A rare hepatic artery variant reporting and a new classification

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Variations of the hepatic artery are very common, but they greatly increase the difficulty of surgery and the risk of complications in perihepatic surgeries such as liver transplantation, liver segmentectomy, and gastroduodenal surgery. Thus, it is important to precisely define the type of hepatic artery variant before surgery. However, there are often rare variants that cannot be defined with existing classifications. For example, the type of hepatic artery variant in the current case could not be classified with conventional classifications, and no such variation has been reported to date, involving two accessory left hepatic arteries from the common hepatic and left inferior phrenic arteries, respectively. Based on the existing 3DCT technology and the CRL classification method, which is applicable to the most common hepatic artery variants, we reviewed many rare variant types and proposed a new classification method (ex-CRL classification) for hepatic artery variations that do not fit the classic scope. The ex-CRL classification can accurately classify the vast majority of rare cases in the literature, greatly compensates for the limitations of current hepatic artery classifications, improves the generalization and understanding of rare cases, and reduces surgical complications.

KEYWORDS accessory hepatic artery, vascular variants, phrenic artery, classification, case report

Introduction

The common hepatic artery (CHA) is one of the main arterial supplies of the liver, gallbladder, lesser omentum, and gastroduodenal region, and its branches and variations are complicated. Interestingly, normal patterns occur only in 42%–75.7% of cases (1, 2). Aberrant arteries may be accidentally injured during surgical procedures resulting in severe haemorrhage and other fatal complications (3–11). A thorough knowledge of possible variations in branching, courses, and distribution of the vessels supplying the liver and gallbladder is essential. Therefore, many variations of the hepatic artery have been described and classified, among which the classic classifications are Michel’s 10 types and Hiatt’s 6 types (1, 8). However, despite the many classifications, more than 10% of hepatic arteries still cannot be classified. In recent years, with the wide application of 3D visualization technology (3DVT) (9–11), Yan et al. (12) presented the CRL classification, which succinctly describes the hepatic artery and its branches and covers a wider range of arterial variants. However, some rare variants still cannot be categorized by the CRL classification, and these variations are more likely to be...
unidentified and injured. These rare types have been reported in the form of individual cases for many years, lacking systematic classification and generalization. The aim of this study was to report a variation of the hepatic artery not in any of the hepatic classifications. Thereafter, based on CRL classification, the study puts forward an extended version (ex-CRL classification) specifically for some rare cases.

Case presentation

The variation of the hepatic artery was found during a routine dissection of an adult male cadaver for teaching purposes at Changzhi Medical College in 2020. In compliance with the confidentiality of the donor, we have not obtained the medical history and related information of the cadaver. The specimens were perfused with 5% formaldehyde and 10% alcohol, and fixed with 4% formaldehyde solution. The structure of the specimen was clear and intact.

CHA, originated from the celiac trunk (CT), which travelled behind the stomach and gave off three branches as follows: the accessory left hepatic artery (aLHA), proper hepatic artery (PHA) and gastroduodenal artery (GDA). The aLHA arose 22.0 mm distal to the origin of the CHA, travelling behind the PHA and finally entering into segment IV, posterior to the common hepatic duct (CHD), and its one bunch ran through the umbilical fissure. The PHA then travelled to the right about 11.4 mm and divided into the right gastric artery and cystic artery, and then ran superiorly about 34.2 mm and entered the liver through the porta hepatis (Figure 1).

In addition, the abdominal aorta gave off the left inferior phrenic artery at about 23.0 mm above the CT and was associated with agenesis of the right inferior phrenic artery. The left inferior phrenic artery provided the second left accessory hepatic artery. The second left accessory hepatic artery partially joined the left interlobar artery of the left lobe through the venous ligament and partially supplied the left lobe of the liver, forming a small branch supplying the falcine ligament and partially supplying the right diaphragm.

