Granular cell myoblastoma of the biliary tree

Ann Savage
M.B., B.S.

P. Devitt
M.B., Ch.B.

Departments of Pathology and Surgery, Southmead Hospital, Westbury-on-Trym, Bristol

Summary

A granular cell myoblastoma of the common bile duct is described which caused episodes of jaundice in a young woman. Histological examination demonstrated a close association of the tumour with nerve fibres, tending to support the theory of origin from Schwann cells. The lesion is benign and after excision carries an excellent prognosis.

Introduction

Granular cell myoblastoma is an uncommon benign lesion of uncertain histogenesis. It was first described in 1926 by Abrikossof who considered that the cells composing the tumour resembled embryonal skeletal muscle. This is no longer generally accepted but the name is still widely used. The lesion occurs most frequently in the oral cavity and skin, but has been described in a wide variety of other situations (Paskin, Hull and Cookson, 1972). Its occurrence in the biliary tree is rare and although fifteen previous cases have been described, there is none in the British literature.

Case history

A female Caucasian was first seen in 1959 at the age of 16 years with jaundice. This was thought to be infective in origin, though the course of the illness was noted to be rather prolonged. Liver function tests at that time were: alkaline phosphatase 89 i.u./l (normal 20–110 i.u./l), SGPT (alanine aminotransferase) 190 i.u./l (normal range less than 35 i.u./l) and bilirubin 22 μmol/l (normal range less than 20 μmol/l).

Ten years later she again became jaundiced, giving a 3-day history of general malaise, pruritus, abdominal pain and dark urine. Liver function tests were: alkaline phosphatase 138 i.u./l, SGPT 104 i.u./l and bilirubin 40 μmol/l. An Australia antigen test was negative. Again the course of the illness was prolonged, an obstructive aetiology considered, but her enzyme levels then started to return to normal and so this was considered unlikely. A cholecystogram, performed during this episode, failed to reveal a functioning gall bladder but this was thought to be consistent with her persistent jaundice.

In April 1975, at the age of 30 years she again became jaundiced, this being associated with nausea, flatulence and heartburn. She had generalized pruritus and noted her urine had turned dark. Liver function tests showed: alkaline phosphatase 338 i.u./l, SGPT 863 i.u./l and bilirubin 44 μmol/l. Again an oral cholecystogram failed to opacify the gall bladder. However i.v. cholangiogram suggested a stricture of the common hepatic duct with proximal dilatation.

At laparotomy in July 1975 a dense fibrous biliary stricture was found high in the porta hepatitis. Operative transhepatic cholangiography demonstrated gross dilatation of the right and left hepatic ducts but failed to outline the distal biliary tree. The mass of fibrous tissue had also occluded the cystic duct with consequent gall bladder distension. Cholecystectomy was performed, the fibrous stricture excised and cholehepatico-jejunostomy carried out with a Roux loop. A wedge biopsy of liver was also taken.

The patient made a satisfactory postoperative recovery and her liver function tests rapidly returned to normal.

Pathological findings

The gall bladder measured 80 × 40 × 30 mm and had a thickened wall up to 5 mm but contained no stones. Histologically there was mild chronic cholecystitis but no evidence of neoplasm. Sections of the wedge biopsy of liver showed only slight portal inflammation without cholestasis.

The material removed from around the common bile duct consisted of an irregular mass of tissue 20 × 25 × 30 mm., with 30 mm of opened bile duct attached to one side. Sections showed that the common bile duct was eccentrically surrounded by fibrous tissue which was diffusely infiltrated by large polygonal or elongated cells with small dark nuclei and abundant granular eosinophilic cytoplasm (Figs. 1 and 2). The granules were markedly positive with the periodic acid-Schiff reaction. Numerous nerve bundles of varying size were present in the excised tissue; the granular cells closely encircled them and in some cases could be seen within the bundle.
FIG. 1. The tumour is composed of large cells with small dark nuclei (H. and E. × 438).

FIG. 2. High power view of tumour showing granular cytoplasm (H. and E. × 875).
Case reports

FIG. 3. Tumour cells are within and closely surrounding a nerve bundle (H. and E. ×263).

