Hemothorax from external rupture of esophageal varices: An unusual fatal complication

Nigel H. Bramwell, BMED SCI BM, BS, Roger W. Byard, BMED SCI MB, BS, CCFP

ABSTRACT: A case of sudden death caused by hemorrhage into the chest cavity from a linear tear of the external esophageal wall following vomiting in a 20-year-old woman with congenital portal vein atresia is described. Compromise of the underlying esophageal wall due to recent sclerotherapy for bleeding varices predisposed to this highly unusual 'adventitial' variant of the Mallory-Weiss syndrome. Can J Gastroenterol 1989;3(2):58-60

Key Words: Esophagus, Variceal hemorrhage

A 20-YEAR-OLD WOMAN WAS ADMITTED WITH HEMATEMESIS. The patient had a long medical history, with an episode of hematemesis at the age of 20 months prompting investigations which led to a diagnosis of esophageal varices and portal hypertension due to congenital portal vein atresia. At the age of 10 years the patient underwent a mesentericaval bypass using a Teflon graft; recovery from which was complicated by a subphrenic abscess. Although the graft had never functioned satisfactorily, with occlusion confirmed by venography, the patient had remained well with no further episodes of bleeding until this admission.

The patient's course in hospital was a difficult one. Bleeding was not adequately controlled with a Blakemore tube and intravenous pitressin. A splenic and mesenteric venogram on the second hospital day showed an atretic portal sys-
Postoperatively, bleeding persisted. On the eighth hospital day esophagogastroscopic sclerotherapy was attempted unsuccessfully. A venogram at this time showed the graft to be functioning well. Liver function tests, including coagulation factors, were normal throughout the hospital course. Further bleeding was managed conservatively until a massive bleed on the 12th hospital day necessitated ligation of varices and devascularization of the stomach with a feeding jejunostomy. Recovery was complicated by septicemia secondary to intra-abdominal abscesses (blood cultures grew Staphylococcus aureus and Streptococcus faecalis) leading to wound dehiscence by the 24th hospital day. Antibiotics were administered and a further laparotomy for lysis of adhesions, revision of jejunostomy and a secondary wound closure were undertaken. Seven days postoperatively the patient had been extubated and appeared to be progressing well. On the morning of the 39th hospital day a chest X-ray was normal. Shortly thereafter the patient had an episode of forceful vomiting, following which she suddenly became obtunded and cyanotic. Resuscitation attempts were unsuccessful.

**AUTOPSY RESULTS**

**Macroscopic:** On opening the thorax, the left pleural cavity was found to contain 1500 mL of fresh blood with a markedly atelectatic lung. After evisceration,
the esophagus, stomach and proximal small bowel were examined separately. Large, prominent varices were easily seen over the distal one-third of the esophagus. The adventitial veins of the esophagus were dilated and tortuous. On close examination of the external surface of the esophagus, a 1.2 cm longitudinal tear was encountered which was 5 cm above the gastroesophageal junction on the left, anteriorly (Figure 2). No mucosal tear, or blood in the esophageal lumen or stomach, was found. The right lung was congested and no evidence of aspiration was encountered in the airways.

The abdomen was easily entered via the recent laparotomy incision. Distended loops of dusky colourd bowel, densely adherent to one another, were found. The adhesions were soft and easily dissected bluntly. Locoluated pockets of pus were widespread and all peritoneal surfaces were coated by florid granulation tissue and purulent debris. Tracing the inferior vena cava proximally from the iliac veins, it was possible to identify the recent mesocaval shunt, which was patent. The older Teflon graft was completely occluded. The portal vein was small, but identifiable and could be traced as far as its small intrahepatic branches. Centrilobular hepatic necrosis and pancreatic fat necrosis were noted.

**Microscopic:** Sections from the esophagus revealed extensive necrosis of the muscular coat in the area adjacent to the varices. Dilated venous channels with degenerative mural changes and perivascular hemosiderin deposition were prominent and in one area one such channel communicated with a breach at the muscularis-adventitial interface corresponding to the linear tear seen macroscopically (Figure 2).

Histopathological findings in the liver and pancreas correlated with the gross changes. The remainder of the macroscopic and microscopic examination revealed no significant pathological changes.

Death was attributed to shock and acute cardiorespiratory compromise due to left hemothorax and lung collapse resulting from external rupture of esophageal varices.

**DISCUSSION**

Spontaneous perforation of the esophagus in patients without varices is a well documented, if infrequent, condition (1) often associated with unusually high intramural pressures induced by vomiting and retching (Mallory-Weiss syndrome) (2) or even by a suppressed sneeze while swallowing food and air (3). Acquired perforations due to foreign bodies are self-explanatory.

Patients with esophageal varices, however, usually bleed on the basis of spontaneous rupture of swollen submucosal veins into the lumen in the absence of causative forces such as vomiting (4,5). Such hemorrhages usually, therefore, lie outside the category which embraces the Mallory-Weiss and Boerhaave syndromes (6). This distinction was made endoscopically by Alvarez and co-workers in their study of children with portal obstruction in whom 79% suffered some form of gastrointestinal hemorrhage, the majority unrelated to a definite precipitating event (7,8).

The present patient represents a highly unusual situation in which therapeutically induced compromise of the structural integrity of the esophageal wall, combined with large adventitial varices, made her vulnerable to external rupture with massive hemorrhage following vomiting. While the Mallory-Weiss phenomenon classically involves the esophageal mucosa, this case demonstrates that increased intraluminal pressure can cause tearing at any level of the esophageal wall, given appropriate predisposing circumstances. The potential seriousness of this rare complication is emphasized by the patient's sudden deterioration and death. This unusual complication, not previously reported to the authors' knowledge, is worthy of note by clinicians who manage these difficult cases.

**ACKNOWLEDGEMENTS:** The authors thank Mrs. K.J. Bramwell for preparing the manuscript.

**REFERENCES**

Submit your manuscripts at http://www.hindawi.com