Fatal Epistaxis Caused by Rupture of an Intratumoral Aneurysm Enclosed by a Large Prolactinoma

—Case Report—

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Abstract

A 72-year-old female presented with episodes of epistaxis. Neuroimaging demonstrated a large prolactinoma totally enclosing a large intracavernous aneurysm of the internal carotid artery. Adjacent bony structures were eroded and destroyed by tumor invasion and extension. Rupture of the intratumoral aneurysm caused fatal epistaxis rather than subarachnoid hemorrhage before surgery. Intratumoral aneurysm is rare and epistaxis caused by rupture of it is extremely rare. Lack of bony protection apparently have contributed to the aneurysmal growth and rupture.

Key words: epistaxis, intracavernous aneurysm, intratumoral aneurysm, pituitary tumor, prolactinoma

Introduction

Epistaxis caused by rupture of intracranial aneurysm usually originates from aneurysms located in the cavernous portion of the internal carotid artery (ICA), and in most cases, was precipitated by head injury.2,4,6,11,12,14) The second most common cause was developmental aneurysm.13) Mycotic aneurysms secondary to the cavernous sinus infection have also caused epistaxis.9) Epistaxis from intratumoral aneurysm in the pituitary tumor is very rare. Association of intracranial aneurysm and pituitary tumor is also well known, but an aneurysm totally enclosed by the pituitary tumor is not common. We describe a very unusual case that rupture of an intratumoral aneurysm in the pituitary tumor causing epistaxis which resulted in death.

Case Report

A 72-year-old female was transferred by ambulance to the Department of Otolaryngology because of sudden epistaxis with headache and nausea on May 18, 1996. Hemostasis was completed with Bellock's tamponade. She was referred to the Department of Neurosurgery. Three months before, she had had two episodes of epistaxis which had subsided without special treatment.

Neurological examination found she was conscious without nuchal rigidity. Her pupils were 3 mm in diameter on both sides. Only light was distinguishable in her right eye. She had not previously realized the presence of decreased visual acuity of the right eye. No visual defect of the left eye was observed. Eye movement was normal, with no nystagmus. No motor or sensory disturbance was found. Her blood pressure was 200/110 mmHg. She had no history of head injury, cerebrovascular disease, diplopia, facial pain, aspirin intake, or anticoagulant use.

Computed tomography (CT) showed a giant mass in the sella turcica extending into the sphenoid and ethmoid sinuses, laterally into the cavernous sinus, and slightly into the suprasellar cistern. Another large round mass of higher density was totally enclosed by this giant mass (Fig. 1). Three-dimensional CT angiography revealed a large aneurysm located in the sella turcica, and destruction of the sphenoid sinus bone (Fig. 2). Cerebral angiography showed a large cavernous aneurysm of the right ICA (Fig. 3). There were no feeding arteries or tumor stain. The balloon occlusion perfomed during cerebral angiography caused left hemiparesis. Her serum prolactin level was extremely high at 6800 ng/ml
The diagnosis of the tumor was prolactinoma.

Trapping of the ICA-cavernous aneurysm with extracranial-intracranial bypass was scheduled. However, sudden massive arterial nasal bleeding due to aneurysmal rupture occurred while she was waiting for surgery. Hemostasis of the massive epistaxis filling in the nostril and the oral cavity was difficult to achieve despite continuous compression of the right carotid artery. Consequently, she lost over 1000 ml blood and died in spite of temporary resuscitation on June 6, 1996. Autopsy was refused.

**Discussion**

Our patient had no history of head injury, and the ICA-cavernous aneurysm and large prolactinoma was incidentally diagnosed after she complained of headache. This aneurysm seemed to be developmental and the several episodes of minor bleeding before admission were probably from the aneurysm rather than the pituitary tumor. Rupture of the intracranial aneurysm possibly caused epistaxis.

The incidence of intracranial aneurysm associated with pituitary tumor is higher than that in the general population or associated with other brain tumors. However, such occurrence is only a chance factor and the risk is no greater than that among the general population.

Most aneurysms associated with pituitary tumor were from adjacent arteries such as the ICA or anterior communicating artery and were located close to or touching the pituitary tumor. Aneurysm totally enclosed by the tumor tissue as in our case is very rare.

Several hypotheses to explain the high frequency of aneurysm associated with pituitary tumor have been proposed, including direct invasion of brain tumor into the cerebral artery, mechanical compression or stretching of the cerebral artery by the brain tumor, rheological stress induced by increased blood supply to the brain tumor, and endocrinological effects on membranous collagen such as growth.
hormone-secreting tumor. However, the exact mechanism has not yet been identified.

The ICA-cavernous aneurysm surrounded by hard tissue such as the dura and the bony structures, so is usually unlikely to rupture. In our case, the bony structures were widely eroded and destroyed by the tumor extending and invading inferiorly and anteriorly into the sphenoid and ethmoid sinuses. Therefore, the wall beneath the aneurysm had become less resistant. The aneurysm had apparently grown in the least resistant direction and had subsequently become large enough to rupture. The tumor might have invaded the aneurysmal wall and caused weakening, so the rupture site was probably the most weakened aneurysmal wall. The resistance to intraneurysmal pressure was also decreased by lack of bony protection. Therefore, she did not suffer subarachnoid hemorrhage although the aneurysm had bled only into the nostril.

Various surgical strategies for the treatment of ICA-cavernous aneurysm are available, including ligation, balloon occlusion, coil embolization of the ICA, trapping of the ICA-cavernous portion, and neck clipping of the aneurysm. Simultaneous neck clipping and tumor resection is often successfully performed in cases of ICA aneurysm associated with pituitary tumor, but not in cases of intratumoral aneurysm. ICA trapping with extracranial-intracranial bypass was scheduled for our patient because of the neurological deficits provided by the balloon occlusion test. Unfortunately she died before surgery. Emergent surgery is recommended in such cases.

References

5) Kagawa R, Shima T, Matumura S, Okada Y, Nishida M, Yamada T, Okita S: [A case of a large prolactinoma complicated with the intratumoral large aneurysm — Findings by magnetic resonance imaging]. Hiroshima Igaku 43: 382-385, 1991 (Jpn)

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