Case report

A 32-year-old male who underwent a straightforward laparoscopic cholecystectomy 5 months earlier, presented with 3-day history of intermittent episodes of haematemesis and melaena. He denied any history of abdominal pain and chronic ingestion of any non-steroidal anti-inflammatory analgesics.

At presentation, his vital signs were stable, and the abdominal examination was unremarkable. The routine blood tests were within the normal range. Gastroscopy showed a deformed duodenal bulb, with narrowing of the 2nd part of duodenum by a smooth convex bulge representing either an external compression or submucosal lesion (Figure 1). Abdominal ultrasound revealed a well-defined (size: 10x6x6 cm) sacculo-tubular mass in the porta hepatis region (Figure 2A). Doppler ultrasound study showed an echogenic thick wall containing a mural thrombus with turbulent arterial flow (Figures 2B). Computed tomography (CT) scan showed a large well-defined hypodense mass with relatively hyperdense periphery arising from the hepatic artery and extending caudally (Figures 2C). A 3D CT-Angiography reconstructed images showed the presence of a huge aneurysm arising from the right hepatic artery (Figure 2D).

The patient was offered selective angiography and embolization, but he refused. He was discharged against medical advice and therefore lost to follow-up.

Discussion

The hepatic artery is the fourth most common site of intra-abdominal aneurysm after infra-renal aorta, iliac artery and the splenic artery. Recently, hepatic artery aneurysms are becoming the most common, due to the explosive increase in the use of percutaneous and laparoscopic hepato-biliary procedures [1]. More than one half of all detected hepatic artery aneurysms are pseudoaneurysms and the extrahepatic artery aneurysms have higher incidence of rupture than the intra-hepatic ones.

Generally, hepatic artery pseudoaneurysm (HAP) is rare (reported incidence is 0.6%) [2] but can lead to serious life-threatening complications. It was first reported after laparoscopic cholecystectomy (LC) in 1994 and attributed to iatrogenic vascular injury [1]. The injury occurs either in association with bile duct injury during LC or in isolation as a result of direct trauma or thermal injury [3]. The diagnosis is often made weeks or months after the laparoscopic procedure [4].

HAP commonly presents with gastrointestinal tract bleeding; commonly haemobilia if the pseudoaneurysm communicates with...
the bile duct. Also, it may present with obstructive jaundice due to
the compression of the biliary tract by the expanding aneurysmal
haematoma [5]. It may also present urgently with massive
intraabdominal bleeding which necessitates emergency surgical
intervention [6]. Contrast CT scan or dynamic CT angiography and
Doppler ultrasound can ‘clinch’ the diagnosis. However, it is best
made by selective angiography. The first line of treatment is non-
operative in the form of selective embolization which offers reasonably
good long-term results [7]. However, coil embolization carries the risk
of aneurysmal rupture, hepatic necrosis and delayed bile duct stricture
due to ischaemia. Sudden rupture of HAP with massive intraabdominal
bleeding carries high mortality rate and hence, necessitates prompt
emergency surgical intervention with ligation of the artery.

Conclusion

- Hepatic artery pseudoaneurysm (HAP) may develop weeks or
  months after laparoscopic cholecystectomy.
- HAP commonly present with intermittent GI bleeding (haematemesis
  or melaena). If left untreated, it ruptures with massive intraabdominal
  bleeding which carries high mortality rate.
- HAP should be suspected in any patient presenting with GI bleeding
  and has a recent history of laparoscopic cholecystectomy.

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