ORIGINAL RESEARCH

Comprehensive analysis of intra-arterial treatment for acute ischemic stroke due to cervical artery dissection

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ABSTRACT

Objective The safety and efficacy of intra-arterial treatment (IAT) in patients with acute ischemic stroke (AIS) due to cervical artery dissection (CeAD) has not been formally studied. The purpose of this study was twofold: first, to define a large series with CeAD treated with IAT, second, to analyze outcomes with CeAD receiving IAT versus (a) CeAD not treated with IAT, (b) CeAD receiving intravenous thrombolysis (IVT) alone, and (c) non-CeAD mechanism of AIS receiving IAT.

Design Demographics, clinical characteristics, treatment, and outcomes were summarized for all CeAD patients treated with IAT from January 2010 to May 2015. Outcomes included favorable 90 day modified Rankin Scale (mRS) score of 0–2, symptomatic intracerebral hemorrhage (sICH), recanalization (Thrombolysis in Cerebral Infarction 2b–3), procedural complications, and mortality. Outcomes were analyzed with χ2 tests and multivariate logistic regression.

Results There were 161 patients with CeAD: 24 were treated with IAT and comprised our target population. Dissections were more common in the internal carotid (n=18) than in the vertebral arteries (n=6). All but one patient had intracranial embolus. IAT techniques included thrombectomy (n=19), IA thrombolysis (n=17), stent (n=14), and angioplasty (n=7). Outcomes included favorable 90 day mRs score of 0–2 in 63%, 4 deaths, 1 sICH, and 3 procedural complications. After adjustment, favorable mRs in our target population was similar to comparison populations: (a) in CeAD, IAT versus no IAT (OR 0.62, p=0.56); (b) in CeAD, IAT versus IVT alone (OR 1.32, p=0.79); and (c) IAT in CeAD versus non-CeAD mechanism of AIS (OR 0.58, p=0.34).

Conclusions IAT is a valid alternative therapeutic option for AIS caused by CeAD due to the low complication rate and excellent outcomes observed in this study.

INTRODUCTION

Cervical artery dissection (CeAD) is a recognized cause of acute ischemic stroke (AIS), characterized by intramural hematoma produced by subintimal tear of the carotid and vertebral arterial walls.1 CeAD is uncommon in the general stroke population accounting for just 2–2.5% of all AIS,2 but is a major cause of stroke in the young, responsible for 10–25% of all AIS.1,4

There is debate as to whether anticoagulation is superior to antiplatelet aggregation agents;1,6 the findings from the first published randomized controlled trial comparing antithrombotic therapies found no difference in efficacy, although this pilot trial’s outcomes were rare occurrences.7 CeAD should not be considered a contraindication to acute treatment with intravenous thrombolysis (IVT) because IVT has been shown to be beneficial across all causes of stroke in inducing recanalization of the thrombosis and distal emboli.

Intra-arterial treatment (IAT) may be a complementary option to IVT and antithrombotic therapy in patients with AIS due to CeAD. There are several case reports and small case series of IAT in patients with CeAD, demonstrating the feasibility of this approach.8–23 However, the safety and efficacy of IAT with CeAD has not been formally studied. In light of the recent groundbreaking randomized controlled trials (RCTs)24–28 demonstrating the superiority of IAT over IVT in selected patients with AIS caused by proximal intracranial occlusion in the anterior circulation, our purpose was to examine the safety and efficacy of IAT for AIS caused by CeAD.

DESIGN

We retrospectively studied all consecutively admitted patients with AIS at our high volume comprehensive stroke center between 1 January 2010 and 31 May 2015. The study was investigator initiated and approved by the institutional review board with a waiver of subject informed consent.

The specific aims of this study were twofold: first, to describe the characteristics and outcomes of a large series of patients with CeAD treated with IAT (n=24); second, to analyze the odds of favorable outcome following IAT in the CeAD population versus three comparison populations: (a) patients with CeAD not receiving IAT (n=137), (b) patients with CeAD receiving IVT alone (n=11), and (c) patients with a non-CeAD cause of AIS receiving IAT (n=421).

