Hemorrhagic colloid cyst

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ABSTRACT

Colloid cysts are benign, thin-walled, cystic lesions that arise from the brain’s endodermal embryonic remnants, and they are usually located in the anterior portion of the third ventricle near the foramen of Monro; these cysts contain colloid material. The clinical presentation of these cysts ranges from incidental findings on brain images to sudden death. The clinical presentation of these cysts is largely dependent on the mass effect on the foramen of Monro. Rarely, hemorrhagic cysts can lead to acute obstructive hydrocephalus or sudden death. We herein report 2 cases of hemorrhagic colloid cysts.

Case Report. Case 1. Patient’s information: A 47-year old male presented with progressive morning headaches for a total of 10 days, which was associated with nausea and vomiting. He had no history of trauma or loss of consciousness.

Clinical findings. On examination, he had an intact level of consciousness. Funduscopic examination revealed bilateral papilledema. The patient’s cranial nerves, as well as his motor and sensory examinations, were normal without any signs of neurological deficits.

Diagnostic assessment. The patient’s initial computed tomography (CT) scan showed a large cystic lesion with peripheral high densities causing acute hydrocephalus. Magnetic resonance (MR) images showed a large intraventricular mass at the foramen of Monro measuring 4.3cm×4.0cm×3.2cm, pushing the septum pellucidum to the right and causing acute lateral ventricular dilatation. The mass was largely cystic and had a thick wall and solid components inferiorly, demonstrating hyperintensity on T1-weighted images and multilayered hypointensity on T2-weighted images with remarkable susceptibility artifacts in the wall and in the inferior part, indicating hemosiderin deposition (Figure 1). The solid component in the inferior part of the lesion demonstrated mild diffusion restriction,
which likely represented bleeding at different ages. The wall of the cystic component of the lesion demonstrated thick enhancement, which was irregular in some places; the solid component did not show enhancement. The perfusion scans did not show increased perfusion. Furthermore, the MR arteriogram study showed normal intracranial arteries with no displacement or occlusion.

Therapeutic intervention. The patient then underwent a frontal craniotomy and an interhemispheric, transcallosal transventricular approach to resect the cyst, as well as left frontal external ventricular drain (EVD). The patient recovered and tolerated the surgery. Pathological examination confirmed a hemorrhage within the colloid cyst.

Follow-up and outcomes. At the one month follow up visit, the patient exhibited enhanced cognitive function and complete resolution of the headaches (figure 2).

Case 2. Patient’s information. A 55-year-old male presented with progressive memory disturbance and unsteady gait for 8 months, which became more profound at the time of presentation.

Clinical findings. A higher cognitive function examination revealed impairments in short-term memory. A fundoscopic examination revealed bilateral papilledema. The patient’s cranial nerve, as well as his motor and sensory examinations, were normal except for bilateral extensor plantar reflexes. Cerebellar examination revealed impaired tandem walking.

Diagnostic assessment. The patient’s initial CT scan showed a heterogeneous mass that adhered to the septum pellucidum at the level of the foramen of Monro; this was associated with bilateral ventricular hydrocephalus laterally and a mild midline shift. Magnetic resonance images showed a heterogeneous,
rounded, intraventricular cystic lesion within the dilated left foramen of Monro and with extension into the third ventricle, causing a shift of the septum pellucidum to the right by 9.4 mm. There was also an obstruction of the ipsilateral foramen of Monro, resulting in obstructive hydrocephalus of the lateral ventricles. The lesion was hyperintense on T1-weighted images, intermediate to hypointense on T2-weighted images with peripheral hemorrhagic susceptibility artifacts, and it did not show any signs of diffusion restriction (Figure 3).

**Therapeutic intervention.** The patient subsequently underwent a left frontal craniotomy and transcallosal resection of the colloid cyst. The patient tolerated the surgery and his unsteady gait improved slightly; however, he developed aphasia postoperatively.

