CASE REPORT

Oculomotor Nerve Palsy in a Patient with a Ruptured Middle Cerebral Artery Aneurysm

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Abstract : We describe a case of acute oculomotor nerve palsy caused by a ruptured middle cerebral artery (MCA) aneurysm. A 59-year-old female presenting with headache and nausea was admitted to our hospital. Her consciousness was alert, and had no other neurological deficit without left oculomotor nerve palsy. A computed tomography (CT) showed SAH extending from left sylvian cistern to basal cistern. CT angiography revealed a left MCA aneurysm which protruded toward internal carotid artery. The patient was successfully treated with surgical clipping. The oculomotor nerve palsy resolved immediately after the surgery. Perioperative radiological evaluation revealed that there were no evidence of midbrain hemorrhage or stroke, vessel anomaly of basilar, posterior cerebral or superior cerebellar artery, vasospasm, and uncal herniation. Furthermore, intraoperative findings revealed that the aneurysm was projected toward the affected carotid cistern and oculomotor nerve. From these findings and time course of oculomotor nerve palsy, it is suggested that the jet flow of bleeding from the ruptured MCA aneurysm caused oculomotor nerve palsy in the patient. J. Med. Invest. 64 : 165-167, February, 2017

Keywords : oculomotor nerve palsy, middle cerebral artery aneurysm, subarachnoid hemorrhage

INTRODUCTION

Patients with acute subarachnoid hemorrhage (SAH) occasionally present with isolated oculomotor nerve palsy. The majority of those patients have ruptured aneurysms of the internal carotid-posterior communicating artery junction or basilar artery-superior cerebellar artery junction, reflecting the anatomical proximity of the affected vessels and the oculomotor nerve (1-3). Conversely, solitary oculomotor nerve palsy due to rupture of an aneurysm that is far from the nerve is extremely rare. Here we describe a case of oculomotor nerve palsy caused by rupture of a middle cerebral artery (MCA) bifurcation aneurysm. The mechanism of this fairly rare condition in association with SAH is discussed.

CASE REPORT

A 59-year-old female experienced sudden headache and nausea and was referred to our institution 14 hours after the onset. She had a history of hypertension and malignant rheumatoid arthritis that had been adequately controlled by medication. She had no history of diabetes mellitus, heart disease, stroke, or cancer. She had no family history of SAH, and no habit of smoking or alcohol. On arrival, she was alert and well oriented. She had no neurological deficit except for left external/internal oculomotor nerve palsy, including ptosis, pupillary dilatation, and loss of direct/indirect light reflexes. Computed tomography (CT) scans showed a diffuse thick SAH, especially in the left sylvian fissure (Figure 1). Three-dimensional CT angiography demonstrated a small, medially projecting, left MCA bifurcation aneurysm with a maximum diameter of 3.5 mm (Figure 2). No other aneurysms were identified. There were no vessel anomalies including the tortuosity or dolichoectasia of the posterior cerebral artery (PCA), superior cerebellar artery (SCA), posterior communicating artery or basilar artery.

The patient underwent an emergent craniotomy and repair of the aneurysm. During surgery, the MCA aneurysm was found to be ruptured and was successfully clipped. A small protrusion at the tip of the aneurysm dome toward the carotid cistern was evident and thought to be the bleeding point. Subsequently, subarachnoid clots in the sylvian and basal cisterns were adequately removed, and careful inspection of the carotid and basilar cisterns around the left oculomotor nerve indicated no carotid or basilar artery
nerve palsy (7). These cases showed prompt recovery of oculomotor nerve palsy due to direct aneurysm compression or stroke recoveries very slowly after repair of the aneurysm (13). Therefore, the clinical course of the present case also supports the idea that oculomotor nerve palsy may be caused by the jet flow of bleeding. Finally, we intraoperatively confirmed that the bleeding point of the aneurysm was directed toward the oculomotor nerve. Accordingly, the oculomotor nerve palsy is suggested to be caused by the jet flow of bleeding from ruptured MCA aneurysm which is directed toward the oculomotor nerve.

Although the small number of reported cases precludes statistical analysis, accumulated clinical data suggest that the characteristic features of oculomotor nerve palsy that is associated with a ruptured aneurysm in a distant location include: 1) no evidence of midbrain hemorrhage or infarction, remarkable elevation of ICP, vasospasm, hematoma compression, anomaly of basilar artery, PCA, SCA or posterior communicating artery, and uncal herniation; 2) aneurysm projection toward the affected carotid cistern and oculomotor nerve; and 3) prompt recovery of the symptom within a few days.

Our case serves as a reminder that oculomotor nerve palsy can be a false localizing sign of a ruptured aneurysm in SAH patients. This unusual occurrence suggests an alternative explanation of oculomotor nerve palsy that is not directly related to aneurysmal compression.

CONFLICTS OF INTEREST

No work resembling the enclosed article has been published for publication elsewhere. We certify that each author has made a substantial contribution so as to qualify for authorship as detailed at the end of the manuscript. We have disclosed all financial support for our work and other potential conflicts of interest.

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