Surgical experience of hepatectomy in a live donor with absent celiac trunk - a case report

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Abstract

Background: Celiac trunk agenesis is the recent and the rarest vascular variation reported. Reported cases in literature were identified either incidentally on imaging or while cadaveric dissections. This is a unique experience of hepatectomy in a patient without celiac artery ever reported.

Case presentation: This is a case of a patient who was scheduled to undergo donor hepatectomy. He was found to have a rare anatomical variation of agenesis of the celiac trunk during preoperative imaging done as a routine transplant surgery workup. He underwent live donor hepatectomy with no intra-operative or post-operative complications. This is the first case to be reported in the literature of a donor hepatectomy for liver transplantation in a patient with this rarest vascular variation of the celiac axis.

Conclusion: Pre-operative imaging to determine hepatic vascular anatomy improves the understanding of the surgeon intraoperatively while dissection and decreases the chance of any iatrogenic damage.

Keywords: Liver transplant, Agenesis of the celiac trunk, Anatomical variation, Hepatic artery, Celiac artery, Case report

Background

Different anatomical variations of the celiac artery have been recognized and reported in the literature with agenesis of the celiac trunk being the rarest anomaly (0.1–2.6%) [1, 2]. The first case of the absent celiac trunk was reported in the year 1832 [3].

Earlier classifications of the celiac artery did not include the missing celiac artery as a variant [4, 5]. As per Morita’s classification [1, 4], this case belongs to type V (typus primitus). Adequate pre-operative workup helped surgeons in understanding the vascular anatomy intraoperatively and therefore, after mutual consensus of transplant team it was decided that no alteration in the standard surgical plan was needed in this donor. A standard hilar dissection for the left hepatic lobectomy was performed. No attempt was thus made to dissect the aortic origin of the three separate vessels replacing the celiac trunk.

Case presentation

A 38-year-old male with no known comorbid, was planned for liver donation, underwent routine pre-operative evaluation before hepatectomy for a liver transplant. CT angiography and volumetry was done to visualize the vascular anatomy of the hepatobiliary system. On reviewing the images, celiac trunk was absent and was replaced by left gastric, common hepatic, and splenic arteries originating separately from the aorta (Fig. 1). The left gastric artery was the first branch to arise, followed by the common hepatic artery (5.8 mm in diameter and 16.5 mm in length) (Fig. 2). Common hepatic artery bifurcated into gastroduodenal artery and hepatic artery proper (4.5 mm in diameter). Hepatic artery proper divided into right hepatic artery (3.7 mm in diameter and 43.9 mm in length) and left hepatic artery...
(1.7 mm in diameter and 41.8 mm in length) (Fig. 3). A branch from right hepatic artery was supplying segment IV (2 mm in diameter and 42.0 mm in length). Venous anatomy was unremarkable except two accessory hepatic veins noted in segment VI (7.3 mm in diameter) and segment VII (2.6 mm in diameter). Total liver volume was 1832.95 cc (right lobe =1386.22 cc, left lobe = 419 cc, caudate lobe = 27.73 cc). The splenic artery originated at the same level as the common hepatic artery, but it was directed to the left toward the spleen. All other relevant pretransplant workup was normal.

The patient had unremarkable donor hepatectomy with no adverse intra-operative event. He remained stable post procedure and was discharged within few days.

Discussion

Pattern of Trikus Halleri (trifurcation) of the coeliac trunk was first described by Haller in 1756. Normally, the celiac artery branches into the left gastric artery, the common hepatic artery, and the splenic artery [1]. Classical trifurcation of the celiac artery has been reported in 87.6% of the cadavers, and 12.4% had variable patterns [3]. Agenesis of the celiac trunk is a rare vascular anomaly with a mean prevalence of 0.38% [6]. The first case of the absent celiac artery was reported in 1832 by Geoffory Saint-Hilaire [3]. Aortic origination of all branches of the celiac axis is seen in 2% of cases [7]. In a systematic review of anatomical variation of celiac artery, 5 cases (41.7%) of the absent celiac trunk were identified out of total of 12 studies [2]. Nearly, 33% of these cases are diagnosed on radiological imaging [1]. 0.19% of 10750 CT scans had absent celiac trunk [4].

Various theories have been explained regarding the missing celiac trunk. It is believed that formative changes of ventral splanchnic arteries that have originated from paired ventral segmental arteries are responsible for morphological types of the celiac axis. The celiac trunk develops from the union of longitudinal anastomoses of segmental arteries. The absence of these anastomoses leads to the remnant of the segmental arteries and subsequent agenesis of the celiac trunk [4].

Initially, anatomical variants of the celiac trunk were classified by Lipshutz (1917) and Adachi (1928). Lipshutz described four variations of the celiac trunk: type I: coeliac artery branches into gastric, splenic, and
hepatic arteries; type II: hepatic and splenic arteries arise from the coeliac axis and the gastric artery originates from the abdominal aorta; type III: the gastric and hepatic arteries arise from the coeliac axis whereas, the splenic artery separately branch from the abdominal aorta; and type IV: the gastric and the splenic arteries branch from the celiac artery and the hepatic artery comes from the abdominal aorta [5]. Adachi classified six types: (i) hepatogastrosplenic, (ii) hepatosplenic, (iii) gastrosplenic, (iv) coeliacomesenteric, (v) hepatosplenomesenteric, and (vi) hepatomesenteric [5].

Agenesis of the celiac trunk was not described in either of these classification systems of the celiac trunk. Morita proposed a modified version of the celiac axis classification which included absent celiac artery as one of the anatomical variants. Morita’s classification included (i) celiac trunk, (ii) hepatosplenic, (iii) gastrosplenic, (iv) hepatogastric, and (v) absent celiac trunk [1, 4].

According to Morita’s classification (1935), the presented case belongs to type V (Typus primitivus). Pre-operative knowledge of abdominal vascular anatomy in this case helped in careful identification and dissection of the left hepatic hilar structures during left donor hepatectomy. That included the left hepatic artery, the separate segment IV hepatic artery, the left hepatic duct, and the left portal vein. Following a careful dissection of these structures and the parenchymal dissection, ligation of the left hepatic artery and segment IV hepatic artery (arising from the right hepatic artery) were key steps in this case of donor hepatectomy. Failure to identify the common hepatic artery can lead to devastating effects on the vascularity of the remaining liver parenchyma in a live donor. The rest of the surgery was carried out without any special intervention or maneuver, with regards to vascular morphology.

This case highlights the importance of pre-transplant imaging as vascular variations when encountered intraoperatively can be challenging for the surgeon; also, CT imaging aids in preventing inaccurate resection and hence reducing donor-related morbidity and mortality.

Conclusion
This case report depicts the importance of pre-operative radiological techniques in patients planned for liver transplant as various anatomical varieties of celiac axis exist and precise details of hepatic vasculature help in the planning of surgery in live hepatic donors for transplant.

Abbreviations
MD-CT: Multi-detector CT scan

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The authors declare that they have no competing interests.

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