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Bilateral parotid secondaries from primary bronchial carcinoid tumour

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Abstract

Three years following a right lower lobectomy for primary carcinoid tumour of the bronchus, a patient presented with bilateral parotid masses. On investigation, both these were shown to be carcinoid tumours, consistent with metastatic spread from the primary bronchial tumour. This unique case is presented together with a discussion of the diagnostic methods employed.

Case report

In 1984, a 64-year-old lady was found on routine chest X-ray, to have a 3 cm opacity in her right lower lobe. Investigation including rigid bronchoscopy, computerized tomography of the chest, liver and bone scans excluded any further disease and she duly underwent a lobectomy. Histology showed features typical of carcinoid tumour.

Three years later, the patient was being followed up in an ENT clinic for chronic otitis media, when a 3 cm left parotid swelling was discovered. Fine-needle aspiration and computerized tomographic (CT) scan were arranged, at which time she was found to have a similar mass on the right side. This too was aspirated.

The scan showed bilateral irregular deposits in the superficial parts of both parotid glands, extending deeply on the left (Fig. 1). Aspirates from each side showed features highly diagnostic of carcinoid tumour (Fig. 2), as described below. The microscopic appearances were identical with those of the earlier lung primary. Twenty four hour urine 5-hydroxyindoleacetic acid (5-HIAA) levels were normal.

Following conventional radiotherapy (4,448 cGy) both lesions have undergone clinical and radiological regression. There has been no recurrence at one year.

Discussion

Carcinoid tumours are thought to arise from endocrine cells, particularly of appendix, rectum, small intestine, and lung (De Lellis et al., 1984). The tumours are classified according to their embryological origin: foregut, midgut and hindgut, with the category being extended more recently to cover extraintestinal neoplasms (Williams and Sandier, 1963). As part of the group of tumours known as APUDomas, they have the potential to secrete various physiologically-active peptides, especially serotonin.

Bronchial carcinoid tumours are particularly slow-growing and thus resection generally leads to an excellent prognosis (Martensson, H. et al., 1987). Malignancy is determined not by histological features, but by the appearance of secondaries which may occur up to ten years after successful treatment of the primary lesion (Bensch et al., 1965). Hence the delay of four years in this instance is by no means unusual.

Diagnosis in this case demonstrated the usefulness of fine needle aspiration biopsy (FNAB), which is used routinely in our department for parotid swellings. Of the last 32 parotid FNAB’s performed, 19 (59 per cent) were diagnostic with a sensitivity of 78 per cent, specificity of 93 per cent, and complete concordance in 74 per cent of patients in whom paraffin section material was later obtained. These figures are similar to those in the larger reported series (Eneroth et al., 1970; Berg et al., 1986; Rodriguez et al., 1989). In addition to reliability, the technique is cheap, safe (Engzell et al., 1971), simple and avoids the risks of an operation, such as possible facial nerve damage in the reported case. The usefulness of CT scanning for parotid masses is well-established and complements the fine-needle findings (Berg et al., 1986; Cameron and Mann, 1989).

The electron microscopic appearances of the cytology obtained were highly specific for carcinoid tumour (Bensch et al., 1965), showing uniform 2000 Angstrom, highly electron-opaque granules, surrounded by a unit membrane, and with otherwise scanty organelles (Fig. 2). This was identical with the original bronchial specimen. Together with the clinical picture, this is convincing evidence that the case illustrated is an example of bilateral parotid secondaries from a bronchial carcinoid, a previously unreported event.

An alternative view would be that this represents multicentric disease. In Godwin’s series, only 6 of 2837 reported carcinoids were multicentric, and these were confined to the small intestine. Hence, this appears less likely (Godwin, 1975).
Transmission electron micrograph from parotid aspirate, showing cellular features characteristic of carcinoid: highly electron-opaque core granules (arrowed) surrounded by a unit membrane and sparse organelles.

Metastases of bronchial carcinoids to the head and neck are very rare. There are only two cases documented in the literature: one to the mandible (Haggerty, 1987), and one to the soft palate (Mintz and Radecki, 1988). The parotid gland is a very uncommon site for metastases from a primary tumour that is not a squamous cell carcinoma (Bergersen et al., 1987).

In view of the patient's age, the hazardous nature of the surgery that would be necessary to remove these tumours, and the indolent nature of the disease, it was decided to refer her for a course of radiotherapy. Although carcinoid tumours are not thought to be very radiosensitive (De Lellis et al., 1984), a recent study (Abrams, 1987) reported an objective response to radiotherapy in 54 per cent of cases and complete regression in 23 per cent of cases. The response of the tumours in the above case is consistent with these findings.

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