Outcomes and covariates

The following demographics and clinical characteristics were described: age, gender, race, transfer status, type of dissection, area of occlusion, history/etiology, presenting signs and symptoms, diagnostic modality, National Institutes of Health Stroke Scale (NIHSS), and treatment type. Outcomes included...
favorable 90 day modified Rankin Scale score (mRS ≤2 at 90 days or at discharge where 90 day mRS scores were unavailable, vs mRS 3–6), recanalization (defined as successful/ unsuccessful for the cervical artery, and defined as successful by Thrombolysis in Cerebral Infarction (TICI) scores 2b–3 for the associated intracranial embolus), procedural complications, rate of symptomatic intracerebral hemorrhage (sICH) defined using European Cooperative Acute Stroke Study criteria, and inhospital mortality, and median change in NIHSS (discharge—initial NIHSS).

Statistical analysis
Analyses were performed with SAS V9.3. $\chi^2$ and Wilcoxon rank sum tests were used to analyze demographic and clinical differences for patients with CeAD treated with IAT with our three comparison populations. The outcomes of favorable 90 day mRS 0–2, sICH, and mortality were univariately analyzed with Pearson and Fisher’s exact $\chi^2$ tests. Favorable 90 day mRS 0–2 was also analyzed with multivariate stepwise logistic regression using entry and exit criteria of $p<0.20$. Change in NIHSS was analyzed with Wilcoxon rank sum tests.

Table 1: Intra-arterial thrombolysis treated patients with acute ischemic stroke due to cervical artery dissection

| ID | Age (years) | Transfer | Baseline NIHSS | CeAD type | Intracranial thrombus | IAT | IAT procedures | CeAD recanalization | Intracranial embolus recanalization | Discharge mRS |
|----|-------------|----------|----------------|-----------|-----------------------|-----|----------------|---------------------|-----------------------------------|--------------|
| 1  | 40–49       | Yes      | 0              | Vertebral | Yes                   | No  | IA lytic, thrombectomy, stent | Successful | TICI 3               | 6             |
| 2  | 50–59       | Yes      | 12             | Carotid   | Yes                   | Prior| Angioplasty, IA lytic, thrombectomy, stent | Successful | TICI 3               | 3             |
| 3  | 60–69       | Yes      | 19             | Carotid   | Yes                   | Prior| Stent | Successful | TICI 3               | 3             |
| 4  | 40–49       | Yes      | 27             | Carotid   | No                    | Prior| Stent | Successful | TICI 3               | 2             |
| 5  | 60–69       | Yes      | 13             | Carotid   | Yes                   | No   | IA lytic, thrombectomy | Non-flow limiting | Unsuccessful | 3             |
| 6  | 80–89       | Yes      | .              | Vertebral | Yes                   | No   | IA lytic, thrombectomy | Left occluded | TICI 3               | 6             |
| 7  | 40–49       | Yes      | 4              | Vertebral |*Yes                   | No   | IA lytic, thrombectomy | Successful | TICI 3               | 2             |
| 8  | 50–59       | Yes      | 15             | Carotid   | Yes                   | No   | Angioplasty, IA lytic, thrombectomy, stent | Successful | TICI 2b              | 4             |
| 9  | 70–79       | No       | 9              | Vertebral |*Yes                   | No   | Angioplasty, IA lytic, thrombectomy, stent | Successful | 2b/Unsuccessful | 6             |
| 10 | 50–59       | Yes      | 12             | Carotid   | Yes                   | Prior| Thrombectomy, stent | Successful | TICI 3               | 2             |
| 11 | 40–49       | No       | 24             | Carotid   | No                    | No   | IA lytic, stent | Successful | TICI 3               | 4             |
| 12 | 50–59       | Yes      | 14             | Carotid   | Yes                   | Yes  | Angioplasty, IA lytic, thrombectomy, stent | Successful | TICI 2b              | 4             |
| 13 | 40–49       | No       | 10             | Carotid   | Yes                   | Yes  | Thrombectomy, stent | Successful | TICI 2b              | 4             |
| 14 | 30–39       | Yes      | 16             | Carotid   | Yes                   | No   | IA lytic, stent, thrombectomy | Left occluded | TICI 2b | 3             |
| 15 | 40–49       | Yes      | 12             | Carotid   | Yes                   | Prior| IA lytic, thrombectomy | Non-flow limiting | TICI 2b | 1             |
| 16 | 60–69       | Yes      | 11             | Carotid   |*Yes                   | No   | Thrombectomy, stent | Successful | TICI 3               | 3             |
| 17 | 20–29       | Yes      | .              | Vertebral | Yes                   | No   | Thrombectomy, stent | Left occluded | TICI 2b | 5             |
| 18 | 60–69       | Yes      | 15             | Carotid   | Yes                   | Prior| IA lytic, thrombectomy, stent | Successful | TICI 2b              | 4             |
| 19 | 40–49       | Yes      | 33             | Vertebral | Yes                   | Prior| Thrombectomy | Successful | TICI 2b              | 3             |
| 20 | 50–59       | Yes      | 6              | Carotid   | Yes                   | No   | Angioplasty, IA lytic, stent | Non-flow limiting | TICI 2b | 4             |
| 21 | 50–59       | Yes      | 23             | Carotid   | Yes                   | No   | Angioplasty, IA lytic, stent | Successful | TICI 3               | 6             |
| 22 | 50–59       | No       | 13             | Carotid   | Yes                   | No   | Angioplasty, IA lytic, stent | Successful | TICI 3               | 3             |
| 23 | 50–59       | Yes      | 15             | Carotid   | Yes                   | Prior| IA lytic, thrombectomy | Non-flow limiting | TICI 2b | 4             |
| 24 | 30–39       | Yes      | 12             | Carotid   | Yes                   | Prior| IA lytic, thrombectomy | Non-flow limiting | TICI 3 | 4             |

