Radial artery access for neuroendovascular procedures: safety review and complications

Kazim H Narsinh,1 Mohammed H Mirza,2 M Travis Caton Jr,1 Amanda Baker,1 Ethan Winkler,3 Randall T Higashida,1 Van V Halbach,1 Matthew R Amans,1,1 Daniel L Cooke,1 Steven W Hetts,1,1 Adib A Abla,3 Christopher F Dowd1

ABSTRACT

Although enthusiasm for transradial access for neurointerventional procedures has grown, a unique set of considerations bear emphasis to preserve safety and minimize complications. In the first part of this review series, we reviewed anatomical considerations for safe and easy neuroendovascular procedures from a transradial approach. In this second part of the review series, we aim to (1) summarize evidence for safety of the transradial approach, and (2) explain complications and their management.

INTRODUCTION

Interest has burgeoned in the use of the radial artery as an access site for neurointerventional procedures because it has been associated with fewer complications, shorter hospital stays, and patient preference. Nevertheless, transradial access (TRA) presents a unique set of considerations to preserve safety, including knowledge of the anatomical variations of the upper extremity arterial supply, access site complications, and closure techniques. Herein, we aim to review radial artery access for neuroendovascular procedures with attention to (1) a literature review of the evidence for safety in neurointervention, and (2) complications and their management.

Literature in interventional cardiology has substantiated certain advantages of the TRA route in the setting of coronary angiography and intervention. Compared with the transfemoral access (TFA) route, TRA has been associated with significantly reduced all-cause mortality, major cardiovascular events, all forms of bleeding, length of hospital stay, and vascular and access site complications (eg, hematoma, pseudoaneurysm, arteriovenous fistula, need for vascular repair). These findings are evidenced by randomized controlled trials, systematic reviews, and meta-analyses cumulatively covering >35 studies and >28,000 participants.1 Of note, however, these studies do not uniformly make use of ultrasound guidance with both access routes, some report slightly lower procedural success rates with TRA, and crossover from TRA to TFA is not infrequent.2 However, a lower complication rate and risk of stroke has been consistently demonstrated with TRA in interventional cardiology, without a difference in contrast media use, procedure duration, fluoroscopy, and door-to-balloon time for ST elevation myocardial infarction (STEMI). As a result, the American Heart Association and European Society of Cardiology recommend a “radial first” strategy for arterial access, particularly in the setting of acute coronary syndromes and myocardial infarction.1,3 This paves the way for interventional neuroradiology/neurointerventional surgery to adopt an alternative access site for other endovascular procedures, although it has yet to be substantiated by the degree of evidence that has supported interventional cardiology’s transition.

RADIAL ARTERY ACCESS SAFETY FOR NEUROENDOVASCULAR PROCEDURES

Methods

Literature search

We performed a literature search through MEDLINE/PubMed and Ovid for papers published in the English literature from 2018 to 2021 using the following search parameters: (radial[Title]) OR (transradial[Title]) AND (["Journal of neurointerventional surgery"] OR [Journal]) OR (neurointerventions OR neuroendovascular). Studies measuring both crossover and access site complication rates as outcomes in patients (n>100) undergoing a neuroendovascular procedure (diagnostic or interventional) with a radial-first approach were included. Exclusion criteria were: (1) case reports/series or studies with n≤100 subjects; (2) studies not reporting either one of the two outcomes of interest; (3) studies published before 2018 (when available wires/catheters/sheaths were more limited); (4) studies not adopting a radial-first approach; and (5) studies not involving neuroendovascular procedures.

Outcome variables

The following TRA outcomes were studied: access site complication rate, overall complication rate, and rate of crossover defined as any conversion from the originally intended access site. Single-arm studies were separated from comparative studies. In the case of comparative studies, the same three outcomes were measured for transfemoral comparison.

RESULTS

A total of 95 unique articles were retrieved after the first round of research, 69 of which were excluded by title and abstract screening. The full texts of the
remaining 26 articles were accessed, and 19 articles matching the inclusion criteria were included: 13 single-arm studies and six comparative studies. Fourteen studies were retrospective cohort, four prospective cohort, and one systematic review and meta-analysis (figure 1).

