Efficacy and safety of recanalization with transseptal needle for chronic total occlusion of the brachiocephalic vein in hemodialysis patients

Xi Yin\textsuperscript{1,2,*}, Xi Shen\textsuperscript{1*}, Zhongxin Zhou\textsuperscript{3}, Qin Chen\textsuperscript{1}, Li Zhou\textsuperscript{1}, Tianlei Cui\textsuperscript{1}

\textsuperscript{1}Department of Nephrology, West China Hospital, Sichuan University, Chengdu, China; \textsuperscript{2}The Second People’s Hospital of Panzhihua, Panzhihua, China; \textsuperscript{3}Department of Vascular Surgery, The Third Affiliated Hospital of Southern Medical University, Guangzhou, China

**Contributions:** (I) Conception and design: T Cui; (II) Administrative support: T Cui; (III) Provision of study materials or patients: T Cui; (IV) Collection and assembly of data: X Yin, X Shen, Z Zhou, Q Chen; (V) Data analysis and interpretation: X Yin, X Shen, L Zhou; (VI) Manuscript writing: All authors; (VII) Final approval of manuscript: All authors.

*These authors contributed equally to this work.

**Correspondence to:** Tianlei Cui, MD. Department of Nephrology, West China Hospital, Sichuan University, 37 Guoxue Alley, Wuhou District, Chengdu, China. Email: tianleicui@163.com.

**Background:** Whilst there are effective techniques for the recanalization of central venous occlusions, recanalization of chronic total occlusions remains particularly challenging. This study aims to evaluate the safety and efficacy of recanalization using a transseptal needle in chronic total occlusions of the right brachiocephalic vein (RBV) in long-term hemodialysis patients.

**Methods:** In this single-center, retrospective study, maintenance hemodialysis patients with chronic total occlusion of the RBV were enrolled between February 2017 to March 2019 from West China Hospital of Sichuan University. Refractory lesions were defined as complete vascular occlusions with failed recanalization using conventional techniques. Oclusions were approached using a transseptal needle to penetrate which offers an alternative strategy with firm support force. Patient data, treatment outcomes and patency rates were collected and analyzed to assess the safety and efficacy of the technique.

**Results:** A total of 16 eligible patients were analyzed. The operation was successful in 13 of the 16 patients and the success rate was 81.25% (13/16). Twelve patients underwent percutaneous balloon dilatation and stent implantation after sharp recanalization, whilst one patient underwent balloon dilatation only. We achieved procedural success in 13 patients without surgical complications, and all of the patients were discharged in a stable condition. The primary patency rates at 3, 6 and 12 months after surgery were 100%, 84.6% and 69.2%, respectively. The primary assisted patency rates were 100%, 84.6% and 76.9%, respectively, and the secondary patency rates were 100%, 84.6% and 76.9%, respectively.

**Conclusions:** This study demonstrates that recanalization of chronic total occlusions to the RBV using a transseptal needle is a safe and effective method after traditional guide wire and catheter techniques fail. It was also found that additional techniques are needed for recanalization in patients with RBV occlusion combined with proximal stenosis or occlusion of the right subclavian vein.

**Keywords:** Transseptal needle; right brachiocephalic vein (RBV); chronic total occlusion; hemodialysis; percutaneous transluminal angioplasty (PTA); percutaneous transluminal angioplasty and stenting (PTAS)

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Introduction

Central venous occlusion is the most common complication of patients undergoing regular hemodialysis treatment. It is caused by mechanical damage to vessel walls from catheter insertion, as well as high flow rate of a preexisting arteriovenous fistula, resulting in intimal hyperplasia, fibrosis, endothelial cell dysfunction, and narrowing and occlusion of blood vessels (1-3). This can reduce the efficacy of hemodialysis by causing ipsilateral limb swelling and vascular access dysfunction, ultimately increasing all-cause mortality (4). In day-to-day clinical practice, effective and safe treatment approaches are gradually developing, involving surgical and endovascular intervention. Although effective, invasive surgery can be difficult for patients with poor health such as long-term hemodialysis patients (5-7). Recently, endovascular intervention has become the first-line treatment for central venous occlusion in patients on hemodialysis (8,9). However, recanalization in patients with long-segment occlusions (more than 2 cm) and/or angular occlusions still remains a challenge (10).