* Bilateral dissections.
† Patient No 9 had successful restoration of flow and then re-thrombosed 2 days later, with unsuccessful recanalization of re-thrombosis.
CeAD, cervical artery dissection; IAT, intra-arterial thrombolysis; IVT, intravenous thrombolysis; mRS, modified Rankin score; NIHSS, National Institutes of Health Stroke Scale; TICI, Thrombolysis in Cerebral Infarction.
angioplasty (n=7). Eleven patients were also treated with IV tissue plasminogen activator.

Regarding stents, 14 patients with at least 99% CeAD occlusion were stented; 7 stents were placed before approaching the intracranial embolus; 4 stents were placed after approaching the intracranial embolus; patient No 14 was unsuccessfully stented both before and after recanalization of the middle cerebral artery (MCA); patient No 4 had an ICAD stent and did not have an associated intracranial embolus; and patient No 3 had an ICAD stent and did not have IAT performed on the intracranial embolus. More than one stent was required in 8 of 14 patients.

Favorable 90 day mRS 0–2 was achieved in 63% (12/19) of patients. The median improvement in NIHSS was −9 points (IQR −11 to 1). There were four deaths; none was attributable to IAT. There was 1 sICH and 3 procedural complications (2 groin pseudoaneurysms, 1 catheter induced acute dissection).

Among the 18 ICADs, 6 internal carotid arteries were left untreated: 5 were non-flow limiting and 1 was left occluded with excellent intracranial collateral circulation following unsuccessful stenting of the right ICAD (table 1). The remaining 12 ICADs were successfully recanalized (100% successful recanalization).

Seventeen of 18 ICADs had associated MCA occlusions; patient No 4 did not have an associated MCA occlusion but was stented due to inadequate collateral flow. Recanalization of the MCA was successful to TICI 2b–3 in 16 patients and was unsuccessful in 1 patient (94% successful recanalization). Patient No 3 was technically successful with prompt restoration of flow to the intracranial circulation after the ICAD was stented, although the small MCA branch occlusions were not approached.

Among the six VADs, two vertebral arteries were left occluded because the risk of opening it was greater than the perceived benefit of revascularization. The remaining four VADs were successfully recanalized (100%) (table 1).

