Approach to Management of Indeterminate Biliary Stricture

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Abbreviations: MRCP: Magnetic Resonance Cholangiopancreatography; ERCP: Endoscopic Retrograde Cholangiopancreatography; DIA: Digital Image Technique; EUS: Endoscopic Ultrasonography; SOC: Single Operator Cholangioscopy; CT: Cross Section Imaging

Introduction

Biliary strictures are traditionally classified as ‘indeterminate’ when basic work up, including transabdominal imaging and endoscopic retrograde cholangiopancreatography (ERCP) with conventional brush cytology, is non-diagnostic for its etiology.

Table 1: Biliary strictures: etiology.

<table>
<thead>
<tr>
<th>Benign</th>
<th>Malignant</th>
</tr>
</thead>
<tbody>
<tr>
<td>Iatrogenic (cholecystectomy, liver transplant)</td>
<td>Cholangiocarcinoma</td>
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<tr>
<td>Chronic pancreatitis</td>
<td>Pancreatic cancer</td>
</tr>
<tr>
<td>Primary sclerosing cholangitis</td>
<td>Gall bladder carcinoma</td>
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<tr>
<td>IgG4-related disease (cholangiopathy, autoimmune pancreatitis)</td>
<td>Metastatic disease (compression by lymph nodes, tumor at hilum)</td>
</tr>
<tr>
<td>Infections (tuberculosis, viral, parasitic, recurrent cholangiohepatitis, AIDS)</td>
<td>Lymphoma</td>
</tr>
<tr>
<td>Cholecystolithiasis</td>
<td></td>
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<tr>
<td>Post biliary sphincterotomy</td>
<td></td>
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<tr>
<td>Trauma</td>
<td></td>
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<td>Post radiation therapy</td>
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<tr>
<td>Vasculitis, ischemic portal biliopathy</td>
<td></td>
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<tr>
<td>Sphincter Oddi dysfunction</td>
<td></td>
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<tr>
<td>Mirizzi syndrome</td>
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<tr>
<td>Extrinsic compression by nodes or vessels</td>
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</table>

Etiology of Biliary Strictures

Majority (70%-80%) of the biliary strictures are malignant, although 20%-30% of it may be benign [1]. The commonest causes of benign strictures are enlisted in Table 1.

Evaluation of Patients with Biliary Strictures

The clinical approach to the patient with indeterminate biliary stricture includes a thorough history and physical examination. By and large, strictures of the bile duct in patients with obstructive jaundice should be considered malignant unless a benign etiology is ascertained. The significance of biliary strictures without jaundice is less certain.
Biomarkers

The most common biomarkers for suspected biliary tract malignancies in clinical use are serum CA 19-9 and carcinoembryonic antigen. In patients with PSC, using cut-off value of 129 U/ml, the sensitivity and specificity of CA 19-9 for the diagnosis of cholangiocarcinoma are 79% and 98% respectively [2]. In patients without PSC, the sensitivity is only 53% with a cutoff value 100 U/ml [3]. Elevated serum carcinoembryonic adenocarcinoma has also been shown to have a sensitivity of 33-68% and specificity of 79-95% for cholangiocarcinoma.

Radiological Work-Up

Ultrasound is helpful in patients with biliary stricture by demonstrating intrahepatic biliary radical dilatation and the level of obstruction. However, the distal part of the common bile duct is not properly evaluated because of the interference of bowel gas. Moreover, it has a very low yield for actual detection of strictures on biliary ductal masses [4]. The development of multi-detector helical scanners, used in conjunction with rapid injection of contrast media, accurate depiction of extension of the tumor, locoregional lymphadenopathy and encasement of blood vessels to determine operability of the tumor can be picked up easily.

Magnetic resonance cholangiopancreatography (MRCP) has a high sensitivity for bile duct lesions and has got comparable diagnostic accuracy in comparison to ERCP. Ductal features at MRCP which may suggest malignancy include long (>10mm), asymmetrical and irregular strictures. The presence of mass lesion is highly suggestive of malignancy, especially in the hilar region. Abrupt cut-off of the CBD in contrast to a smooth tapering has traditionally been considered to be a sign of malignancy (Figure 1). The sensitivity and specificity of MRCP to differentiate malignant from benign strictures are reported to be 38%-90% and 70%-85% respectively [5]. MRCP is 88-96% accurate in predicting the extent of involvement of the bile duct in cholangiocarcinoma.

Role of Endoscopy

Endoscopic retrograde cholangiopancreatography (ERCP) remains the first line approach for tissue sampling of biliary strictures. The reported sensitivity of conventional brush cytology is 25% to 55% [6]. Different techniques have been employed to improve the sensitivity of conventional brush cytology including novel brushes, biliary stricture dilation with subsequent brushings and repeated brushings. The pluricellular nature and submucosal pattern of tumor growth in cholangiocarcinoma or extrinsic malignancy involving the bile ducts attributes to the low sensitivity of biliary brushings. Inadequate biliary cytology specimens are one of the main reasons for non-diagnostic samples. This may be overcome by the presence of an onsite cytopathologist. Several other strategies include cutting the entire brush, creation of slides by the endoscopy team and placing them in a fixative solution prior to submission to pathology are being used to overcome inadequate sampling. The fluorescein in situ hybridization analysis detects chromosomal polysomy using fluorescent probes, whereas the Digital Image Technique (DIA) quantifies nuclear DNA with special stains to assess the presence of aneuploidy.