**Follow-up and Outcomes.** At the 2-month follow-up visit, the patient was walking independently, his memory showed signs of improvement, and his aphasia demonstrated improvements in speech. A pathological examination of the excised lesion confirmed the presence of a hemorrhage within the colloid cyst (figure 4).

**Discussion.** The reported estimated incidence of third ventricle colloid cysts (TVCC) is approximately one per one million. Colloid cysts account for 0.5%-1% of all intracranial tumors. Generally, colloid cysts are benign and most cases are asymptomatic, with both genders being affected at equal rates. Symptomatic intracystic hemorrhage within these colloid cysts is rarely encountered. A PubMed search was conducted for this literature review, and the following keywords were used: “Hemorrhagic colloid cyst” and “Colloid cyst and

**Figure 2** - A 47-year-old male (Case 2) presented with progressive morning headaches for a total of 10 days, which were associated with nausea and vomiting. He had no history of trauma or loss of consciousness.
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There were only 13 clinically diagnosed cases of these cysts reported in the literature,\textsuperscript{2,3,9-19} and 5 additional cases were reported at the time of autopsy.\textsuperscript{4,5,20-22} Of these 5 cases, only 3 had associated risk factors,\textsuperscript{4,5,11} - 2 cases had arterial hypertension and one case demonstrated bleeding tendencies. Although it is unknown whether these factors contributed to colloid cyst bleeding, arterial hypertension and coagulation abnormalities may have played a role and should be carefully considered in patients with colloid cysts in the future.\textsuperscript{14}

The mechanisms underlying sudden death in patients with colloid cysts are poorly described in the literature, with several authors postulating that hydrocephalus and brain herniation serve as the direct causes of the rapid deterioration that precedes mortality.\textsuperscript{23,24} Moreover, intracystic hemorrhage should be considered as a possible mechanism for sudden deterioration in these patients as well.\textsuperscript{14} Table 1 summarizes the characteristics, clinical presentation, imaging features, pathological confirmation, treatment methods, and outcomes of all clinically reported cases.

Interestingly, the ages of the reported cases ranged from 9-77 years old, with a mean age of 39.67 years; there was only one pediatric case. In all, there were 8 males and 5 females. Most of the clinical cases reportedly presented with signs of raised intracranial pressure due to obstructive hydrocephalus.

Six of the 13 cases reportedly demonstrated rapid deterioration of the patient’s clinical picture following the initial presentation.\textsuperscript{2,3,9-11,19} Of those 6 cases, 4 had poor outcomes,\textsuperscript{2,10,11,19} illustrating that the degree of rapidity with which clinical deterioration occurs

Figure 3 - Radiological images of case 1. A) Axial CT scan showing a heterogeneous mass adherent to the septum pellucidum at the level of the foramen of Monro associated with bilateral ventricular hydrocephalus laterally and a mild midline shift. B) An axial T2-weighted image showing an intermediate to hypointense heterogeneous intraventricular cystic lesion resulting in obstructive hydrocephalus of the lateral ventricles. C and D) Axial and sagittal T1-weighted images showing hyperintensity of the lesion. E) An axial SWI demonstrating multiple peripheral, hemorrhagic susceptibility artifacts. F) An axial postoperative T1-weighted image showing complete excision of the colloid cyst, improvements in hydrocephalus, a small intraventricular hemorrhage, and a small amount of air.
**Table 1** - The characteristics, clinical presentation, imaging features, pathological confirmation, treatment methods, and outcomes of all clinically reported cases.