All six VADs had a basilar artery occlusion; recanalization of the basilar artery was 100% successful to TICI 2b–3 (table 1). In the two VADs where the vertebral artery was left untreated, the basilar artery embolus was extracted through the contralateral patent vertebral artery. In patient No 9, the vertebral artery re-thrombosed 2 days after the initial restoration of flow and the vertebral artery was unsuccessfully recanalized.

Outcomes analysis

Overall there were 3671 patients with AIS. There were 445 patients treated with IAT, including 421 patients with a non-CeAD mechanism of AIS and 24 patients with a CeAD mechanism.

Thirty-five patients with CeAD were acutely treated with IAT (n=24; 11 with and 13 without IVT) or IVT only (n=11) (figure 1). The remaining 125 patients with CeAD were acutely treated with either anticoagulation or antiplatelet therapy.

First we analyzed patients with CeAD who received IAT versus those who did not receive IAT. There were significant differences in demographics, clinical characteristics, and outcomes (table 2). In particular, patients with CeAD who were treated with IAT had significantly more severe stroke compared with those not receiving IAT (median NIHSS 13 vs 3, p<0.001). As a result, unadjusted outcomes were significantly worse in patients treated with IAT than patients who did not receive IAT, with the exception of a greater improvement in NIHSS (−9 vs 1, p=0.02). However, after adjustment for baseline NIHSS and age, the odds of a favorable 90 day mRS 0–2 were not significantly different for patients receiving IAT versus Those not receiving IAT (OR 0.62 (0.12–3.14), p=0.56) (figure 2).

Next we analyzed patients with CeAD treated with IAT versus IVT alone. There were no significant differences between groups in demographics, clinical characteristics, or unadjusted outcomes, except for patients receiving IAT who had more severe stroke than those receiving IVT alone (median NIHSS 13 vs 10, p=0.04) (table 2). After adjustment for baseline NIHSS, there was no difference in the odds of a favorable 90 day mRS 0–2 with IAT versus IVT alone (OR 1.32 (0.16–10.72), p=0.79) (figure 2).

Finally, we analyzed outcomes in 445 patients receiving IAT: 24 with CeAD and 421 with another cause of AIS. Patients with CeAD were younger (median age 52 vs 70, p<0.001) with less severe stroke (median NIHSS 13 vs 18, p=0.03) than those with another cause of AIS (table 2). Despite these differences, unadjusted outcomes were similar by AIS cause (table 2). After adjustment for baseline NIHSS and age, there was no difference in the odds of a favorable 90 day mRS 0–2 with CeAD versus another cause of AIS (OR 0.58 (0.19–1.78), p=0.34) (figure 2).

DISCUSSION

Groundbreaking findings of several recently published RCTs demonstrated the superiority of IAT over standard medical therapy, resulting in updated American Heart Association/ American Stroke Association guidelines recommending endovascular treatment in selected patients with AIS.31 32 The role of IAT in patients with CeAD is empirically demonstrated because there are no controlled trials; data are confined to 16 published case series,8–23 the largest of which reported on 26 patients with CeAD and IAT,33 34 and two systematic reviews by Pham and Zinkstok.33 34 The review by Pham et al33 reported a 99%
technical success rate and 1.3% complication rate with IAT in 140 patients with CeAD; data on mRS were not summarized. The systematic review by Zinkstok et al reported safety and efficacy of thrombolysis (IAT and IVT) in 180 patients with CeAD, of whom 59 received IAT. Overall rates were 3% sICH, 9% mortality, and 56% with mRS ≤2. These rates are similar to our case series of CeAD patients treated with IAT, with 4% sICH, 17% mortality, and 63% with mRS ≤2.