In patients with PSC these chromosomal abnormalities can be seen without the presence of malignancy. Thus, the specificity of FISH in this setting is lower than routine cytology. Thus, FISH increases the sensitivity of brush cytology of indeterminate biliary strictures without much improvement in the specificity. FISH should be reserved for patients with high pre-test probability for malignant strictures. Using endobiliary forceps, the malignancy detection rates ranges from 44% to 89% for cholangiocarcinoma and 33% to 71% for pancreatic cancer [7]. Triple sampling with brushing, transpapilary biopsy and endoluminal FNA has shown the highest sensitivity. Risk of biliary ductal perforation after endobiliary biopsy, however, remains a concern.

Endoscopic ultrasound (EUS) has emerged as an important method for evaluating indeterminate biliary stricture. It provides an excellent alternative method for visualizing and sampling the extra-hepatic biliary tree, hilar masses, gallbladder and peri-hilar lymph nodes and vessels. Sensitivity is significantly better in distal compared to proximally located tumors.

Cholangioscopy

Endoscopic retrograde cholangiopancreatography (ERCP) remains the first line approach for tissue sampling
The technique has recently gained attention with the development of a single operator cholangioscopy (SOC) system known as the SpyGlass (Boston Scientific, Natick, MA, USA). Visually directed biopsies can be obtained using biopsy forceps (SpyBite). Overall sensitivity and specificity of SOC examination for differentiating malignant and benign ductal abnormalities have been seen to be 78% and 82% respectively, higher than the 51% and 54% of ERCP alone [8]. Among the cholangioscopic features, the presence of abnormal tumor vessels due to neovascularization within the biliary stricture is suggestive of biliary malignancy. Intraductal nodules and masses can be visualized during cholangioscopy and are indicative of malignancy (Figure 2). Using these features, good concordance has been seen between cholangioscopic appearance and histopathology.

Intraductal ultrasound

ERCP with IDUS improves the diagnostic yield of biliary strictures. A small and high-frequency ultrasound probe provides high resolution images of ductal and periductal tissues. IDUS features which suggest malignancy include eccentric wall thickening with an irregular surface, a hypoechoic mass, heterogeneity of the internal echo pattern, a papillary surface, disruption of the normal three-layer sonographic structure of the bile duct, presence of lymph nodes, and vascular invasion. It is, however, not commonly available and expertise is needed for a successful outcome.

Chromoendoscopy, autofluorescence and narrow-band imaging

For better characterization of biliary strictures, several techniques have been employed during cholangioscopy. In chromoendoscopy, different stains are topically applied to the surface of the mucosa. Methylene blue can successfully differentiate malignant lesions and ischemic strictures from normal mucosa. Biliary narrow band imaging enhances the vascular pattern of the mucosal surface and delineates tumor extent effectively. Initial cholangioscopic studies with autofluorescence have been less promising; poor specificity and high rates of false positivity were observed [9].

Confocal laser endomicroscopy (The cellvizio system)

Cellvizio is a probe-based CLE system which generates optical biopsies, providing physicians with microscoping images of tissue instantaneously and in a minimally invasive manner. This technique produces specific patterns that correlate with standard histology and differentiate between malignancy, inflammation and normal mucosa. In a recent multicentre study CLE was found to provide significantly higher diagnostic accuracy for malignant biliary strictures than standard ERCP (90% vs 73%) [10].

The Miami classification system has been proposed to characterize pCLE findings for biliary strictures. The presence of thick white bands (>20 micrometer), thick dark bands (>40 micrometer), dark clumps, epithelial structures and contrast leakage were the factors which could differentiate malignant from benign strictures [11]. Another newer classification system called the Paris classification was recently described. This includes additional features such as vascular congestion, dark glandular patterns, increased interglandular space and thickened reticular structures [12].

Suggested Approach to Indeterminate Biliary Strictures (Figure 3)

The choice of diagnostic workup should be individualized, depending to a great extent on local expertise and availability of the particular technology. Cross section imaging (CT or MRI) is useful to assess the respectability in patients with suspected malignancy. Table 2 outlines differences between benign and malignant strictures.
**Table 2**: Differences between benign and malignant biliary stricture.

<table>
<thead>
<tr>
<th></th>
<th>Benign Biliary Stricture</th>
<th>Malignant Biliary Stricture</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>Any age, usually younger</td>
<td>Usually age &gt; 50 yrs</td>
</tr>
<tr>
<td>Loss of weight, appetite</td>
<td>Less common</td>
<td>Significant loss</td>
</tr>
<tr>
<td>Jaundice</td>
<td>Deep jaundice unusual</td>
<td>Deep jaundice usual</td>
</tr>
<tr>
<td>Features of cholangitis</td>
<td>More common</td>
<td>Less common</td>
</tr>
<tr>
<td>Presence of lump in abdomen</td>
<td>Not a feature</td>
<td>Favors malignancy</td>
</tr>
<tr>
<td>Radiology MRCP/ERCP</td>
<td>Smooth stricture, no mass</td>
<td>Eccentric, irregular stricture, abrupt cut-off, presence of mass</td>
</tr>
<tr>
<td>Ca 19.9</td>
<td>Normal except in cholangitis</td>
<td>High</td>
</tr>
<tr>
<td>Cholangioscopy features</td>
<td>Smooth mucosa, no mass or tumor vessel</td>
<td>Tumor vessels present, nodules, mass</td>
</tr>
<tr>
<td>Cytology, biopsy</td>
<td>Not suggestive</td>
<td>Suggestive</td>
</tr>
</tbody>
</table>

**References**


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