All neurointerventional studies of TRA suffer from a lack of randomization (tables 1 and 2), which introduces patient selection bias, except for one recent small study which assigned access site based on the day of presentation. A recent single arm study of 121 patients who underwent diagnostic and interventional neuroendovascular procedures after adoption of a ‘radial-first’ approach at a single institution had a transfemoral crossover rate of about 25%, while an older retrospective study of 49 patients undergoing transradial flow diversion had a transfemoral crossover rate of about 20%. In contrast, crossover rates in non-randomized neurointerventional cases are significantly lower, approaching approximately 5%. These data suggest that patient selection bias has strongly affected the complication rates of most prior studies of TRA (tables 1 and 2). Randomized controlled clinical trials are needed to establish that TRA has a decreased complication rate relative to TFA for neuroendovascular procedures, although current non-randomized studies suggest that the complication rate may be lower for selected patients.

COMPLICATIONS OF TRA

Common complications of transradial procedures include radial artery spasm (further complicated by catheter entrapment and radial artery eversion), radial artery occlusion (further complicated by hand ischemia), and radial artery perforation (further complicated by forearm hematoma and compartment syndrome). Rare complications include radial artery pseudoaneurysm, granuloma or sterile abscess formation, and arteriovenous fistula formation. Below we provide a more detailed discussion of the former three conditions that arise more frequently (and their associated sequelae), followed by the latter three rare complications.

Radial artery spasm

The radial artery is a muscular artery rich in α-1 receptors, making it highly spasmotic compared with other vessels. Thus, radial artery spasm causes a significant reduction in lumen diameter. This can be prevented by using intra-arterial 'cocktails' that include heparin, nitroglycerine, and verapamil diluted in normal saline. They induce rapid, prolonged radial artery vasodilatation and reduce vasospasm during transradial procedures. Preprocedural administration of a combined topical gel of verapamil–nitroglycerine–lidocaine can also significantly increase the size of the radial artery and reduce pain during puncture. The incidence of radial artery spasm can also be decreased by use of a long 23 cm hydrophilic sheath because they protect the full length of the radial artery and diminish repetitive friction forces of the catheter against the vessel wall. Although prevention of radial artery spasm is the preferred approach, once radial artery spasm has been detected, several techniques can be used to relieve spasm including forearm heating with forearm compresses, deepening of sedation or induction of general anesthesia, and administering additional intra-arterial vasodilators. A small randomized controlled trial (RCT) demonstrated inflating normal saline through the sheath (10 mL/s for 10mL) to induce pressure-mediated vasodilation induces greater radial artery intraluminal diameter acute gain when compared with intra-arterial administration of a combination of nitroglycerin and verapamil, while causing significantly less reduction in blood pressure. Another small RCT demonstrated inflating a blood-pressure cuff above systolic blood pressure for 5 minutes before rapid deflation to induce flow-mediated vasodilation leads to similar median time to return of radial pulse—compared with sublingual nitroglycerin—with less prevalence of headache and decreased blood pressure. Isolated cases have also been reported wherein administering a radial nerve or brachial plexus anesthetic block under ultrasound guidance with the help of anesthesiologist as a rescue measure provides a pharmacologic symptomatic relief.

Catheter entrapment

Severe radial artery spasm, worsened by excessive catheter manipulation or kinking or radial artery tortuosity, can lead to catheter entrapment. Any kinks in catheters should be untied slowly and carefully by rotating in the reverse direction or by introducing guidewires within catheters. Any fragments of catheters retained within the arterial system may lead to hand ischemia, thrombus formation, or infection.

Radial artery eversion

Radial artery eversion (figure 2) is an extremely rare complication caused by severe radial artery spasm, which may prevent catheter and sheath removal. As above, when encountering severe radial artery spasm, analgesia, vasodilators, deep sedation or even general anesthesia are administered. While these measures are usually successful, instruments must never be forcefully extracted, and careful attention to preventing or treating radial artery spasm is the preferred approach.

