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Comparison of alternative arterial anastomosis site during liver transplantation when the recipient's hepatic artery is unusable

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Background: Few studies have analyzed outcomes of liver transplantation (LT) when the recipient hepatic artery (HA) was not usable.

Methods: We retrospectively evaluated the outcomes of LT performed using the different alternative sites to HA.

Results: Between 2002 and 2017, 1,677 LT were performed in our institution among which 141 (8.4%) with unusable recipient HA were analyzed. Four groups were defined according to the site of anastomosis: the splenic artery (SA group, n=26), coeliac trunk (CT group, n=12), aorta using or not the donor's vessel (Ao group, n=91) and aorta using a vascular prosthesis (Ao-P group, n=12) as conduit. The median number of intraoperative red blood cell transfusions was significantly increased in the Ao and Ao-P groups (5, 5, 8.5 and 16 for SA, CT, Ao and Ao-P group respectively, $P=0.002$), as well as fresh frozen plasma (4.5, 2.5, 10, 17 for the SA, CT, Ao and Ao-P groups respectively, $P=0.001$). Hospitalization duration was also significantly increased in the Ao and Ao-P groups (15, 16, 24, 26.5 days for the SA, CT, Ao and Ao-P groups respectively, $P<0.001$). The occurrence of early allograft dysfunction (EAD) ($P=0.07$) or arterial complications ($P=0.26$) was not statistically different. Level of factor V, INR, bilirubin and creatinine during the 7th postoperative days (POD) was significantly improved in the SA group. No difference was observed regarding graft ($P=0.18$) and patient ($P=0.16$) survival.

Conclusions: In case of unusable HA, intraoperative and postoperative outcomes are improved when using the SA or CT compared to aorta.

Keywords: Liver transplantation (LT); hepatic artery (HA); arterial anastomosis; splenic artery (SA)

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Introduction

Liver transplantation (LT) is the best treatment for end-stage liver disease and primary liver tumor. Despite surgical, medical and radiological improvements in diagnosis and treatment, arterial anastomosis remains a challenge with an incidence of hepatic artery thrombosis (HAT) varying from 4% to 15% (1-3) with a significant impact on graft and patient outcomes (4,5).

Arterial anastomosis is usually performed between the graft and the recipient's hepatic artery (HA). However, when the recipient's HA is unusable (owing to complete thrombosis, intimal dissection, small size or inadequate flow), alternative sites such as aorto-hepatic anastomosis with or without conduits (6,7), coeliac trunk (CT) (8,9) or splenic artery (SA) (10-12) are used.

To date, few studies have compared the intraoperative and postoperative outcomes of these different alternative sites. Consequently, the choice usually depends on the surgeon's experience or habits.

The aim of this study was to analyze and compare the outcomes of alternative anastomosis sites in case of unusable recipient HA during LT. We present the following article in accordance with the STROBE reporting checklist (available at <https://hbsn.amegroups.com/article/view/10.21037/hbsn-20-10/rc>).

Methods

Patient selection

From January 2002 to December 2017, all adult LT or retransplantations performed in a high-volume LT center in which native recipient HA was not used (n=146) were retrospectively analyzed. Patients with associated transplantation of another organ (n=1), transplanted with a split liver graft (n=2), pediatric donor (n=1) or donor after cardiac death (n=1) were excluded from the analysis (*Figure 1*).

Depending on the recipient site of arterial anastomosis and reconstruction technique, 4 groups were defined: (I) anastomosis on the SA (SA group) usually in a side-to-end fashion; (II) on the CT (CT group); (III) on the aorta using or not the donor's vessel (Ao group) or (IV) using a vascular prosthesis (Ao-P group) as conduit. No patients had previous splenectomy.

Patients transplanted using the recipient HA (n=1,531) were also analyzed as the reference group after excluding transplantations with associating another organ (n=67),

split liver graft (n=55), pediatric donor (n=2) or donor after cardiac death (n=12).

The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study received approval from local ethics committee (comité d'éthique du CHU de Rennes NO21.49) and non-opposition consent was obtained from all participants.

Surgical procedure and postoperative care

After standard (with systematic donor vessels) procurement, liver grafts were usually preserved with Celsior® (before 2016) or custodiol® (after 2016).

All patients had orthotopic LT with inferior vena cava preservation. Briefly, after standard wound incision and exposition, the liver pedicle was first dissected. The native liver was removed and careful hemostasis was performed. Graft implantation started with a side-to-side caval anastomosis followed by an end-to-end portal vein anastomosis. The graft was then vascularized prior to arterial anastomosis which was usually an end-to-end anastomosis between the recipient's proper HA (at the junction between the left and right branch) and the graft's common artery in order to have the largest anastomosis surface and the shortest length. The biliary anastomosis was usually performed by an end-to-end anastomosis between the main bile duct of the recipient and the graft intubated with a T-tube.