Discussion

To the best of our knowledge, there has been no case report of the hepatic artery arising from the inferior phrenic artery and replacing the right inferior phrenic artery. Only Covey et al. (13) reported the origin of the hepatic artery from the phrenic artery. They collected arterial anatomical data from 600 cases of visceral angiography from May 1996 to October 2000, and found that one case of rRHA and aRHA each originated from the right phrenic artery. Thereafter, Liang (14) searched the previous literature and analysed 21 studies, including that of Covey et al., with a total of 10,966 cases, and only 2 cases reported by Covey et al. noted that the hepatic artery originated from the phrenic artery. Jin (15) reviewed 10,211 cases in 21 articles and found that only 2 cases originating from the phrenic artery were reported by Covey et al. Therefore, the origin of the accessory hepatic artery from the phrenic artery reported in this case is extremely rare and noteworthy.

In addition, another accessory hepatic artery in the current case originated from the common hepatic artery. Futara (16) dissected 110 cases and found that in 5 cases (4.5%), the left accessory hepatic artery originated from the common hepatic artery.
artery, while Pai (17) collected autopsy data from 72 cadavers between 2000 and 2005 and found only one (1.38%) variant of the accessory hepatic artery originating from CHA, which was much lower than the expected incidence (4.5%).

However, we think about reasons why this variant has not been previously described, except that it has an extremely low incidence and may also be associated with limited 2D CT techniques. 2D technologies very test the spatial imagination of surgeons (18), except for conventional variants such as left gastric artery and superior mesenteric artery, other rare variants are difficult to find. In this case, the accessory left hepatic artery originating from the inferior phrenic artery is short and concealed, and we only exposed its course completely after excising part of the left lobe, so it is easily overlooked.

In the current case, the hepatic artery that replaced the right inferior phrenic artery was thicker than the left inferior phrenic artery, and has a wider range of blood supply. This variation is likely to cause ischaemia and dysfunction of the right diaphragm during left hepatic lobectomy, lymph node dissection around a gastric cancer, infusion of chemotherapeutic drugs for hepatocellular carcinoma, and...
vascular embolisation. In addition, the accessory hepatic artery may not be fully visualised during selective hepatic arteriography, and such vascular variants interfere with effective tumour control by transcatheter arterial chemoembolisation (19). If a brain-dead donor of this variant type is encountered during liver transplantation, all blood vessels entering the liver should be preserved as long as possible. During vascular reconstruction, it can be flexibly determined according to the recipient’s vascular condition, and blood vessels with an appropriate diameter and in good condition are selected for end-to-end anastomosis with the donor. When the recipient has poor vascular conditions such as the common hepatic artery and gastroduodenal artery due to the underlying liver disease, but the splenic artery is in good condition, an anastomosis with the easily separated splenic artery is selected; if the donor liver cannot obtain arterial blood supply in situ, The donor iliac vessels were selected to bypass the hepatic artery with the abdominal aorta (20). If this type of variant is a living donor, due to the presence of the double accessory left hepatic arteries, a right half liver can also be considered for living donor liver transplantation. Of course, it is still controversial as to whether multiple arteries need routine reconstruction, so full surgical protocol should also be made with adequate preoperative evaluation (21).

Although the variants of the two combinations in this case are extremely rare, these variants are complex and easy to ignore and could lead serious intraoperative complications. Therefore, the accumulation of knowledge regarding this type of variant has important anatomical significance for surgery and interventional therapy.

In recent years, with the development of 3DCT technology, 3DVT is clearer and more accurate in imaging blood vessels compared to 2D imaging (22). 3DVT is widely used in preoperative evaluation and provides a clearer understanding of the origin and course of CHA, LHA, and RHA. Therefore, there is also a need for a new type of classification that is more compatible with 3DCT to describe and classify hepatic arteries more concisely. The CRL classification is a new classification developed based on 3DCT technology, with each hepatic artery described in detail. It conforms to the current requirements for preoperative evaluation and is suitable for most routine variants. However, it is currently not specific for rare cases. Therefore, we propose a new classification, the extra-CRL classification (Figure 2), on the basis of the CRL classification to name and classify variants that cannot be classified by the CRL classification.

