A tale of two arteries: dual posterior cerebral arteries with vascular bridges. A possible protective pattern?

Authors: Y. Mansour, R. Kulesza

DOI: 10.5603/FM.a2020.0070

Article type: CASE REPORTS

Submitted: 2020-06-19

Accepted: 2020-07-01

Published online: 2020-07-08

This article has been peer reviewed and published immediately upon acceptance. It is an open access article, which means that it can be downloaded, printed, and distributed freely, provided the work is properly cited. Articles in "Folia Morphologica" are listed in PubMed.
A tale of two arteries: dual posterior cerebral arteries with vascular bridges. A possible protective pattern?

A tale of two arteries: dual posterior cerebral arteries with vascular bridges

Y. Mansour, R. Kulesza
Department of Anatomy, Lake Erie College of Osteopathic Medicine, Erie, PA 16509, United States

Address for correspondence: Dr. Yusra Mansour, Department of Anatomy, Lake Erie College of Osteopathic Medicine, 1858 West Grandview Blvd, Erie, PA 16509, USA, tel: (419) 442-1354, e-mail: ymansour49902@med.lecom.edu

Abstract

Stroke is a common morbidity and a frequent cause of disability and even death. The impact of cerebrovascular events are dictated by the brain region involved and can be complicated by anatomical variations. One of the most common variations impacting the cerebrovasculature is the presence of a fetal posterior cerebral artery. This vessel arises from the internal carotid artery instead of the basilar artery and is often associated with more extensive injury in cerebrovascular events. Herein, we report the case of a 60-year-old male who had numerous arterial abnormalities, including a kink and a coil of the left internal carotid, two posterior communicating arteries on the right and two posterior cerebral arteries (PCA) on the left, one arising from the internal carotid (fetal PCA) and one from the basilar. The fetal PCA supplied the thalamus, splenium of the corpus callosum and primary visual cortex. The basilar PCA supplied the midbrain and parts of the occipital lobe. These PCAs were connected to each other by a vascular bridge and the fetal PCA was connected to the middle cerebral artery by an additional vascular bridge. This vascular pattern would appear to provide collateral support around blockages in the internal carotid and main stem middle and posterior cerebral arteries.

Key words: variation, cerebral, vasculature, fetal
INTRODUCTION
Stroke is a leading cause of death and a major cause of disability [1]. Stroke, its treatment and subsequent mortality can be complicated by variations in arterial patterns. One of the most common intracranial arterial variations is persistence of the fetal posterior communicating artery (PCOM), which occurs in approximately 20% of subjects [2, 3, 4, 5]. Therefore, clinical knowledge of both normal and variant cerebrovascular patterns is important in predicting vascular risk, recognizing common and atypical stroke syndromes and planning neurointerventions [2, 6, 7]. Herein, we report a rare case of a subject with two posterior cerebral arteries (PCA) on the left and two PCOMs on the right side. Additionally, this subject had two abnormalities involving his left internal carotid artery [8, 9]. We discuss the embryological origin of these vessels and their potential impacts/consequences.

CASE REPORT
Case presentation
A 60-year-old male presented to the emergency department with the chief complaints of seeing flashing lights in his right eye and left arm weakness. His neurological exam was non-focal. He underwent CT and CTA imaging to rule out possible stroke. The CT series was imported into Amira (Version 6.7.0) and a slice by slice approach was taken to create a 3D rendering of both his carotid and vertebral circulations.

Imaging findings
Examination of the 3D model (figure 1A, B) of this subject's cerebrovasculature revealed multiple variations. On the left internal carotid artery (ICA) there was a cervical kink (figure 1, white arrowhead) and a cavernous coil (figure 1, yellow arrowhead) [6]. On the left side there was a PCA that arose at the basilar bifurcation (“basilar PCA” - bPCA; figure 1, blue arrowheads) and a fetal PCA (fPCA; figure 1, green arrowheads) that continued directly from the internal carotid (ICA). The CT and 3D reconstruction clearly showed both of these arteries wrapping around the midbrain and distributing within the occipital lobe. The fPCA had a vascular bridge to the middle cerebral artery (MCA; figure 1, black arrowhead) and
bPCA (figure 1, cyan arrowhead). On the right side there were two PCOMs (figure 1, orange arrowheads). A video of the 3D reconstruction can be found here: https://drive.google.com/file/d/1AfJl9WwRe4LNhH2gNECcjPqQa75hRtiY/view?usp=sharing