Methods for recanalization are traditional recanalization including regular guide wire and 21-guage needle or the sheath in combination with the dilatator or the stiff end of hydrophilic wire and sharp recanalization including PTCD puncture needle, TIPS needle and transseptal needle. Transseptal needle is used in transseptal puncture which is a critical step in achieving left atrial access for a variety of cardiac procedures. It is thinner and harder than other puncture needles and guide wires, as well as its small curved tips, making it a potentially superior choice for recanalization in this population of patients. Figure 1 showed the transseptal needle. Aside from a case report utilizing transseptal needle to recanalize total occluded segment (4 mm), transseptal needle have not been studied for successful use in long-segment occlusions (more than 2 cm) and/or angular occlusions (11). Therefore, the purpose of this study was to evaluate the efficacy and safety of sharp recanalization using a transseptal needle after failure using standard guide wire and catheters in hemodialysis patients with chronic total occlusion of the right brachiocephalic vein (RBV).

We present the following article in accordance with the STROBE reporting checklist (available at http://dx.doi.org/10.21037/atm-20-5369).

Figure 1 The Transseptal Needle Set consists of a transseptal needle, obturator, and transseptal guiding introducer.

Methods

Study population

We retrospectively enrolled hemodialysis patients with chronic total occlusion of the brachiocephalic vein between February 2017 to May 2019 from West China Hospital of Sichuan University. Patients who underwent conventional recanalization were included in this study. The inclusion criteria were as follows: patients with occlusion of the RBV leading to dysfunction of the arteriovenous internal fistula; patients were male or non-pregnant females >14 years old; patients with confirmed complete occlusion of the RBV through digital subtraction angiography (DSA) or three-dimensional (3D) reconstruction of chest CT. Patients with a history of tumors and/or had an intravascular device such as a pacemaker were excluded. We also excluded patients with an allergy to the contrast medium, swelling of limbs caused by skin soft tissue infection, and coagulation dysfunction. Patient demographic information and treatment outcomes were recorded. All procedures performed in this study involving human participants were in accordance with the Declaration of Helsinki (as revised in 2013). This study was approved by the ethics committee of West China Hospital of Sichuan University (2017204). Because of the retrospective nature of the research, the requirement for informed consent was waived.

Surgical technique

Thoracic contrast-enhanced CT scans and 3D virtual model reconstructions were performed prior to the procedure
for all patients. All procedures were performed with local analgesia. Under DSA guidance with the patient in a supine position, 9F, 5F (or 6F) vascular sheaths were inserted at the femoral and forearm (or upper extremity fistula) sites respectively. A contrast medium was administered to delineate the anatomy of the adjacent vessels and occlusion to identify the degree and the accurate length of the occluded segment. Multiple approaches were made from both sides to penetrate the occlusion with the soft or hard end of a 0.035 inch (1 inch ≈ 0.025 m) hydrophilic guide wire (Terumo, Tokyo, Japan). If this failed, the upper guide wire was advanced to the distal end of the occlusion as a target. Aided by two orthogonal projections in the anterior-posterior and lateral view under DSA, a transseptal needle BRK (St. Jude Medical; St. Paul, MN, USA) was used to retrogradely cross through the femoral vein. Confirmation was obtained that the wires crossed in an appropriate route consistent with the occlusion segment. Next, balloons of different sizes were sequentially passed over the guide wire to produce graded dilations (Cordis Corporation, Milpitas, CA, USA). After dilation, contrast medium was administered in order to rule out extravasation and verify the patency of the occlusive segment. If necessary, stent implantation was performed. Figure 2 shows the surgical procedure. Technical success was defined as successful recanalization followed by balloon angioplasty and stent implantation without major complications. All procedures were performed by the same experienced interventional nephrologists. At follow-up, the primary and secondary patency rates, along with primary assisted patency rates at 3, 6 and 12 months were recorded.

Statistical analysis

SPSS 26.0 (Chicago, IL, USA) was used for statistical analysis. The continuous variables conforming to a normal distribution were expressed as mean ± standard deviation. And the qualitative data were expressed as frequencies or percentages (%). Patency rates were assessed using the Kaplan-Meier method.

Results

Patient characteristics

A total of 16 patients were included in this study (Table 1), including 11 males (71.4%) and 5 females (28.6%). The average age was 56.4±17.2 years. The patients had received hemodialysis for an average of 54.0±33.4 months. The average occlusion length in the RBV was 28.7±5.1 mm. There were 9 patients with occlusion of the RBV only, whilst other patients had several other comorbid lesions including occlusions to the right internal jugular vein, right subclavian vein and the superior vena cava, or stenoses. A summary of patient demographics is presented in Table 1.