Regarding the arterial anastomosis, the decision to not use recipient HA was made preoperatively in case of known thrombosis or decided intraoperatively by the surgeon after evaluation of the quality of the HA and its flow. The choice of an alternative arterial site was also made by the surgeon according to his or her preferences. The SA anastomosis was performed in a side-to-end fashion while the CT anastomosis was performed in an end-to-end fashion with preservation of the splenic and gastric vessel when possible (*Figure 2*). When use of a conduit was required, allograft (usually the donor's iliac vessels) was preferred. In case of massive arteriosclerosis or insufficient length, a vascular prosthesis (GoreTex®) was used. After arterial declamping, arterial flow was systematically evaluated by intraoperative Doppler ultrasonography with calculation of resistivity index.

After the procedure, aspirin was systematically administered as soon as possible as well as standardized immunosuppression (associating calcineurin inhibitor,

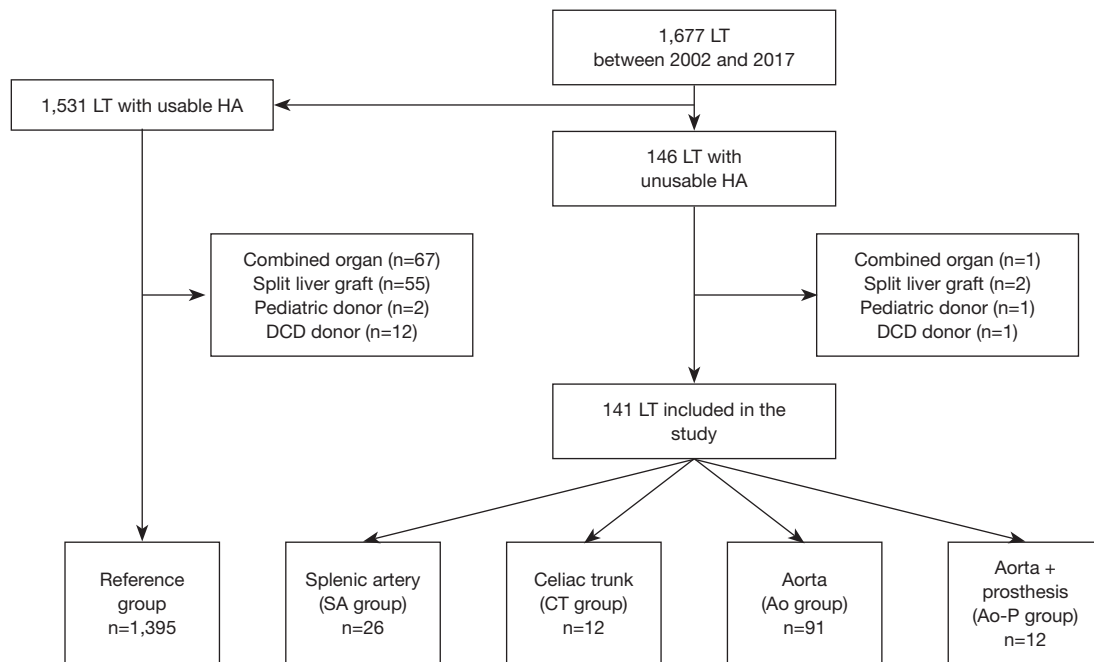


Figure 1 Flow chart. DCD, donation after circulatory death; HA, hepatic artery; LT, liver transplantation.

