

Isolation of *Klebsiella pneumoniae* from an hepatic inflammatory pseudotumour

DICKRAN A MALATJALIAN, MD, JOHN MORRIS, MD, AUDLEY BODURTHA, MD

DA MALATJALIAN, J MORRIS, A BODURTHA. Isolation of *Klebsiella pneumoniae* from an hepatic inflammatory pseudotumour. *Can J Gastroenterol* 1992;6(2):84-86. Inflammatory pseudotumours of the liver are rare, solid mass lesions (often mistaken for malignancy) and although inflammatory in nature, are obscure in etiology and pathogenesis. An hepatic inflammatory pseudotumour is presented in which cultures obtained from the lesion intraoperatively grew *Klebsiella pneumoniae*. This is the first case of an hepatic inflammatory pseudotumour in which a microorganism has been identified. The isolation of *K pneumoniae* suggests that inflammatory pseudotumours may arise from a low grade infection with chronic inflammatory tissue response.

Key Words: Etiology, Hepatic tumours, Inflammatory pseudotumours, *Klebsiella pneumoniae*

Isolement de *Klebsiella pneumoniae* à partir d'une pseudotumeur inflammatoire hépatique

RESUME: Les pseudotumeurs inflammatoires du foie sont des masses solides rares, qu'on croit souvent malignes et dont on méconnaît les causes ou la pathogénèse. On rapporte ici le cas d'une pseudotumeur inflammatoire hépatique, où la culture de prélèvements peropératoires a mis en évidence *Klebsiella pneumoniae*. Il s'agit du premier cas de pseudotumeur inflammatoire hépatique qui ait donné lieu à l'isolement d'un germe pathogène. On conclut que les pseudotumeurs inflammatoires pourraient résulter d'une subinfection accompagnée d'une réaction inflammatoire chronique des tissus.

Departments of Pathology and Surgery, Victoria General Hospital and Dalhousie University, Halifax, Nova Scotia

Correspondence and reprints: Dr DA Malatjalian, Department of Pathology, Victoria General Hospital, Halifax, Nova Scotia B3H 2Y9

This study was first presented at the Regional Meeting of Canadian Association of Gastroenterology, June 1990, Halifax, Nova Scotia

Received for publication September 23, 1991. Accepted January 10, 1992

INFLAMMATORY PSEUDOTUMOURS (IPT) are rare solid inflammatory mass lesions which have been described in several organs and given different names, including plasma cell granuloma (1). Hepatic IPT are very rare with only 23 case reports in English language literature (2-9). The etiology and pathogenesis of IPT has not been determined. Janigan and Marrie (10) isolated *Coxiella burnetii* from an IPT of the lung. However, until the case here presented, no microorganisms have been identified from hepatic IPT. The authors present the case of a patient with hepatic IPT from which *Klebsiella pneumoniae* was isolated.

CASE PRESENTATION

A 42-year-old male coal miner with chronic obstructive lung disease and controlled hypertension presented to his family physician with left chest pain and fever of several days duration. Clinical and radiological findings confirmed bilateral lower lobe pneumonia. Sputum cultures grew *K pneumoniae* and blood cultures were negative on three occasions. Despite two weeks of treatment with tetracycline, clinical improvement was slow and the patient

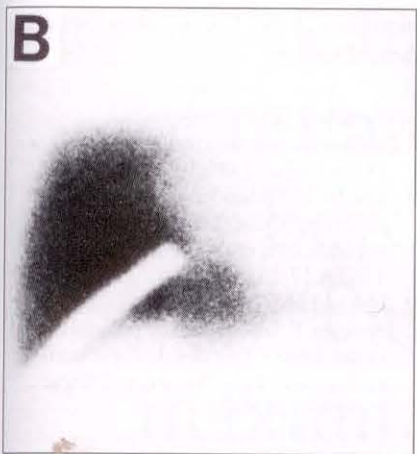
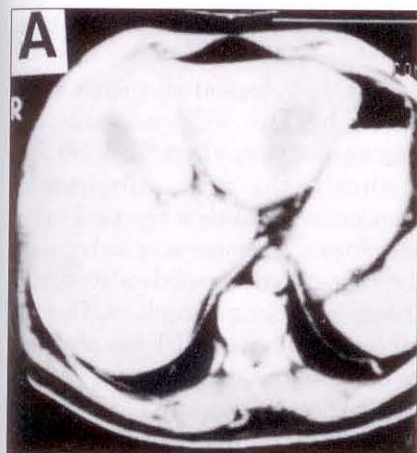