Surgical success

Sharp recanalization of the brachiocephalic vein was performed successfully in 13 patients, with a success rate of 81.25% (13/16) and without any serious complication. Only 1 patient underwent percutaneous balloon angioplasty exclusively, whilst 12 patients underwent both percutaneous balloon angioplasty and stent implantation after sharp recanalization. Two of the remaining patients who suffered from occlusion of the RBV combined with proximal right subclavian vein, accepted percutaneous superior vena cava puncture straightly to placement tunneled cuffed catheters with excellent success after failed in sharp recanalization. In one of the patients, a 14 years old female with occlusion of the RBV and superior vena cava, recanalization was not performed as her family did not consent to another operation. Details of occlusion length and treatment outcomes are presented in Table 2.

Clinical outcomes

The median follow-up was 12 (range, 3–24) months. The primary patency rates at 3, 6, and 12 months were 100%, 84.6%, and 69.2%, respectively. Primary assisted patency rates were 100%, 84.6%, and 76.9%, respectively, whilst the secondary patency rates were 100%, 84.6%, and 76.9%, respectively (Figure 3). Two patients were lost at follow-up, and 3 patients developed stenosis, occlusion and thrombosis at follow-up. At 9-month follow-up, 1 patient had stenosis of the right basilic vein, stenosis with thrombosis at the entrance of right subclavian vein stent, and stenosis of the right innominate vein with thrombosis. At 14 months post-surgery, severe stenosis occurred in the proximal part of the right innominate vein and the distal part of the subclavian vein, with stent thrombosis. In the second patient, the right subclavian vein was occluded 8 months post-surgery, losing the access point. At 13 months post-surgery, stenosis of the right cephalic arch was accompanied by thrombosis. At 17 months post-surgery, there were multiple stenoses in

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the right axillary vein, subclavian vein and the stent. In the third patient, stenoses occurred in the right axillary vein, subclavian vein and brachiocephalic vein 14 months post-surgery. These three patients were treated with balloon dilation and stent implantation for stenosis and occlusion, restoring blood flow.

**Discussion**

In this case series, recanalization with a transseptal needle was performed in 16 patients who had chronic total occlusion in the RBV. The success rate of this technique in the present study was higher than previous reports, reaching 81.25%. The surgery was able to successfully...
penetrate the occlusion of the RBV, overcoming the failures of conventional recanalization methods.

Central vein occlusion is a serious problem facing long-term hemodialysis patients, often exhausting all alternative sites of access, leading to reduced long-term survival rates. It has been reported that the prevalence of central venous occlusion in these patients varies considerably, ranging from 4.3% to 41% (12,13). For these patients, even in the absence of central vein catheterization, using the arteriovenous fistula as an access site for hemodialysis may also eventually lead to narrowing or occlusion of the central vein. The mechanisms of central vein occlusion involve extrinsic and intrinsic factors. The former are attributed to compression by adjacent anatomical structures, and the latter is mainly attributed to smooth muscle hyperplasia, mural thrombosis or fibrosis due to intravascular device insertion and high flow rate of the arteriovenous fistula (14).

The RBV, also known as the innominate vein, originates from the merging of the right subclavian vein and right internal jugular vein, and converges with the RBV to form the superior vena cava. The length is usually 2 to 3 cm (15). Occlusion of the RBV could decrease the ipsilateral upper extremity hemodialysis access flow rates and lead to upper limb swelling (16).

Intravascular interventional therapy, namely percutaneous balloon angioplasty, is a standard treatment for central vein

Table 1: Patient demographics and baseline characteristics

<table>
<thead>
<tr>
<th>Demographics</th>
<th>Patients (n=16)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age [years]</td>
<td>56.4±17.2 [14–77]</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>11 (71.4)</td>
</tr>
<tr>
<td>Female</td>
<td>5 (28.6)</td>
</tr>
<tr>
<td>Dialysis duration</td>
<td>54±33.4 [12–120]</td>
</tr>
<tr>
<td>Primary disease</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>5 (31.3)</td>
</tr>
<tr>
<td>Diabetes</td>
<td>3 (18.8)</td>
</tr>
<tr>
<td>Arthrolithiasis</td>
<td>2 (12.5)</td>
</tr>
<tr>
<td>Polycystic kidney</td>
<td>1 (6.25)</td>
</tr>
<tr>
<td>Allergic purpura</td>
<td>1 (6.25)</td>
</tr>
<tr>
<td>Unknown</td>
<td>7 (43.8)</td>
</tr>
</tbody>
</table>

Data is expressed as mean ± SD (range) or n (%).