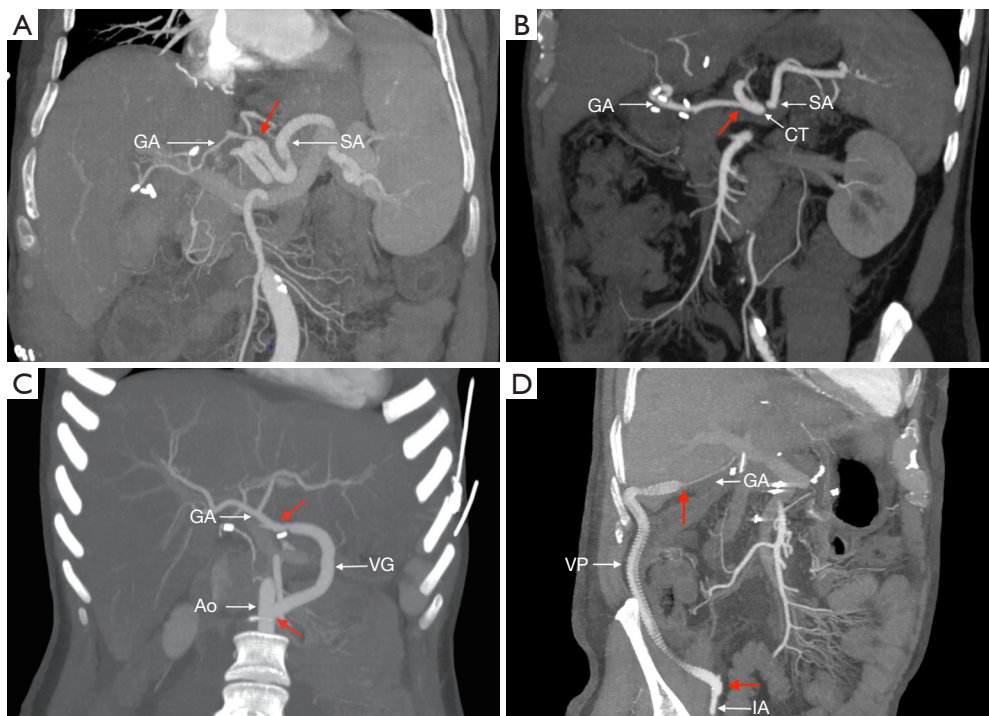


Figure 2 Arterial anastomosis using alternative implantation site on recipient. (A) Side-to-end anastomosis on recipient's splenic artery; (B) end-to-end anastomosis on celiac trunk with preservation of left gastric and splenic artery; (C) implantation on recipient's aorta using vascular graft (donor's iliac vessel); (D) implantation on recipient's right iliac artery using a vascular prosthesis. Red arrow: anastomosis site. Ao, recipient aorta; CT, coeliac trunk; GA, graft artery; IA, iliac artery; SA, splenic artery; VG, vascular graft (donor's iliac vessel); VP, vascular prosthesis.

usually tacrolimus, mycophenolate mofetil and a short course of corticosteroids).

Systematic doppler ultrasonography was performed on postoperative days (POD) 1 and 7 and if hepatic dysfunction or vascular complication was suspected. When suspected, computed tomography was performed to confirm vascular complications and treatment (medical, radiological or surgical) was discussed in multidisciplinary meetings. Arterial thrombosis occurring before POD7 were operated for urgent revascularization, while thrombosis occurring after this period were followed (when asymptomatic) and listed for retransplantation when they became symptomatic (mostly biliary complications). Arterial stenosis was treated by radiological intervention.

No significant modification regarding the surgical procedure or postoperative medical care was observed during the study period.

Clinical and biological parameter collection

Clinical and biological data were retrospectively collected from a prospective database. The following parameters were analyzed:

- ❖ Donor and recipient demographic data (age, sex and BMI), MELD score, primary LT or retransplantation, use of an extended criteria donor (ECD) graft (defined by the presence of at least one of the following criteria: age >65 years; BMI >30; ICU stay >7 days prior procurement; natremia >155 mmol/L; liver enzymes 3 times higher the normal value (i.e., aspartate aminotransferase (ASAT) blood level >150 IU/mL, alanine aminotransferase (ALAT) blood level >170 IU/mL) and cause of donor death;
- ❖ Intraoperative data: cold ischemia time, delay to arterialization (defined as the time between portal and arterial revascularization), operative time, transfusion, and cause of unusable recipient HA;
- ❖ Postoperative data: incidence of arterial complications, duration of hospitalization, Clavien-Dindo classification, patient and graft survival;
- ❖ Biological data: incidence of early allograft dysfunction (EAD) defined according to Olthoff *et al.* (13) and evolution of liver function and creatinine level during the 7th POD.

Statistical analysis

Comparative statistical analysis was made between the four

study groups in order to compare them.

Quantitative variables were expressed as median with extreme values and compared using the Kruskal-Wallis test. Qualitative variables were expressed as number and percentage and compared using the chi-squared test. Survival analysis was performed with Kaplan-Meier curves and compared with the log-rank test. Repeated-measures analysis (i.e., evolution of biological parameters) during the early postoperative course was performed using a linear mixed-effects model (14).

All statistical analyses were performed with R software version 3.4.3. using the lme4 package version 1.1.15 for linear mixed-effects analysis. A $P < 0.05$ value was considered to be statistically significant.

Results

Population characteristics (Table 1)

During the study period, 1,677 LT were performed at our center. After the selection process, 141 (8.4%) LT performed with unusable recipient HA, were analyzed. The SA (SA group) was used in 26 (18.4%) cases, the CT (CT group) in 12 (8.5%) cases, the aorta (Ao group) in 91 (64.5%) cases and the aorta with use of a vascular prosthesis (Ao-P group) in 12 (8.5%) cases (*Figure 1*). Median age was 54 [17–73] years with 65.2% (n=92) men without differences between the groups.

In 74 cases (52.5%), the procedure was a retransplantation (occurring after a median delay of 7.6 months and in 20 (37%) cases within the first month) and was secondary to arterial thrombosis in 36 cases (49%).

The median MELD score was 18.5 without significant differences between the groups.

The cause for not using recipient HA was reduced caliber or insufficient flow in 54 (35.1%) cases, HAT in 79 (59.4%) cases and intimal dissection in 8 (5.5%) cases.

In the Ao group, the implantation site on the aorta was infrarenal in 57% (n=52), supraceliac in 36% (n=33) and on the iliac artery in 6% (n=6). A conduit was used in 61.5% (n=56). In the Ao-P group, implantation was on the infrarenal aorta in 50% (n=6), supraceliac in 8% (n=1) and on the iliac artery in 42% (n=5). Median follow-up was 61 months for the entire cohort.

The reference group (i.e., using the recipient HA) consisted on 1,395 patients and as expected present major differences compared to the four study groups especially regarding rate of retransplantation.

