

An Autopsic Study on Bilateral Hypoplasia of the Posterior Communicating Arteries in Cerebrovascular Diseases

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Abstract

Objective: To identify the clinical consequences of bilateral hypoplasia of the posterior communicating arteries (PCoAs) in order to provide a clue to neurosurgeons and neurologists in their work to diagnose and treat patients with stroke symptoms.

Material and methods: We performed an anatomical retrospective study on gross morphology of 98 circles of Willis (CW) that were identified at the time of the autopsies made on patients who have died in the "Prof. dr. N. Oblu" Emergency Clinical Hospital Iasi, Romania, the main neurological and neuro-surgical healthcare Centre in the region of North-Eastern Romania, for a period of three years. Demographical and autopsic data, as well as photographic images were taken from the registers and archives of the Pathology Department. Only CWs with bilateral hypoplasia of PCoA without any other associated anatomical abnormalities were studied.

Results: In 5 cases (5.12%) we identified bilateral hypoplasia of PCoAs. Male: Female ratio was 3: 2. The average age was 63.5 years. 80% of cases died of ischemic or haemorrhagic stroke, but 10% died of heart disease. In 2 (40%) cases, one of the two PCoAs presented aplasia and the other was hypoplastic, an event related to frontal lobe strokes on the same side. In the other two (40%) cases in which both arteries were hypoplastic, the cause of death was ischemic stroke located in the brainstem. In only one case (20%), even in the presence of bilateral hypoplasia of PCoAs, the cause of death was non-neurological.

Conclusion: Even if literature claims that PCoA hypoplasia becomes a risk factor for ischemic stroke only in the presence of ipsilateral internal carotid artery (ICA) occlusion, in our study we found that aplasia of one of the two PCoAs associated with hypoplasia of the controlateral artery caused ischemic stroke in the brainstem, thus revealing the role of PCoAs in ensuring the posterior circulation in case of occlusion of the vertebro-basilar system.

Keywords: Posterior communicating artery; Bilateral hypoplasia; Autopsy

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Introduction

Circle of Willis (CW) or circulus arteriosus cerebri is an anastomotic arterial structure located at the base of the brain, around the optic chiasm and other structures of the interpeduncular fossa.

The posterior communicating artery (PcoA) has its origin in the intradural part of the internal carotid artery (ICA), below the point of emergence of the anterior choroidal artery and it follows a posterolateral path, above the oculomotor nerve, to join the posterior cerebral artery, which is a branch of the basilar trunk. PCoA serves as an anastomotic channel between the anterior and posterior cerebral circulation as it extends posteriorly and

join the primary segment of the ipsilateral Posterior Cerebral Artery, thus completing the circle of Willis. In its path, PCoA sends branches that irrigate the optic tract, optic chiasm, posterior hypothalamus, the pituitary stalk, the anterior and ventral nuclei of the thalamus [1].

Therefore, this artery provides a major blood supply to the ipsilateral cerebral hemisphere if there is a hypoplasia or an obstruction of the ICA, basilar artery or vertebral arteries.

On the other hand, occlusion, aplasia, or hypoplasia of the ACoP can significantly compromise brain irrigation, which can lead to the development of a stroke.

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In the literature, autopsic reports of hypoplastic PCoAs are quite rare, but articles with bilateral hypoplasias are even rarer, either as isolated cases [2] or as a number of cases included in a large series of anatomical variants of CW [3]. So far, only 24 studies have been published. However, there are articles on this issue, but they were realized as angiographic studies [4].

The aim of our research was to identify the clinical consequences of bilateral hypoplasia of the posterior communicating arteries (PCoAs) in order to provide a clue to neurosurgeons and neurologists in their work to diagnose and treat patients with stroke symptoms.

Materials and Methods

This research was done in the Department of Pathology, "Prof. dr. N. Oblu" Emergency Clinical Hospital, Iasi, Romania, mentioning the fact that this represents the main neurological and neuro-surgical healthcare Centre in the region of north-eastern Romania.