Figure 1) A Ultrasound showing hypochoic mass. B Technetium scan demonstrating a cold solitary lesion occupying most of the left lobe

was referred to hospital for further evaluation. There was no history of diabetes and physical examination on admission was normal.

The following were the abnormal laboratory findings, with the normal ranges (where appropriate) shown between parentheses: white blood cells $11.9 \times 10^9/L$; erythrocyte sedimentation rate 115 mm/h; alkaline phosphatase 562 U/L (30 to 104); alanine aminotransferase 97 U/L (1 to 41); aspartate aminotransferase 48 U/L (8 to 29); lactate dehydrogenase 335 U/L (117 to 259); gamma glutamyltransferase 585 U/L (0 to 40). The rest of the routine biochemical and hematological profiles were normal.

Chest x-ray films demonstrated bilateral lower lobe segmental atelectasis. Ultrasound of abdomen revealed a hypochoic mass in the left lobe of



Figure 2) Cut surface of left hepatic lobectomy specimen showing a well circumscribed 6 x 5 x 3 cm yellowish mass surrounded by partially fixed normal liver parenchyma

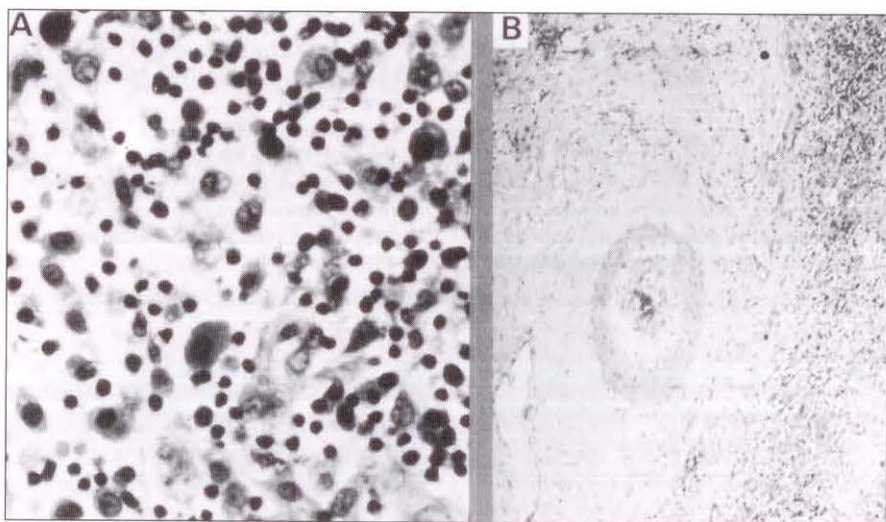


Figure 3) A Inflammatory pseudotumour composed of a mixture of lymphocytes, plasma cells, macrophages, fibroblasts and capillaries. Hematoxylin and eosin. B Sclerosing phlebitis with partial obliteration of a portal vein. Hematoxylin and eosin.

liver. A computed tomography scan demonstrated the liver's left hepatic lobe to be enhancing along the periphery but not in the centre (Figure 1A). A technetium scan (Figure 1B) identified the lesion as 'cold' while the gallium scan depicted it as 'hot'. Selective left hepatic angiography suggested increased vascularity of the rim with relative hypovascular centre. The radiological findings were considered inconclusive. As the lesion was suspected to be malignant, a left hepatic lobectomy was

performed. Intraoperative biopsy for cultures taken from the lesion grew *K pneumoniae*. The patient had an uneventful recovery and is alive seven years later.