Table 1 Patient and donor characteristics

Variables	SA group (n=26)	CT group (n=12)	Ao group (n=91)	Ao-P group (n=12)	P value	Reference group (n=1,395)
Recipient characteristic						
Gender (male)	16 (61.5%)	9 (75%)	59 (64.8%)	8 (66.7%)	0.87	1,079 (77.3%)
Age (years)	58 [19–66]	56.5 [32–67]	52 [17–73]	54.5 [38–65]	0.10	57 [15–73]
BMI	24.6 [13.6–33.6]	27.1 [19.8–34.4]	25.3 [17.4–40.4]	25.8 [19.7–37.1]	0.90	26.3 [13.6–47.2]
MELD	18 [6.5–37.9]	13.7 [6.4–40]	19 [5.8–40]	22.1 [17.1–37.1]	0.1	15 [5.4–40]
Creatinine (μmol/L)	81 [43–241]	69 [59–175]	89 [37–636]	91 [53–200]	0.32	74 [28–787]
Dialysis	1 (3.8%)	0 (0%)	7 (7.7%)	1 (8.3%)	0.70	29 (2.1%)
Retransplantation rate	13 (50%)	3 (25%)	54 (59.3%)	4 (33.3%)	0.07	86 (6.1%)
Cause of unusable HA					<0.001	
Small caliber-low flow	7 (26.9%)	8 (66.7%)	33 (36.3%)	6 (50%)		NA
Thrombosis	13 (50%)	4 (33.3%)	56 (61.5%)	6 (50%)		NA
Dissection	6 (23.1%)	0	2 (2.2%)	0		NA
Donor characteristics						
Gender (male)	15 (57.7%)	9 (75%)	53 (58.2%)	8 (66.7%)	0.68	811 (58.1%)
Age	44.5 [20–86]	53 [11–82]	48 [14–80]	58.5 [16–79]	0.37	55 [10–93]
BMI	23.44 [16.9–39.8]	24.20 [18.1–31.6]	23.05 [16.1–32]	27.21 [19.6–32.5]	0.37	24.69 [13.9–54.4]
ECD graft	16 (61.5%)	6 (50%)	51 (56%)	8 (66.7%)	0.81	881 (63.2%)
Cause of donor death					0.83	
Traumatic	8 (30.8%)	5 (41.7%)	34 (37.4%)	5 (41.7%)		365 (26.2%)
Vascular	16 (61.5%)	6 (50%)	44 (48.4%)	7 (58.3%)		818 (58.6%)
Anoxic	2 (7.7%)	1 (8.3%)	8 (8.8%)	0 (0%)		168 (12%)
Other	0 (0%)	0 (0%)	5 (5.5%)	0 (0%)		43 (3.1%)
Study period					0.1	
2002–2007	10 (15%)*	2 (3%)*	46 (71%)*	7 (11%)*		472 (100%)
2008–2012	9 (19%)*	4 (8.5%)*	30 (64%)*	4 (8.5%)*		393 (100%)
2013–2017	7 (24%)*	6 (21%)*	15 (52%)*	1 (3%)*		530 (100%)
Follow up (months)	70 [0–188.5]	32.6 [0.1–184.3]	62 [0–192.3]	30.1 [0.9–192.9]	0.59	61.3 [0–205.9]

*, % calculated for each study period. Ao, aorta group; Ao-P, aorta + vascular prosthesis; BMI, body mass index; CT, celiac trunk; ECD, extended criteria donors; SA, splenic artery.

Intraoperative outcomes (Table 2)

Median operative time was 405 minutes and was close to being significantly increased in the Ao-P group with a median duration of 437.5 min ($P=0.05$). No differences

were noted regarding cold and warm ischemia times.

The median number of transfused red blood cell units was significantly increased in the Ao and Ao-P groups (8.5 and 16 respectively, $P=0.002$). In the same way, the

Table 2 Intraoperative and postoperative outcomes

Variables	SA group (n=26)	CT group (n=12)	Ao group (n=91)	Ao-P group (n=12)	P value	Reference group (n=1,395)
Intraoperative outcomes						
Operative duration (min)	413 [240–862]	307 [185–500]	405 [160–830]	437.5 [285–800]	0.05	360 [134–1,020]
Cold ischemia (min)	568 [297–1,046]	414.5 [251–679]	540 [227–1,405]	605.5 [319–942]	0.12	526 [166–1,393]
Delay to arterialization (min)	48 [20–140]	45 [28–84]	48 [21–253]	53 [32–73]	0.58	48 [0–190]
Transfusion						
Red blood cells	5 [0–23]	5 [0–12]	8.5* [0–60]	16* [2–31]	0.002	5 [0–60]
Fresh frozen plasma	4.5 [0–21]	2.5 [0–16]	10* [0–60]	17* [3–54]	0.001	5 [0–40]
Platelet count	0.5 [0–2]	0 [0–2]	0.5 [0–7]	1 [0–4]	0.13	0 [0–4]
Postoperative outcomes						
ICU stay (days)	4 [0–25]	4.5 [2–15]	6 [1–160]	7 [3–63]	0.05	4 [0–199]
Hospital stay (days)	15 [1–72]	16 [3–33]	24* [1–270]	26.5* [3–63]	<0.001	19 [0–368]
Clavien-Dindo grade >2	7 (26.9%)	5 (41.7%)	38 (41.8%)	7 (58.3%)	0.30	340 (24.4%)
Arterial complications	6 (23.1%)	5 (41.7%)	20 (22%)	5 (41.7%)	0.26	126 (9%)
Stenosis	2 (7.7%)	3 (25%)	4 (4.4%)	1 (8.3%)	0.07	56 (4%)
Thrombosis	4 (15.4%)	2 (16.7%)	15 (16.5%)	4 (33.3%)	0.53	70 (5%)
EAT	2 (7.7%)	2 (16.7%)	8 (8.8%)	2 (16.7%)	0.68	35 (2.5%)
EAD	6 (23.1%)	5 (41.7%)	41 (45.1%)	8 (66.7%)	0.07	328 (23.5%)
Survival outcomes						
90-day graft survival	22 (84.6%)	9 (75%)	74 (81.3%)	7 (58.3%)	0.26	1,250 (89.6%)
1-year graft survival	21 (80.8%)	9 (75%)	65 (71.4%)	6 (50%)	0.27	1,182 (84.7%)
5-year graft survival	18 (69.2%)	6 (50%)	50 (54.9%)	4 (33.3%)	0.21	1,033 (74.1%)
90-day patient survival	23 (88.5%)	11 (91.7%)	79 (86.8%)	8 (66.7%)	0.24	1,292 (92.6%)
1-year patient survival	22 (84.6%)	10 (83.3%)	72 (79.1%)	7 (58.3%)	0.30	1,224 (87.7%)
5-year patient survival	19 (73.1%)	8 (66.7%)	60 (65.9%)	5 (41.7%)	0.30	1,082 (77.6%)

