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rial and tuberculous orchitis, syphilitic infection of the testis, sperm granuloma and testicular tumour. There were no clinical or histological features of these disorders in our case. An association between this condition and local trauma and urinary tract infection have been described; the significance of these remain problematical. The differential diagnosis from cancer is not easy, hence the importance of orchidectomy and early rapid diagnosis.

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ACTINOMYCOSIS OF THE LIVER

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Abstract. We present our first case of actinomycosis in Trinidad and we believe this is also the first reported case of hepatic actinomycosis in the West Indies

Key words: actinomycosis; liver; Trinidad

Case Report

In February 1984 a 35-year-old Trinidadian of Spanish descent presented to the Port-of-Spain General Hospital with right-sided upper abdominal pain and fever. His symptoms started 4 weeks before and became progressively worse. He was a labourer by occupation and was involved in raising common fowl.

On admission, he had a temperature of 38°C. Clinical examination was essentially normal except for tender hepatomegaly extending 6 cm below the right costal margin. He had no previous abdominal surgery. Laboratory investigations were all normal except for an elevated white blood cell count of 12.000 with 70% polymorphonuclears. Blood cultures were negative. Repeated stool examinations and serologic tests for amoebiasis were negative. Roentgenograms of the chest and abdomen were normal.

At exploratory laparotomy, all viscera and organs were normal except for a hard, nodular 12×8 cm size mass in the right lobe of the liver; it was adherent to the anterior abdominal wall and diaphragm. Biopsy of this hepatic lesion showed classical actinomycotic granules (Figure 1). Aerobic and anaerobic cultures of the

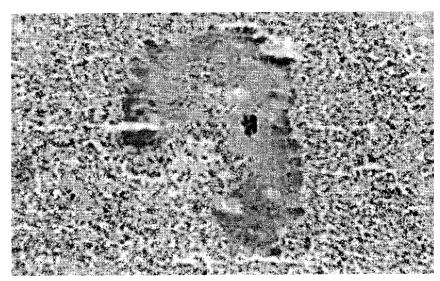


Figure 1a. Liver biopsy showing actinomycotic granule and dense polymorphonuclear infiltrate (H & Eimes 100).

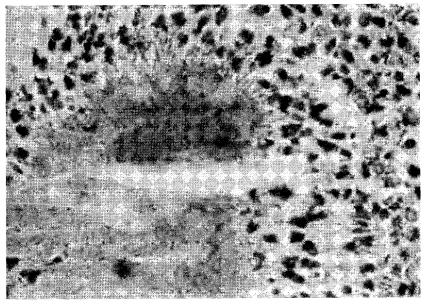


Figure 1b. Higher magnification showing the details of actinomycotic granule (H & E \times 200).

tissue did not yield growth. Postoperatively, the patient was treated with crystalline penicillin, 5 million units intravenously, 4 times a day for one week.

Two months later the patient was readmitted because of mild persistent pyrexia. An ultrasound scan demonstrated sub-hepatic fluid collection. At re-exploration, a serosanguineous sub-hepatic accumulation of fluid (400 ml) was drained and a shrunken hepatic nodule at the previous biopsy site was excised. Cultures

of this fluid were sterile while histology of the hepatic nodule showed fibrosis with no evidence of residual actinomycosis. The patient recovered uneventfully and six months later remained afebrile and free of symptoms.

Discussion

Actinomycosis is usually an endogenous infection in humans and presents in cervicofacial, thoracic and abdominal regions. Abdominal actinomycosis usually begins in the intestine as a complication of inflammatory disease or surgery [1]. The liver is the primary site of infection in 15% of patients with abdominal actinomycosis [2]. Hepatic actinomycosis may develop after appendicitis or surgery for non-inflammatory diseases [3]; it is probably conveyed to the liver by the portal vein. However, the pathogenesis remains unclear [4].

The most common hepatic lesion is a solitary abscess, though multiple abscesses may occur resulting in a honey-comb appearance. These abscesses are usually confined to the liver, but may extend to involve the adjacent organs [5]. Pre-operative diagnosis of hepatic actinomycosis is difficult. The commonest clinical findings are fever and tender hepatomegaly [6]. Serologic techniques are not conclusive in the diagnosis of actinomycosis; but crossed immuno-electrophoresis may provide greater sensitivity and specificity [7]. The specific diagnosis however, can only be made on histology or anaerobic cultures of the material.

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