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Duplicated middle cerebral artery associated with aneurysm at M1/M2 bifurcation: a case report

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Abstract

Background: A duplicated middle cerebral artery arises from the internal carotid artery and supplies blood to the middle cerebral artery territory. A duplicated middle cerebral artery is sometimes associated with an intracranial aneurysm. Most aneurysms associated with duplicated middle cerebral artery are located at the origin of the duplicated middle cerebral artery. An aneurysm located at the distal middle cerebral artery is not common.

Case presentation: We encountered a 62-year-old Asian man with duplicated middle cerebral artery associated with aneurysms at the M1/M2 junction of the duplicated middle cerebral artery and top of the internal carotid artery.

Conclusions: In cases of duplicated middle cerebral artery, association with a distal aneurysm on the duplicated middle cerebral artery is rare. However, the aneurysm may be formed on the thicker middle cerebral artery due to hemodynamic stress.

Keywords: Anomaly, Duplicated middle cerebral artery, Aneurysm, Distal, M1/M2

Background

Some anomalies of the middle cerebral artery (MCA) have been reported on autopsy and radiological examinations [1]. Among them, a duplicated MCA (DMCA), accessory MCA, and fenestration are well known [1–3]. DMCA arises from the internal carotid artery (ICA) and supplies blood to the MCA territory. The incidence of DMCA has been reported to be 0.7–2.9% on autopsy and 0.24–1.5% on angiography [4].

There have been some reports describing cases of an anomalous MCA associated with an aneurysm [3, 5, 6]. Approximately 35 cases of DMCA associated with aneurysms have been reported in the literature [6–11]. Most of the cases had an aneurysm at the origin of the DMCA, and an aneurysm located at the distal DMCA was not common. To the best of our knowledge, only five cases of DMCA associated with distal MCA aneurysms have been reported

[12–15]. Although the incidence is low, we should be aware of the association of DMCA and aneurysm.

We encountered a patient with DMCA and an aneurysm at the M1/M2 junction of the DMCA. This patient also had a small aneurysm at the top of the ipsilateral ICA. In this case report, we describe the patient with DMCA, and aneurysms on the distal DMCA and top of the ICA, and discuss the radiological findings and characteristics of the rare vascular anomaly and associated aneurysms.

Case presentation

A 62-year-old Asian man with a medical history of diabetes mellitus and pancreatitis due to alcohol experienced speech disturbance. At the age of 58 years, he was treated with insulin for his diabetes mellitus. After that, his blood sugar level was well controlled by diet therapy. There were no relatives with intracranial aneurysms. His symptom was transient and had completely improved when he presented to our institution. He had no neurological abnormalities when he underwent radiological examinations. Magnetic resonance images showed no abnormality in his brain including hemorrhage or cerebral infarction. Magnetic resonance angiography (MRA) revealed a left DMCA that originated from the ICA distal to

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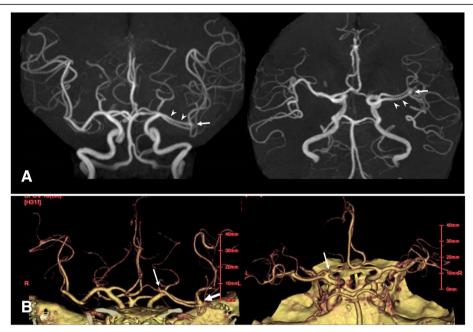


Fig. 1 a Magnetic resonance angiography showing duplicated middle cerebral artery (*arrowheads*) on the left and an aneurysm at the M1/M2 junction of duplicated middle cerebral artery (*arrow*). **b** Three-dimensional computed tomographic angiography also demonstrating the left duplicated middle cerebral artery and aneurysms at the M1/M2 junction of duplicated middle cerebral artery (*arrow*) and internal carotid artery top (*small arrow*)

the anterior choroidal artery (Fig. 1a). An aneurysm at the M1/M2 junction of the DMCA was found (Fig. 1a). Three-dimensional computed tomographic angiography (CTA) also demonstrated the left DMCA associated with aneurysms at the M1/M2 junction and left ICA top (Fig. 1b). In our patient, the aneurysm was located on the DMCA, and not the main trunk of the MCA. The diameter of the DMCA was almost the same as that of the main MCA. The diameters of the DMCA and ICA top aneurysms were both less than 5 mm. Therefore, the aneurysms were not surgically treated, and periodic examinations by magnetic resonance images (MRI) and MRA were planned.