Table 2: Occlusion characteristics and treatment outcomes

<table>
<thead>
<tr>
<th>Patient</th>
<th>Occluded vein</th>
<th>Occlusion length (mm)</th>
<th>Balloon (mm)</th>
<th>Stent (mm)</th>
<th>Successful</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>RBV, RIJV</td>
<td>37</td>
<td>6,10,12</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>2</td>
<td>RBV</td>
<td>23</td>
<td>8</td>
<td>8</td>
<td>Yes</td>
</tr>
<tr>
<td>3</td>
<td>RBV</td>
<td>39</td>
<td>6,8,10,12</td>
<td>–</td>
<td>Yes</td>
</tr>
<tr>
<td>4</td>
<td>RBV</td>
<td>28</td>
<td>4,8,10</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>5</td>
<td>RBV</td>
<td>23</td>
<td>4,8,10</td>
<td>10</td>
<td>Yes</td>
</tr>
<tr>
<td>6</td>
<td>RBV</td>
<td>29</td>
<td>10,12</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>7</td>
<td>RBV, RSV</td>
<td>25</td>
<td>6,8,10</td>
<td>10</td>
<td>Yes</td>
</tr>
<tr>
<td>8</td>
<td>RBV</td>
<td>36</td>
<td>6,10</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>9</td>
<td>RBV, RIJV</td>
<td>24</td>
<td>4,6,8</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>10</td>
<td>RBV, RSV</td>
<td>33</td>
<td>10,12</td>
<td>6,12</td>
<td>Yes</td>
</tr>
<tr>
<td>11</td>
<td>RBV</td>
<td>24</td>
<td>4,5,6</td>
<td>10</td>
<td>Yes</td>
</tr>
<tr>
<td>12</td>
<td>RBV</td>
<td>25</td>
<td>8,10</td>
<td>11</td>
<td>Yes</td>
</tr>
<tr>
<td>13</td>
<td>RBV</td>
<td>28</td>
<td>8,12</td>
<td>12</td>
<td>Yes</td>
</tr>
<tr>
<td>14</td>
<td>RBV, SVC</td>
<td>26</td>
<td>–</td>
<td>–</td>
<td>No</td>
</tr>
<tr>
<td>15</td>
<td>RBV, RSV</td>
<td>28</td>
<td>–</td>
<td>–</td>
<td>No</td>
</tr>
<tr>
<td>16</td>
<td>RBV, RSV</td>
<td>31</td>
<td>–</td>
<td>–</td>
<td>No</td>
</tr>
</tbody>
</table>

RBV, right brachiocephalic vein; RSV, right subclavian vein; RIJV, right internal jugular vein; SVC, superior vena cava.
occlusion. This technique utilizes the anterior or posterior end of a guide wire, coaxially supporting the catheter and vessel sheath, whilst delivering continuous balloon dilation followed by stent implantation (5). Although primary patency rates of percutaneous balloon angioplasty and stents vary, this technique still has limitations, and secondary patency can be obtained with repeated surgery. In a study of 70 patients with hemodialysis-related central venous stenosis, Quaretti et al. (17) reported a primary patency rate of 58% at 12 months post-percutaneous balloon angioplasty. Similarly, in a study of 74 patients with symptomatic central vein occlusion, Saleh et al. (18) reported a primary patency rate of 51.7%±7.4% for the percutaneous balloon angioplasty group, and 70.6%±11.1% in the stent group. In the previous study, covered stent grafts resulted in a primary patency rate of 91.7% at 12 months post-surgery, with an excellent secondary patency rate of 95.8% when used for central venous stenosis in hemodialysis patients (19). The primary patency rate in our study was higher, but the long-term patency rate was still not encouraging. As a result, repeated interventions may be required to maintain patency. Surgery success rates and long-term patency rates are influenced by several factors, including comorbidities, severity of lesions, fistula flow, types of instruments used and interventional experience of the surgeons.