Radial artery occlusion

Radial artery occlusion is a common complication of transradial catheterization, with most operating physicians underestimating its incidence. It occurs at an incidence of 7.7% within 24 hours, declining to 5.5% at 1 month due to spontaneous recanalization. Radial artery occlusion has been associated with larger sheath sizes, smaller radial arteries, and diabetes mellitus. Other predictors such as patient age, body habitus, procedure duration, and hemostasis duration have shown inconsistent findings among different clinical studies. Radial artery occlusion proximal to the anatomic snuffbox can be decreased by accessing...
<table>
<thead>
<tr>
<th>Ref</th>
<th>Study population</th>
<th>N</th>
<th>Study design</th>
<th>Access site complication rate</th>
<th>Overall complication rate</th>
<th>Crossover rate</th>
<th>Follow-up length and method</th>
<th>Weaknesses</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>Patients undergoing 'radial first' diagnostic/interventional neuroendovascular procedure</td>
<td>121</td>
<td>Prospective cohort</td>
<td>Single center</td>
<td>7 (5.79%)</td>
<td>8 (6.61%)</td>
<td>30 (24.8%)</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>6</td>
<td></td>
<td></td>
<td>Retrospective cohort</td>
<td>Multi-center</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
<td>30 day follow-up with Allen test and/or Doppler</td>
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<tr>
<td>71</td>
<td>Diagnostic and intervention (aneurysm, flow diversion, stroke)</td>
<td>2203</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>7 (5.3%)</td>
<td>N/A</td>
<td>7 (5.3%)</td>
<td>Follow-up at each repeat procedure</td>
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<td>72</td>
<td>Patients undergoing repeat TRA at same access site</td>
<td>133</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>73</td>
<td>Intracranial aneurysm coiling in anterior circulation</td>
<td>103</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>74</td>
<td>Anterior circulation interventions</td>
<td>130</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>0%</td>
<td>0%</td>
<td>0%</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>75</td>
<td>Carotid artery stenting</td>
<td>723</td>
<td>Systematic review and meta-analysis</td>
<td>37 (5.12%)</td>
<td>61 (8.44%)</td>
<td>66 (9.13%)</td>
<td>No uniform protocol for follow-up</td>
<td>Patient selection bias (non-randomized)</td>
</tr>
<tr>
<td>29</td>
<td>Reaccessing occluded radial artery for neurointervention</td>
<td>106</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>0%</td>
<td>N/A</td>
<td>6 (5.66%)</td>
<td>Short-term (unspecified) follow-up for complications</td>
</tr>
<tr>
<td>76</td>
<td>Thrombectomy in acute stroke</td>
<td>309</td>
<td>Systematic review and meta-analysis</td>
<td>5/280 (1.4%)</td>
<td>N/A</td>
<td>7.2% (mean)</td>
<td>No uniform protocol for follow-up.</td>
<td>Patient selection bias (non-randomized)</td>
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<tr>
<td>77</td>
<td>Diagnostic and interventional (aneurysm, stroke)</td>
<td>225</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>3 (1.33%)</td>
<td>N/A</td>
<td>5 (2.22%)</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>78</td>
<td>Cerebral angiography</td>
<td>148</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>10 (6.75%)</td>
<td>10 (6.75%)</td>
<td>7 (4.7%)</td>
<td>Barbeau test at discharge; telephone interviews with 80 patients during postoperative period</td>
</tr>
<tr>
<td>79</td>
<td>Angiography, intracranial/ head and neck intervention</td>
<td>328</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>8 (2.4%)</td>
<td>8 (2.4%)</td>
<td>26 (7.9%)</td>
<td>30-day follow-up outpatient visit: evaluation of access site and radial pulse</td>
</tr>
<tr>
<td>80</td>
<td>Diagnostic angiography (93%)</td>
<td>121</td>
<td>Retrospective cohort</td>
<td>Single center</td>
<td>0%</td>
<td>1 (0.8%)</td>
<td>6 (5%)</td>
<td>Not collected</td>
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<td>81</td>
<td>Diagnostic and intervention (embolization, coiling)</td>
<td>506</td>
<td>Prospective cohort</td>
<td>Single center</td>
<td>4 (0.8%)</td>
<td>4 (0.8%)</td>
<td>33 (6.5%)</td>
<td>No uniform protocol for follow-up</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td>5156</td>
<td></td>
<td></td>
<td></td>
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</tr>
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</table>