| Authors | Age/Gender | Comorbidities | Clinical course | Radiological features of CC | Pathology confirmation | Treatment | Outcome/Follow-up period |
|---------|------------|---------------|----------------|-----------------------------|------------------------|-----------|--------------------------|
| (2)     | 35/F       | A known case of a colloid cyst, as well as progressive headaches and memory disturbances, followed by progressive deterioration and 2 seizure episodes. | +               | +                           | Emergency ventriculostomy followed by endoscopic subtotal excision of the cyst. | Poor; after few months post-op, was still fully dependent on others. |
| (3)     | 9/F        | Headache, nausea, and sudden deterioration in mental status. | +               | +                           | Emergency ventriculostomy, followed by craniotomy and cyst excision. | Tolerated the procedure well and was discharged on postoperative day 3 without any neurological complications. |
| (9)     | 48/M       | Headaches and a memory deficit for 2 weeks, followed by sudden deterioration. | +               | NA                          | Stereotactic aspiration with limited evacuation. Followed by ventricular drainage due to herniation. Renewed stereotactic aspiration with endoscopic guidance and subtotal excision of the cyst. | Residual cyst with a slow increase in cyst size, radiologically. |
| (10)    | 29/F       | Mental status deterioration, Glasgow Coma Scale 4 with respiratory insufficiency. | +               | +                           | Emergency ventricular drainage followed by endoscopic removal. | The radiological results were excellent, but there were poor clinical outcomes (vegetative state). |
| (11)    | 47/M       | Hypertension, Acute right-sided hemiparesis and speech impediment, followed by rapid deterioration of consciousness. | +               | +                           | Emergency EVD followed by elective total resection of the lesion via a transcallosal route. | Right-sided hemiparesis (3/5) with impaired proprioception and a mixed-type sensitive and motor dysphasia (1 year post-op). A neuropsychological assessment revealed Bradipsychia and anosognosia, as well as attention, concentration, and memory deficits. |
| (12)    | 48/M       | A 4-day progressive headache followed by the sudden onset of 24-hour anterograde amnesia. (Normal neurological examination without papilloedema.) | +               | +                           | Emergency neuro-endoscopic total excision of the cyst. | The patient's amnesia completely resolved within 24 hours post op, and he was discharged home 2 days later. Returned to full-time employment (2 months post-operatively). |
| (13)    | 28/M       | Known case of a colloid cyst with unilateral hydrocephalus and a right ventriculoperitoneal shunt for 8 years. Presented with severe headache and multiple episodes of vomiting for 1 day. | +               | +                           | Right frontal craniotomy, with total excision of the cyst (an anterior interhemispheric, transcortical–transventricular approach). EVD for 3 days. | At 1-year follow-up, he has no neurological deficits and the headache has resolved. |
| (14)    | 77/M       | Type II diabetes mellitus. On warfarin for atrial fibrillation; INR: 1.2 Unsteady gait, incontinence, and gradually worsening confusion over 3 weeks. The GCS was 14, with mild right-sided weakness; power 4/5. The cranial nerves were intact. | +               | +                           | Right frontal craniotomy with total excision of the cyst. (transcortical approach). Ventriculoperitoneal shunt for delayed hydrocephalus. | Satisfactory with rehabilitation. |

M - male, F - female, EVD - external ventricular drain, ND - no data.
### Table 1 (continued) — The characteristics, clinical presentation, imaging features, pathological confirmation, treatment methods, and outcomes of all clinically reported cases.

| Authors | Age/Gender | Comorbidities | Clinical course | Radiological features of CC | Pathology confirmation | Treatment | Outcome/Follow-up period |
|---------|------------|---------------|----------------|-----------------------------|------------------------|-----------|--------------------------|
| (15)    | 20/F       |               | Progressive headache, three episodes of generalized tonic clonic seizures, gait disturbances, impaired short-term memory, bilateral papilloedema, extensor plantar reflex and (+) Romberg’s sign. | + + | Rightfrontoparietal craniotomy with total excision of the cyst (interhemispheric transcrallosal approach). | Seizure-free and improved memory function in 3 weeks. |
| (16)    | 47/ F      |               | Headache, N/V, papilledema, GCS of 13. | + + | Ventriculostomy followed by craniotomy and total excision of the cyst. | Uneventful recovery. |
| (17)    | 35/M       |               | Progressive increasing headaches over months; nausea, transient loss of consciousness, and incontinence. | + + | Total removal of the cyst (transcallosal approach). | No neurological complications. |
| (18)    | 43/M       |               | Known case of colloid cyst of the third ventricle without evidence of ventriculomegaly for 6 years; presented with sudden onset of headache. Intact level of consciousness with no neurological deficits. | + + | Elective endoscopic partial removal of the cyst. | Resolved headache postoperatively without any neurological deficits. Follow-up imaging demonstrated a gradual reduction in the residual cyst size and normalization of the size of the lateral ventricle. |
| (19)    | 45/M       |               | Headache and malaise initially, then rapid deterioration of consciousness; GCS of 3. | + + | EVD initially on each side. Intensive care unit admission where brain death was certified. This was followed, however, by craniotomy with excision of the cyst (transcortical frontal approach). | Brain death. |

