Unilateral agenesis of the internal carotid artery with intracavernous carotid-carotid anastomosis

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CASE SUMMARY
A 91-year-old right-handed man presented to the emergency department with left-sided weakness, which started approximately five hours prior to presentation. Pertinent medical co-morbidities included atrial fibrillation and a prosthetic aortic valve, not on anticoagulation, as well as hypertension. His initial National Institute of Health Stroke Scale (NIHSS) score was 18, and his clinical presentation was consistent with a right hemispheric infarct. Symptoms included dysarthria, left lower facial droop, flaccid left arm, weak left leg, and left hemineglect. As part of the routine stroke protocol at our institution, a non-contrast CT head, CT perfusion of the brain, and CT angiography of the head and neck were obtained.

IMAGING FINDINGS
CT angiography of the neck demonstrates a normal course of the left common carotid artery, which terminates/continues as the external carotid artery; there is no bifurcation and no internal carotid artery (Figure 1). At the level of the skull base, the left internal carotid artery is absent (Figure 2A) and there is also absence of the left carotid canal (Figure 2B). CT angiography of the head demonstrates a vessel branching from the cavernous segment of the right internal carotid artery, which crosses the cavernous sinus, and assumes the course of the intracranial left internal carotid artery terminus (Figure 3).

DIAGNOSIS
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DISCUSSION
The internal carotid artery (ICA) arises from the embryonic third aortic arch.1 Congenital absence of the ICA is classified as a spectrum of rare anomalies, which occur in less than 0.01% of the population, and range from agenesis (complete failure of ICA development) to hypoplasia (incomplete ICA development).2,3 Lie4 developed a classification system with six types of collateral circulation patterns, which can be seen with carotid absence (Figure 4):

Type A: Unilateral ICA agenesis; flow into the ipsilateral anterior cerebral artery (ACA) via the anterior communicating artery (AComm), and flow into the ipsilateral middle cerebral artery (MCA) via flow through the ipsilateral posterior communicating artery (PComm).

Type B: Unilateral ICA agenesis; flow into the ipsilateral ACA and MCA via the AComm.

Type C: Bilateral ICA agenesis; flow
Type C:   Bilateral ICA hypoplasia; flow into the MCAs via enlarged PComms.

Type D:   Unilateral ICA agenesis; flow into the ipsilateral carotid siphon via an intracavernous carotid-carotid anastomosis from the posterior circulation via the PComms or other patent carotid-vertebrobasilar anastomoses.

Type E:   Bilateral ICA hypoplasia; flow into the MCAs via enlarged PComms.

FIGURE 1. Curved reformatted image demonstrates the left common carotid artery (black arrows) continuing as the external carotid artery (white arrows), which then branches as expected. There is no carotid bifurcation and no internal carotid artery.

FIGURE 2. (A) Axial CT image at the level of the skull base demonstrates the right internal carotid artery (white solid arrow) and bilateral vertebral arteries (black arrows); however, the left internal carotid artery is absent (white dotted arrow). (B) Just superiorly, the right internal carotid artery is present within the right carotid canal (white arrow), however, the left internal carotid artery and left carotid canal are absent.

FIGURE 3. (A) Axial, (B) sagittal, and (C) coronal images demonstrate a vessel traversing the sella (arrows), which arises from the right internal carotid artery and continues as the intracranial left internal carotid artery terminus, consistent with intracavernous carotid-carotid anastomosis.
FIGURE 4. (A) Normal (conventional anatomy). (B) Depiction of Lie’s classification system with six types of collateral circulation patterns, which can be seen with carotid absence or hypoplasia.
Type F: Bilateral proximal ICA agenesis; flow into the distal ICAs via the rete mirabile (transcranial anastomoses arising from the internal maxillary arteries).

A simplification of Lie’s classification system describes three types of collateral pathways in the setting of ICA absence: 1) flow through the circle of Willis, 2) persistent fetal vessels, and 3) branches from the external carotid artery.2,3

This case is an example of type D, in which the cervical, petrous, and cavernous segments of the ICA are absent, with the supraclinoid segment arising from an intracavernous vessel from the contralateral ICA. There are two theories regarding the origin of this anastomotic vessel. One is that there is an intracavernous connection between the two primitive trigeminal arteries, which do not connect to the posterior circulation.4 The other is that there is compensatory hypertrophy of a clival or sellar branch off the contralateral ICA, which eventually becomes the main blood supply for the intracranial ICA territory.5 In this case, as in other reports of intracavernous anastomosis,2,3,6-8 the intracavernous vessel continues as the middle cerebral artery (MCA), while the A1 segment of the anterior cerebral artery (ACA) ipsilateral to the absent ICA is aplastic or hypoplastic; this is slightly different than Lie’s original description in which the intracavernous anastomosis supplies both the MCA and ACA territories. Thus, the MCA territory is supplied by the intracavernous anastomosis, while the ACA territory is supplied via a patent anterior communicating artery. This is important because the relative isolation of the MCA territory leaves the cerebral hemisphere ipsilateral to the absent ICA susceptible to injury, due to lack of collateral flow from the circle of Willis. For example, in this case, occlusion of the right ICA could potentially cause infarct in the bilateral MCA territories.

Congenital absence of the ICA in itself is a benign finding and often found incidentally;9 however, there is an increased incidence of concurrent intracranial aneurysms upwards of 34%,10 as compared to the 2-4% incidence in the general population. It is therefore important to look carefully for aneurysms; none were found in this case.

CONCLUSION

Congenital absence of the internal carotid artery is a rare congenital anomaly. Collateral circulation to the ipsilateral cerebral hemisphere may be provided by flow through the circle of Willis, persistent fetal vessels, or collaterals from the external carotid artery. This case is an example of unilateral internal carotid agenesis with intracavernous anastomosis, with collateral flow from both the circle of Willis and from a persistent fetal vessel. This entity may be benign, but can have profound implications regarding collateral circulation to the ipsilateral cerebral hemisphere in the event of ischemia, as well as an increased incidence of intracranial aneurysms.

REFERENCES


Prepared by Dr. Montoya while in the Department of Imaging Sciences; Dr. Towner while in the Department of Neurosurgery; and Dr. Bhatt while in the Department of Imaging Sciences at the University of Rochester Medical Center, Rochester, NY.