The study was started by undertaking the institutional ethical approval. We performed an anatomical retrospective study on gross morphology of 98 circles of Willis that were identified at the time of the autopsies, which were performed at the request of the attending physician to establish the diagnosis of death in patients who died in same hospital in a three years period (January 1, 2013-December 31, 2015). At the time of the each autopsy, the pathologist (AS) carefully separated CW from brain tissue for better view. The pathologist (AS), assisted by two neurosurgeons (DMT and AIC), made detailed morphological and morphometric analysis of each circle of Willis. All constituting arteries of the CW were observed, measuring the outer diameter that was and note in a autopsy register. Gross morphological variations of CW were photographed and later archived.

We included in this study only those circles of Willis with bilateral hypoplasia (less than 0.5mm in the outer diameter) of the posterior communicating arteries (PCoAs) [2] and without any other associated anatomical abnormalities. The circles of Willis that showed normal anatomy or other gross morphological variations were excluded from the study. Any vessel with an outer diameter less than 0.5 mm was considered hypoplastic.

We recorded from the register of the Pathology Department of the same hospital the following data: age and gender of each deceased patient, the outer diameters of both PCoAs (left and right) for each circle of Willis, which were measured with a ruler and recorded during the autopsy by the pathologist, but also the medical cause of the death for these patients. All demographical data and autopsic findings were recorded in the patient data sheet.

Results

Of the 98 deceased who were autopsied in the hospital Prosecutor's Office during the three years studied period, only in 5 cases (5.12%) we identified bilateral hypoplasia of the PCoAs (Figure 1).

Of these 5 cases, three were male and two were female (Male-Female Ratio=3:2) (Figure 2). The average age of the deceased

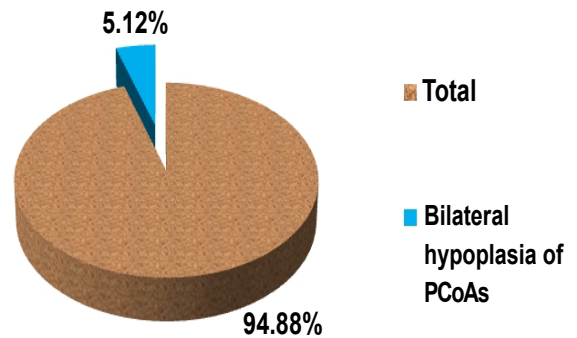


Figure 1 Percentage representation of bilateral PCoA hypoplasia in our study.

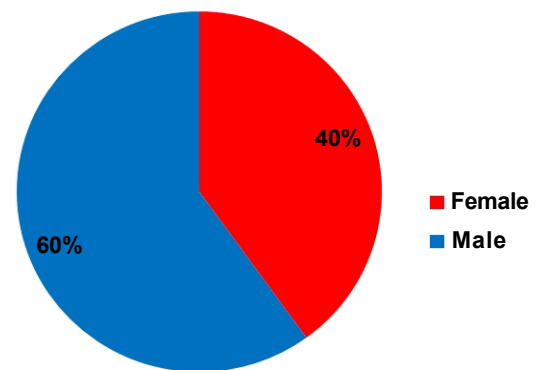


Figure 2 The Male to Female Gender Ratio in our study (M:F=3:2).

patients that were included in our study was 63.5 years.

Table 1 highlights the variety of causes of death in bilateral hypoplasia/aplasia of PcoAs. Most of the cases (4/5 cases) revealed a stroke located either supratentorially, in the frontal or fronto-parietal area, or in the brainstem, and manifested as hematoma, hemorrhage or infarction. However, in each case, other causes of death were associated, such as hypertension or systemic atherosclerosis. Only one case died of non-neurological causes (Table 2).

In two (40%) cases, we found a strokes in frontal lobe. In two (40%) cases, the cause of death was an ischemic stroke of the brain stem.

In only one case (20%), even in the presence of bilateral hypoplasia of PCoAs, the cause of death was non-neurological.

Discussions

Normal PCoA presents an external diameter of 1.00-1.5 mm and an average length of 1.33-1.36 cm, regardless of its origin [5].

