Left Renal Vein Division for Juxtarenal Aortic Exposure: Influence on Renal Function and Role of the Communicating Lumbar Vein

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Abstract

Background In this study, we evaluate the outcome of renal function in patients undergoing juxtarenal abdominal aortic aneurysm repair with or without division of the left renal vein with special focus on the role of the communicating lumbar vein.

Methods A retrospective analysis of prospectively collected data of 110 patients undergoing elective juxtarenal abdominal aortic aneurysm repair between 2000 and 2018 was performed. The demographic characteristics and comorbidities were reviewed in detail and the renal function was analysed pre- and post-operatively. The cohort of patients was split into group A (left renal vein divided) and B (left renal vein mobilised). Group A was further sub-analysed regarding the presence of a communicating lumbar vein on preoperative imaging data (group A+ = vein present, group A− = no communicating lumbar vein present).

Results The patients were matched well regarding their demographic characteristics and comorbidities. In the analysis of renal function, no statistically significant difference could be detected between group A and B. In the sub-analysis of group A, the group with a communicating lumbar vein (group A+) turned out to have a significantly better renal function in the long term (sCrea 0.87 vs. 1.51; p = 0.016).

Conclusion Ligation of the left renal vein is a safe procedure in surgery of juxtarenal aortic aneurysms regarding the outcome of the renal function. A communicating lumbar vein between the left renal vein and the left ascending lumbar vein seems to play a key role to provide venous drainage after division of the left renal vein.

Introduction

Juxtarenal aortic aneurysms are rare with an incidence of 2.2/100,000 and account for 10.5% of abdominal aortic aneurysms [1]. During open repair of juxtarenal aortic aneurysms, the left renal vein (LRV) is dissected and frequently divided to achieve adequate exposure of the juxtarenal aortic segment. According to the literature, the left renal vein division (LRVD) rate ranges from 1.3 to 18.8% during this procedure [2,3]. After LRVD, a significant increase in serum creatinine values is reported one day and one month after an elective juxtarenal aneurysm repair [4]. Also, West et al. [5] documented a significant association between renal impairment and LRVD after elective surgery.
aortic aneurysm repair. Moreover, AbuRahma et al. [6] described increased perioperative levels of creatinine. Wang et al. found decreased glomerular filtration rates (GFR) and, as observed by Mehta et al. [2], increased creatinine values immediately after surgery which recovered in the long-term. Samson [7] described no changes in serum creatinine and GFR during the first postoperative days, at discharge and at 12 months postoperatively compared to preoperative values.

The literature seems quite contradictory and no reasonable explanation for deteriorated renal function after LRVD in one group of patients vs. normal function in the other group has been identified so far.

Typically, the venous system is characterised by a large number of collateral vessels, which provide alternative pathways to allow blood flow if vessels are obstructed or ligated like the left renal vein during juxtarenal aneurysm repair. Gonadal, adrenal and left ascending lumbar veins are the venous collaterals of the left renal vein [8–10]. A communicating vein, linking the ascending lumbar vein to the left renal vein is described by some authors [8,11]. The incidence of this communicating lumbar vein (CLV) in contrast-enhanced CT studies is approximately 35% [8]. In case of division or obstruction of the LRV, venous collaterals (including the CLV) and the left renal stump may enlarge due to the diversion of blood flow from the left kidney [8]. However there is no data on changes in renal function taking into consideration this anatomical variation [12].

The aim of this study was to investigate the perioperative renal function of patients undergoing elective open juxtarenal aortic aneurysm repair. The follow-up period was two years. The effects of LRVD were compared to mobilization only or reconstruction of the LRV. In addition, a sub-analysis on a potentially protective effect of a communicating lumbar vein on the renal function after LRVD was performed. The images shown in this manuscript are provided in agreement with the respective patient.

