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Relationship of vascular variations with liver remnant volume in living liver transplant donors

Vascular variants and liver volumes

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Abstract

Background: In this study, we investigated the relationship between the portal vein and hepatic artery variations and the remaining liver volume in living donors in liver transplantation.

Materials and methods: In the study, triphasic abdominal CT images of 180 live liver donor candidates were analyzed retrospectively. Portal veins were divided into four groups according to the Nakamura classification and seven groups according to the Michels classification. The relationship between vascular variations and remnant liver volume was compared statistically.

Results: According to the Nakamura classification, there were 143 (79.4%) type A, 23 (12.7%) type B, seven (3.9%) type C and seven (3.9%) type D cases. Using the Michels classification, 129 (71%) type 1, 12 (6.7%) type 2, 24 (13%) type 3, two (2.2%) type 4, 10 (5.6%) type 5, one (0.6%) type Six and two (1.1%) type 7 cases were detected. There was no significant difference in the percentage of the remaining volume of the left liver lobe between the groups ($p = 0.055$, $p = 0.207$, respectively).

Conclusions: Variations in the hepatic artery and portal vein do not affect the remaining liver volume in liver transplantation donors.

Key words: portal vein, hepatic artery, anatomic variations, liver remnant volume, liver transplantation, living donor

INTRODUCTION

An accurate evaluation of potential living donors before liver transplantation is important in preventing postoperative liver failure. (1) There are many factors related to donor safety, such as obesity, age, liver volume, fatty liver, medical problems, anatomical variations, surgical process, and operative procedures.(2) The correct assessment of liver vascularity and volume is very important for both the donor and the recipient.(1, 3) Posttransplantation liver failure has been reported in the right lobe donors at around 10%.(3) On the other hand, the low liver volume of the recipient is an important problem that affects the recovery rate.(3, 4) Triphasic abdominal CT imaging has minimized errors in vascular and volume assessment.(3) The determination of vascular variations is essential for the operative procedure because they can affect decisions regarding the resection line or the use of grafts (2, 5). A minimum of 30% of liver remnant volume after posttransplantation is considered as the critical limit.(1, 4) Therefore, researchers have explored the relationships between liver volume and vascular structures.(6-9) Similarly, in this study, we examined the relationship of the portal and hepatic artery variations, which are surgically important parameters before liver transplantation, with the remnant liver volume in liver transplant donors.

MATERIAL AND METHODS

Patient selection

This study was approved by the local ethics committee (Number file: B.30.2.ATA.0.01.00/163). All donors were informed about the examination and study procedure, and their written consent was obtained. Of the healthy volunteers who presented to the radiology department of our hospital between July 2018 and January 2020 as potential transplant donors, those aged 18 to 45 years were included in the evaluation. A routine laboratory evaluation, hemogram analysis, and liver ultrasonography, and triphasic abdominal computed tomography (CT) were performed. After the laboratory and radiological evaluation, 25 patients with fatty liver, two with an abdominal aortic aneurysm, and 13 with diffuse atherosclerosis in vascular structures were excluded from the study. The data of the remaining 180 volunteers were evaluated in the study.

CT Examination

In this study, a 320-row multi-detector CT device (Aquillion ONE Vision; Toshiba Medical Systems Corporation, Otawara, Japan) was used for liver imaging. All CT scans were performed using the parameters recommended by the manufacturer (slice thickness, 0.5 mm; rotation time, 0.5 sec; and scan interval, 240 mm [480 slices, 0.5 mm]). Using a pressure injector, 1.5 mL/kg contrast enhancement (300 mg/mL iohexol) was applied at a rate of 3.5 ml/sec. Triphasic images were obtained in the arterial, portal, and hepatic vein phases. The images were evaluated on the radiological workstation (Syngo Via Console, software version 2.1, Siemens AG Medical Solutions, Erlangen, Germany) by a single radiologist (B.Y.C.) with 10 years' experience in the field. The volumetric volume assessment of the liver was undertaken using another workstation (Myrian Pro; Intrasure, Montpellier, France)

Image evaluation

The images were divided into groups according to Nakamura et al.'s anatomic classification of portal veins(10) and Michels et al.'s classification of hepatic arteries(11).

