Intraventricular cavernous hemangiomas are uncommon. Among them, those occurred at the foramen of Monro in the third ventricle may be of particular interest because of its rarity, development of hydrocephalus, being differentiated from other brain lesions. We present a rare case of intraventricular cavernous hemangioma at foramen of Monro which was resected through microsurgery and also review the relevant literatures.

Key Words: Cavernous hemangioma · Intraventricular · Third ventricle · Foramen of Monro · Hydrocephalus · Arteriovenous malformations.

INTRODUCTION

Cavernous hemangioma constitutes 5-10% of vascular malformations occurring in the entire central nervous system (CNS). Intraventricular location constituting 2.5-10.8% of cerebral cavernous hemangioma is uncommon because most of them are usually found in the parenchyma of CNS. So far, the intraventricular cavernous hemangioma has been reported to be less than 90 cases to our best knowledge. Those located in the third ventricle account for approximately half of all intraventricular cavernous hemangiomas. In particular, cavernoma at the foramen of Monro which have been reported to be about 13 cases are very rare and may be difficult to differentiate from other brain tumors. Herein, we present an additional case of cavernous hemangioma located in the foramen of Monro of the third ventricle and review the relevant literatures.

CASE REPORT

A 30-year-old female patient who was pregnant presented with intermittent non-throbbing headache for a month before admission. She has often suffered from short-term memory deficits gradually. She complained the projectile vomiting and fever without consciousness disturbance which were observed shortly after normal vaginal delivery. She was referred to department of neurology with the impression of meningitis. On physical examinations, definite neurological deficits were not found. Computed tomogram (CT) showed enlargement of both lateral ventricles which may be caused by 3 cm sized multi-lobular calcified mass with slightly peripheral rim enhancement at the foramen of Monro in the third ventricle. The lesion seemed to involve the left thalamus parenchyme partially. Magnetic resonance image (MRI) revealed a relatively well-delineated mass at the foramen of Monro with heterogeneous signal intensity on both T1 and T2 weighted images. The surrounding rim of low signal intensity around mass was not delineated definitely on T2-weighted image. A peri-lesional edema in the left thalamus was observed on the FLAIR image and only minimal peripheral rim enhancement around mass was found on enhanced T1-weighted image (Fig. 1). Conventional cerebral angiograms revealed no vascular abnormalities. Anterior interhemispheric trans-callosal approach was performed to remove the intraventricular lesion. Operative findings showed that the mass with xanthochromic and lobular surface obstructing the foramen of Monro was located in the third ventricle. The left posterior margin of the mass was close contact to the thalamus while resection. The mass was removed totally without significant bleeding (Fig. 2). The histological examination revealed large hyalinized vessels filled with organizing thrombus. Immunostaining for smooth muscle actin (SMA) was positive in the vessel wall (Fig. 3). Those findings were compatible with characteristics of cavernous hemangioma. After operation, the patient's symptoms including vomiting and headache
were recovered gradually (Fig. 4).

**DISCUSSION**

Cavernous hemangiomas known as cavernous malformations, cavernous angiomas or cavernomas are a kind of vascular malformation which is angiographically occult unlike arteriovenous malformations. These benign lesions are composed of sinusoidal vascular channels which are lined with a single layer endothelium predisposing to hemorrhage and have perilesional surrounding gliosis. Hemorrhages at different stages due to repeated bleeding, occlusion and thrombosis in the vascular channels are found within the lesion. Organization of the hematoma results in chronic granulation and scar formation. Progressive pseudotumorous evolutions are induced by repeated bleeding which occur in the immediate vicinity of cavernous hemangioma, leading to hemosiderin deposits and gliosis. Most of them occur in the intra-axial and supratentorial regions and they are typically located in the subcortical deep white matters, basal ganglia, brain stem and cerebellum. However, they have rarely seen in the intraventricular or extra-axial locations. Kivelev et al. summarized intraventricular cavernomas of 89 cases including their own cases and reported that 34 cases (38%) of them were located in the third ventricle. Third ventricular cavernomas can be classified into suprachiasmatic, foramen of Monro, lateral wall, and floor regions depending on the location. Those located at the foramen of Monro may have particular clinical interests because of hydrocephalus due to foraminal obstruction and have been reported to be only 13 cases so far. It is notable that our case can be additional one as foramen of Monro cavernomas of third ventricle. Clinical symptoms of intraventricular cavernomas may be elicited by mass and increased intracranial hypertension due to acute bleeding or obstruction of cerebrospinal fluid pathway. In our case, headache and vomiting resulted from increased intracranial pressure due to obstructive hydrocephalus induced by cavernoma located at the foramen of Monro. Fourteen intraventricular cavernomas located at the foramen of Monro including our case are summarized in the Table 1. There were 5 males and 9 females whose age distribution is ranged from 11 years to 64 years. The clinical findings were

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**Fig. 1.** Computed tomogram (CT) shows enlargement of both lateral ventricles which is caused by 3 cm sized multi-lobular calcified mass with slightly peripheral rim enhancement. It is located at the foramen of Monro in the third ventricle and seems to be close to the left thalamus parenchyme partially (A). Magnetic resonance image (MRI) reveals a relatively well-delineated third ventricular mass with heterogeneous signal intensities in the center of lesion on T1-weighted image (B). Typical peripheral hemosiderin rim of low signal intensity is not delineated on T2-weighted image (C). Minimal peripheral enhancement at the right posterior portion of the mass is seen on enhanced T1-weighted image (D). There is a minimal perilesional edema in the left thalamus on the FLAIR image (E). The mass around foramen of Monro is seen on enhanced sagittal T1-weighted image (F).

