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Case Report

A bi-foraminal craniometric-guided approach to endoscopic third ventriculostomy and biopsy of a pineal tumour

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ABSTRACT

Background: Pineal tumors are very rarely encountered, with an incidence of <1% of intracranial lesions in adults. Life-threatening hydrocephalus due to obstruction of the third ventricle can result from the location of these tumours. Endoscopic third ventriculostomy (ETV) and tumor biopsy is a safe and feasible strategy, particularly if the tumor appears benign. This mitigates the high risks of uncontrollable venous bleeding from open and stereotactic biopsies. While typically performed using either ipsilateral single or dual bur holes, the location of the tumor may require modifications to the standard endoscopic techniques.

Case Description: A 55-year-old male presented with signs of intracranial hypertension and was found to have obstructive hydrocephalus due to a pineal tumour. The tumour displayed a right-sided dominance when the pre-operative imaging was assessed, which would risk forniceal injury if biopsied via a right-sided burr hole. Craniometric measurements revealed a superior trajectory to the tumour via the left foramen of Monro. A biforaminal approach was performed, with a traditional ETV using a right coronal bur hole and biopsy via a left frontal bur hole. This minimized forniceal stretching and allowed a safe biopsy.

Conclusion: The bi-foraminal approach has not been widely described in the literature but can potentially avoid morbidity with biopsy in patients with right-sided pineal tumours. We believe this technique should be considered, particularly in low-resource settings where neuroendoscopy is not commonly done, and where the use of ipsilateral single or dual-bur holes may lead to forniceal injury.

Keywords: Biforaminal approach, Craniometrics, Endoscopic third ventriculostomy, Pineal tumour

INTRODUCTION

Tumours of the pineal gland are very rarely encountered, with a quoted incidence of less than 1% of all intracranial lesions in adults. Life-threatening hydrocephalus from obstruction of the third ventricle may occur, due to the location of these tumours. Endoscopic third ventriculostomy (ETV) and tumour biopsy is known to be a safe and feasible strategy, particularly if the tumour appears benign. This mitigates the high risks of uncontrollable venous bleeding from open and stereotactic biopsies.

It is typically performed using either ipsilateral single or dual bur holes, which are placed with the guidance of neuronavigation. However, the location of the tumour may require modifying of the standard endoscopic techniques. This is particularly true in settings where neither neuronavigation

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nor flexible endoscopes are available, and craniometric measurements must be used to facilitate safe and accurate tumour biopsy. We report on a case of a pineal tumour with obstructive hydrocephalus, in whom a bi-foraminal approach to ETV and biopsy was utilized, with a good neurological outcome.

CASE REPORT

A 55-year-old male presented to the outpatient clinic with generalized headaches and blurred vision for 1-month's duration. He noted progressive worsening of the headaches, and they were particularly harsh on mornings, graded as 7/10 severity. They were not associated with nausea/vomiting, fever or seizures. His blurred vision prevented him from driving, and he noted that it had progressively declined. He had no known medical comorbidities, and the remainder of his history was unremarkable.

Physical examination revealed a fit-looking man who was haemodynamically stable. He was oriented to time, place and person and had a normal gait. Sensorimotor examination did not reveal any deficits. Pupillary assessment was within normal limits, at 3 mm and reactive to light and accommodation bilaterally. There were no gaze palsies or ophthalmoplegia, and no evidence of Parinaud's syndrome.

Visual acuity was reduced to 20/40, but visual fields were spared. Fundoscopy was positive for mild disc oedema.

The patient had an urgent magnetic resonance imaging (MRI) brain done [Figure 1] which demonstrated non-communicating hydrocephalus. There was obstruction of cerebrospinal fluid (CSF) flow through the aqueduct of Sylvius secondary to a pineal region tumour measuring 20 × 15 × 25 mm. It was isointense to the surrounding parenchyma on T1-weighted imaging, demonstrating homogenous contrast enhancement post-gadolinium administration.

