGY OF ASCITES: CLASS II, Rivalta (-)
	RA (+), CRP (++), ASLO 80
	1

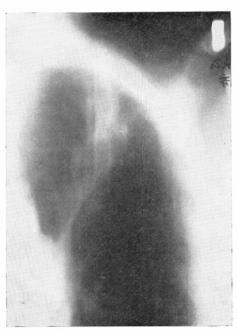


Fig. 3. Tomography following FAMT chemotherapy of about 2 months duration.

昭和50年12月25日 1095—(27)

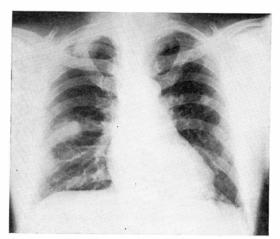


Fig. 4. Chest X-ray film in October 1972 showing nodular densities in the apex of the right lung and middle lung field. Mottling shadow and scar in the apex of the left lung are seen.

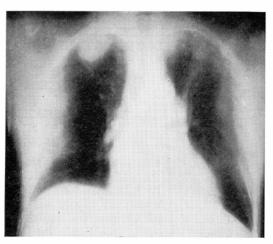


Fig. 5. Tomography at the same period as Fig. 4 (9cm distal to the back) showing shadow in the apex of right lung. Scar and mottling shadows in the apex of the left lung.

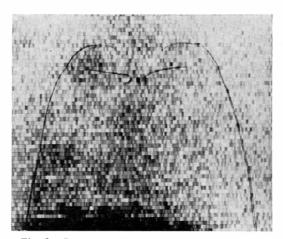


Fig. 6. Lung scanning was performed with the rectilinear scanner after the injection of 2mCi of 67Ga-citrate into the antecubital vein. Max counting rate was 100 counts per second, intensity 80, time constant 2 seconds, line spacing 4 mm and speed 60cm per minute.

Scintigram shows positive uptakes of radioisotope in the right apex and right middle lung.

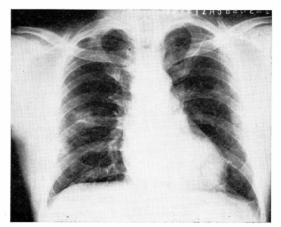


Fig. 7. Chest X-ray film following the 2nd FAMT chemotherapy showing only scars in the right apex, right middle lung and the left apex.

Following the above described treatment the pulmonary nodular densities were disappeared after about 2 months as shown in Fig. 3 and the patient was discharged.

In October 1972, the patient again complained of backache. Chest radiogram revealed multiple nodular shadows as shown in Fig. 4 and 5. ⁶⁷Ga-citrate scanning revealed increased uptakes in the areas corresponding to the shadow as shown in Fig. 6. Laboratory findings at this stage is also indicated in



Fig. 8. Histology of the lung showing scarred granulomatous fibrosis with histiocytes containing hemosiderin pigment or lipid in the bilateral lungs by Elastical-Masson Goldner modification stain.

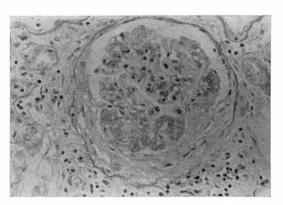


Fig. 9. Glomerular histology suggestive of severe focal glomerulonephritis. The portion of the tuft to the bottom shows increased cellularity with some breaking up of noclei, whereas the top of the glomerulus is spared.

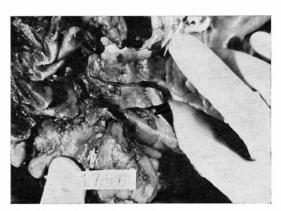


Fig. 10. Advanced necrotizing and phlegmonous colitis resulted probably from thrombosis of the inferior mesenteric vein. Arrow displays the thrombosis of the inferior mesenteric vein.

Table 1. FAMT therapy was then performed almost the same amounts as previously treated. It was also effective to the pulmonary lesions as shown in Fig. 7. However, the patient gradually became anorexia, and was readmitted in January 1973. Laboratory findings on admission were shown in Table 1. He showed the symptoms of stomatitis, conjunctivitis, blood stool with diarrhea, followed by ileus and then peritonitis. Ascites was repeatedly aspirated, he lost consciousness and died on January 24, 1973.