N/A, not available; TRA, transradial access; US, ultrasound.
Table 2 Recent comparative studies of transradial versus transradial access for neuroendovascular procedures

<table>
<thead>
<tr>
<th>Ref</th>
<th>Study population</th>
<th>N (radial arm)</th>
<th>Study design</th>
<th>Access site complication rate (radial vs femoral)</th>
<th>Overall complication rate (radial vs femoral)</th>
<th>Crossover rate (radial vs femoral)</th>
<th>Follow-up interval</th>
<th>Weaknesses</th>
</tr>
</thead>
<tbody>
<tr>
<td>82</td>
<td>Interventional (thrombectomy, thrombolysis, carotid) and diagnostic</td>
<td>206 (20%)</td>
<td>Retrospective cohort, Single center</td>
<td>3 (1.5% vs 6%)</td>
<td>4 (2.0% vs 7.0%)</td>
<td>3 (1.5% vs 1.3%)</td>
<td>Unclear</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no uniform follow-up protocol)</td>
</tr>
<tr>
<td>83</td>
<td>Flow diversion</td>
<td>134 (6%)</td>
<td>Retrospective cohort, Multicenter</td>
<td>0 (0% vs 2.48%)</td>
<td>5 (3.73% vs 9.02%)</td>
<td>12 (8.63%)</td>
<td>Unclear</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no uniform follow-up protocol)</td>
</tr>
<tr>
<td>84</td>
<td>Anterior circulation mechanical thrombectomy</td>
<td>130 (35%)</td>
<td>Retrospective cohort, Single center</td>
<td>0 (0% vs 6.5%)</td>
<td>3 (2.3% vs 11.4%)</td>
<td>6 (4.6% vs 1.6%)</td>
<td>90 days</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no US as part of follow-up)</td>
</tr>
<tr>
<td>4</td>
<td>Cerebral arteriography with right TRA</td>
<td>158 (51%)</td>
<td>Prospective cohort, Single-center</td>
<td>4 (2.5% vs 5.8%)</td>
<td>4 (2.5% vs 5.8%)</td>
<td>5 (3.2% vs 1.3%)</td>
<td>Unclear</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no uniform follow-up protocol)</td>
</tr>
<tr>
<td>85</td>
<td>Diagnostic angiograms performed by two new fellows</td>
<td>169 (58%)</td>
<td>Prospective cohort, Single center</td>
<td>2 (1.2% vs 1.6%)</td>
<td>2 (1.2% vs 1.6%)</td>
<td>6 (2.1% vs not reported)</td>
<td>Unclear</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no uniform follow-up protocol)</td>
</tr>
<tr>
<td>86</td>
<td>Neurointerventional treatments (embolization, thrombectomy)</td>
<td>162 (28%)</td>
<td>Retrospective cohort, Single center</td>
<td>2 (1.2% vs 6.7%)</td>
<td>5 (3% vs 10%)</td>
<td>7 (4% vs 3.1%)</td>
<td>Unclear</td>
<td>Non-randomized study (patient selection bias) Possible misclassification bias (no uniform follow-up protocol)</td>
</tr>
<tr>
<td></td>
<td>Large interventional cardiology RCTs for comparison:</td>
<td></td>
<td></td>
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<td></td>
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</tr>
<tr>
<td>42</td>
<td>ACS (RIVAL)</td>
<td>3507 (50%)</td>
<td>Multicenter, 32 countries, 1:1 randomization, open label</td>
<td>49 (1.4% vs 3.7%)</td>
<td>167 (4.8% vs 7.4%)</td>
<td>265 (7.6% vs 2.0%)</td>
<td>30 days</td>
<td>Follow-up lacks US assessment</td>
</tr>
<tr>
<td>87</td>
<td>ACS (MATRIX)</td>
<td>4917 (50%)</td>
<td>European multicenter, 1:1 randomization, open label</td>
<td>73 (1.8% vs 5.2%)</td>
<td>460 (11% vs 13.6%)</td>
<td>243 (5.8% vs 2.3%)</td>
<td>30 days and 1 year</td>
<td>Follow-up lacks US assessment</td>
</tr>
</tbody>
</table>

