A Case Report of Pontine Infarction as an Initial Manifestation of Systemic Lupus Erythematosus: Diagnostic Clues from MRI and Digital Subtraction Angiography

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Brainstem infarction due to vertebrobasilar insufficiency is a rare initial presentation of systemic lupus erythematosus (SLE) patients and small-vessel dissection as the direct cause of infarction has not been reported. We report the case of a 20-year-old female with acute infarction on the right side of the pons due to a small artery (pontine perforator) dissection, identified on digital subtraction angiography and high-resolution vessel wall MRI (vwMRI). She was diagnosed with SLE based on the presence of neurologic disorders and relevant laboratory findings. The pontine perforator-dissecting aneurysm had occluded and the right distal vertebral artery had resolved on subsequent vwMRI. She had a modified Rankin Scale (mRS) score of 1 at discharge with mild symptom improvement, and exhibited no further aggravation of symptoms at 3 or 12 months, maintaining an mRS score of 1.

Index terms Stroke; Systemic Lupus Erythematosus; Dissecting Aneurysm; Magnetic Resonance Imaging; Digital Subtraction Angiography
INTRODUCTION

Systemic lupus erythematosus (SLE) is an autoimmune disorder characterized by microvascular inflammation within multiple organ systems secondary to the production of autoantibodies. Stroke represents one of the most severe complications of SLE with an incidence rate of 3%–20% (1) and results in increased morbidity and mortality (2). SLE is associated with a twofold higher risk of ischemic stroke compared to the general population and a > 10-fold higher risk in patients < 40 years old (3). There are several potential mechanisms for SLE-related ischemic stroke, including diffuse cerebral small vessel disease, accelerated atherosclerosis, antiphospholipid syndrome, thrombosis, cerebral vasculitis, dissection, and infection (2, 4).

Ischemic stroke is very rare as the first manifestation of the disease and can often be misdiagnosed. A few cases have been reported with an acute posterior circulation infarction as the initial manifestation of SLE. These cases showed stenosis, occlusion, or normal appearance in the large vessel adjacent to the infarcted region on digital subtraction angiography (DSA), CT, and MRI (5). No studies have reported small vessel dissection as the direct cause of infarction. Here, we report the first case of pontine perforator dissection (identified on DSA and vessel wall MRI [vwMRI]) resulting in acute brainstem infarction as the initial manifestation of SLE.

CASE REPORT

A 20-year-old female presented to the emergency department complaining of headache, dizziness, and blurred vision. She had no history of stroke or autoimmune disease and the family history was unremarkable. There was no evidence of skin lesions, musculoskeletal abnormalities, lymphadenopathy, or splenomegaly. On neurologic examination, there was left-side hypesthesia (admission National Institutes of Health Stroke Scale score was 1). Complete blood count, electrolytes, thyroid, and kidney function tests were all normal; however, C-reactive protein (7.7 mg/L) and erythrocyte sedimentation rate (22 mm/hr) were elevated. Rheumatoid factor was negative and there were no renal abnormalities. However, the patient showed fever (> 38.3°C), thrombocytopenia (34 × 10^9/L), low complement C3 (78.6 mg/dL), positive antinuclear antibodies (1:320) and anti-dsDNA antibodies (1:320) on examination several days later and was diagnosed as SLE (Total score: 15) (6).

The initial diffusion-weighted imaging (DWI) MRI depicted a subtle high signal intensity area in the right pons, suggesting acute infarction and there was old lacunar infarction in right upper pons (Fig. 1A). Cerebral DSA revealed a vascular structure with a fusiform appearance on the right side of the basilar artery, which showed irregular margins and sluggish flow. Time-of-flight (TOF) MR angiography (MRA) showed two hemispherical aneurysms in nonbranching site of right cavernous internal carotid artery and distal vertebral artery (Fig. 1B). At day one follow-up, the DWI and vwMRI revealed the increased extent of infarction in the pons and the fusiform dilated vascular structure originated from the basilar artery and intramural hematoma with wall enhancement and susceptibility artifact (i.e. pontine perforator dissecting aneurysm) (Fig. 1C, D). There was segmental concentric wall enhancement (i.e. vasculitis) distal to the dissecting aneurysm of right pontine perforator. Bilateral anterior ce-
rebral artery, middle cerebral artery, and external carotid artery were intact on TOF MRA. At day nine follow-up, the vwMRI revealed occlusion of dissecting aneurysm in right pontine perforator by an intramural thrombus as a sequela of the dissection (Fig. 1E). The patient

![Fig. 1. Pontine Infraction as an initial manifestation of systemic lupus erythematosus in a 20-year-old female.](image)

A. Initial MRI. DWI (left and middle) and B0 image of DWI sequence (right) depict an acute infarction (black arrows) and an old lacunar infarction (white arrow) in the right pons.

B. Digital subtraction angiography images (left and middle) reveal the dilated vascular structure (black arrows) on the right side of the basilar artery. Time-of-flight MR angiography (right) reveals two hemispherical aneurysms in the right cavernous internal carotid artery and distal vertebral artery (white arrows).

C. DWI and vwMRI at day 1 follow-up. DWI (left end) reveals the increased extent of acute infarction in the right pons. Proton density vwMRI images (right series) confirm the dilated vascular structure’s (black arrows) origin from the basilar artery (white arrows), that is, the pontine perforator.