Materials and methods

From January 2000 to December 2018, 110 patients underwent elective aortic open surgery for juxtarenal abdominal aortic aneurysm repair at our institution. Data were prospectively collected and retrospectively analysed. Patients with infrarenal abdominal aortic aneurysms and patients with ruptured abdominal aortic aneurysms were excluded from analysis. Patients were divided in two groups: group A (n = 81 patients) with suprarenal aortic cross clamping and division of the LRV; group B (n = 29 patients) with suprarenal aortic cross clamping and LRV mobilisation without division. The cross-clamping times were comparable between the groups. The left renal vein was divided close to the inferior vena cava, far from the collateral veins (adrenal, gonadal and communicating vein to the left ascending lumbar vein) to preserve venous drainage. If the LRV was mobilised, the suprarenal and gonadal veins were divided. The patients were matched well according to risk factors, interestingly there was a trend towards smoking in group A. In group B coronary artery disease was (insignificantly) more common (Tables 1, 2).

Furthermore, to identify a possible variable to stratify the risk for deterioration of renal function after suprarenal cross clamping and division of the LRV, we focused on the
anatomy of the left renal vein in the preoperative contrast enhanced CT scan images. Special attention was given to the presence of a communicating lumbar vein between the left renal vein and the left ascending lumbar vein that could serve as an additional drainage pathway after division of the left renal vein.

Group A (n = 81 patients) was further divided into two subgroups: group A+ (communicating lumbar vein present on CT-scan preoperatively) with n = 25 patients and group A– (no communicating lumbar vein present on CT-scan preoperatively) with n = 56 patients.

For both comparisons of groups A and B as well as of groups A+ and A–, patient characteristics, risk factors and perioperative parameters including intensive care unit stay and 30 day mortality were analysed. The following parameters were calculated: mean values of serum creatinine preoperatively and at postoperative days 10 and 15 as well as at discharge and 12 and 24 months after the operation. In addition to that, the GFR was calculated using the CDK-EPI formula preoperatively, at discharge and 12 as well as 24 months postoperatively [3,13,14]. Dichotomous variables were compared using the chi-square test or Fisher’s exact test where indicated, continuous variables were compared using the student’s t-test or Mann Whitney U Test as appropriate. For all statistical analysis a p-value <0.05 was considered significant. All statistical analyses were performed using the statistical analysis software package SPSS (26.0, IBM, Boston, USA version).

Ethical statement

All procedures performed in this study were in accordance with the ethical standards of the institutional research committee and with the 1964 Helsinki Declaration and its later amendments. The present study was approved by the local ethics review board with the reference number EC 1309/2017. The study is registered on ClinicalTrials.gov, trial identification number: NCT05054972.

Results

Detailed patients’ comorbidities and demographics are described in Tables 1 and 2. No significant differences were found between groups A and B (Table 1) or groups A+ and A– (Table 2).

The evaluation of the renal outcome showed similar results between groups A and B during the early postoperative course (days 1–15) as well as in the long-term follow-up (12 and 24 months postoperatively), details are described in Table 3.

The sub-analysis of group A showed that 25 patients (group A+) had a communicating lumbar vein detectable in the preoperative CT scan (diameter ≥ 2 mm), whereas in 56 patients this vein was absent (group A–). With regard to renal function between group A+ and group A–, both sCr and eGFR had an initial overlap during the early postoperative period and after 12 months. Slightly lower mean sCrea values were observed in group A+, but no statistical significance could be detected (1.01 vs. 1.23; p = 0.262). After 24 months of follow-up patients who did not present with a communicating lumbar vein in the preoperative CT scan showed a significant deterioration of renal function: sCr at 24 months was 0.87 mg/dl in group A+ and 1.51 mg/dl in group A–, p = 0.016; the eGFR at 24 months was 96.2 ml/min/1.73 m² in group A+ and

