# CARCINOSARCOMA OF THE OESOPHAGUS

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Carcinosarcoma of the oesophagus is a rare condition, for there appear to be but 30 cases recorded in the literature.

In 1949 Stout, Humphreys and Rottenberg<sup>1</sup> reviewed 20 reported cases when reporting their own. Six years later King and Koerner<sup>2</sup> added a case and referred to 6 further

cases, published in the interim. Since then Poppens, Nicolas and Szanto's case<sup>3</sup> is the only one listed.<sup>9</sup>

Sarcoma of any type is a rarity in the oesophagus for, in the course of making 12,000 barium-meal examinations in White and non-White patients, this is the first case I have

encountered, while carcinoma of the oesophagus is relatively common in non-Whites in South Africa.

### CASE REPORT

A Coloured farm labourer aged 45 years, from a fruit district in the Western Cape Province, was referred to the Karl Bremer Hospital, Bellville, early in June 1958 for investigation of difficulty with swallowing and a sensation of obstruction in the oesophagus in the middle of the chest of one month's duration. Regurgitation of food was experienced for the same period, during which time the difficulty in swallowing solids became worse and semi-fluid food only could be taken sparingly, with consequent rapid loss of weight. A cough and hoarseness were troublesome.

### Examination

The patient was emaciated. No anaemia, jaundice or cyanosis was noted. The cardiovascular, genito-urinary and central nervous

systems revealed no abnormality.

Respiratory system. A laryngo-tracheo-bronchitis was present. X-ray changes were apparent in both lung fields, mainly in the mid-zones and bases, suggestive of a spill-and-aspiration phenomenon with increased broncho-vascular striation, as is commonly seen with obstructive oesophageal lesions. An oval softtissue mass in the long axis of the mediastinum was visible posterior to the tracheal bifurcation, with a fluid level in a dilated upper third of the oesophagus on a level with the aortic arch.

# Barium Meal

There was a high-grade obstruction in the upper third of the The upper edge was at the level of the aortic arch, anterior to D5. An oval mass extended downwards for 9 cm.

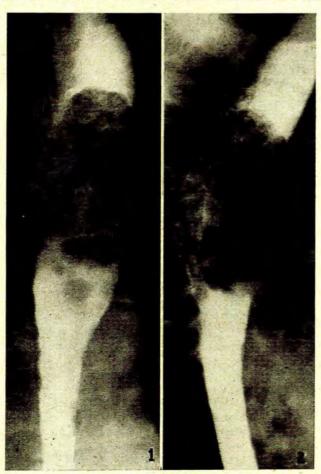


Fig. 1. Barium meal, postero-anterior position.

Fig. 2. Barium meal, oblique position.

from here and expanded the lumen of the oesophagus to a width of 6 cm. The oesophagus above the mass was dilated to about twice the average normal diameter. A cupola effect at the upper and lower margins of the mass was seen in the erect and Trendelenburg positions. The barium trickled slowly past the mass, showing a coarse net-like thin layer of barium smeared on its surface but no mucosal pattern. (See Figs. 1 and 2.) The rest of the upper alimentary tract was normal. The appearance was unlike a carcinoma and suggested a benign lesion such as a leiomyoma but the fairly rapid development of symptoms favoured sarcomatous change.

Biopsy was performed on 7 June 1958 and the following report from the Department of Histo-pathology (Prof. H. W. Weber) was furnished: 'The specimen consists of 4 small pieces of greyish tissue. The histology is that of a malignant anaplastic tumour consisting of spindle-shaped and round cells with many mitoses. I cannot decide with confidence whether it is a sarcoma or a carcinoma but a spindle-celled sarcoma seems to be more probable."

At operation, on 17 June, there was no sign of malignant spread to the mediastinum. The tumour was resected with the lower two-thirds of the oesophagus. The stomach was mobilized and anastomosed to the upper third of the oesophagus, and a pyloroplasty was carried out.

Post-operatively the patient progressed favourably for 5 days and then developed fever, delirium and pulmonary changes. The anastomosis showed no abnormality or leak on radiological investigation. The patient's condition deteriorated and he died

on the 12th post-operative day.

The pathological report (20 June) on the specimen stated that there was a large greyish tumour 10 cm. in diameter, which appeared to be a carcinosarcoma consisting of a squamous-cell carcinoma and a spindle-cell sarcoma filling the oesophagus.

Of special interest was the X-ray appearance in this case, which suggested the intramural, extramucosal origin of the tumour by 4 features, viz. (a) the cupola effect with sharply defined margins above and below, (b) distension of the oesophagus with absence of mucosal pattern, (c) smear effect resulting in a web-like pattern as barium passed over the tumour mass, and (d) constancy of shape on respiration unlike a cystic tumour.

The short duration of the symptoms indicated a malignant process but the absence of ulceration and irregularity of contour

were quite unlike a carcinoma.

# DISCUSSION

The illustrations of reported cases reveal that it is impossible to make the diagnosis on the X-ray signs alone, and that myomata<sup>5-7</sup> and other sarcoma types, e.g. melanosarcoma,<sup>4</sup> and leiomyosarcoma may all present as extramucosal lesions,8 exhibiting sharply defined rounded defects, some with ulceration. With a short clinical history the diagnosis of sarcoma is a reasonable certainty, for it is very unlike carcinoma.

# **OPSOMMING**

'n Geval van karsino-sarkoom van die slukderm by 'n nie-Blanke pasiënt word beskryf. Die röntgenologiese kenmerke met 'n barium sluk is 'n skerp begrensde tumor wat die slukderm uitsit met koepel fatsoen bo en onder, 'n smeereffek en spinnerak patroon waar die barium daaroor loop, sonder enige slymvliespatroon of ulserasie en 'n kort geskiedenis van slukbesware wat spoedig toeneem.

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### REFERENCES

- 1. Stout, A. P., Humphreys, G. H. and Rottenberg, L. A. (1949): Amer, J.
- Radiol., 61, 461.
- 2. King, H. A. and Koerrer, T. A. (1957); Surgery, 42, 389. 3. Poppens, A. D., Nicolas, A. and Szanto, P. B. (1957): Quart. Bull. Northw.

- Burnett, J. M. and Elmer, St. J. (1951): Radiology, 57, 868. 5. Zaslow, J. and Krasnoff, S. O. (1953); J. Thorac, Surg., 26, 83.
- 6. Rose, J. D. (1937); Brit. J. Surg., 24, 297.
- 7. Golden, T. and Stout, A. P. (1941): Surg. Gynec. Obstet., 73, 784. 8. Schatzki and Howes, L. (1942): Amer. J. Radiol., 48, 1.
- 9. Current List of Medical Literature (1958), vol. 34. Washington, D.C.:
  - U.S. Department of Health, Education and Welfare.