CRL classification rules

CRL describes the origin of CHA, RHA, and LHA, with CRL being the abbreviation of their initials. CRL describes normal origins; that is, CHA originates from CT, while RHA and LHA originate from PHA. Thereafter, the origins of other arteries are marked with capital letters corresponding to their arteries. For example, CRL means that CHA has changed, not from CT but from SMA, while the origins of RHA and LHA remain changed. The replaced or accessory hepatic arteries need to be represented by lowercase letters “r” or “a”. For example, CRL means that CHA originates from CT, rRHA originates from GDA, and aLHA originates from the aorta. The CRL classification is detailed in Table 1.

| Classification | Description |
|----------------|-------------|
| Type 1 (CRL) | CHA from coeliac trunk; LHA and RHA from PHA |
| Type 2 (CRaL) | rRHA only |
| 2a (CRaL) | rRHA from aorta |
| 2b (CRaL) | rRHA from coeliac trunk |
| Type 3 (CRaL) | aLHA from other arteries |
| 3a (CRaL) | aLHA from aorta |
| 3b (CRaL) | aLHA from coeliac trunk |
| 3c (CRaL) | aLHA from SMA |
| 3d (CRaL) | aLHA from other arteries |
| Type 4 (CRL) | rLHA |
| 4a (CRL) | rLHA from GDA |
| 4b (CRL) | rLHA from other arteries |
| Type 5 (CRL) | aLHA |
| 5a (CRL) | aLHA from GDA |
| 5b (CRL) | aLHA from LGA |
| 5c (CRL) | aLHA from other arteries |
| Type 6 (CRL) | rRHA and rLHA (type 2 + type 4) |
| CRaL | rRHA from aorta, coeliac trunk, GDA, SMA or other arteries; rLHA from GDA, LGA or other arteries |
| Type 7 (CRaL or CRaL) | rRHA and aLHA (type 2 + type 5), or aRHA and rLHA (type 3 + type 4) |
| 7a (CRaL) | rRHA from aorta, coeliac trunk, GDA, SMA or other arteries; aLHA from GDA, LGA or other arteries |
| 7b (CRaL) | aLHA from coeliac trunk, GDA, SMA or other arteries; rLHA from GDA, LGA or other arteries |
| Type 8 (CRL) | aRHA and aLHA (type 3 + type 5) |
| CRaL | aRHA from aorta, coeliac trunk, GDA, SMA or other arteries; aLHA from GDA, LGA or other arteries |
| Type 9 (CRL) | rCHA not from coeliac trunk |
| 9a (CRL) | CHA from aorta |
| 9b (CRL) | CHA from GDA |
| 9c (CRL) | CHA from LGA |
| 9d (CRL) | CHA from SMA |
| 9e (CRL) | CHA from other arteries |
The ex-CRL classification is suitable for a small number of variants outside the CRL classification. Other arteries in the CRL classification are uniformly represented by "O". For example, CRaOL indicates that aRHA originates from other arteries other than AA, CT, PHA, LGA, GDA, or SMA, while ex-CRL subdivides other arteries that have appeared in the past, such as "OR", which indicates origin from the renal artery; "OP", from the phrenic artery; "OS", from the splenic artery; "OC", from CHA; "OD", from the pancreaticoduodenal artery; "OM", from the inferior mesenteric artery; and "ODP" from the dorsal pancreatic artery. Detailed rules for the ex-CRL classification are shown in Tables 2, 3.

TABLE 2 Terminology and Nomenclature of CRL and ex-CRL classification.

| Label | Description |
|-------|-------------|
| C     | origin of CT |
| R     | origin of RHA |
| L     | origin of LHA |
| O     | other arteries |
| OR    | originating from the renal artery |
| OP    | originating from the phrenic artery |
| OS    | originating from the splenic artery |
| OC    | originating from the CHA |
| OD    | originating from the pancreaticoduodenal artery |
| OM    | originating from the inferior mesenteric artery |
| ODP   | originating from the dorsal pancreatic artery |

 TABLE 3 ex-CRL classification.