| Characteristics | Total cohort (n = 81) | Suprarenal clamp + LRV division patients + communicating lumbar vein present (group A+; n = 25) | Suprarenal clamp + LRV division patients + communicating lumbar vein missing (group A–; n = 56) | p-Value |
|-----------------|----------------------|---------------------------------------------------------------------------------|---------------------------------------------------------------------------------|---------|
| Sex, male, n (%) | 65 (80.2%)           | 17 (68.0%)                                                                      | 45 (80.4%)                                                                      | 0.225   |
| Age, years, mean ± SD | 68.8 ± 9.3         | 68.9 ± 7.7                                                                      | 68.8 ± 10.0                                                                     | 0.959   |
| ASA, median     | 3.04                 | 2.84                                                                           | 3.13                                                                           | 0.064   |
| Dialysis, n (%) | 1 (1.2%)             | 0                                                                               | 1 (1.8%)                                                                        | 0.507   |
| CDK stage ≥ 3b, n (%) | 9 (11.1%)          | 2 (8.0%)                                                                        | 7 (12.5%)                                                                       | 0.114   |
| Dyslipidemia, n (%) | 48 (59.3%)         | 16 (64.0%)                                                                      | 32 (57.1%)                                                                      | 0.567   |
| Diabetes mellitus, n (%) | 14 (17.3%)        | 3 (12.0%)                                                                        | 11 (19.6%)                                                                      | 0.407   |
| Hypertension, n (%) | 68 (84.0%)          | 21 (84.0%)                                                                      | 47 (84.0%)                                                                      | 0.994   |
| Smoker, n (%)   | 53 (65.4%)           | 15 (60.0%)                                                                      | 38 (67.9%)                                                                      | 0.498   |
| CAD, n (%)      | 30 (37.0%)           | 9 (36.0%)                                                                        | 21 (37.5%)                                                                      | 0.899   |
| Peripheral artery disease, n (%) | 29 (35.8%) | 8 (32.0%)                                                                       | 21 (37.5%)                                                                      | 0.638   |
| Pulmonary disease, n (%) | 23 (28.4%)  | 5 (20.0%)                                                                        | 18 (32.1%)                                                                      | 0.269   |

CKD chronic kidney disease; CAD coronary artery disease

Table 2 Demographic characteristics of patients (group A+ vs. group A–)
64.3 mL/min/1.73m² in group A, \( p = 0.041 \). Detailed results are described in Table 4.

Two patients died peri-operatively and both belonged to group A. They were both ASA 4 patients, affected by chronic ischaemic cardiac disease, chronic obstructive pulmonary disease, second stage chronic kidney disease and peripheral artery disease. One patient had an intraoperative cardiac arrest, the other patient died after 19 days due to congestive heart failure. Despite this, 30 day mortality was insignificantly different between the groups (2.5% in group A and 0% in group B, \( p = 0.15 \)). No significant difference was observed concerning the mean duration of the ICU stay (6.14 days in group A and 4.2 days in group B, \( p = 0.19 \)).

The sub-analysis of group A showed a 30 day mortality similar between the groups (3.6% in group A and 0% in group A, \( p = 0.16 \)) and no statistically significant difference emerged in mean ICU stay between group A+ and A– (8.64 days in group A+ and 5.18 days in group A–, \( p = 0.22 \)).

No patient was lost to follow-up.

**Discussion**

Despite the advances and increasing frequency of endovascular techniques performed, juxtarenal and complex aortic aneurysms frequently require open surgical treatment. At our institution, we prefer the transperitoneal approach. For adequate exposure of the pararenal aortic segment LRV division or extensive mobilisation of the LRV is required.

In our series, we compared renal function outcomes between patients who required division of the LRV (group A) with patients in whom the aortic cross clamp could be placed safely without division but mobilisation of the LRV (group B). We subsequently performed a sub-analysis of group A with regard to the presence of a communicating lumbar vein (CLV) as detected on preoperative CT imaging (group A+: CLV present; group A–: CLV not present).

We observed an elevation of serum creatinine values throughout postoperative day 15, but no significant differences between groups A and B or A+ and A–. This observation is consistent with early postoperative data from Wang et al. [15] and Samson et al. [7]. We therefore cannot confirm the findings of Huber et al. West et al. and AbuRahma et al. who described a relevant deterioration of renal function after division of the left renal vein, compared to mobilization only [2,4–6]. Interestingly the collective of patients in these studies was not as uniform as in our study. Huber et al. [4], Mehta et al. [2] and Wang et al. [15] included a large number of patients with ruptured aortic aneurysms into their analyses (29%, 39% and 37% of patients; respectively). In patients with ruptured aortic aneurysms shock due to blood loss represents an important factor contributing to the deterioration of renal function. In our opinion, these patients should be analysed as an individual cohort.