**Correlative anatomy**

Developmentally, the arterial supply of the brain is derived from vertebral and internal carotid arteries and the PCOM initially connects these two circulations (figure 2A). The PCA is normally derived from the PCOM and supplies the midbrain, diencephalon and caudal telencephalon. We propose that the arterial variations identified in this subject arose from the altered embryonic vascular pattern shown schematically in figure 2B. Careful study of the 3D reconstruction and CTA series allowed us to develop a vascular map (figure 2C, D). In the CTA, we carefully traced branches from both the bPCA and fPCA and constructed maps of their vascular territories. In this subject, the bPCA supplied the central aspect of the midbrain and the lateral aspect of the PCA territory (figure 3). The fPCA supplied the lateral aspect of the midbrain, the entire thalamus, splenium of the corpus callosum and the medial aspect of the occipital lobe including the calcarine sulcus.

**DISCUSSION**

The PCA normally develops as a direct continuation of the embryonic PCOM and supplies the midbrain, thalamus and occipital lobe. The most common variant of the PCOM/PCA is a fPCA where this vessel forms as a direct branch from the internal carotid artery (ICA) [2, 3, 4, 5]. When a fPCA is present, the ICA provides arterial supply to the entire cerebral hemisphere, thalamus and midbrain via anterior, middle and posterior cerebral arteries. Accordingly, if a subject with a fPCA has a blockage of the ICA, this can impair blood flow to the entire thalamus and cerebral hemisphere with often catastrophic results [10, 5]. However, the presence of a fPCA is not considered an additional risk factor for stroke [11].

The subject of this case presentation has both a bPCA and a fPCA and these vessels made major connections to other vessels in the left hemisphere through vascular bridges. In our 3D reconstruction and study of the CTA series, we found the fPCA and bPCA were
connected by a small vascular bridge near their origin [4], but there were no additional connections between the two PCAs. Further, we found an additional vascular bridge connecting the fPCA and MCA. Together, we believe this arterial architecture may provide a protective support pattern. Specifically, if the subject of this case presentation were to have a thrombus in the ICA or in the main stem of fPCA, bPCA or MCA on the left, the bridging vessels could provide collateral support and potentially prevent ischemia or infarct in PCA or MCA territories.

CONCLUSIONS

The fPCA is a fairly common anatomical variation involving arterial support of the midbrain, thalamus and occipital lobe. Vascular compromise involving the anterior circulation in a subject with a fPCA is often associated with extensive ischemia and poor outcomes. However, the presence of vascular bridges connecting the fPCA to surrounding mainstem vessels may provide protective anastomoses.

Acknowledgements

The authors would like to thank the University of Pittsburgh Medical Center Hamot, Department of Neurology for the CT and CTA data.

REFERENCES

1 Stroke. (2020, April 7). Centers for Disease Control and Prevention. https://www.cdc.gov/stroke/index.htm

2 Zampakis P, Panagiotopoulos V, Petsas T, Kalogeropoulou C. Common and uncommon intracranial arterial anatomic variations in multi-detector computed tomography angiography (MDCTA). What radiologists should be aware of. Insights Imaging. 2015 Feb; 6(1): 33–42.

3 Uchino A, Saito N, Takahashi M, Okano N, Tanisaka M. 2016. Variations of the posterior cerebral artery diagnosed by MR angiography at 3 tesla. Neuroradiology 58:141–146
4 Vasović L, Trandafilović M, Jovanović I, Antović A, Stojanović J, Zdravković M, Milić M. 2010. An excess vessel in the posterior part of the human cerebral arterial circle (CAC): a case series. BMC Neurol. Jun 23;10:53.