In a 3D volumetric image processor, the liver parenchymal volume was distinguished from vascular structures (Figure 1A). Along the Cantlie line used during transplantation (Figure 1B), the liver was volumetrically divided into two lobes as right and left. The percentage of the remnant left lobe relative to the total liver volume was determined after liver transplantation.

Statistical evaluation

Statistical evaluation was performed using Medcalc statistics (v. 12, Mariagerke, Belgium). The D'Agostino-Pearson test was used to determine whether the data was parametric. The left liver lobe percentages of the four groups formed according to the portal vein classification were compared using the Kruskal-Wallis test. The left liver lobe percentages of the five arterial variation groups formed according to the hepatic artery classification were compared by the Kruskal-Wallis test. The demographic data between the groups were comparatively evaluated using the t-test for age and the chi-square test for gender. P values of <0.05 were considered as statistically significant.

RESULTS

The mean age of the 180 volunteers evaluated was 28 ± 8.5 years. The number of female volunteers was 76 (41.1%). According to the Nakamura classification, there were 143 (79.4%) type A, 23 (12.7%) type B, seven (3.9%) type C, and seven (3.9%) type D patients. There was no statistically significant difference in age and gender distribution between the groups ($p = 0.54$ and 0.096 , respectively). The data of the left liver lobe volume were non-parametric and did not show normal distribution. In the portal vein groups (types A, B, C, and D), the median values [95% confidence interval (CI)] of the left liver lobe volume were found to be 36 (35-37), 35 (33-37), 33 (30.5-35.5), and 32 (30.5-38.5), respectively. There was no significant difference between the Nakamura groups in terms of the left lobe remnant percentage relative to the liver transplantation resection line ($p = 0.055$) (Figures 2 and 3). The related data and statistical results are shown in Table 1.

According to the Michels classification, the following seven groups were observed: type 1 (n = 129; 71%), type 2 (n = 12; 6.7%), type 3 (n = 24; 13%), type 4 (n = 2; 2.2%),

type 5 (n = 10; 5.6%), type 6 (n = 1; 0.6%), and type 9 (n = 2; 1.1%). There was no statistical difference in age and gender distribution between these groups (p = 0.341 and 0.132, respectively). In the hepatic artery groups (types 1, 2, 3, 4, 5, 6 and 9), the median values of the left liver lobe volume (95% CI) were found to be 36(35-37), 34.5(30.3-38.8), 35(32-37), 39, 36.5(33-42), 42, and 30. No significant difference was determined between the Michel's groups in relation to the percentage of the left lobe remnant volume relative to the liver transplantation resection line (p = 0.207) (Figures 4 and 5). Table 2 presents the related data and statistical results.

DISCUSSION

In this study, we found that the volume of the left liver lobe was not affected by the portal vein and hepatic artery variations which are important pre-transplantation parameters that determine donor safety and success of the procedure. In the literature, there are some studies that has investigated the effects and relationship of vascular structures concerning liver volume (6, 8, 9). Since 3D software providing preoperative volumetric evaluation is not available in every center, researchers have attempted to perform this evaluation using various formulas (12). In this study, we examined the relationship of the liver remnant volume with the portal vein and hepatic artery anatomical variations, which, to the best of our knowledge, has not been previously evaluated in the literature.

Some external anatomic landmarks can be used to divide the liver into the lobes however this approach is not sufficient when planning a liver surgery. Further division of the liver into the segments based on the biliary and vascular trees was introduced previously (13,14). On the other hand, various anatomical variations of the liver were reported (15) which can affect the volumetric evaluation. The development of the portal and hepatic venous system is completed at the end of 6th week of development. Embryological origin of the liver veins can conflict with Couinaud's model and segmental anatomy is closely related with an adaptive mechanism of the liver which varies with metabolic demand and perfusion (14).

Abdalla et al. performed the volumetric evaluation of the left lobe and left lobe segmentation and found differences between the patients with a certain standard deviation

(16). It was reported that these differences might be due to anatomical differences (16). Therefore, they may result from either physical differences or vascular variations between patients. In another study, Altunkaynak et al. showed that the body mass index of the patients was associated with their liver volume (17). Kokuda et al. detected differences between study groups in terms of liver volume and attributed it to the thoracic width (18). Such studies demonstrate that liver volume can vary according to the populations examined and their anatomical differences. The current study was conducted in a Turkish population, and no significant difference was observed between the groups in terms of age and gender. Therefore, we focused on the effects of vascular variations on the liver volume in our study.