Differential diagnoses of pineocytoma, germ cell tumour and metastasis were considered. Computed tomography scans of the chest, abdomen and pelvis did not reveal any evidence of malignancy. Hematological investigations including and serum germ-cell markers (alpha-fetoprotein, beta-human chorionic gonadotropin, placental alkaline phosphatase) were within normal reference ranges. The goals of management at this time were to divert CSF and obtain a tissue diagnosis. A decision was made to proceed with endoscopic third ventriculostomy (ETV) and pineal tumor biopsy. This was due to the high ETV success score calculated in this patient (over 90%).

Craniometric measurements were taken to determine the simplest route to the tumor, since neuronavigation is not

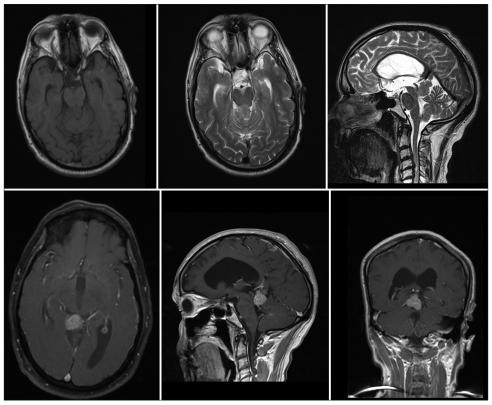


Figure 1: Top row-axial T1 (left), T2 (middle) and sagittal T2 (right) magnetic resonance imaging (MRI) showing pineal tumour dorsal to the midbrain, and resultant aqueductal stenosis. Bottom row-axial (left), sagittal (middle) and coronal (right) T1 post-contrast MRI showing homogenous enhancement of lesion, obstructive hydrocephalus and surrounding vascular structures.



Figure 2: T1 axial (left, sagittal (middle) and coronal (right) magnetic resonance imaging showing craniometrics for endoscopic third ventriculostomy (ETV) at the floor of the third ventricle, and ideal trajectory for tumor biopsy. As noted on the sagittal image, there was a significant distance between the ideal trajectories for the ETV and biopsy, necessitating dual bur holes. The left frontal pre-coronal bur hole was placed 5.19 cm from the glabella, with a measured distance of 10.789 cm to the tumour. The right coronal bur hole for the ETV was placed 11 cm from the nasion, with a measured distance of 8.15 cm to the floor of the third ventricle.

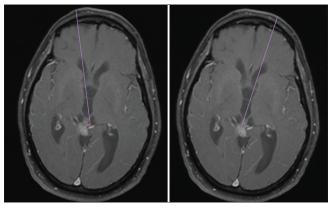


Figure 3: T1 post-contrast axial images showing trajectory through the foramen of Monro on right (left image), which does not target tumour, and the approach via the left foramen (right image) which directly enters into the pineal tumor.

available in our setting. The ETV would be performed through a standard right coronal bur hole, however, a separate left frontal-precoronal bur hole was decided to be appropriate for the biopsy. This tumor had a right-sided predominance, and possible trajectories through the right foramen were not leading to the tumour. A direct and simple trajectory would however be achieved by the use of a left-sided bur hole [Figures 2 and 3]. While a flexible endoscope and neuronavigation may have facilitated a single bur hole ETV and biopsy, only rigid endoscopes are available in our setting. The potential risk of causing multiple haemorrhages in the patient's dominant frontal lobe, as well as likely forniceal stretching and contusions, deterred us from the use of either a single or ipsilateral dual-bur holes.

The patient was placed under general anaesthesia and positioned supine. The head was kept neutral and slight neck flexion was done, and then secured in the Mayfield head clamp. A right Kocher's point bur hole was placed, and the right lateral ventricle cannulated with an external ventricular drain. CSF samples were taken at this time, to assess for germ-cell tumour markers and cytology. The ventricular drain was removed, and a rigid 0-degree endoscope then introduced. Confirmation of correct position was done by identifying landmark anatomic structures (septal vein, choroid plexus, thalamostriate vein). The foramen of Monro was entered and care taken to avoid injury to the fornices. In the third ventricle, the basilar artery was identified and the ventriculostomy then performed safely away from it. Flow of CSF through the ventriculostomy was noted, and there was no vascular injury.