*, significantly different. Ao, aorta group; Ao-P, aorta + vascular prosthesis; CT, celiac trunk; ICU, intensive care unit; EAD, early allograft dysfunction; EAT, early artery thrombosis (<30 days); SA, splenic artery.

median number of transfused frozen plasma units was significantly increased in the Ao and Ao-P groups (10 and 17 respectively, $P=0.001$).

Evolution of biological parameters (Figure 3)

While the evolution of liver enzyme levels was not statistically different between the four study groups, linear mixed-effect analysis revealed that bilirubin levels were significantly increased in the Ao group ($P=0.03$)

compared with the SA group and INR and factor V level were significantly increased in the Ao-P group ($P=0.04$ and $P=0.007$ respectively) compared with the SA group (Table 3).

Accordingly, the EAD rate was 23.1% in the SA group, 41.7% in the CT group, 45.1% in the Ao group and 66.7% in the Ao-P group, but without a significant difference ($P=0.07$). However, the creatinine level [which was similar preoperatively (Table 1)] was significantly increased in both the Ao ($P=0.007$) and Ao-P groups ($P=0.008$) compared with the SA group (Table 3).

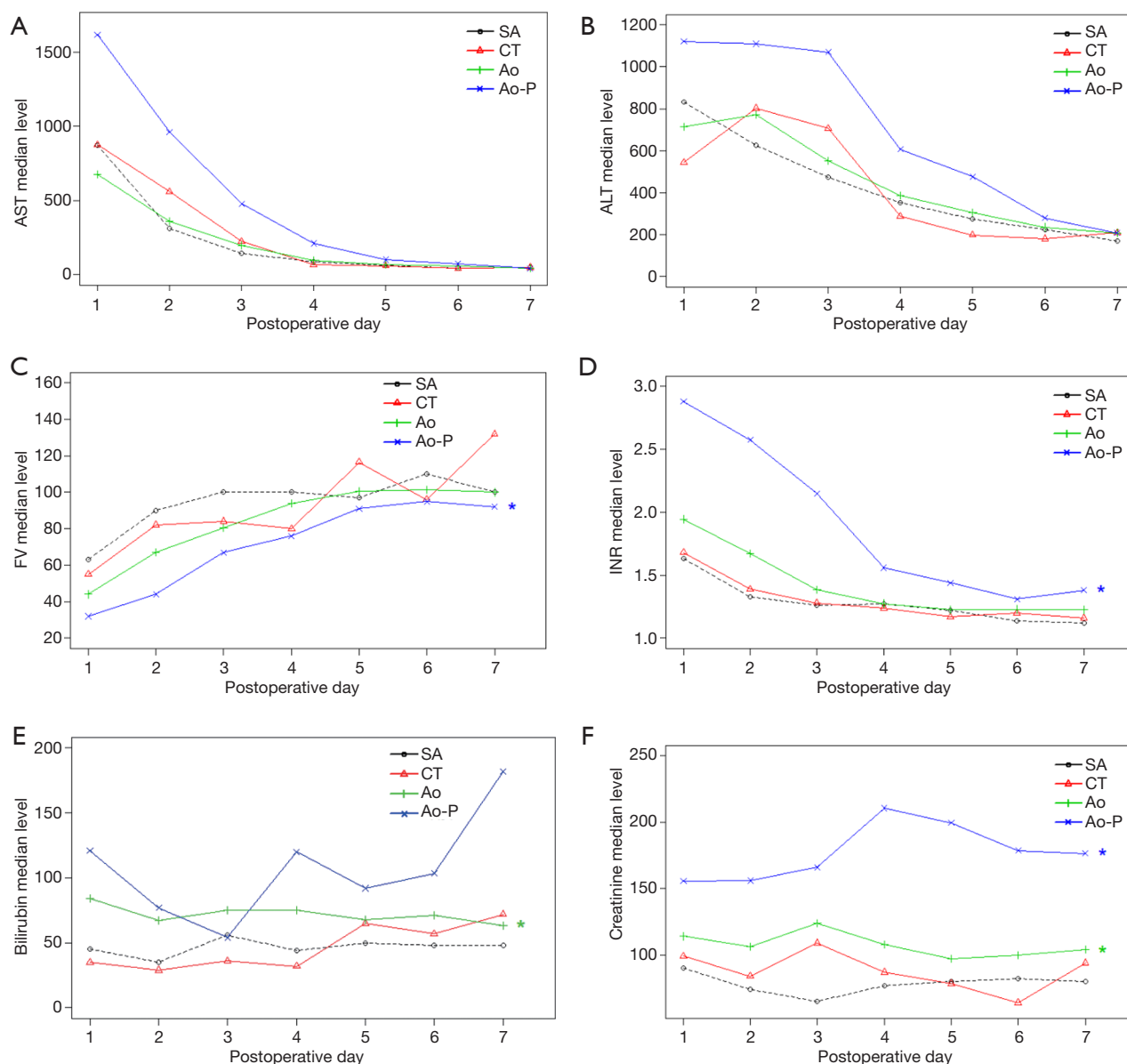


Figure 3 Biological parameter evolution. (A) AST level; (B) ALT level; (C) factor V level; (D) INR level; (E) total bilirubin level; (F) serum creatinine level. The * symbol means a significant difference compared with the splenic artery group. SA, splenic artery; CT, coeliac trunk; Ao, aorta group; Ao-P, aorta + vascular prosthesis group; AST, aspartate aminotransferase; ALT, alanine aminotransferase; INR, international normalized ratio.

Postoperative outcomes (Table 2)

Hospitalization duration was significantly higher in the Ao and Ao-P groups (24 and 26.5 days respectively, $P < 0.001$). The incidence of significant complications (i.e., Clavien-Dindo grade > 2) was observed in 26.9% in the SA group, 41.7% in the CT group, 41.8% in the Ao group and 58.3% in the Ao-P group, without significant differences ($P = 0.3$).

Overall, 36 (25.5%) recipients had an arterial event (stenosis and/or thrombosis) without a difference between groups ($P = 0.26$). Early HAT (i.e., < 30 POD) rate was 7.7%, 16.7%, 8.8% and 16.7% for the SA, CT, Ao and Ao-P groups respectively ($P = 0.68$).

