Paracardiac lipomatosi in exogenous Cushing's syndrome

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Paracardiac lipomatosi in exogenous Cushing's syndrome. Histologically proven lipomatosi presenting as paracardiac masses on the chest radiograph is described in a patient with a renal graft. This lesion was caused by the prednisone therapy and diminished after lowering the dosage.

Review of 90 renal transplant patients revealed one likely example of this syndrome and four patients had other radiographic abnormalities suggestive of intrathoracic lipomatosi.

It is pointed out that intrathoracic lipomatosi can produce a variety of abnormalities on the chest radiograph and that sometimes these changes need to be differentiated from primary tumours.

Deposition of fat in the trunk, face, and neck is a characteristic feature of Cushing's syndrome. Unusual episternal (Lucena, Bennett, and Pierre, 1966) and presacral (Sowerbutts, 1959) fat accumula- tions have also been recorded. Several recent case reports mention superior mediastinal widening due to lipomatosi in endogenous (Santini and Williams, 1971) and exogenous (Koerner and Sun, 1966; Bodman and Condemi, 1967; Price and Rigler, 1970; Teates, 1970; Fraser and Paré, 1970) Cushing's syndrome. The condition is rare and has sometimes led to surgical exploration.

In addition to mediastinal widening, prominent epicardial fat pads may be another manifestation of intrathoracic lipomatosi (Koerner and Sun, 1966; Bodman and Condemi, 1967; Price and Rigler, 1970; Teates, 1970).

The patient to be reported developed puzzling paracardiac masses whilst receiving corticosteroids after renal transplantation. Subsequent histological examination showed the lesion to be a form of drug-induced lipomatosi. At a later date one likely example of this syndrome was discovered after reviewing all the patients who had undergone renal transplantation.

CASE REPORT

L. H. M., a 51-year-old woman, underwent intermittent haemodialysis because of renal insufficiency due to hereditary nephritis. A cadaver renal transplantation was performed in August 1971. Two rejection episodes were treated by high doses of prednisone, up to 150 mg daily, for short periods. Four months after the operation a routine chest radiograph revealed bilateral paracardiac masses (Fig. 1) not present before transplantation (Fig. 2). Superior mediastinal widening and thickening of the thoracic wall were also noted.

Apart from a marked Cushingoid appearance, the physical examination was unremarkable. There was no significant increase in weight compared to the pre-transplantation period. Extensive laboratory investigations gave no indication as to the nature of the radiographic abnormalities. Finally, a left thoracotomy revealed a mass of adipose tissue (Fig. 3). Biopsy showed normal fat tissue covered by a monolayer of flat epithelial cells, presumably the parietal pleura. Over a period of several months, as the dose of prednisone was gradually decreased to 10 mg daily, the Cushingoid appearance and the paracardiac shadow diminished simultaneously.

The above findings then prompted us to examine the available material in our renal transplantation series (90 patients), since all the patients were receiving
FIG. 1. Chest radiographs taken four months after transplantation showing the paracardiac masses (arrows) in the posteroanterior (a) and lateral (b) views.
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FIG. 2. Posteroanterior (a) and lateral (b) chest radiographs before transplantation.
corticosteroid therapy. One likely case was discovered and in four patients we found radiographic abnormalities suggestive of intrathoracic lipomatosis—mediastinal widening, prominent epicardial fat pads, or both. All the patients showed a marked Cushingoid appearance.

**DISCUSSION**

Although paracardiac lipomatosis has been noted in the obese (Holt, 1947), we were unable to find any report of intrathoracic lipomatosis in Cushing's syndrome. Lipomatosis was suspected in the present case because of the associated Cushing’s syndrome, the lack of other symptoms, the symmetrical localization of the abnormal shadows on the chest radiographs, and the additional radiographic abnormalities—thickening of the thoracic wall and mediastinal widening. As mentioned in earlier reports (Bodman and Condemi, 1967; Teates, 1970), the disappearance of the abnormalities when corticosteroids are withdrawn also seems to support this diagnosis. However, the possibility of a malignancy, in a patient on immunosuppressive therapy, made further investigation desirable.

Knowledge of this benign abnormality may prevent unnecessary investigation and even surgical exploration.

**REFERENCES**