Case 1 47/M Type II diabetes mellitus, long-term smoker. Progressive morning headaches for 10 days associated with nausea and vomiting. Bilateral papilledema. + + Frontoal craniotomy and interhemispheric transcalsal transventricular approach for cyst excision. Left frontal EVD. Resolved headache.

Case 2 51/M Progressive memory disturbance and unsteady gait for 8 months. Bilateral papilledema. + + Left frontal craniotomy and transcalsal resection of the colloid cyst. Aphasia postoperatively, which improved with speech therapy. Unsteady gait demonstrated slight improvements.

M - male, F - female, EVD - external ventricular drain, ND - no data.

may play a role in the subsequent outcomes of these patients. All cases demonstrated the typical imaging features associated with a hemorrhagic colloid cyst, including hypointense areas along the wall of the cyst on T2-weighted images and susceptibility artifacts, which are suggestive of a bleed; these were histopathologically confirmed.\(^2,3,9-19\)

The first clinically diagnosed case of a hemorrhagic colloid cyst, reported by Beems et al.,\(^2\) was initially treated by an emergency ventriculotomy followed by
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elective endoscopic cyst removal. Additionally, Carrasco et al.\textsuperscript{11} reported a case of a hemorrhagic colloid cyst where a safe surgical strategy was employed, consisting of elective microsurgical removal following EVD.

The only pediatric case, reported by Farooq et al.\textsuperscript{3} was of a 9-year-old girl with a hemorrhagic colloid cyst; this patient had initially presented with sudden deterioration in her level of consciousness. She was treated with a similar approach with good clinical outcomes. In fact, most of the reported cases were also treated with a similar approach (Table 1).\textsuperscript{10,13-19} Only one case presented with abrupt anterograde amnesia, which was treated by direct emergency neuro-endoscopic excision of the cyst; this patient demonstrated complete resolution of the anterograde amnesia following surgical removal of the cyst. This case emphasized the importance of timely surgical excision on rapid improvements in memory disturbances.\textsuperscript{12} Similarly, one of our cases presented with the gradual onset of memory disturbances; this patient was treated with direct surgical removal and had similar outcomes.

Four out of the 13 reported cases were treated endoscopically.\textsuperscript{2,10,12,18} Only one reported case was treated by stereotactic aspiration followed by ventricular drainage, then stereotactic aspiration with endoscopic guidance and subtotal excision of the cyst.\textsuperscript{9} The rest of the cases were treated by an open approach.\textsuperscript{3,11,14,15,19} Our 2 cases were treated by an open transcallosal approach.

Figure 4 - A 55-year-old male (case 2) presented with progressive memory disturbance and unsteady gait for 8 months, which became more profound at the time of presentation.
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In both of our cases, there were severe adhesions at the surgical field which can make the endoscopic approach and stereotactic aspiration very difficult. So, in theory, we suggest to go with an open approach in hemorrhagic cases.

The outcomes were poor in 4 cases, satisfactorily in 3, and good with complete resolution in 5. All 4 cases that experienced poor outcomes tended to undergo rapid clinical deterioration following their initial presentation.

In conclusion, a hemorrhagic colloid cyst is a rare entity; however, bleeding tendencies and risk factors for bleeding should be carefully considered in patients with these cysts. The degree of rapidity in which clinical deterioration occurs may play a major role in the preferred treatment approach and in subsequent outcomes.

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