The following anatomical variants of PCoA are described in the literature: 1). hypoplasia of one or both PcoAs (i.e. the outer diameter should be less than 0.5 mm, regardless of its length); 2). aplasia of one or both PcoAs (a complete or partial failure of the arteru to grow); 3). atresia of one or both PcoAs (i.e. the outer diameter should be less than 0.5 mm and lack of a vascular lumen), 4). bilateral absence of PcoAs; 5). fenestration; 6). persistent fetal pattern [3,6-8].

Table 1 Demographic and morphological characteristics of our cases.

No.	Gender/Age	Right PCoA	Left PCoA	Medical Cause of Death
1.	M/50	Aplasia	Hypoplasia	Right frxonto-parietal hemorrhagic infarction Occlusion of right ICA Systemic hypertension
2.	F/71	Hypoplasia	Hypoplasia	Ponto-mesencephalic hemorrhage Systemic atherosclerosis Arterial hypertension
3.	M/76	Hypoplasia	Aplasia	Left F hematoma Occlusion of left ICA Systemic atherosclerosis
4.	M/40	Hypoplasia	Hypoplasia	Bulbar and pontine infarction Occlusion of basilar trunk Arterial hypertension
5.	M/67	Hypoplasia	Hypoplasia	Acute myocardial infarction of the left ventricle

PCoA=Posterior Communicating Artery; ICA=Internal Carotid Artery; F=Frontal; P=Parietal; T=Temporal; IV=intraventricular

Table 2 Comparison of the data obtained in our study with those from literature.

Authors	Country	Material	Hypoplasia
Kapoor et al [9]	India	1000 brains	13,2%
Sinha et al [10]	India	80 brains	17%
Gunnal et al [8]	India	150 formalized brains	27,33%
Siddiqi, Tahir and Lone [11]	Pakistan	51 fresh brains	15,6% unilateral hypoplasia 7,8% bilateral hypoplasia
Macchi et al [12]	Italia	100 healthy patients	21%
Paşcalău et al [13]	Romania	10 formalized brains with CW with anatomical variants	41%
Our study	Romania	98 fresh brains	5,12% (bilateral)

In the studies published so far, there are not many reported variations of PCoAs. More over, there are very few studies on the proposed topic, usually as isolated case reports (Nagawa et al). There are also a few reports on large number of cases with hypoplasia of PcoA, bur usually they presented only the unilateral variant [3,8]. Only Gunnal et al, in India, analysing 170 human cadaveric CW, found numerous types of variations of PCoA in the form of its aplasia, unilateral or bilateral hypoplasia, fenestration, and persistent fetal pattern. From all this types of variants, unilateral and bilateral hypoplasia were identified in a quater of the cases [3].

However, all studies published until now, realized on autopsied brains or on cerebral angiographs, in Asia or Europe, showed that hypoplasia is the most common anatomical variation of PCoAs [9-13].

The reduction of the diameter of the PCoA is associated with reducing the required blood supply to the brain because the volume of blood flow is inversely proportional to the length of the artery and directly proportional to its diameter. Thus, the shorter and wider one of the PCoA, the more efficient the transmission of blood to the nervous tissue dependent of that artery will become. Conversely, the longer and narrower the lumen of one or both PCoAs, the weaker the irrigated area of the brain will be [2].

This fact explains the development of extensive stroke identified

by us in bilateral hypoplasia of PCoA, especially if there is an associated systemic pathology as well as the occlusion of an important artery that ensures brain irrigation, such as ICA or basilar trunk.

Even if literature claims that PCoA hypoplasia becomes a risk factor for ischemic stroke only in the presence of ipsilateral ICA occlusion [14], in our study we found that aplasia of one of the two PCoAs associated with the hypoplasia of the controlateralPCoA causes ischemic stroke of the brainstem, revealing their role in ensuring posterior circulation in case of occlusion of the vertebro-basilar system, too.

Conclusions

We have found that the bilateral hypoplasia of the two Posterior Communicating Arteries led to a stroke, either haemorrhagic or ischemic, which ultimately caused death of that individual. Although bilateral hypoplasia of both Posterior Communicating Arteries may be asymptomatic and neurologically uncomplicated, in most cases this anatomical variant increases the risk of hemorrhagic or ischemic stroke and plays a significant role in planning neurosurgical interventions and in diagnoses of neurological diseases.

Disclosure of Conflicts of Interest

None

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