Endoscopic Removal of Third Ventricular Colloid Cyst: A Case Report

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Abstract:
Third ventricular colloid cyst is slow-growing benign tumor comprising <1% of intracranial tumors. Previously it was removed by interhemispheric transcallosal or transcortical approach. Now a days, endoscopic removal of colloid cyst is a popular option because it is less invasive, panoramic view helps in total removal of tumor. Here we are reporting a 8 years old boy presented with occasional headache, vomiting, seizure and loss of consciousness for 3 months. Neuro-imaging revealed anterior third ventricular colloid cyst. We removed the tumor completely with rigid neuroendoscope which was confirmed by histopathology. After three months follow up patient was quiet healthy.

Key words: Third ventricular Colloid cyst, rigid endoscope, Transcortical, Transcallosal

Introduction
Colloid cyst of the third ventricle is an uncommon lesion and the reported incidence is 3.2 per 100,000 population. The natural history is varied and most of the patients have an insidious course. However, some of the patients present with sudden neurological deterioration: drop attacks; life-threatening acute hydrocephalus; and sudden death. Even though the treatment approach for asymptomatic incidentally detected colloid cyst is debatable, most neurosurgeons advise intervention to avoid catastrophic neurological deterioration. However, long-term follow-up study of 68 subjects with asymptomatic colloid cysts suggests that only a few (8%) subjects go on to develop symptoms at 10 years.

The surgical approach for symptomatic colloid cysts is varied. The operative procedures usually consist of open microsurgical excision techniques. Less often stereotactic aspiration or stereotactic-guided surgery have been described. Endoscopic approach for excision of colloid cysts is relatively recent and offers the advantage of excellent visualization and minimally invasive corridor through the naturally dilated ventricular system. A more controlled resection of the cyst is possible through this approach avoiding damage to critical structures in the anterior third ventricle. We have presented this case because of minimal invasiveness of the procedure.
Case report
A 8 years old boy has presented with occasional headache, vomiting for three months. Intensity of headache increased at very early in the morning and was global in nature. His father also complaints of occasional seizure for several times which was generalized, clonic, tonic in nature. Seizure has persisted for few minutes. There was history of tongue bite, frothing with seizure. His father also complaints of occasional loss of consciousness for a brief period for several times. Computerized tomography scan (CT scan), Magnetic Resonance Imaging scan (MRI) with contrast showed colloid cyst at the anterior part of the third ventricle. We operated the case with rigid neuroendoscope and with the help of flexible neuroendoscope.

Surgical technique
The patient underwent the procedure under general anesthesia. Patient was positioned supine with the

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**Fig.-1:** (1a,1b,1c): Preoperative MRI of brain shows colloid cyst at third ventricle

**Fig.-2:** Postoperative patient with external ventricular drainage

**Fig.-3:** Postoperative CT scan of brain with shunt tube and ventricular haemorrhage

**Fig.-4:** Postoperative CT scan after 3 months
The opening. Thereafter, the cyst contents were inserted through the cyst wall and inflated to enlarge scissors. Once this was done, a Fogarty balloon was at all times. The cyst wall was cut open by endoscopic contact of the probe with normal ventricular structures to shrink towards the probe, avoiding then coagulated carefully allowing the vascular fibrous coagulator was then introduced. The cyst wall was the ipsilateral lateral ventricle. The endoscopic was seen bulging through the foramen of Monro into and the choroid plexus. In this case the colloid cyst of Monro, the septum pellucidum, thalamostriate vein and the endoscope introduced to visualize the foramen after ventricular puncture, the obturator was withdrawn and the endoscope introduced to visualize the foramen of Monro, the septum pellucidum, thalamostriate vein and the choroid plexus. In this case the colloid cyst was seen bulging through the foramen of Monro into the ipsilateral lateral ventricle. The endoscopic coagulator was then introduced. The cyst wall was then coagulated carefully allowing the vascular fibrous outer layer to shrink towards the probe, avoiding contact of the probe with normal ventricular structures at all times. The cyst wall was cut open by endoscopic scissors. Once this was done, a Fogarty balloon was inserted through the cyst wall and inflated to enlarge the opening. Thereafter, the cyst contents were aspirated.

We have introduced an external ventricular drainage for five days. Postoperatively patients developed small lateral ventricular haemorrhage. Ventricular haemorrhage become clear after five days. Postoperatively patient was uneventful. After 3 months follow up with CT scan there was no residual tumour and there was no clinical signs & symptoms. Patients used to take the anticonvulsant drug.

Discussion
Colloid cysts occur most commonly in the fourth through seventh decade. The symptom profile in our patient was similar to the other reports. Mishra et al. reported about 22% of patients in their series had triad of memory loss, gait abnormality and incontinence, symptom complex of normal pressure hydrocephalus. The mean age of these patients at presentation was 44 years. In our patient had gait disturbance, headache, vomiting and seizure. The optimal surgical management of colloid cysts continues to be a matter of debate. Colloid cysts are benign lesions with low risk of recurrence after radical resection. Open microsurgical excision was traditionally considered the "gold standard" treatment. However, operative approaches are not without risks. Of the 38 patients who underwent transcortical-transventricular resection of the colloid cyst, 30% of patients had postoperative complications. In Transcallosal approach avoids the potential risks of a cortical incision, but still is associated with occasionally serious morbidity. The reported complications of this approach include cortical venous infarct, fornical damage, injury to the deep venous system, subdural hematoma, disconnection syndromes, ventriculitis and meningitis. In this case there was ventricular haemorrhage and was managed by external ventricular drainage.

The quest for less invasive approaches for deal these lesions, led to the introduction of aspiration techniques for decompression of the cyst. Free-hand aspiration of colloid cysts was attempted in 1975 by Gutierrez-Lara, et al. To increase the accuracy of the procedure, stereotactic guidance was employed first by Bosch and co-workers and it gradually gained popularity, because the procedure is associated with negligible morbidity. However, it was soon realized that not all colloid cysts could be suitable for aspiration and high viscosity of the cyst contents seriously limited the evacuation of cyst material. Kondziolka and colleagues had found that only cysts which are hypodense or isodense on plain CT may be amenable to successful aspiration.

It is to be acknowledged that there is a definitive learning curve associated with the endoscopic treatment of colloid cysts. Presently we have attempted to perform a gross total excision of tumour.

Conclusion
We would prefer to approach such lesion through endoscope. Endoscopically resecting such lesions would prevent involve excessive manipulation of the fornix which can potentially cause short-term memory loss.

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