Paracardiac lipomatosis in exogenous Cushing’s syndrome

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van de Putte, L. B. A., Wagenaar, J. P. M., and San, K. H. (1973). Thorax, 28, 653–656. Paracardiac lipomatosis in exogenous Cushing’s syndrome. Histologically proven lipomatosis 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 lipomatosis. It is pointed out that intrathoracic lipomatosis 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 accumulations have also been recorded. Several recent case reports mention superior mediastinal widening due to lipomatosis 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 lipomatosis (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 lipomatosis. 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 thoracoscopy 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 paracardiad 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.
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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