| Classification | Description |
|----------------|-------------|
| C0RL (CRL Type 9) | CHA from other arteries |
| C0RL | CHA from the renal artery |
| C0RL | CHA from the splenic artery |
| C0RL | CHA from the pancreaticoduodenal artery |
| C0RL | CHA from the inferior mesenteric artery |
| C0RL | CHA from the dorsal pancreatic artery |
| CRaOL (CRL Type 2o) | aRHA from other arteries |
| CRaOL | aRHA from the renal artery |
| CRaOL | aRHA from the splenic artery |
| CRaOL | aRHA from the CHA |
| CRaOL | aRHA from the pancreaticoduodenal artery |
| CRaOL | aRHA from the inferior mesenteric artery |
| CRaOL | aRHA from the dorsal pancreatic artery |
| CRaOL (CRL Type 3o) | aLHA from other arteries |
| CRaOL | aLHA from the renal artery |
| CRaOL | aLHA from the splenic artery |
| CRaOL | aLHA from the CHA |
| CRaOL | aLHA from the pancreaticoduodenal artery |
| CRaOL | aLHA from the inferior mesenteric artery |
| CRaOL | aLHA from the dorsal pancreatic artery |
| CRaOL | aLHA from the renal artery |
| CRaOL | aLHA from the splenic artery |
| CRaOL | aLHA from the CHA |
| CRaOL | aLHA from the pancreaticoduodenal artery |
| CRaOL | aLHA from the inferior mesenteric artery |
| CRaOL | aLHA from the dorsal pancreatic artery |
| CRLaO (CRL Type 5o) | aLHA from other arteries |
| CRLaOP | aLHA from the renal artery |
| CRLaOP | aLHA from the phrenic artery |
| CRLaOS | aLHA from the splenic artery |
| CRLaOC | aLHA from the CHA |
| CRLaOP | aLHA from the pancreaticoduodenal artery |
| CRLaOM | aLHA from the inferior mesenteric artery |
| CRLaODP | aLHA from the dorsal pancreatic artery |

TABLE 3 Continued

| Classification | Description |
|----------------|-------------|
| Combinatorial variation | Two or more accessory artery, replaced artery or middle hepatic artery |
| CRa1_a2_L | two aRHAs; eg: "CRa1A,a2LL", means "aRHA1 from aorta, aRHA2 from LGA" |
| CRr1_a2_L | two rRHAs; eg: "CRr1A,a2LL", means "rRHA1 from aorta, rRHA2 from LGA" |
| CR_La1_a2 | two lHAs; eg: "CRLa1A,a2L", means "lLHA1 from aorta, lLHA2 from LGA" |
| CR_Lr1_a2 | two lHAs; eg: "CRLr1A,a2L", means "lLHA1 from aorta, lLHA2 from LGA" |
| C_R_L | CHA, aRHA or rRHA, aLHA or rLHA from three different arteries, and CHA not from coeliac trunk |
| C_R_M_L | MHA is present and not from PHA |

ex-CRL classification rules

The ex-CRL classification is suitable for a small number of variants outside the CRL classification. Other arteries in the CRL classification are uniformly represented by "O". For example, CRaOL indicates that aRHA originates from other arteries other than AA, CT, PHA, LGA, GDA, or SMA, while ex-CRL subdivides other arteries that have appeared in the past, such as "OR", which indicates origin from the renal artery; "OP", from the phrenic artery; "OS", from the splenic artery; "OC", from CHA; "OD", from the pancreaticoduodenal artery; "OM", from the inferior mesenteric artery; and "ODP" from the dorsal pancreatic artery. Detailed rules for the ex-CRL classification are shown in Tables 2, 3.

In addition, ex-CRL complements multiple replaced or accessory arteries, as well as in the case of MHA. It uses "a1", "a2", "r1", and "r2" to represent multiple replaced or accessory hepatic arteries. As reported in this case, aLHA1 originates from CHA, and aLHA2 originates from the phrenic artery, and is represented by CRa1OC, a2ODP, which succinctly and clearly illustrates the variation of the two aLHAs. In addition, MHA is not uncommon in variants. When MHA originates from PHA, RHA, or LHA, it does not need special notation, because it is close to the hepatic portal and bifurcates in advance before entering the liver to supply the fourth segment of liver. However, when MHA originates from other arteries and is difficult to be classified as RHA or LHA,
special labeling is required. For example, Gündoğdu et al. (23) described MHA originating from the pancreaticoduodenal artery, and aRHA originating from the dorsal pancreatic artery in 2021, which can be represented by CRaODPMOPL.

Finally, cases with variations in CHA, RHA, and LHA have been supplemented. For example, CHA from AA, aLHA from LGA, aRHA from SMA, were reported by Gruttadauria et al. (24) in 2001. However, the current case cannot be classified with the CRL classification, but can be expressed as CArAOCL based on the ex-CRL classification.

