A 61-year-old woman with advanced alcoholic liver cirrhosis (Child–Pugh score: C, 10 points; MELD score: 24 points) developed evident jaundice with prevalent conjugated hyperbilirubinemia (bilirubin: total 12.0 mg/dL, direct 7.5 mg/dL). Aminotransferases were only mildly elevated (AST 76 U/L, ALT 40 U/L), while there was a predominant elevation of alkaline phosphatase (162 U/L) and γ-glutamyl transpeptidase (103 U/L). Routine laboratory analysis also showed anemia (hemoglobin 9.8 g/dL), thrombocytopenia (platelets 25,000/μL), and prolonged prothrombin time (PT-INR 2.01). Ultrasonography showed dilated bile ducts. Magnetic resonance cholangiopancreatography (MRCP) revealed an abnormal biliary tree with multiple strictures and focal dilatations characterized by irregular contrast enhancement, potentially suggestive of intraductal malignancy, although no prominent obstruction was observed (▶Fig. 1a, b). Radiographic features of portal hypertension with perigastric and perisplenic collaterals were also observed. Neither esophageal nor gastric varices were found by esophagastroduodenoscopy, while congestive gastropathy was present. Endoscopic retrograde cholangiopancreatography (ERCP) was carried out with the aim of defining the diagnosis and facilitating biliary drainage. ERCP documented multiple narrowing of intra- and extrahepatic bile ducts without major stenosis (▶Fig. 1c). The still unexplained biliary strictures prompted performance of peroral cholangioscopy (▶Fig. 1d;▶Video 1). Biliary sphincterotomy was performed to be able to pass the cholangioscope. Peroral digital single-operator cholangioscopy (SpyGlass DS; Boston Scientific) revealed multiple bile duct varices (BDV) with red spots and microbleedings localized to the common bile duct, thereby defining a diagnosis of portal biliopathy. Neither a critical (i.e., clinically significant) stenosis nor blood clots causing obstruction were found. Therefore, no endoscopic treatment was performed. The patient remained clinically stable on follow-up, although no substantial improvement of jaundice was observed (bilirubin: total 13.2 mg/dL, direct 7.7 mg/dL). She accepted an offer to participate in a program of alcohol counseling and, finally, after 6 months of abstinence from alcohol, was considered for liver transplantation.

The term portal biliopathy refers to biliary obstruction associated with cavernous transformation of the portal venous system. Jaundice is its main clinical manifestation, but cholangitis and hemobilia may also be present. The diagnosis of portal biliopathy requires three criteria to be fulfilled: (i) presence of portal cavernoma and/or hypertension, (ii) typical cholangiographic changes (e.g., irregular ductal contour, strictures and dilatations) on ERCP/MRCP, and (iii) absence of other conditions that cause similar changes (e.g., neoplasms, primary sclerosing cholangitis, choledocholithiasis) [1]. Although BDV have been known about for decades [2,3], their diagnosis is still a clinical challenge for which a high index of suspicion is crucial [4]. BDV should be considered in the differential diagnosis of obstructive jaundice, especially in patients with known portal cavernoma and/or hypertension when medical im-
aging is inconclusive. Peroral cholangioscopy, allowing direct visualization of the biliary tract, may be a useful diagnostic tool for uncovering the causes of indeterminate biliary anomalies [5].

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Competing interests

The authors declare that they have no conflict of interest.

The authors

Gloria Tacchella1, Odencja Gjermeri1, Stefano Francescon Crinò2, Armando Gabrieleli2, Nicola Martinelli1
1 Department of Medicine, University of Verona, Policlinico G. B. Rossi, Piazzale L. A. Scuro 10, 37134 Verona, Italy
2 Gastroenterology and Digestive Endoscopy Unit, The Pancreas Institute, Policlinico G. B. Rossi, University Hospital, Verona, Italy

Corresponding author

Nicola Martinelli, MD, PhD
Department of Medicine, University of Verona, Policlinico G. B. Rossi, Piazzale L. A. Scuro 10, 37134 Verona, Italy
nicola.martinelli@univr.it

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