Autopsy findings were as follows: 1) several scars of granulomatous fibrosis with histiocytes containing hemosiderin pigment or lipid by Elastical-Masson Goldner modification stain in the bilateral lungs (Fig. 8) 2) several focal glomerulonephritis (Fig. 9) and 3) advanced necrotizing and phlegmonous colitis resulted probably from thrombosis of the inferior mesenteric vein (Fig. 10).

From above findings, the final pathological diagnosis was made to be Wegener's granulomatosis of the lung with focal glomerulonephritis.

Discussion

In this case, the large nodular densities appeared intermittently in the both lung fields. ⁶⁷Ga-citrate scanning revealed positive images in the same region. Cytology of sputum showed histiocytes. Leucocytosis and proteinuria was also observed (Table 1). Initially, differential diagnosis of pulmonary carcinoma, metastatic pulmonary carcinoma and granuloma was discussed from the above described clinical data and history. Thereafter, the postmortem examination revealed the scarred granulomatous lesions in the bilateral lungs and severe focal glomerulonephritis. Therefore, the final diagnosis was Wegener's granulomatosis.

As observed in this case, no clinical symptom in the upper respiratory tract was noted and therefore, a suspicion remained on the diagnosis of Wegener's granulomatosis. However, Sharnoff and Schneider reported a case of Wegener's granulomatosis of the lung with kidney involvement¹⁰. Furthermore, according to the recent report of Gonzalez and Van Ordstrand, the upper respiratory involvement was evidenced only in 6 out of 11 cases with Wegener's granulomatosis⁶.

⁶⁷Ga-citrate was known to be uptaken not only in neoplastic disease⁹⁾, but also in sarcoidosis²⁾⁷, pulmonary abscess, osteomyelitis, chronic pneumonia, pulmonary tuberculosis²⁾, pulmonary silicosis¹¹⁾ and pneumoconiosis¹⁸⁾. Therefore it seemed possible that the pulmonary lesions of Wegener's granulomatosis would also show positive scan. The results supported this expectation.

References

- Carrington, C.B. and Leibow, A.A.: Limited forms of angitis and granulomatosis of Wegener's type. Amer. J. Med. 41 (1966), 497-527.
- Dige-Petersen, H., Heckscher, T. and Hertz, M.: ⁶⁷Ga scintigraphy in nonmalignant lung diseases. Scandinav. J. Resp. Dis. 53 (1972), 314–319.
- 3) Felson, B. and Braunstain, H.: Noninfectious necrotizing granulomatosis. Radiology. 70 (1958), 326-333.
- 4) Fraser, R.G. and Pare, J.A.P.: Diagnosis of disease of the chest. W.B. Saunders, Philadelphia (1970), p. 874.
- 5) Garland, L.H. and Sisson, M.A.: Roentgenfindings in the collagen diseases. Radiology 71 (1954), 581-598.
- Gonzalez, L. and Ordstrand, H.S.V.: Wegener's granulomatosis review of 11 cases. Radiology 107 (1973), 295–300.
- Machida, K., Yasukochi, H., Tada, S., Kimura, F. and Ohshima, M.: ⁶⁷Ga-citrate scanning of sarcoidosis. Diagnosis of malignant tumors by gallium scanning (Yasukochi, H. edit.), Igakutosho-Shuppan, Tokyo (1973), 71–73.
- 8) Roghair, G.D. and Ross, P.M.B.: Wegener's granulomatosis. Brit. J. Radiol. 43 (1970), 216-217.
- 9) Schoot, J.B., Groen, A.S. and Jong, J.: Gallium-67 scintigraphy in lung diseases. Thorax 27 (1972), 543-546.
- 10) Scharnoff, J.G. and Schneider, L.: Wegener's granulomatosis. Amer. Rev. Resp. Dis. 86 (1962), 553-556.
- Shida, T., Chiyotani, K., Mishina, R., Saito, K., Kozuka, N., Yasukochi, H. and Tada, S.: Gallium schintigram of pulmonary silicosis. Diagnosis of malignant tumors by gallium scanning (Yasukochi, H. edit.), Igakutosho-Shuppan, Tokyo (1973), 62–70.
- Wegener, F.: Über generalisierte, septische Gefässerkrankungen. Verh. Deutsch. Gas. Path. 29 (1936), 202-210.
- 13) Winsor, D.W., Grebe, S.F., Sargent, N. and Siemsen, J.K.: Pulmonary concentration of ⁶¹Ga in pneumoconiosis. J. Nucl. Med. 14 (1973), 466.