5 Klimek-Piotrowska W, Kopeć M, Kochana M, et al. Configurations of the circle of Willis: a computed tomography angiography based study on a Polish population. *Folia Morphol (Warsz).* 2013;72(4):293-299. doi:10.5603/fm.2013.0049

6 Brzegowy P, Polak J, Wnuk J, Łasocha B, Walocha J, Popiela TJ. Middle cerebral artery anatomical variations and aneurysms: a retrospective study based on computed tomography angiography findings. *Folia Morphol (Warsz).* 2018;77(3):434-440. doi:10.5603/FM.a2017.0112

7 Zurada A, Gielecki JS. A novel formula for the classification of blood vessels according to symmetry, asymmetry and hypoplasia. *Folia Morphol (Warsz).* 2007;66(4):339-345.

8 Benson JC, Brinjikji W, Messina SA, Lanzino G, Kallmes DF. 2020. Cervical internal carotid artery tortuosity: A morphologic analysis of patients with acute ischemic stroke. Interv Neuroradiol. 2020 Apr;26(2):216-221.

9 Griessenauer CJ, Yalcin B, Matusz P, Loukas M, Kulwin CG, Tubbs RS, Gadol AA. 2015. Analysis of the tortuosity of the internal carotid artery in the cavernous sinus. Childs Nerv Syst. Jun;31(6):941-4.

10 Yu J, Qu L, Xu B, Wang S, Li C, Xu X, Yang Y. Current Understanding of Dolichoarteriopathies of the Internal Carotid Artery: A Review. Int J Med Sci. 2017; 14(8):772-784.

11 de Monyé C, Dippel DW, Siepman TA, Dijkshoorn ML, Tanghe HL, van der Lugt A. Is a fetal origin of the posterior cerebral artery a risk factor for TIA or ischemic stroke? A study with 16-multidetector-row CT angiography. J Neurol. 2008 Feb;255(2):239-45.
Figure 1. Images of 3D reconstruction (A-F) and CTA (G) of this patient’s arterial pattern. The views in A and B show the carotid and vertebral arteries (V) from their origin in the neck. The formation of the basilar artery (B) appeared normal. In the neck, the left ICA had both a coil (yellow arrowhead; observed also in D) and a kink (white arrowhead). The ICAs were found to give rise to the middle (MC) and anterior (AC) cerebral arteries. In C, E and F the abnormal PCAs and PCOMs are identified. Figure C shows a top-down view. The blue arrowheads indicate the bPCA, and the orange arrowhead indicates the PCOM. On the patient’s right side there were two PCOMs. On the patients left, there were two PCAs: one from the ICA (fPCA) and one from the basilar (bPCA). There were two vascular bridges on the left: one from the fPCA to the MCA (black arrowhead) and one from the fPCA to the bPCA. Figure E shows the two PCAs from the patients left side. Figure F shows the double PCOM from the patient’s right side. Figure G shows a slice from the CTA study. The fPCA is indicated by the green arrowhead and the bPCA is indicated by the blue arrowhead.

Figure 2. Proposed arterial patterns. Shown in A is the typical embryologic pattern that forms the PCOM and PCA and the pattern we propose developed in this subject in B. Shown in C is the typical arterial pattern and in D is the arterial pattern discovered in this case study. Key to arrowheads: green-fPCA, blue-bPCA, orange-PCOM, black-fPCA to MCA vascular bridge, cyan-fCPA to bPCA vascular bridge.

Figure 3. Shown in A-B are transverse slices through the brain at the level of the thalamus (A) and midbrain (B). Shown in C is a coronal slice through the brain at the level of the thalamus. On the left side of each image (the patient’s right), the territory of the bPCA is shown in red. On the right side of each image (the patient’s left) the territory of both the bPCA and fPCA are shown. Note that in the subject of this case study, the fPCA supplies the lateral and posterior midbrain, thalamus, splenium of the corpus callosum and primary visual cortex. The inset in the bottom right shows the approximate levels and locations of the slices shown in A-C.
