A 78-year-old man was admitted to hospital due to massive, bright red hematemesis. As he was hemodynamically unstable, vigorous resuscitation procedures were initiated. Esophagogastroduodenoscopy (EGD) showed a large Mallory–Weiss lesion and a necrotic polypoid lesion about 4 cm below the upper esophageal sphincter (Figure 1). No active bleeding was visible. Six hours later, there was again an abrupt onset of torrential, bright red hematemesis. The bleeding did not stop before the insertion of a Sengstaken–Blakemore tube into the upper esophagus. Angiography of the thoracic aorta revealed a ruptured aneurysm in an aberrant right subclavian artery (ARSA).

Because of the expected high mortality of a surgical intervention in this situation, all treatment efforts were withdrawn, and the patient died 14 h after admission. The autopsy showed a ruptured aneurysm of an ARSA with an arterioesophageal fistula (Figure 2).

Rupture of an ARSA aneurysm into the esophagus is rare, with only 20 reported cases [1]. Most fistulas appear as a spontaneous rupture of a preexisting atherosclerotic aneurysm in the ARSA, but there have been several reports of arterioesophageal fistulas after prolonged nasogastric esophageal intubation in young patients with an ARSA who did not have an aneurysm [2]. Survival following a ruptured ARSA with an arterioesophageal fistula has only been documented in four patients.

Fistulas between an ARSA (or another large artery) and the esophagus should be suspected when there is massive, bright red hematemesis [2]. EGD may exclude other causes of bleeding, but the sensitivity of EGD for detecting arterioesophageal fistulas is only 38% [2]. Definitive diagnosis in stable patients is achieved by computed tomography and catheter angiography [3].