The 1-year graft and patient survival rate was 80.8% and 84.6% for the SA group, 75% and 83% for the CT group, 71.4% and 79.1% for the Ao group and 50% and 58.3%

Table 3 Analysis of biological parameter evolution during the first 7 postoperative days

Variables	Mixed linear model (P value)
AST level	
SA group (= reference)	
CT group	0.15
Ao group	0.29
Ao-P group	0.51
ALT level	
SA group (= reference)	
CT group	0.14
Ao group	0.43
Ao-P group	0.51
Bilirubin level	
SA group (= reference)	
CT group	0.87
Ao group	0.03
Ao-P group	0.1
INR level	
SA group (= reference)	
CT group	0.54
Ao group	0.1
Ao-P group	0.04
Factor V level	
SA group (= reference)	
CT group	0.93
Ao group	0.05
Ao-P group	0.007
Creatinine level	
SA group (= reference)	
CT group	0.06
Ao group	0.007
Ao-P group	0.008

SA, splenic artery; CT, celiac trunk; Ao, aorta group; Ao-P, aorta + vascular prosthesis group.

for the Ao-P group ($P=0.27$ and $P=0.30$, respectively). The cause of graft loss at 1-year was an hemorrhagic or vascular complication in 15% ($n=4$) for SA group (resulting

in patient death in 1 case), 25% ($n=3$) for CT group (resulting in patient death in 0 case), 11% ($n=10$) in Ao group (resulting in patient death in 2 cases) and 16.7% ($n=2$) in Ao-P group (resulting in patient death in 0 case) and a severe sepsis or multivisceral failure in 0% for SA group, 0% for CT group, 11% ($n=10$) for Ao group (resulting in patient death in 10 cases), 25% ($n=3$) for Ao-P group (resulting in patient death in 3 cases), $P=0.27$.

The 5-year graft and patient survival rate was 69.2% and 73.1% for the SA group, 50% and 66.7% for the CT group, 54.9% and 65.9% for the Ao group and 33.3% and 41.7% for the Ao-P group ($P=0.21$ and $P=0.30$, respectively). Long-term graft ($P=0.18$) and patient ($P=0.16$) survival was also not different (*Figure 4*).

When comparing with the reference group, the graft survival was significantly worst for CT group ($P=0.02$), Ao group ($P=0.001$) and Ao-P group ($P<0.001$) while it was not different for SA group ($P=0.31$). Regarding the patient's survival, there was a significant difference between the reference group and the Ao-P group ($P=0.004$) while it was not different for Ao group ($P=0.096$), CT group ($P=0.24$) and SA group ($P=0.61$).