This study also provides class III evidence that the rate of functional independence, defined as mRS 0–2 at 90 days, was similar for our target population of patients with CeAD treated with IAT as the rates observed in our three comparison populations, after adjustment for relevant covariates (namely stroke severity and age). However, caution must be taken when interpreting these findings because the study was not powered to detect significant differences between the groups due to small sample size. First, we examined IAT versus no IAT in 161 patients with CeAD, reporting non-significant adjusted odds of functional independence in favor of no IAT (OR 0.62, 0.12–3.14). Only one other study performed a similar analysis: Engelter et al used the Cervical Artery Dissection and Ischemic Stroke Patients (CADISP) registry to examine patients with CeAD and compared any thrombolysis (n=68, 9 of whom had IAT and 55 had IVT) with no treatment (n=548). In their study, the odds of functional independence were virtually identical between groups, both after adjustment (OR 0.95, 0.45–2.00) and after propensity matching (OR 1.00, 0.49–2.00). Second, we examined patients with CeAD and compared IAT versus IVT alone, reporting adjusted odds of functional independence of 1.32 (0.16–10.72). We identified two meta-analyses that performed a similar analysis, both showing nearly identical odds of functional independence to our study, all non-significantly favoring IAT to IVT alone. Engelter’s 2015 meta-analysis examined 111 patients (32 received endovascular therapy) and reported an OR of 1.41 (0.45–3.45). Zinkstok et al examined 180 patients (59 received IAT), reporting an OR of 1.45 (0.61–3.45). Lastly, we compared IAT in patients with CeAD versus another cause of AIS. Two meta-analyses examined IAT in CeAD versus a non-CeAD mechanism. The first meta-analysis did not report efficacy outcomes but rather showed the sICH rate was no different following IAT in patients with CeAD compared with other causes of AIS identified via the Safe Implementation of Thrombolysis in Stroke-Monitoring Study (SITS-MOST) registry (1.9% vs 1.7%). While our rates of sICH were higher, we also found no significant difference in sICH rates between groups, at 4% versus 6.4%. The second meta-analysis found no difference in the rates of functional independence after IAT in patients with CeAD compared with other causes of AIS identified via the Safe Implementation of Thrombolysis in Stroke-International Stroke Thrombolysis (SITS-ISTR) registry (56% vs 52%). Our reported rates of functional independence after IAT in those with versus Those without CeAD mechanism were comparable (65% and 54%, respectively).

Our results, taken together with the published meta-analyses, suggest three key findings: (1) IAT appears to be as safe and effective when used in patients with ischemic stroke due to

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### Table 2  Clinical and demographic characteristics and unadjusted outcomes

|                         | CeAD and IAT (n=24) | CeAD, No IAT (n=137) | CeAD, IVT alone (n=11) | Non-CeAD mechanism (n=421) |
|-------------------------|---------------------|----------------------|------------------------|-----------------------------|
| Age (years)             | 52 (43–60)          | 44 (35–54)†          | 52 (37–59)             | 70 (61–79)†                 |
| Female sex              | 25.0 (6)            | 41.6 (57)*           | 36.4 (4)               | 54.16 (228)†                |
| White race              | 83.3 (20)           | 65.0 (89)            | 100.0 (11)             | 77.4 (326)                  |
| Transferred in          | 83.3 (20)           | 69.3 (95)            | 72.7 (8)               | 71.7 (302)                  |
| Initial NIHSS           | 13 (12–16)          | 3 (1–7)†             | 10 (7–12)†             | 18 (13–22)†                 |
| Discharge NIHSS         | 6 (3–13)            | 1 (0–3)†             | 9 (3–16.5)             | 1 (0–3)†                    |
| Change in NIHSS         | –9 (–11–1)          | –1 (–3–0)†           | –4 (–5.5–5)            | –10 (–15–5)                 |
| Discharge mRS           | 4 (3–4)             | 2 (1–3)              | 3.5 (3–4)              | 2 (1–3)†                    |
| 90 day mRS 0–2§         | 65.0 (13/20)        | 93.0 (93)†           | 60.0 (3/5)             | 53.7 (145/270)              |
| Symptomatic ICH         | 4.0 (1/25)          | 0 (0)§               | 0.0 (0/11)             | 6.4 (27/421)                |
| Inhospital mortality    | 16.0 (4/25)         | 3.7 (5)†             | 9.1 (1/11)             | 16.4 (69/421)               |

Values are % (n) or median (IQR). Change in NIHSS: discharge NIHSS—initial NIHSS (negative values indicate improvement). Pearson χ² p value versus CeAD and IAT.