The lesion was approximately 6 x 5 x 3 cm, well demarcated, yellowish, rubbery and contained small areas of necrosis and hemorrhage (Figure 2). The surrounding liver parenchyma was normal. Histologically, the lesion was composed of lymphocytes, plasma cells, foamy histiocytes, hemosiderin-laden

macrophages and fibroblasts with collagen (Figure 3). Scattered foci of necrosis were noted but there was no cavitating abscess. Some veins within the lesion showed sclerosing phlebitis (Figure 3B).

DISCUSSION

The etiology of IPT is unknown. There is, however, strong clinical circumstantial evidence indicating it is infectious in origin. Patients often present with fever, weight loss, leukocytosis, elevated erythrocyte sedimentation rate and positive C-reactive protein (7,9). Microorganisms from the gut may be responsible for hepatic IPT (9) but

until the case under discussion, no microorganisms have been isolated from an hepatic IPT. The mechanism of IPT formation remains obscure.

Histologically, the lesion consists of a chronic inflammatory process in which lymphocytes, plasma cells, macrophages and fibroblasts abound while suppuration with liquefactive necrosis is minimal or absent. The isolation of *K pneumoniae* from the patient's sputum and subsequently from the hepatic IPT suggests a low grade infection with chronic inflammatory tissue response predominating. The concept of chronic low grade infection may explain why previous attempts to isolate microor-

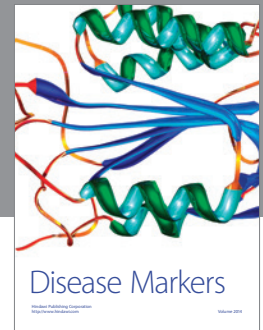
ganisms from hepatic IPT have been unsuccessful.

The histological distinction between hepatic IPT and resolving hepatic abscess may be difficult (9). Accordingly, the term 'inflammatory pseudotumour' likely is best used in the radiological and macroscopical context for space-occupying solid inflammatory lesions mimicking a neoplasm. This latter aspect of hepatic IPT has often led to the diagnosis of malignancy and extensive hepatic resection (7,9). Needle biopsy under ultrasonography guidance would be the procedure of choice for identifying the lesion, thus avoiding radical surgery.

ACKNOWLEDGEMENTS: The authors are grateful to Ms P Morgan for typing the manuscript.

REFERENCES

1. Bahadori M, Liebow AA. Plasma cell granuloma of lung. *Cancer* 1973;31:191-208.
2. Pack GT, Baker HW. Total right hepatic lobectomy: Report of a case. *Ann Surg* 1953;138:253-8.
3. Hertzler NR, Hawk WA, Hermann RE. Inflammatory lesions of the liver which simulate tumor: Report of two cases in children. *Surgery* 1971;69:839-46.
4. Someren A. 'Inflammatory pseudotumor' of the liver with occlusive phlebitis: Report of a case in a child and review of the literature. *Am J Clin Pathol* 1978;69:176-81.
5. Chen KTK. Inflammatory pseudotumor of the liver. *Hum Pathol* 1984;15:694-6.
6. Heneghan MA, Kaplan CG, Priebe CJ, Partin J, Partin JS. Inflammatory pseudotumor of the liver: A rare cause of obstructive jaundice and portal hypertension in a child. *Pediatr Radiol* 1984;14:433-5.
7. Anthony PP, Telesinghe PU. Inflammatory pseudotumor of the liver. *J Clin Pathol* 1986;39:761-8.
8. Kessler E, Turani H, Kayser S, Bar-Ziv J, Chaimoff CH. Inflammatory pseudotumor of the liver. *Liver* 1988;8:17-23.
9. Horiuchi R, Uchida T, Kojima T, Shikata T. Inflammatory pseudotumor of the liver. Clinicopathologic study and review of the literature. *Cancer* 1990;65:1583-90.
10. Janigan DT, Marrie TJ. An inflammatory pseudotumor of the lung in Q-fever pneumonia. *N Engl J Med* 1983;308:86-8.



Hindawi
Submit your manuscripts at
<http://www.hindawi.com>