**Evaluation of the ex-CRL classification**

Different classifications were used to describe rare cases in the previous literature (Table 4). We reviewed the literature on hepatic artery variants from 1994 to 2021, and finally retained 16 studies (2878 cases) describing rare variants, including 17 rare hepatic arterial variants that cannot be described by common classification methods. We found that the classic Hiatt and Michel classifications were unsuitable for these rare and complex classifications, and their coverage by the CRL classification was relatively improved, but the specificity for these rare cases was not high. For example, rRHA described by Abdullah et al. (33) in 2006 originated from the inferior mesenteric artery, which is classified as CRrOML by ex-CRL, and rRHA described by Imam et al. (25) in 2021 was derived from the pancreaticoduodenal artery, which was classified as CRrODL by ex-CRL, but both cases are represented by CRL classification as “2o (CRrOL)”. The CRL classification does not clearly describe this part of “other arteries”, making it necessary to use ex-CRL for reclassification.

The CRL classification can classify some rare cases, but it lacks specificity and cannot describe them accurately. When the case base is large, the number of rare cases will increase; thus, attention is required for these rare cases. The application of ex-CRL, as an extension of the CRL classification, is conducive to improving the understanding and classification of rare variants and reducing the occurrence of surgical complications.

However, all classification methods, including ex-CRL classification, describe the origin of the variant artery, but lack the description of the path of the variant artery. Even if they originate from the same artery, their bifurcation planes and paths may be different. For example, two cases of aRHA originating from the renal artery reported by Darsan (27) in
2019 originated from the proximal end of the renal artery near the aorta and the distal end of the renal artery near the renal hilum, and their ascending paths were also different. This requires careful observation while performing an analysis of the detailed path of the artery to ensure that rapid and precise judgements are made in a limited surgical field of view.

Variations of the hepatic artery and surrounding blood vessels are associated with increased surgical difficulty and complications (36). For example, the accessory hepatic artery varies in liver transplant donors. If the accessory hepatic artery is ligated, it will lead to local arterial ischaemia, which may compromise liver function. If the accessory hepatic artery is preserved, it will lead to an increase in the number of arterial reconstructions and anastomosis, and a longer warm ischaemia time, which has been proven to be a risk factor for hepatic artery thrombosis (37). However, if the recipient has a complex hepatic artery vascular variation, due to the small and complex blood vessels of the hepatic artery, it will increase the incidence of hepatic artery thrombosis, hepatic artery stenosis, hepatic artery haemorrhage, or other nonvascular complications, including bile duct stenosis after liver transplantation as well as liver abscesses (38). Additionally, in pancreaticoduodenectomy, perihepatic arterial variation increases the risk of compromised hepatic arterial supply, which may lead to unintended bleeding or ischaemia, as well as an increased risk of biliary anastomotic leakage, transient liver function disorders, and, in some patients, liver failure (39).

In addition to invasive angiographic techniques, and noninvasive vascular imaging techniques, 3D reconstruction techniques such as volume rendering technique (VRT) and curved planar reformation (CPR) have rapidly evolved, they allow stereoscopic imaging of splanchnic arteriovenous and other conduits, reflecting the complete vessel diameter, length, course, and positional relationships of surrounding soft tissues and organs (40, 41). We expect that 3D visualization technology and CRL classification will become more widespread and routinely used for preoperative evaluation and surgical simulation of perihpatic surgery. Improved understanding of vascular anatomy, imaging, and classification of hepatic arteries will facilitate the development of individualized surgical protocol and avoid complications, while also expecting that more rare types of new hepatic artery variants will be discovered.

**Conclusion**

Comprehensive knowledge, accumulation, and classification of hepatic arterial variant types can greatly reduce surgical complications, and the ex-CRL classification can be used for hepatic arterial variants that cannot be classified with conventional methods.

**Data availability statement**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

**Ethics statement**

The studies involving human participants were reviewed and approved by Ethics Committee of Changzhi Medical College. The patients/participants provided their written informed consent to participate in this study. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

**Author contributions**

XW conceptualized the topic and wrote the manuscript. JK and YL consulted the literature. JN and YS conceptualized the topic and supervised and facilitated the conduct of the study. GS modified the edited and made significant revisions to the manuscript. All authors read and approved the final manuscript. All authors contributed to the article and approved the submitted version.

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**Conflict of interest**

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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