ACS, acute coronary syndrome; MATRIX, Minimizing Adverse haemorrhagic events by Transradial access site and systemic Implementation of angioX, RCTs, randomized controlled trials; RIVAL, RadIal Vs femorAL access for coronary intervention; TRA, transradial access; US, ultrasound.
Some measures to decrease the risk of radial artery occlusion include higher intra-arterial heparin doses ($\geq 5000$ IU), shorter compression times ($\leq 15$ min), and patent hemostasis defined as hemostasis with enough pressure applied to the vessel to avoid bleeding but not enough to prevent blood flow.\textsuperscript{30–36} Patent hemostasis technique should ensure radial artery patency during closure with a reverse Barbeau’s test.\textsuperscript{38} Another technique to reduce radial artery occlusion is ipsilateral ulnar artery compression during hemostasis, which increases radial artery flow, and can be used simply and effectively for reducing the risk of radial artery occlusion after transradial catheterization.\textsuperscript{39–40}

Hand ischemia

Hand ischemia rarely presents as a complication of radial artery occlusion due to the dual blood supply of the hand and an extensive network of collaterals. In fact, some suggest that pre-procedural examination for a complete palmar arch using Barbeau’s test may not be necessary because it is not predictive of ischemic complications.\textsuperscript{35} The RIVAL (Radial Vs femoral access for coronary intervention) RCT comparing TFA and TRA in acute coronary syndrome patients undergoing percutaneous coronary intervention showed a 0.2% rate of symptomatic radial artery occlusion requiring medical attention.\textsuperscript{40} Raynaud’s disease puts patients at higher risk of hand ischemia and is a contraindication to TRA.\textsuperscript{4} Management of hand ischemia is mainly conservative with possible short-term use of aggressive anticoagulation, with a success rate of 87%.\textsuperscript{41} Percutaneous and surgical management have been described in cases of acute limb-threatening ischemia (figure 3).\textsuperscript{42–45}

Radial artery access closure

Prevention of radial artery occlusion and hand ischemia can be achieved through use of the smallest possible sheath size, intraarterial vasodilators and heparin, as well as proper attention to closure technique. Following TRA, several techniques are available to achieve hemostasis of the radial artery including manual compression, mechanical compression, and hemostatic agents.

Manual compression remains a viable option for achieving hemostasis, especially in diagnostic catheterizations with short durations. The superficial course of the radial artery makes manual compression easily achievable with two to three fingers and intermittent ulnar artery compression every 2 min. Pulse oximetry is monitored to ensure adequate perfusion without compromising hemostasis, and pressure over the radial artery is decreased in case of absent antegrade flow waveforms. Manual compression can be more effective than mechanical compression in performing patent hemostasis without any difference in the incidence of radial artery occlusion, and also achieves hemostasis in a shorter duration (14–22 min vs 2–3 hours) and at a lower cost than mechanical compression.\textsuperscript{46–48} Despite its advantages, manual compression is not widely adopted as first line because it usually demands prolonged post-procedural involvement of the interventional team.

Mechanical compression has historically used a balloon compression bracelet typified by the TR Band (Terumo Medical, Tokyo) with similar iterations including the VascBand (Teleflex, Wayne, PA), Tracelet (Medtronic, Dublin, Ireland), Safeguard Radial (Merit, South Jordan, UT), and Zephyr (Cordis, Miami Lakes, FL). The bracelet is applied at the arterial puncture site, then the bladder inflated, and the radial sheath is removed from under the band. The bladder is then decompressed until some mild bleeding occurs, and then the bladder is re-inflated with 1–2 mL of air. Then, antegrade radial artery flow is confirmed with a reverse Barbeau test (compression of the ulnar artery.
New devices and techniques