Discussion

During LT, HA reconstruction remains a technical challenge and complex arterial reconstruction (typically when recipient HA cannot be used) is a well-known risk factor of arterial thrombosis (14-17).

In the present study, we found that using the recipient SA or CT was associated with lower intraoperative transfusion rates and better graft function recovery, resulting in lower durations of hospitalization. On the other hand, implantation on the aorta using a vascular prosthesis as conduit was associated with higher intraoperative transfusions and worse postoperative graft and renal function resulting in longer stays in the intensive care unit and the hospital. However, we did not find a significant difference regarding the rate of arterial complications although the occurrence of arterial stenosis increased in the CT group (25%, $P=0.08$) and thrombosis in the Ao-P group (33.3%, $P=0.53$). In the same way, we did not find a significant difference regarding graft or patient survival between the four study groups although visual analysis of the Kaplan-Meier curves suggests a worse outcome with a vascular prosthesis. This absence of statistical significance is probably due to the limited number of patients in the different groups, resulting in a lack of power. However,

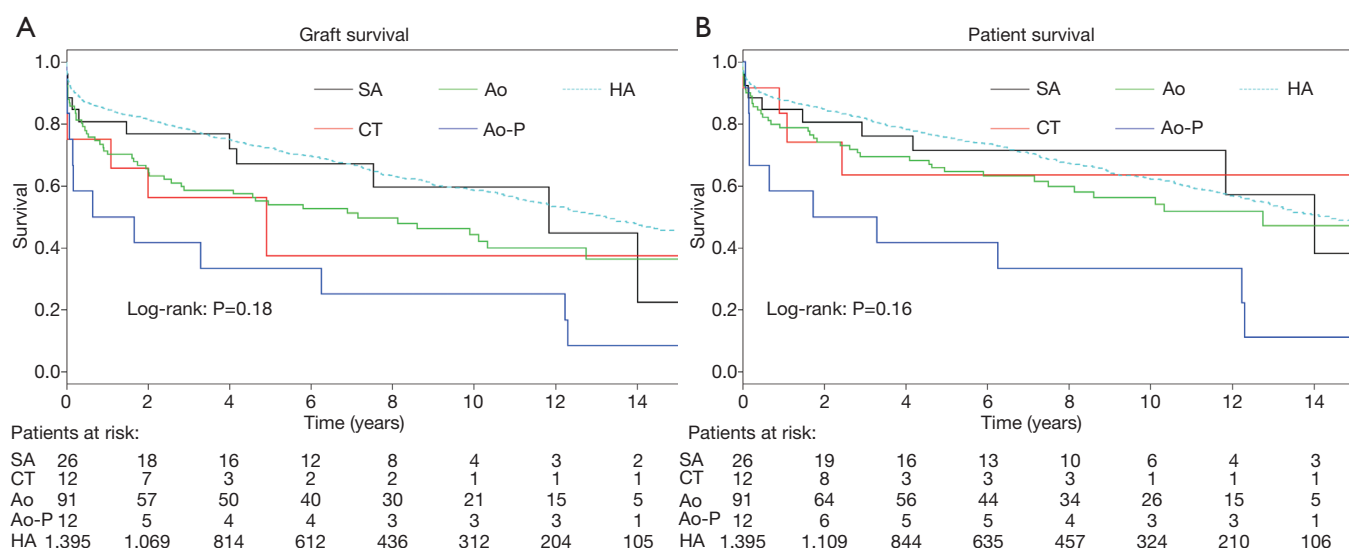


Figure 4 Long-term graft (A) and patient (B) survival. SA, splenic artery; CT, coeliac trunk; Ao, aorta group; Ao-P, aorta + vascular prosthesis group; HA, hepatic artery.

when comparing with the reference group's (n=1,395) outcome, only patients in the SA group shown similar outcomes. Indeed, the CT, Ao and Ao-P group presented worst graft survival while the Ao-P group also presented worst patient survival.

Typically, when a recipient HA is unusable, the preferred arterial implantation site is the aorta with use of the donor iliac vessels as conduits if needed (18-20). Recently in a systematic review, Reese *et al.* (21) found that this option was associated with a higher incidence of early arterial thrombosis (7% *vs.* 2%, $P=0.001$) as well as worse grafts ($HR=1.38$, $P<0.001$) and patient survival ($HR=1.57$, $P=0.009$) compared with the non-conduit groups. In their study, they reported 5-year survival at 54% and 67% for grafts and patients respectively which is similar to our findings with 5-year survival at 54.9% and 65.9% for grafts and patients respectively. Moreover, we found that these results were even worse with vascular prosthesis instead of allograft with 5-year survival at 33.3% and 41.7% for grafts and patients respectively. Even if it did not reach significance, we believe that our findings plead for avoiding this method if possible and explain the decrease rate of Ao or Ao-P anastomosis performed during the study period contrary to SA and CT (Table 1).

