

# Recurrent Extracranial Internal Carotid Artery Vasospasm Diagnosed by Serial Magnetic Resonance Angiography and Superselective Transarterial Injection of a Calcium Channel Blocker

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Recurrent vasospasm of the extracranial internal carotid artery (ICA) is extremely rare, and optimal management is unclear. A 25-year-old woman developed transient dysarthria and left-sided hemiparesis. Initial magnetic resonance (MR) imaging showed spotty acute infarction in the right temporal lobe, and MR angiography revealed right ICA occlusion. ICA occlusion was spontaneously resolved within 6 days of its onset, whereas transient left ICA narrowing was evident at 12 days. Because recurrent occlusion of the right ICA occurred at 14 days when the contralateral ICA was still narrowed, we attempted a local intra-arterial injection of a calcium channel blocker based on the diagnosis of recurrent extracranial ICA vasospasm. The local injection of 1 mg of nicardipine partially dilated the affected ICA, which confirmed the diagnosis of vasospasm. After the introduction of oral medication with benidipine hydrochloride, bilateral ICA vasospasm was completely resolved 23 days after its onset, as shown by MR angiography. In conclusion, we recommend intensive radiologic follow-up at the acute stage and therapeutic catheter angiography when the bilateral lesion is evident because bilateral occlusion of the ICA could lead to a catastrophic condition. **Key Words:** Recurrent extracranial internal carotid artery vasospasm—magnetic resonance angiography—interventional neuroradiology—calcium channel blocker.

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## Introduction

Vasospasm of the extracranial internal carotid artery (ICA) has generally been attributed to mechanical stress such as catheter examination and trauma, whereas idiopathic recurrent vasospasm of the extracranial ICA is extremely rare, with only 8 cases being reported to date.<sup>1-7</sup> Here, we report a case of bilateral recurrent extracranial ICA vasospasm that underwent serial magnetic resonance (MR) angiography and the interventional transarterial injection of a calcium channel blocker to the affected vessel.

## Case Report

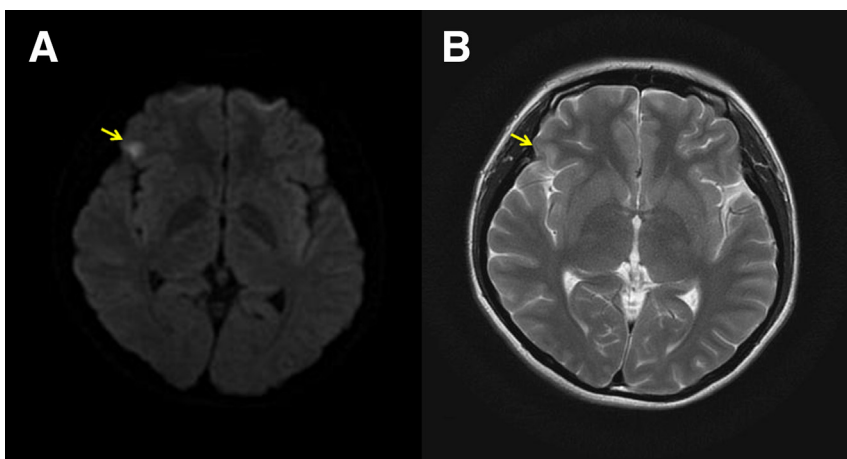
A 25-year-old woman developed transient dysarthria and left-sided hemiparesis. Diffusion-weighted MR imaging revealed a spotty high intensity area in the right frontal operculum (Fig 1, A), and MR angiography demonstrated right ICA occlusion (Fig 2, A). She was subsequently transferred to our hospital with the initial diagnosis of right ICA dissection. Although her neurologic status was unremarkable on admission, she had a headache. She did not have any risk factors for cardiovascular disease, except for migraine-like headaches. She had a habit of active dancing but had no history of trauma to the head or neck.

Laboratory data, x-rays of the chest, and screening for thrombophilia and vasculitis found no apparent abnormalities. *N*-isopropyl-*p*-[<sup>123</sup>I]iodoamphetamine single-photon emission computed tomography (CT) demonstrated no hemodynamic compromise (data not shown); therefore, she was treated conservatively with the initial diagnosis of idiopathic ICA dissection. Occlusion of the right ICA was relieved 6 days after its onset, as shown by MR angiography (Fig 2, B). Repeated MR angiography 12 days after the onset demonstrated no steno-occlusive changes in the right ICA, but segmental narrowing at the contralateral extracranial ICA at the left C1-2 junction (Fig 2, C, arrow). Three-dimensional CT angiography revealed occlusion of the right ICA again 2 days later when the contralateral ICA was still narrowed (Fig 2, D). In light of the dynamic courses of bilateral extracranial ICA stenosis, we suspected recurrent extracranial ICA vasospasm, and attempted therapeutic catheter angiography 14 days after the onset. Bilateral carotid angiography revealed that the left ICA narrowing was almost relieved and cross flow from the left ICA supplied the vascular territory of the right ICA (Fig 2, G). Because the right ICA was still occluded at the cervical portion (Fig 2, H), we attempted interventional therapy to the affected vessel. Although a local intra-arterial injection of 15 mg of fasudil hydrochloride (15 mg) was not

effective, an intra-arterial injection of 1 mg of nicardipine partially dilated the vasoconstricted right ICA, suggesting the efficacy of the calcium channel blocker to this intrinsic pathology (Fig 2, I, arrow). We then administered oral medication of benidipine hydrochloride (4 mg/day) in addition to low-dose aspirin (100 mg/day). Right ICA vasospasm was ameliorated 2 days later (Fig 2, E), and the patient completely recovered at 23 days (Fig 2, F). She did not exhibit neurologic deterioration during the follow-up period.

## Discussion

We demonstrated the temporal profile of recurrent extracranial ICA vasospasm with bilateral lesion by neuroimaging, such as MR angiography, and revealed a dynamic course during a short period. Recurrent extracranial ICA vasospasm is characterized by reversible steno-occlusive changes in the ICA 2-4 cm distal to the common carotid artery bifurcation, and has been reported in 9 cases including the present case (Table 1). All 9 cases were relatively younger adults (24-48 years),<sup>1-7</sup> and repeatedly exhibited temporary segmental vasospasm during a short period ranging from 3 days to 42 months. Eight of the 9 cases had bilateral ICA vasospasm including our case.<sup>1-7</sup> None of the 9 cases showed involvement of the external carotid artery. Our case also showed dynamic changes in the bilateral extracranial ICA during the 20 days from admission, although the site of vasospasm of the left ICA appeared to be more distal than that of the typical location. Eight of the 9 cases developed cerebral infarction.<sup>1-7</sup> Our patient also developed spotty cerebral infarction in the right frontal operculum. Although we failed to detect apparent hemodynamic compromise in the initial flow study in our case, development of cerebral infarction could be because of the hemodynamic ischemia at the acute stage after vasospastic occlusion of the extracranial ICA, in light of the absence of any atherosclerotic background in



**Figure 1.** Diffusion-weighted imaging (A) and T2-weighted imaging (B) of initial magnetic resonance study demonstrating spotty acute infarction in the right frontal operculum (arrows).

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