Vascular variations are important anatomical markers that determine liver segmentation (19). In particular, portal and hepatic variations are vascular components used in liver segmentation. Therefore, variational changes can affect segmentation (including the right-left lobe separation) (19). Besides, the proper functioning of the portal vein, hepatic artery, and hepatic vein structures that provide tissue vascularization allows for the volumetric and functional development of the related tissues (20-22). Due to the changes in liver function caused by these functional variations in vascular structures, the differences in anatomical structures and their features may also affect the liver. Similarly, Choi et al. reported that the portal vein flow was related to the volume ratios of the right and left liver lobes; thus, they concluded that vascular drainage might affect the liver volume and functional capacity (6). However, in the same study, it was shown that the portal vein area was not associated with liver volume (6). Although no variational assessment was undertaken in this study, the portal vein area was not associated with liver volume which was an important finding revealing that the differences in anatomical structures did not affect the volume. Our results confirmed that the anatomical variations did not result in any changes in the liver volume percentage.

Our study had several limitations. First, the number of our patients was not sufficient. Because some vascular variations were not observed, and some were few in statistical terms. Our second limitation was not looking at the body mass index of our patients. However, we think that there will be no important difference in the level to affect the result since there is no difference in age and gender distribution.

CONCLUSIONS

In conclusion, variational changes in the hepatic artery and portal vein are important markers that affect decisions concerning the surgical procedure; however, they do not affect the remnant liver volume in liver transplant donors.

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Table 1. Nakamura groups, the left lobe remnant percentage relative to the liver transplantation resection line

Portal vein statistics data	Nakamura type A	Nakamura type B	Nakamura type C	Nakamura type D	p
Number of cases	143	23	7	7	
Age	30.4 ± 8.4	30.4 ± 8.7	31.2 ± 10.4	26.1 ± 6.8	0.54
Male gender (%)	85 (59%)	9 (39%)	5 (71%)	5 (71%)	0.096
Left lobe volume percentage Median (95% CI)	36 (35-37)	35 (33-37)	33 (30.5-35.5)	32 (30.5-38.5)	0.055

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CI: Confidence interval

Table 2. Michels groups, the left lobe remnant percentage relative to the liver transplantation resection line

Hepatic artery statistics data	Michels type 1	Michels type 2	Michels type 3	Michels type 4	Michels type 5	Michels type 6	Michels type 9	p
Number of cases	129 (71%)	12 (6.7%)	24 (13%)	2 (2.2%)	10 (5.6%)	1 (0.6%)	2 (1.1%)	
Age	30.8 ± 8.4	23.5 ± 5.7	30.5 ± 8.8	21 ± 2.8	30.9 ± 9.6	26 ± 0	24 ± 2.8	0.341
Male gender (%)	76 (58.9%)	5 (41.7%)	13 (32.5%)	2 (100%)	6 (60%)	1 (100%)	1 (50%)	0.132
Left lobe volume percentage Median (95% CI)	36 (35-37)	34.5 (30.3-38.8)	35 (32-37)	39 (-)	36.5 (33-42)	42 (-)	30 (-)	0.207

Nakamura T, Tanaka K, Kiuchi T, Kasahara M, Oike F, Ueda M, Kaihara S, Egawa H, Ozden I, Kobayashi N, Uemoto S: Anatom-ical variations and surgical strategies in right lobe living donor liver. Transplantation 2002;73:1896–1903.

CI: Confidence interval

Figure 1. The operation resection line (Arrows) is made to pass right of the middle hepatic vein (Arrowheads)(A). Liver volume is calculated by distinguishing it from vascular structures through the 3D volume program(B).

Figure 2. Statistical graph of percentages of left lobe percentage volume according to Nakamura classification

Figure 3. Computed tomography(A) and 3D volume rendering imaging(B) show the Nakamura type A portal bifurcation structure. The 3D volumetric program(C) demonstrated liver left lobe volume values in the donor with Nakamura Type A.

Figure 4. Statistical graph of percentages of left lobe percentage volume according to Michel's classification

Figure 5. Computed tomography(A) and 3D volume rendering imaging(B) show the Michel's type 1 hepatic artery variation. The 3D volumetric program(C) demonstrated liver left lobe volume values in the donor with Michel's type 1 hepatic artery variation.









