Hepatic portal venous gas (PVG) is a rare and often puzzling clinical entity, with widely varying features regarding presentation [1].

A 77-year-old white woman with a history of primary biliary cirrhosis presented complaining of hematemesis and underwent an esophagogastroduodenoscopy and band ligation. Three days later, CT of her abdomen was performed and showed the presence of gas in the portal venous system (Fig. 1) without any evidence of pneumatosis, ischemia, or bowel thickening. The Department of Surgery was consulted for further evaluation of the PVG, but given the patient’s debilitated condition and the benign abdominal exam findings, it was elected to continue to monitor her. She did very well, without any complaints of abdominal pain or fever, and was eventually discharged home in a satisfactory condition.

First described amongst infants with necrotizing enterocolitis, PVG has been regarded as a sign of an abdominal emergency requiring urgent exploratory surgery [2]. With the ever-expanding role of CT scanning in the diagnosis and treatment of abdominal complaints, however, the finding of PVG has been demonstrated in a wider array of abdominal conditions ranging from bowel necrosis to chronic conditions such as ulcerative colitis and Crohn’s disease [4].

It is not entirely clear what caused the PVG in our patient. In the absence of any abdominal findings and with a normal serum lactate level, an abdominal emergency is less likely. In theory, the esophagogastroduodenoscopy may have been responsible for causing vascular breach, resulting in PVG without any evidence of bowel necrosis.

The patient in this case was at high risk for perioperative mortality with underlying cirrhosis, and it was determined that surgical intervention would be more harmful than beneficial.

In conclusion, in selected patients in whom PVG has been demonstrated on imaging, such as the one described here, supportive care may often be all that is required.

References