A second low left frontal bur hole was then made, allowing the shortest safe linear trajectory to the pineal tumour. The third ventricle was entered, and the tumor was identified as a red, vascular lesion on the posterior wall of the third ventricle [Figure 4]. Biopsies were taken and the endoscopic bipolars used for haemostasis. A ventricular drain was left, and the left frontal bur hole was covered with titanium micro-plates. No defect was palpable or visible.

Post-operatively, the patient reported immediate cessation of headaches and resolution of his blurred vision. The drain was removed on postoperative day 2, and he was discharged on day 5. All CSF samples were negative for germ-cell tumour markers and cytology. Final histology was consistent with a pineocytoma, with conservative management and serial imaging as the management strategy. At a 4-month followup, the patient remains asymptomatic.

DISCUSSION

Pineal region tumors have an incidence of 1% of intracranial lesions in adults, and as high as 10% in the pediatric population.[3] Pineal parenchymal tumors, germ cell tumors,



Figure 4: Clockwise from top left-endoscopic entry into the left foramen of Monro, with choroid plexus, septal vein and thalamostriate vein visible. Top right-view within foramen of Monro. Bottom right-view of vascular appearing pineal tumor from the third ventricle. Bottom left-forniceal columns inferior to tumor.

neuroepithelial tumors and metastases comprise some of the pathology encountered.[3,9] The deep-seated location of the pineal gland, near the brainstem, makes traditional surgical approaches daunting and fraught with potential morbidity.

Up to 90% of these tumors present with potentially lifethreatening obstructive hydrocephalus, which requires urgent management prior to addressing the tumor. [6-8] Even if a stereotactic biopsy is available, this should not be done prior to treating the hydrocephalus, as post-biopsy edema may result in brain herniation.[3] While external ventricular drainage (EVD) or ventriculo-peritoneal shunting (VPS) are options, ETV is a well-described and safe method of CSF diversion in cases of obstructive hydrocephalus. EVD is only a temporary method of diversion, and both EVD and VPS both carry a significant risk of infection. VPS can promote tumor seeding into the abdomen, and there is a higher failure/malfunction rate of VPS in pineal tumors, up to 20%.[8]

The role of endoscopy

The use of the endoscope in the management of these tumors serves two purposes, CSF diversion and histological diagnosis. Treatment of hydrocephalus by CSF diversion can be safely and effectively done via ETV. The ETV success score predicts the likelihood of a good outcome in these patients and has been validated in adults. [4,5] It assesses the patient's age, cause of hydrocephalus and previous shunt surgery, and assigns a score to each. Our patient had a success score between 90% and 100% given his age over 50 years, hydrocephalus due to tumour and no past shunting.

Simultaneous biopsy of the tumor can be done, either using the same bur hole as for the ETV or a separate one. The availability of neuronavigation can aid in determining the ideal bur hole placement, or this trajectory can be calculated based on imaging studies.

When both procedures are planned to be done simultaneously, the ETV takes precedence. The rationale for this is to both address the more pressing concern of hydrocephalus, and prevent obscuring of anatomy if there is bleeding from the biopsy. [6,7] The main advantage of endoscopic biopsy compared to stereotactic biopsy is visualization of the nearby venous structures, and therefore a reduced risk of injury and venous bleeding.

Single versus dual bur holes, and the bi-foraminal approach

ETV is traditionally done using a right-sided coronal bur hole which allows visualization of the floor of the third ventricle and avoids potential injury to the eloquent speech area in the left hemisphere. The ideal trajectory for a pineal tumour biopsy is done by an ipsilateral frontal/pre-coronal bur hole that aims at the posterior third ventricle. [2,8] While some neurosurgeons favour a single bur hole placed between these two points, the method used should be based on the relative position of the tumor, ventricular size, dimension of the massa intermedia and goal of surgery.^[6,7] A flexible endoscope, along with neuronavigation, can facilitate the use of a single bur hole in these cases, with a lower risk of forniceal contusions when compared to the use of a rigid endoscope.[10]

In settings where neuro-endoscopic procedures are not commonly done, and where neuronavigation is unavailable, the use of ipsilateral single or dual-bur holes in cases such as these may unnecessarily risk intraoperative cortical bleeding, forniceal contusions and basilar artery injury.[11] Neurocognitive deficits post-endoscopic ventricular surgery have been described by Soleman and Guzman in their review article in 2020, with a 2% transient and 1% permanent rate of neurological impairment.[10] It was noted that the use of the non-dominant hemisphere for the endoscopic approach to the ventricles was associated with a reduced risk of deficits.

