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Fenestrations of the Middle Cerebral Artery Associated with Aneurysms

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Fenestration of the vertebrobasilar artery is well known, but that of the middle cerebral artery (MCA) has been reported rarely [1–8]. We report two cases with fenestration of the MCA associated with cerebral aneurysm. The pathogenesis and clinical significance of fenestration of the MCA in relation to cerebral aneurysms is discussed.

Case Reports

Case 1

A 65-year-old man experienced a sudden, severe headache followed by left hemiparesis. On neurologic examination somnolence, nuchal rigidity, left spastic hemiparesis, and bilateral papilledema were present. Computed tomography (CT) demonstrated an intracerebral hematoma in the right temporal lobe, and right carotid angiography showed an aneurysm arising from the bifurcation of the right MCA and a fenestration at the origin of the same MCA (fig. 1). At surgery marked arteriosclerosis in the vicinity of the aneurysm was found. The aneurysm was clipped at its neck. The fenestrated artery at the origin of the MCA was not exposed surgically.

Case 2

A 45-year-old man had a subarachnoid hemorrhage. On admission bilateral papilledema with retinal hemorrhage and an abducens nerve paresis on the left were present. Cerebral angiography demonstrated a persistent primitive trigeminal artery and four cerebral aneurysms: on the left vertebral artery close to the origin of the posterior inferior
cerebellar artery, on the left MCA, and on the right internal carotid artery. An aneurysm was also present in the axil of a fenestrated right MCA (fig. 2). CT demonstrated high-density areas in the basal cisterns, both sylvian fissures, and the cavum septi pellucidi and cavum vergae. All the aneurysms were successfully clipped or coated. The aneurysm on the right MCA had originated from the proximal end of the fenestration.

Discussion

Fenestration of the MCA has been rarely reported. The incidence is 0.28% at autopsy [1] and 0.26% on angiography [3]. To our knowledge, 13 such cases have been reported in the literature [1-8] with nine cases in the Japanese. Seven of the nine cases were male and two were female. Cerebral aneurysms were present in six of the 13 cases. In all cases the aneurysms arose in the ipsilateral artery at the fenestration or from the anterior communicating artery. In our case 1 the fenestration and aneurysm were in the same MCA.

Crompton [1] stated that aneurysms could occur at the proximal end of arterial fenestrations independent of any branches. Our case 2 appears to confirm this.

Embryologically a primitive vascular network is formed between the anterior cerebral artery and the MCA, and variations of the cerebral arteries such as fenestrations are explained by the persistence of embryologic transition branches and anastomoses [9]. Kwak et al. [4] considered that the association of fenestration and cerebral aneurysm is incidental. Other investigators [2, 5, 10] reported that fenestrations were apt to accompany cerebral aneurysms. In our case 2, four aneurysms were present in association with fenestration of the MCA and a persistent primitive trigeminal artery. We believe that the association of the middle cerebral artery fenestrations and aneurysms is from concurrent congenital anomalies.

REFERENCES

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