**Table 3** Serum creatinine (sCr) and eGFR values (group A vs. group B)

| Characteristics | Total cohort (\( n = 110 \)) | Suprarenal clamp + LRV division patients (group A: \( n = 81 \)) | Suprarenal clamp + LRV mobilization patients (group B: \( n = 29 \)) | \( p \)-Value |
|-----------------|------------------------------|---------------------------------------------------------------|---------------------------------------------------------------|--------------|
| Preoperative    |                              |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.2                          | 1.17                                                         | 1.27                                                         | 0.271        |
| eGFR (ml/min/1.73m²) | 73.32                      | 72.31                                                       | 61.18                                                       | 0.699        |
| Postoperative day 10 |                    |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.46                         | 1.57                                                         | 1.17                                                         | 0.086        |
| Postoperative day 15 |               |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.48                         | 1.59                                                         | 1.18                                                         | 0.119        |
| Discharge       |                              |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.38                         | 1.46                                                         | 1.16                                                         | 0.126        |
| eGFR (ml/min/1.73m²) | 67.39                      | 66.91                                                       | 68.73                                                       | 0.726        |
| 12 months postoperative |                 |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.2                          | 1.16                                                         | 1.24                                                         | 0.646        |
| eGFR (ml/min/1.73m²) | 70.61                      | 72.35                                                       | 65.2                                                         | 0.303        |
| 24 months postoperative |                |                                                               |                                                               |              |
| sCr (mg/dl)     | 1.29                         | 1.31                                                         | 1.17                                                         | 0.619        |
| eGFR (ml/min/1.73m²) | 73.92                      | 74.01                                                       | 73.50                                                       | 0.975        |
In other studies, mixed cohorts of patients who received both supra- and infrarenal aortic clamps were analysed negligible of the position of the aortic cross clamp [2,4,7,16]. Using this approach the effect of LRVD on renal function in patients with suprarenal aortic cross clamping cannot be determined sufficiently. Therefore, findings of the aforementioned studies need to be interpreted with caution [2,4,17]. A recent publication compared renal outcome with regard to LRVD versus mobilisation only in a cohort in which 68.4% of the patients included underwent infrarenal aortic cross clamping [16]. In our opinion it seems impossible to draw conclusions concerning the impact of LRVD on renal function when the position of the aortic cross clamp is not standardised.

A third factor that affects renal function, besides emergency surgery and the exact position of the aortic cross clamp is additional treatment to the renal arteries (bypass, reimplantation, endarterectomy) [5].

For these aforementioned reasons, we only included patients undergoing aortic cross clamping at the same level (suprarenal) who were not operated under emergency conditions and who did not receive additional treatment of the renal arteries.

Our data show that there is no statistically significant difference in short-term perioperative renal function in patients with LRVD compared to patients with LRV mobilization only. The data become interesting when follow-up is evaluated. When groups A and B are compared no differences are observed, however, after following-up for two years, we could identify a cohort of patients with better renal outcome after ligation of the left renal vein. In this group we could confirm a communicating lumbar vein ≥ 2 mm between the left renal vein and the left ascending lumbar vein on pre-operative CT scans. Renal function was significantly better compared to the group without a communicating lumbar vein (p = 0.016). There was no difference in patient management between the groups. We suggest that this result is probably linked to better venous drainage from the left kidney offered by a stronger network of venous collaterals that prevents damage to the organ due to congestion (Fig. 1).

Our data suggest that the presence of the communicating lumbar vein may provide an alternative way of drainage that allows restoration of the renal function to preoperative levels as seen in group A+, we could not observe this recovery in group A−. This has not been described so far and we suppose that recent studies did not find this difference because the communicating lumbar vein may not have been considered as a relevant individual factor [7,15].

The accuracy of the venous phase in contrast-enhanced CT to detect venous collaterals or anomalies ranges from 96 to 100% which is excellent and underlines the validity of our observation [8,9,12]. Furthermore, in our cohort of patients, the incidence of the communicating lumbar vein in preoperative contrast-enhanced CT studies was similar to previously reported data (30.9% [8]).