Other authors have described alternative implantation sites such as the CT (8,9,22) or SA (5,10,11,23,24) with interesting results (Table 4). Indeed, El-Hinnawi *et al.* (9) reported a series of 9 cases (5 pediatric and 4 adults)

using the CT without major postoperative complications. Dokmak *et al.* (8) reported a series of 7 patients using CT. In their study, one patient died in the early postoperative course and among the 6 remaining patients, patent arterial flow was found in 100% with a follow-up of 6 to 15 months. Stange *et al.* (5) reported a series of 133 anastomoses on the SA with 0.8% HAT. Figueras *et al.* (11) reported a series of 23 patients using the SA and reported 3-year graft and patient survival of 72% and recently, Llado *et al.* (26) reported a series of 54 patients using SA and found comparable outcomes to those having a standard HA reconstruction. Our results are in line with previous reports with similar favorable intraoperative and postoperative outcomes when using SA or CT (Table 4). However, the long-term outcomes in our study appear to be in favor of using SA instead of CT in term of arterial complications.

Indeed, the SA artery is usually dilated owing to frequent splenomegaly in cirrhotic patients and can be easily controlled after its emergence above the pancreatic head, thereby representing a safe and good alternative for arterial anastomosis. On the contrary, control of the supraceliac or infrarenal aorta could be difficult in cirrhotic patients owing to concurrent portal hypertension especially in obese recipients. Therefore, avoiding aortic clamping and unnecessary dissection (leading to less bleeding and transfusion) could represent an additional benefit by preserving recipient hemodynamics, which could already be unstable after graft revascularization, and reducing

Table 4 Main studies reporting alternative arterial implantation site outcome

Reference	Implantation site	EAT (%)	Graft survival	Patient survival
Cherqui <i>et al.</i> , (10) [1994]	SA: 7	0	–	100% at 30-month
Figueras <i>et al.</i> , (11) [1997]	SA: 23	0	78% at 1-year; 72% at 3-year	78% at 1-year; 72% at 3-year
	Ao: 12	17	74% at 1-year; 74% at 3-year	80% at 1-year; 80% at 3-year
Stange <i>et al.</i> , (5) [2003]	SA: 133	0.8	–	–
	Ao: 95	9.5	–	–
D’Albuquerque <i>et al.</i> , (24) [2007]	SA: 4	0	–	–
Vanderlan <i>et al.</i> , (12) [2008]	SA: 7	14	86% at 9-month	100% at 9-month
El-Hinnawi <i>et al.</i> , (9) [2013]	CT: 9	0	83% at 1, 3, 5 years	83% at 1, 3, 5 years
Dokmak <i>et al.</i> , (8) [2015]	CT: 7	0	–	86% at 15-month
	SA: 2	0	–	100% at 14-month
	Ao: 2	0	–	100% at 14-month
Kazemi <i>et al.</i> , (25) [2017]	SA: 17	–	88% at 1-year; 82% at 5-year	94% at 1-year; 88% at 5-year
	Ao: 76	–	78% at 1-year; 62% at 5-year	97% at 1-year; 69% at 5-year
Llado <i>et al.</i> , (26) [2019]	SA: 54	3.7	64% at 5-year	67 % at 5-year
Beaurepaire <i>et al.</i> , [2019]	SA: 26	7.7	80% at 1-year; 69% at 5-year	84% at 1-year; 73% at 5-year
	CT: 12	16.7	75% at 1-year; 50% at 5-year	83% at 1-year; 67% at 5-year
	Ao: 91	8.8	71% at 1-year; 55% at 5-year	79% at 1-year; 65% at 5-year
	Ao-P: 12	16.7	50% at 1-year; 33% at 5-year	58% at 1-year; 41% at 5-year

Ao, aorta group; Ao-P, aorta + vascular prosthesis; CT, celiac trunk; EAT, early artery thrombosis (<30 days); SA, splenic artery.

delay between portal and arterial revascularization as well as operative time (25). These benefits could explain the improved graft and kidney function observed in the SA and CT groups during the postoperative course.

Thus, if preoperative imaging is compatible, we believe that SA or CT should be considered firstly, instead of implantation on aorta like stated by other authors (7).

Our results must be interpreted with caution. First, the retrospective nature of the study and the arbitrary decision of the surgeon regarding the choice of the arterial site inevitably induce a methodological and selection bias although there was no difference regarding the demographic data. Moreover, the limited population size (with a large majority in group Ao) as well as the analysis of 4 different groups led to a lack of power that could explain the absence of significant differences especially regarding survival outcomes or arterial complications.

To our knowledge, the present study is the largest monocentric study (n=141) that compares all alternative

arterial sites of anastomosis along with the largest series using a aorto-hepatic conduit with a vascular prosthesis (n=12) (6,27–29). Therefore, our study represents the “real-life” practice of a high volume LT center. Moreover, our results are in line with the other reports from previous studies which strengthened our findings (*Table 4*). In addition, our study is the first to analyze and report the postoperative biological parameters of graft and kidney function which should also be interpreted with cautious since it may be influenced by multiple factors.

In any case, our results must be confirmed in a larger multicentric series since a prospective randomized study may be difficult to set up.

In conclusion, we believe that using the SA or the CT (despite an elevated stenosis rate in our series) is a safe and efficient alternative. When an aorto-hepatic anastomosis should be performed, use of a vascular prosthesis must be avoided if possible. Our results must be confirmed in a larger series.

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Footnote

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