Figure 3  Hand ischemia. (A) Radial artery occlusion (dashed arrow) with distal flow supplied by the palmar carpal branch of the interosseous artery (solid arrow) in an adult patient on day 1 after transradial access who presented with significant wrist pain and fourth digit paresthesias with palor on exam. Note the ulnar artery filling defects (dotted arrow) despite the patient’s pre-procedural Barbeau type A waveform. (B) Follow-up angiogram after both radial and ulnar intra-arterial tPA lysis demonstrates marked improvement in both radial (dashed arrow) and ulnar (dotted arrow) artery flow with minimal reliance of collateral supply from the interosseous artery (solid arrow). tPA, tissue plasminogen activator.

with observation of the plethysmography waveform to ensure ‘patent hemostasis’; 3 mL of air is withdrawn from the bracelet after 30 min, then an additional 3 mL every 15 min. If bleeding occurs, the removed air is re-introduced. There is no significant difference in bleeding between mechanical and manual compression.46 However, compared with pressure dressings, mechanical compression reduces hemostasis duration, radial artery occlusion incidence, puncture site oozing, and patient discomfort.49

Gauze pads impregnated with a hemostatic agent such as kaolin (QuickClot Radial, Teleflex, Wayne, PA), chitosan (ChitoClot, Anscare, Taiwan), or potassium ferrate (StatSeal, Biolife, Sarasota, FL) can lead to shorter hemostasis durations.50–52 Kaolin activates factor XII, initiating the coagulation cascade, while chitosan attracts platelets, promoting platelet plug formation. Such agents are applied tightly at the puncture site with the sheath still in place. The sheath is then removed, and a small amount of blood is allowed to come into contact with the gauze activating the hemostatic agent. Manual pressure is applied to the gauze for 5 min, and then an elastic band is used to apply pressure for an additional 25–55 min. Compared with mechanical compression with the TR Band, hemostatic agents usually achieve hemostasis in significantly less time (30–60 min versus ~150 min). In addition, chitosan-based hemostatic pads have demonstrated a lower incidence of radial artery occlusion at 24 hours when compared with the TR Band.53 Furthermore, adding chitosan-based hemostatic pads to both mechanical compression devices (TR Band, rotary compression device) significantly reduces hemostasis duration with no difference in vascular complications.51–54

Some hemostasis techniques achieve significantly shorter hemostasis times, such as use of the VasoStat (Forge Medical, Bethlehem, PA), a focused compression device,55–57 or hemo-

Radial artery perforation

Radial artery perforation is a rare complication of TRA with a 0.1–0.1% incidence. Risk factors include female sex, old age, short stature, hypertension, excess tortuosity, excessive anticoagu-

Forearm hematoma

Hematoma formation at the site of radial puncture or in the forearm is an uncommon and usually benign condition. Forearm hematomas are classified into four grades: grade 1 (local, superficial) <5 cm, grade 2 (moderate muscular infiltration) <10 cm, grade 3 reaches distal to the elbow, and grade 4 extends proximal to the elbow. The incidence of grade 1 is <5%, grade 2 <3%, grade 3 <2%, and grade 4 <0.1%.62 Forearm hematomas are associated with radial artery occlusion, significant access site bleeding, improper hemostasis, radial artery perforation, use of non-coated sheaths, and female sex.

Asymptomatic, non-expanding forearm hematomas are managed conservatively with observation, elevation, and compression. Early detection is key and reversal of anticoagulation (eg, using protamine) should be considered. Grade 1 and 2 hematomas are managed with analgesia, additional compression, and local ice, while grade 3 and 4 hematomas may benefit from blood pressure cuffs inflated to 20 mmHg less than systolic blood pressure and deflated every 15 min with pulse oximetry monitoring.44–62

An expanding hematoma or one causing neurovascular compromise must be urgently examined to confirm patency of both radial and ulnar arteries, evaluate digital perfusion, and measure compartment pressures. Forearm hematomas rarely progress to compartment syndrome requiring surgical hema-

Compartment syndrome

Forearm compartment syndrome is a rare but devastating complication following TRA. The etiology is the same as for forearm hematomas, with some authors considering it a grade 5 hematoma, and has been associated with female sex and low body surface area.63–64