<table>
<thead>
<tr>
<th>Case</th>
<th>Author</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Race</th>
<th>Situation</th>
<th>Symptoms +</th>
<th>Treatment and Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Coggins 1952</td>
<td>25</td>
<td>F</td>
<td>B</td>
<td>CBD</td>
<td>Liver failure, Congestive cardiac failure</td>
<td>Diagnosed at post-mortem</td>
</tr>
<tr>
<td>2</td>
<td>Fialho 1952 and Hilarco, 1957</td>
<td>21</td>
<td>F</td>
<td>B</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Excised. Well at follow-up</td>
</tr>
<tr>
<td>3</td>
<td>Duncan 1957</td>
<td>30</td>
<td>F</td>
<td>B</td>
<td>CBD</td>
<td>Obstructive jaundice</td>
<td>Excised. Discharged well</td>
</tr>
<tr>
<td>4</td>
<td>Serpe et al., 1960</td>
<td>34</td>
<td>F</td>
<td>B</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Excised. Well at follow-up</td>
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<tr>
<td>5</td>
<td>Goldman et al., 1967</td>
<td>15</td>
<td>F</td>
<td>B</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Excised</td>
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<td>6</td>
<td>Mackay et al., 1968</td>
<td>45</td>
<td>F</td>
<td>NS</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Excised. Well 9 months later</td>
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<tr>
<td>7</td>
<td>Whitmore et al., 1969</td>
<td>37</td>
<td>F</td>
<td>B</td>
<td>CBD</td>
<td>Obstructive jaundice</td>
<td>Excised. Discharged well</td>
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<tr>
<td>8</td>
<td>Whitmore et al., 1969</td>
<td>61</td>
<td>F</td>
<td>B</td>
<td>CBD</td>
<td>Incidental finding at post-mortem</td>
<td>Chronic alcoholic with chronic lung disease</td>
</tr>
<tr>
<td>9</td>
<td>Christensen 1970</td>
<td>34</td>
<td>F</td>
<td>NS</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>NS</td>
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<tr>
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<td>NS</td>
<td>Cystic duct</td>
<td>Epigastric pain</td>
<td>NS</td>
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<td>44</td>
<td>F</td>
<td>B</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
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<tr>
<td>12</td>
<td>Li Volsi et al., 1973</td>
<td>40</td>
<td>F</td>
<td>NS</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Died of myocardial infarction 4 years after excision. No evidence of tumours</td>
</tr>
<tr>
<td>13</td>
<td>Li Volsi et al., 1973</td>
<td>30</td>
<td>F</td>
<td>NS</td>
<td>CBD</td>
<td>Diabetic. Abnormal liver function tests</td>
<td>Excised. Postoperative biliary fistula. Otherwise well 1 year later. Lesion on ear lobe, also granular cell myoblastoma</td>
</tr>
<tr>
<td>14</td>
<td>Whisnant et al., 1974</td>
<td>16</td>
<td>M</td>
<td>B</td>
<td>CBD</td>
<td>Obstructive jaundice</td>
<td>Excised. Well 1 year later. Neurofibroma adjacent to tumour. Lesion on anterior abdominal wall also granular cell myoblastoma</td>
</tr>
<tr>
<td>15</td>
<td>Reul et al., 1975</td>
<td>39</td>
<td>F</td>
<td>B</td>
<td>Cystic duct</td>
<td>RUQ pain</td>
<td>Excised</td>
</tr>
</tbody>
</table>

* B, black; W, White; NS, not stated.
† CBD, common bile duct.
‡ RUQ, right upper quadrant.
itself (Fig. 3). The epithelium of the common bile duct was extensively ulcerated probably owing to trauma, but surviving islands were normal with no evidence of hyperplasia. Normal glands were present in the periductal tissue. The lesion was a characteristic granular cell myoblastoma.

Discussion
Fifteen cases of granular cell myoblastoma of the biliary tree have been reported previously (Table 1). Including the present case, seven were situated in the common bile duct and the remainder (nine) in the cystic duct. When the lesion was in the common bile duct the clinical picture was usually of obstructive jaundice, whereas with cystic duct lesions right upper quadrant (RUQ) pain was the usual symptom. Except for the case described by Coggins (1952) and one in Whitmore's series (Whitmore et al., 1969) which was diagnosed incidently at postmortem, all cases were successfully treated surgically. Of the reported cases only one was male, and the age range (excluding Whitmore's case) was 15-45 years. The race was not stated in five cases; all the remainder except the present case were Negro. This predilection for black females has been found in granular cell myoblastomas at other sites (Vance and Hudson, 1969). Two of the reported cases had granular cell myoblastomas elsewhere and in one of these (Whisnant et al., 1974) there was also a neurofibroma adjacent to the lesion in the common bile duct.

The histogenesis of granular cell myoblastoma is controversial. In his original paper Abrikossoff (1926) believed that the lesion was a neoplasm arising from myoblasts or embryonal striated muscle cells. This view was supported by the tissue culture work of Murray (1951). Azzopardi (1956) and others have proposed that the lesion is a metabolic disorder rather than a neoplasm and that the granular cells are histiocytes. Fust and Custer (1949) considered that the lesions were neoplasms arising from nerves. The neurogenic origin was supported by the histochemical and electron microscopic studies of Fisher and Wechsler (1962), who considered that the lesion arose from Schwann cells but that it was more likely to be histiocytic than neoplastic. They proposed the term 'granular cell Schwannoma', which has not however been widely accepted. Other workers have accepted a Schwann cell origin but conclude that a neoplastic nature is more likely (Sobel et al., 1971).

A number of granular cell myoblastomas of the biliary tree have been described showing a close association with nerve fibres and in the present case granular cells were present around and within nerve bundles. This feature tends to support the theory of origin from Schwann cells.

Granular cell myoblastoma is rare in the extra-hepatic biliary tree but the lesion is readily curable. It should especially be considered in the differential diagnosis of obstructive jaundice or chronic cholecystitis in young black women.

Acknowledgments
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References
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A. Savage and P. Devitt

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