Discussion

An aneurysm at the DMCA origin was initially reported in 1962 [8]. Most reported aneurysms related to DMCA were located at the DMCA origin [6, 7]. Aneurysms at locations other than the DMCA origin have also been reported [4, 6]. DMCA-related aneurysms have been suggested to be formed due to hemodynamic stress. To the best of our knowledge, 35 cases of DMCA origin aneurysm have been reported [6–11]. And almost half of the cases of DMCA aneurysms have multiple aneurysms due to hemodynamic condition [6]. Among them, only five cases of DMCA with distal MCA aneurysms have been reported [8–11].

These previously reported five cases and our case are summarized in Table 1. Among them, four aneurysms were located on the distal portion of the main MCA [12, 13, 15], and one aneurysm was located on the distal portion of the DMCA itself [14]. In our case, the aneurysm was found at the M1/M2 junction of DMCA, and not the main trunk of the MCA. The diameter of the DMCA was almost the same as that of the main MCA. Among the four reported cases with distal aneurysms on the main MCA, the main MCA trunk was thicker than that of the DMCA in three cases. In the other case, the diameters of the main MCA and DMCA were almost the same. On the other hand, in a case of an M1/M2 aneurysm on DMCA, the diameter of the main MCA was almost the same as that of the DMCA. These previous observations and our experience suggest that an aneurysm might be formed at the thick MCA due to hemodynamic factors. All the previously reported five cases were ruptured ones. The risk of rupture of distal DMCA aneurysm seems to be high. If the aneurysm is not small, radical treatment should be considered. For the three ruptured aneurysms, common treatment such as clipping was performed, and two patients showed good recovery. Furthermore, the distal ICA was tortuous in our case. The ICA top might be stressed by blood flow. Therefore, an aneurysm was formed at the

 Table 1
 Summany of cases of duplicated middle cerebral artery accompanied with distal aneurysms

Case	Age	Ethnicity	Size of MCA (mm)	CA (mm)	Location of	Size of aneurysm (mm)	Rupture	Rupture Treatment	Outcome	Associated aneurysm	Reference
	(years), sex		MCA	DMCA	aneurysm					or vessel anomaly	
_	67, F	Asian (Japanese)	0.71	0.37	M1 of main MCA	QN	+	ND	ND	I	[12]
2	99 W	Asian (Japanese)	0.92	0.32	M1/M2 of main MCA	ND	+	Q	Q.	AComA aneurysm	[12]
m	55, F	Q	Both arteries were similar in size	ries were size	Sylvian portion of the main MCA	ΩN	+	Clipping	Death	Contralateral MCA aneurysm (superior trunk)	[13]
4	45, F	Asian (Japanese)	Both arteries were similar in size	ries were size	Distal trunk of DMCA	ND	+	Clipping	GR	Bilateral accessory MCA	[14]
5	34, F	Black	DMCA we than the I	DMCA was smaller than the main MCA	Inferior division of main MCA	ND (> 10 mm in figure)	+	Clipping	GR	I	[15]
Present case	62, M	Asian (Japanese)	Almost the same	ne same	DMCA	3	1	Observation		No change Ipsilateral ICA top aneurysm	

AComA anterior communicating artery, DMCA duplicated middle cerebral artery, F female, GR good recovery, ICA internal carotid artery, M male, MCA middle cerebral artery, ND not described

top of the ICA due to hemodynamic stress. In the six cases of DMCA with distal MCA aneurysms, half of them had another aneurysm.

Conclusions

In cases of DMCA, association with a distal aneurysm on DMCA is rare. However, the aneurysm may be formed on the thicker MCA due to hemodynamic stress.

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Availability of data and materials

Data sharing is not applicable to this article as no datasets were generated or analyzed during the current study.

Authors' contributions

KM drafting of manuscript, critical revision, treatment of patient. AT drafting of manuscript, critical revision, treatment of patient. SS drafting of manuscript, critical revision, treatment of patient. YI drafting of manuscript, critical revision, treatment of patient. YK drafting of manuscript, critical revision, treatment of patient. TN drafting of manuscript, critical revision, treatment of patient. MN study conception and design, drafting of manuscript, critical revision, treatment of patient. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Ethical approval was obtained from the Human Research Ethics Committee of Yokohama Sakae Kyosai Hospital.

Consent for publication

Written informed consent was obtained from the patient for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Competing interests

The authors declare that they have no competing interests.

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