### Table 4  Serum Creatinine (sCr) and eGFR values (group A+ vs. group A−)

| Characteristics                  | Total cohort (n = 81) | Suprarenal clamp + LRV division patients + communicating lumbar vein present (group A+; n = 25) | Suprarenal clamp + LRV division patients + communicating lumbar vein missing (group A−; n = 56) | p-Value |
|----------------------------------|----------------------|------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------------|---------|
| **Preoperative**                 |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.17                 | 1.12                                                                                           | 1.19                                                                                            | 0.471   |
| eGFR (ml/min/1.73m²)            | 72.31                | 79.64                                                                                            | 69.04                                                                                            | 0.063   |
| **Postoperative day 10**        |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.57                 | 1.36                                                                                           | 1.66                                                                                            | 0.295   |
| **Postoperative day 15**        |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.59                 | 1.46                                                                                           | 1.64                                                                                            | 0.591   |
| **Discharge**                   |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.46                 | 1.37                                                                                           | 1.49                                                                                            | 0.605   |
| eGFR (ml/min/1.73m²)            | 66.91                | 72.18                                                                                            | 64.56                                                                                            | 0.209   |
| **12 months postoperative**     |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.16                 | 1.01                                                                                           | 1.23                                                                                            | 0.262   |
| eGFR (ml/min/1.73m²)            | 72.35                | 76.13                                                                                            | 70.71                                                                                            | 0.487   |
| **24 months postoperative**     |                      |                                                                                                |                                                                                                |         |
| sCr (mg/dl)                      | 1.31                 | 0.87                                                                                           | 1.51                                                                                            | 0.016 sig. |
| eGFR (ml/min/1.73m²)            | 74.01                | 96.23                                                                                            | 64.29                                                                                            | 0.041 sig. |

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Limitations

The limitation of our study is the moderate number of patients, due to the low incidence of the disease and a mindful selection of patients. However, this is in large part due to the strict selection criteria in this study as we did not include infrarenal abdominal aortic aneurysms, ruptured abdominal aortic aneurysms or patients who received additional treatment to the renal arteries during juxtarenal aortic repair. We exclusively analysed patients with juxtarenal abdominal aortic aneurysms in whom a suprarenal aortic cross clamp was applied. All our patients were treated via a transperitoneal approach, so we do not have data on the postoperative renal function in patients with juxtarenal aortic aneurysms treated by a retroperitoneal approach, which per se can eliminate the need of LRV division. In case of insufficient exposure after extensive mobilization of the LRV, our centre’s policy is to divide the LRV rather than to divide the adrenal and gonadal and lumbar vein as this procedure enhances the exposure only to a limited extent.

The limited number of patients is also relevant to the observation of a trend towards a better renal function in the early postoperative period in patients with juxtarenal aortic aneurysms treated by a retroperitoneal approach, which per se can eliminate the need of LRV division. In case of insufficient exposure after extensive mobilization of the LRV, our centre’s policy is to divide the LRV rather than to divide the adrenal and gonadal and lumbar vein as this procedure enhances the exposure only to a limited extent.

The limited number of patients is also relevant to the observation of a trend towards a better renal function in the early postoperative period in patients in whom the left renal vein could be preserved ($p = 0.09$ and 0.12 on days 10 and 15). With a higher number of patients, this result may reach significance.

Future studies with more patients and longer follow-up periods are needed to evaluate the impact of our findings on the long-term.

Conclusion

In patients treated for juxtarenal abdominal aortic aneurysms, LRV division appears not to affect renal function in the short-term. The absence of a communicating lumbar vein $\geq 2$ mm in the preoperative CT scan in patients requiring LRV division seems to correlate with a deterioration of renal function in the long-term. Therefore, we suggest reserving this procedure if possible to patients presenting a good communicating lumbar vein in the preoperative CT scan, to have an additional route of good venous drainage for a better renal outcome. Otherwise reconstruction of the LRV may be advisable, especially in patients that would be more affected in the long-term from a renal deterioration such as patients with one kidney or already increased creatinine preoperatively. Further data are necessary to corroborate our results.

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Declarations

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval This study complies with ethical standards according to the 1964 Helsinki Declaration and its later amendments. The reference number of the local ethics review board is EC 1309/2017.

Fig. 1 Example of a patient with a strong communicating lumbar vein pre- (a) and 5 years post-operatively (b)
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