* <0.15.
† <0.05.
‡ <0.01.
§ Defined as mRS 0–2 at 90 days or discharge, or discharge home.

CeAD, cervical artery dissection; IAT, intra-arterial thrombolysis; ICH, intracerebral hemorrhage; IVT, intravenous thrombolysis; IV tPA, intravenous tissue plasminogen activator; mRS, modified Rankin Scale; NIHSS, National Institutes of Health Stroke Scale.

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### Figure 2  Adjusted odds of a favorable 90 day modified Rankin Scale score (mRS) (defined as mRS 0–2 at 90 days or discharge, or discharge home). Point estimates are noted with a diamond; estimates to the right of 1.0 favor the referent population. CeAD, cervical artery dissection; IAT, intra-arterial thrombolysis, IVT, intravenous thrombolysis.
CeAD compared with other causes; (2) IAT may be favored over IVT alone based on the direction of the adjusted ORs of 1.32, 1.41, and 1.45 across published studies (the CIs for all ORs were wide, which can be partially explained by the small sample sizes); and (3) despite the IAT group being a select group with more severe baseline deficits and presence of occlusions, the adjusted odds of achieving functional independence was similar in patients with CeAD treated with IAT to those not treated with IAT.

Our comprehensive stroke center has an active interventional neuroradiology program, with approximately 150 IAT procedures performed in the past year. Our stroke center does not have explicit guidelines, but as a general rule patients are treated with IAT if there is distal embolus or thrombus extension or if there is a decline despite receiving standard medical therapy. CeAD is not considered a contraindication to treatment; approximately 15% of the CeAD population were treated with IAT, which is similar to 12% treatment in the broader AIS population. Patients with CeAD who were treated with IAT were likely selected because of the presence of large deficits and MCA or basilar artery occlusions.

The primary limitation to this analysis was that this study was retrospective and carries with it the limitations of such a study design, including reliance on existing medical records and lack of controls. Second, this study was conducted at a single center and, despite reporting a large series of CeAD, the subgroups for analysis were small. While our analysis was performed in a smaller sample size than the published meta-analyses and the study using the CADISP registry, we were able to control for potentially confounding factors, such as population and treating physician and hospital, in addition to stroke severity.

CONCLUSION
These data suggest IAT is an alternative therapeutic option for AIS due to CeAD. Outcomes appear to be similar to those presented in large RCTs with stroke from all causes, and to the comparison populations at our high volume comprehensive stroke center. Following the major advances in IAT for AIS in the general population, IAT should be considered in patients with AIS due to CeAD. Given the small numbers in our series and the paucity of data in the literature, we encourage others to present their experience with IAT in the CeAD population. We believe randomized trials of IAT in CeAD are appropriate and timely, particularly in patients with intracranial occlusions.

Contributors All authors gave final approval of the published version and agree to be 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. Specifically, JJ contributed to conception and design, acquisition and interpretation of the data, and drafting the manuscript. KS contributed to conception and design, analysis and interpretation of the data, and drafting the manuscript. DF contributed to acquisition and interpretation of the data and critical revision. DL contributed to design of the study and critical revision. KMCh contributed to acquisition of the data and critical revision. JW contributed to interpretation of the data and critical revision. MW contributed to acquisition of the data and critical revision. RB contributed to interpretation of the data and critical revision. DB-O contributed to interpretation of data and critical revision.

Competing Interests DF reports a consulting relationship with MicroVention, Siemens, Penumbra, Stryker, and Codman. DF owns options to purchase Penumbra equity. JW is a speaker for Genentech. MW reports a consulting relationship with Clinical Data Management and Lombardi Hall, and is a speaker for Activate. RB is a speaker for Penumbra.

Ethics approval The study was approved by HealthOne IRB.

Provenance and peer review Not commissioned; externally peer reviewed.

Data Sharing statement Anonymized data and statistical code are available from the corresponding author (dbaror@ampiopharma.com).

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