**Fig. 2.** Operative finding shows that the mass with xanthochromic and lobular surface obstructing the foramen of Monro is located in the third ventricle. The left posterior margin of the mass is close to the thalamus.

**Fig. 3.** The histological examinations reveal a large hyalinized vessels filled with organizing thrombus (A). Immunostaining for smooth muscle actin (SMA) is positive in the vessel wall (B).
| Reference        | Age (yrs), sex | Symptoms                                                                 | Hydrocephalus | Size (mm) | CT findings                                                                 | MRI findings                                                                 | Angiogram       | Surgical approach |
|------------------|----------------|--------------------------------------------------------------------------|---------------|-----------|--------------------------------------------------------------------------------|--------------------------------------------------------------------------------|-----------------|-------------------|
| Britt, 1980      | 11, F          | Nausea, vomit                                                           | Present       |           | Iso-high density minimal enhance                                               |                                                                                | Transcortical    |                   |
| Pozzati, 1981    | 31, F          | Nausea, vomit                                                           | Present       |           | Iso density                                                                   |                                                                                | Transcortical    |                   |
| Harbaugh, 1984   | 44, F          | Headache, nausea, meningismus, hemorrhage                              | Present       |           | Calcified, iso density no enhance                                              |                                                                                | Negative         | Transcallosal     |
| Katayama, 1994   | 50, F          | Headache, nausea, meningismus, hemorrhage                              | Present       |           |                                                                                  |                                                                                | Transcallosal    |                   |
| Kaim, 1997       | 64, M          | Headache, tinnitus, ataxia, memory disturbance                         | Present       | 15        | Slight enhance                                                                | Heterogenous                                                                 | Enlarged vein   | Transcallosal     |
| Crivelli, 2002   | 38, M          | Short term memory loss, headache, vomit, nausea, ataxia, disorientation | Present       | 30        | Partially calcified slight enhance                                             | Heterogenous                                                                 | Transcortical    |                   |
| Suess, 2002      | 36, F          | Short term memory loss                                                  | Present       | 15        | Calcium, slight enhance                                                        | Heterogenous                                                                 | Transcallosal    |                   |
| Chen, 2006       | 51, F          | Headache, ataxia, vomit, conscious change, disorientation              | Present       | 15        | Calcium, slight enhance                                                        | Heterogenous                                                                 | Transcortical    |                   |
| Longatti, 2006   | 35, M          | Headache, vomit                                                         | Present       | 12        | High density                                                                  | Heterogenous                                                                 | Endoscopic       |                   |
| Sato, 2006       | 47, F          | Headache                                                                | Present       | 15        | High                                                                          | Iso                                                                          | Strong homogenous| Enlarged vein     |
| Prat, 2008       | 56, M          | Headache, confusion, hemorrhage, familiar multiple cavernomatosis      | Present       | 15        |                                                                                | Low                                                                          | Endoscopic       |                   |
| Kivelev, 2010    | 52, M          | Headache, nausea, vomit                                                | Absent        | 25        |                                                                                |                                | Transcallosal    |                   |
| Present case     | 30, F          | Headache, short term memory loss, vomit                                | Present       | 30        | Multiple calcification                                                        | Heterogenous                                                                 | Transcallosal    |                   |

**FLAIR** : fluid-attenuated inversion recovery, **GE** : gradient echo
symptoms associated with hydrocephalus in most cases, hemorrhages in 3 cases, focal neurological deficits in 5 cases. In all patients, surgical removal was performed completely without significant neurological deficits using the microsurgery or/and endoscopic surgery through the trans-callosal or trans-cortical approaches. Postoperative shunt surgery was not required in most cases. Most of them often showed calcification on computed tomogram (CT) and heterogeneous signal intensities and mild enhancement on magnetic resonance image (MRI). Typical CT findings of cavernous hemangiomas in the third ventricle show moderately hyper-dense lobular lesions with mild contrast enhancement. Unlike arteriovenous malformations, they are angiographically occult because cavernous hemangiomas usually do not have feeding or draining vessels obviously. Therefore, MRI has been useful in diagnosis of clinically silent and angiographically occult cavernomas. Common MRI features include a hyper to iso-intense center correlated to the hemorrhages at different stage on T1-weighted images and a surrounding hemosiderin rim of low intensity caused by calcification or fibrosis on T2-weighted images. Perilesional edema is unusual and contrast enhancement is various. And also, it has been reported that typical perilesional hemosiderotic rim might be thinner or absent and core of lesion could be less heterogeneous.

Therefore, MRI has been useful in diagnosis of clinically silent and angiographically occult cavernomas. Common MRI features include a hyper to iso-intense center correlated to the hemorrhages at different stage on T1-weighted images and a surrounding hemosiderin rim of low intensity caused by calcification or fibrosis on T2-weighted images. Perilesional edema is unusual and contrast enhancement is various. And also, it has been reported that typical perilesional hemosiderotic rim might be thinner or absent and core of lesion could be less heterogeneous.

CONCLUSION

We present a very rare case of intraventricular cavernous hemangioma located at the foramen of Monro that may mimic tumor lesions.

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