Biliorenous fistula related to self-expandable biliary metallic stent placement: a rare complication of endoscopic retrograde cholangiopancreatography

An 87-year-old woman was admitted to our hospital with cholangitis caused by a cholangiocarcinoma (Figure 1). She refused surgery, so a self-expandable metallic stent (SEMS) (Memotherm; Bard Inc. Billerica, Massachusetts, USA) was placed endoscopically. One year later, she presented with cholangitis related to tumor progression. We placed a second and a third coaxial "Y"-shaped SEMS (Luminexx, Bard Inc. Billerica, Massachusetts, USA) in the right and left bile ducts. Over the following 4 months she was admitted three times with sepsis of biliary origin, which resolved after cleansing of biliary sludge from the lumen of the stents. In the last episode, when contrast was injected at the proximal end of the right bile duct stent, a leak of contrast with rapid and turbulent flow was clearly seen, indicative of a biliovascular fistula (Figure 2). We cleared out the bile duct once again and her symptoms resolved. She died 4 months later after general deterioration in her condition.

Bilhemia occurs when the hepatic vessels and the bile ducts are in communication and the pressure is higher in the latter, leading to flow of bile into the bloodstream [1,2]. In our case, taking into account the location of the fistula and the temporal relationship between the onset of bacteremia and stent placement, we believe that the fistula was created after the opening of the proximal end of the right bile duct stent, which led to focal necrosis and created a communication with the vascular tree at this level. When the bile duct pressure became progressively higher as a result of tumor growth and sludge formation, small amounts of infected bile entered the bloodstream, causing bacteremia. The endoscopic management of bilhemia is based on reducing the pressure in the bile duct by placement of a biliary stent or by nasobiliary drainage. However, the patients who have been treated endoscopically for this complication previously were young and had benign disease [1,3,4]. The therapeutic decisions we made in this case were guided by our patient’s age, poor general condition, and prognosis.

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