The first step in percutaneous balloon angioplasty is feeding the guide wire through the occlusion intraluminally. This process is called “crossing” and is a key step towards the success of the surgery. However, there are several studies reporting difficulties in crossing the total occlusion segment during conventional revascularization. The first clinicians to describe sharp recanalization were Gupta et al. (20) and Farrell et al. (21) in 1998 and 1999. A hollow-core puncture needle was used to pass through the occluded segment with a target balloon, then a conventional guide wire and catheter are used. The 21-gauge micro-puncture kit gradually replaced the first 18-gauge puncture needle (22). In 2005, Honnef et al. (23) reported a successful case which traversed a 2 cm occlusion of the RBV with transjugular intrahepatic portosystemic shunt (TIPS) set directed by the right brachiocephalic stent. A Rosch-Uchida needle was previously used in 33 patients with a subclavian vein or brachiocephalic vein occlusion with a revascularization success rate of 93.9%. The primary patency rates at 3-, 6-, and 12-month were 43.6%, 24%, and 8%, respectively, and the secondary patency rates were 77.4%, 68.8% and 55.9%, respectively (24). The patency rates in our study compared with the Rosch-Uchida needle technique were significantly higher. Several studies using radiofrequency wires (RF), which use radiofrequency energy to generate heat and vaporize cells in contact with the wire tip during sharp recanalization, have been reported in chronic central venous occlusion patients (25-28). However, this technique also penetrates the adjacent soft tissue, potentially resulting in fatal complications. Subsequently, in a study by Arabi et al. (29), transseptal needles were used for sharp central venous recanalization in 6 patients.

It is easier in recanalization of partly RBV occlusion, and
there were many methods to make it come true, including mother-child technique, extra stiff guidewire, through-and-through wire technique, segmented sharp recanalization strategy, as well as using transseptal needle (30-33). Nevertheless, there are limited data and explicit procedure on sharp recanalization of complete RBV occlusions before, our study focused on total RBV occlusion and didn't include partly one. We performed recanalization using a transseptal needle in 16 patients who all had chronic total occlusion of the RBV, confirming that the technique is safe and effective in a larger sample size. Malik et al. (11) reported the use of a transseptal needle to cross the RBV through the femoral vein, with an occlusion degree of 4 mm. The local intima hyperplasia and dense fibrosis of the dysfunctional vessels form fibrous cord tissue. The guide wire or puncture needle is able to easily pass through the dense tissue, breaking through the weak site in the normal position to the outside of the vascular cavity, resulting in rupture and bleeding (8,34). The main advantages of the transseptal puncture needle over other puncture needles include that the needle can adjust its curvature according to the angiographic path, controlling degree and direction. In addition, the diameter of the transseptal needle is 21G, but it is 16G of Rosch-Uchida needle and 20G of TIPS, so that the severity of the perforation complications is reduced (23,24,29). For long segment (length >2 cm), it is advisable to use the transseptal puncture needle kit to perform the recanalization operation. In the present study, we found that recanalization of the RBV is easier than recanalization of angular (such as subclavian vein combined with brachiocephalic vein) occlusions. The brachiocephalic vein and the superior vena cava are adjacent to the pericardium, lungs and other vital organs. Serious complications such as pneumothorax, hemothorax, hemopneumothorax, pericardial tamponade or major artery injury may occur if the needle direction deviates. It is difficult to anterogradely puncture chronic vascular occlusions, particularly superior vena cava occlusions. Combining anterograde and reverse puncture from upper limb veins, neck vein and femoral vein is a more effective method. Apart from the initial catheter, a support catheter or vascular sheath is also required in order to provide sufficient stiffness and orientation. Most of the occlusion can be crossed through different angles and bidirectional exploration under X-ray fluoroscopy. After that, percutaneous balloon angioplasty is necessary. Because of the chronic fibrosis characteristics of the occlusion, a balloon dilation is carried out step-by-step until it restores blood flow (35).

In summary, we describe a successful technique using a transseptal puncture needle in the treatment of chronic total occlusion of the RBV. This technique was found to be safe and feasible when carried out by a multidisciplinary team of experienced physicians.

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**Footnote**

Reporting Checklist: The authors have completed the STROBE reporting checklist. Available at http://dx.doi.org/10.21037/atm-20-5369

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at http://dx.doi.org/10.21037/atm-20-5369). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All procedures performed in this study involving human participants were in accordance with the Declaration of Helsinki (as revised in 2013). This study was approved by the ethics committee of West China Hospital of Sichuan University (2017204). Because of the retrospective nature of the research, the requirement for informed consent was waived.

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