D. Pre and post-contrast MRI images at day 1 follow-up reveal intramural hematoma (double arrow) and wall enhancement in the dissecting aneurysm (arrow) (left two). In the dissecting aneurysm of the right distal vertebral artery, there is a corresponding intimal flap (arrow) on the pre-contrast T1 vwMRI and wall enhancement (double arrow) on the contrast-enhanced T1 vwMRI (right two).

DWI = diffusion-weighted imaging, vwMRI = vessel wall MRI
was diagnosed with SLE and dissecting aneurysm in the right pontine perforator could be explained by it. The mural wall thrombus in the pontine perforator likely led to the right pontine infarction. The patient was started on an anti-platelet agent, steroid pulse therapy, and hydroxychloroquine 400 mg qd. She was given azathioprine 50 mg qd to treat the underlying autoimmune condition. She had a modified Rankin Scale (mRS) score of 1 at discharge with mild symptom improvement, and she showed no further aggravation of symptoms at 3 months or 12 months, maintaining an mRS score of 1.

Ethics approval and consent to participate. In accordance with the Declaration of Helsinki, informed consent was obtained from the patient.

DISCUSSION

Although it is not completely understood, SLE-related ischemic stroke is known to be caused by various mechanisms including cardioembolism, large artery stenosis (of either non-atherosclerotic or premature atherosclerotic etiology), arterial dissection, hypercoagulable states, and cerebral vasculitis (2, 6). Large vessel occlusion due to premature atherosclerosis is unusual in the early stages of SLE, when cardioembolism, hypercoagulable states, and arterial dissection are more often the etiologies (7). Some reports hypothesize that the most significant cause of cerebral ischemia is the narrowing or occlusion of small vessels. Antiphospholipid antibodies and premature atherosclerosis all play roles in these processes along with cerebral vasculitis (4, 7). In our patient, the acute infarction was due to a pontine perforator...
dissection that might have been caused by underlying vasculitis. Patients with SLE and other connective tissue diseases have a higher risk of vascular dissection than the general population (8). Multiple factors are involved in the activation of autoimmune inflammation of the arterial walls such as immune complex formation and deposition, intimal damage, and medial extracellular matrix damage (9). According to a case report and literature review of 11 patients who presented with ischemic stroke as the initial manifestation of SLE, most patients (7/11), had ischemic infarcts in the vertebrobasilar artery and the pons was the most frequently affected region. The pathophysiology of infarction could be suspected in only three cases (two dissections and one with an inflammatory lesion) but eventually in none of them it was clearly identified. However, most of the cases were reported before 1999 i.e. image modalities to identify vasculitis or dissection were very limited. All patients had at least one autoantibody, which probably caused an underlying inflammatory or hypercoagulable state. Arterial occlusion or stenosis was identified in only six of the 11 cases (5).

We suggest vwMRI as diagnostic investigation tools to reveal the direct cause of ischemic infarcts, such as small vessel dissection or underlying vasculitis in the early phase of SLE and DSA can cross-check the lesion depicting an intimal flap, focal vessel wall irregularity, pre-aneurysmal narrowing, and fusiform dilatation. The predominant vwMRI characteristics of dissection are eccentric or combined eccentric/concentric shape, T1 high signal intensity with susceptibility artifact due to intramural hematoma, a double lumen caused by intimal flap, and aneurysmal dilatation with or without wall enhancement. Differential diagnosis of intramural hematoma is intraplaque hemorrhage, vasculitis and Moyamoya disease. Intraplaque hemorrhage usually shows irregular eccentric T1 high signal and no change. The main differentiation point of intramural hematoma are change by time and smooth luminal margin, longer segment than intraplaque hemorrhage. Vasculitis usually shows concentric strong enhancement in the active stage that resolves in the chronic stage but does not show intimal flap or aneurysmal dilatation on vwMRI (10).

In a young patient presenting with acute stroke of unknown origin, it is necessary to investigate for rarer etiologies such as SLE and other autoimmune diseases. High-resolution vwMRI and DSA might provide important clues.

Author Contributions
Conceptualization, B.J.S.; data curation, all authors; investigation, Y.Y.; methodology, C.M.S.; project administration, B.J.S.; resources, all authors; supervision, B.J.S.; visualization, B.J.S.; writing—original draft, C.M.S.; and writing—review & editing, B.J.S.

Conflicts of Interest
The authors have no potential conflicts of interest to disclose.

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전신성 홍반성 루푸스의 초기 증상으로 나타난 교뇌경색의 증례 보고: 자기공명영상 및 디지털감산 혈관조영술에서의 진단 단서

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척추기저동맥의 혈류공급감소로 인한 뇌간 경색은 매우 드문 전신성 루푸스 환자의 초기 증상으로, 경색의 직접적인 원인으로 매우 작은 기저동맥 분지인 교뇌공급혈관의 박리성 동맥류는 보고된 사례가 없다. 이에 저자들은 디지털감산 혈관조영술과 고해상도 혈관벽 자기공명영상을 이용하여 작은 교뇌공급혈관의 박리성 동맥류의 진단과 추적관찰 중 치유된 20세 여성의 사례를 보고하고자 한다. 전신성 루푸스의 진단은 신경학적장애인의 유무와 혈액학적 검사 결과를 바탕으로 하였다. 추적 고해상도 혈관벽 자기공명영상에서 환자의 교뇌천공지의 박리성동맥류는 폐색되어 있었고 우측 척추기저동맥의 박리성동맥류는 치유되어 보이지 않았다. 환자는 퇴원시 수정랭킨척도 점수가 1점으로 증상 개선을 보였으며 3개월과 12개월 추적관찰에서도 증상이 약화되지 않고 1점을 유지하였다.

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