Normal compartment pressures in the forearm are <10 mmHg, with an increase to >30 mmHg considered diagnostic of acute compartment syndrome. However, the diagnosis should rather
be based on clinical presentation that can be quite rapid with acute forearm swelling, pallor, and pain following the procedure that can further progress to ischemia of the muscles and nerves, with disability and contracture if not managed in a timely fashion.\(^{63-65}\)

Prevention involves objectively examining blood flow to the palmar arch before the procedure, using an appropriate anticoagulant dose, and proper management of intraprocedural tortuosity and radial artery spasm. After the procedure, the hemostatic device and puncture site should be continuously checked.\(^{44-63}\)

Management includes inflation of a blood pressure cuff at the site of pain or swelling at 15–20 mmHg below systolic blood pressure to ensure distal flow confirmed by pulse oximetry for 15 min and repeated twice on average. Blood pressure should also be tightly controlled with anticoagulation reversal or discontinuation. Most cases respond to conservative management, but surgical fasciotomy may occasionally be required.\(^{44,63}\)

**Radial artery pseudoaneurysm**

Radial artery pseudoaneurysm is a very rare complication following TRA with an incidence rate of 0.03%.\(^{66}\) Presentation is within days to weeks as localized, pulsating swelling that may be painful. Diagnosis is confirmed by Doppler ultrasound or angiography. Radial artery pseudoaneurysm can be caused by excessive anticoagulation, anomalous radial artery perforation, excessive puncture attempts, and inadequate compression hemostasis.\(^{67}\)

Management can be escalated sequentially from firm compression to ultrasound-guided compression of the pseudoaneurysm neck to ultrasound-guided thrombin injection. Surgery is indicated in rare cases of infection, compressive ischemic symptoms, or failure of non-surgical management.

**Granuloma and sterile abscess formation**

Granuloma and sterile abscess formation are rare, self-limiting complications induced by a reaction to introducer sheath coating that can be stripped off and retained within the skin. Of note, the Cook vascular sheaths implicated in causing granulomas are no longer in use.\(^{68}\) They form within 2–3 weeks as a tender mass near the puncture site. Due to its late presentation, patients present to their primary care physician rather than their interventionalist, frequently leading to its misdiagnosis as a radial artery pseudoaneurysm. Thus, Doppler ultrasound is important to confirm diagnosis and avoid unnecessary intervention.\(^{69}\)

Management is limited to observation. Antibiotics are not indicated in the absence of fever and leukocytosis. Acute granulomas resolve within weeks and chronic granulomas within months. Wiping the vascular sheath with damp gauze before insertion may remove excess hydrophilic material and reduce the incidence of granulomas.

**Arteriovenous fistula**

Arteriovenous fistulae rarely present as a complication—with an incidence <0.03%—due to the lack of large veins near the radial artery. It presents as a mass or palpable thrill near the puncture site, and Doppler ultrasound confirms the diagnosis. Management may be conservative with prolonged compression, interventional using covered stents, or surgical repair.\(^{14,70}\)

**CONCLUSION**

The radial artery provides an alternative—and in some ways more favorable—access site for the neurointerventionalist. Nonetheless, it demands the physician be well versed in identifying anatomical variants, weighing the risks and benefits of different closure techniques, and recognizing access site complications. While the advantages of transradial over transfemoral catheterization has been established in the field of interventional cardiology with randomized controlled clinical trials, further study is warranted to directly establish its advantage in specific realms of interventional neuroradiology/neurointerventional surgery.

**Contributors**

KHN wrote the manuscript and supervised the review. MHH performed the literature review and analysis and provided with figures. MTC, AB, EAW, RTH, VVH, MRA, DCL, SWH, AAA, CFD provided edits to the manuscript.

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**ORCID iDs**

Kastim H Narsinh http://orcid.org/0000-0002-2019-5461
Mohammed H Mirza http://orcid.org/0000-0003-3784-5894
Matthew R Amans http://orcid.org/0000-0002-8209-0534
Steven W Hetts http://orcid.org/0000-0001-5885-7259

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New devices and techniques


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New devices and techniques