In our case, given the lack of flexible endoscopes and the relative right-sided location of the tumour [Figure 3], we opted for an ETV with a right pre-coronal bur hole, and tumour biopsy using a left low-frontal bur hole. Based on detailed imaging analysis, we decided that the risk of forniceal stretching and potential injury was inevitable if biopsy was attempted through a bur hole on the same side as the ETV. Additionally, the performance of two bur holes in the dominant frontal lobe (with two separate trajectories) and the risk of causing cortical haemorrhages and subsequent neurological deficits were a major concern.

A separate, contralateral left-sided bur hole, therefore, afforded a superior trajectory to the tumour, while reducing the above-mentioned risks. While this technique has not been specifically described in the literature, it should be considered as an option in tumours that may not be accessible via an ipsilateral bur hole to the ETV. The value of the biforaminal approach increases when only rigid endoscopes are available, and even more so when there is no accessible neuronavigation.

Biopsy success rate

Balossier et al. compared several series in 2015 and found a range of 60-100% positive endoscopic biopsies of pineal region tumours.[3] However, a more recent series of 64 patients by Samadian et al. in 2019 had a negative biopsy rate of only 3%. A higher number of samples (6-10) increases the positivity rate, and all studies with 100% success rate had a minimum of 8 samples.[9]

Risks and complications

If there is a failure of resolution of hydrocephalus, patients may require eventual VPS. However, the ETV success score allows prediction this, and some authors have reported a 1% chance of requiring shunting after ETV.[9]

Hemorrhage from intraventricular surgery reported rate of 4.8%.[3,11] The majority of bleeding during intraventricular surgery can be arrested with steady irrigation, bipolar coagulation and patience. If severe, abandoning the procedure and placement of a ventricular drain, or switching to microsurgery may be necessary. [9] Coagulation of the tumour prior to sampling may reduce the risk of intraoperative haemorrhage. However, this can alter the cellular architecture of the sample, and lead to difficulty in histological interpretation.

Short-term memory loss from forniceal contusions is documented after this procedure, but fortunately quite rare. [2,7,9,10] Infection, transient diabetes insipidus, intraventricular haemorrhage requiring temporary ventricular drainage and tumour seeding are other complications that may arise from this procedure.[1-3,9]

CONCLUSION

The bi-foraminal approach to combined ETV and biopsy for pineal tumours can be utilized to enable safe tumour biopsy. The tumour location should be studied carefully on preoperative imaging, to guide the placement of bur holes using craniometric measurements. While typically ipsilateral single- or dual-bur holes are done, this may unnecessarily risk forniceal injury, especially if only rigid endoscopes are available. This technique is an additional strategy available to neurosurgeons, particularly in regions where there is a lack of readily accessible neuronavigation. This necessitates careful study of preoperative imaging and accurate craniometric measurements, for a safe and successful procedure.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent.

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Conflicts of interest

There are no conflicts of interest.

REFERENCES

- Abbassy M, Aref K, Farhoud A, Hekal A. Outcome of singletrajectory rigid endoscopic third ventriculostomy and biopsy in the management algorithm of pineal region tumors: A case series and review of the literature. Childs Nerv Syst 2018;34:1335-44.
- Amer MA, Elatrozy HI. Combined endoscopic third ventriculostomy and tumor biopsy in the management of pineal region tumors, safety considerations. Egypt J Neurosurg 2018;33:4-9.
- Balossier A, Blond S, Touzet G, Lefranc M, de Saint-Denis T, Maurage CA, et al. Endoscopic versus stereotactic procedure for pineal tumour biopsies: Comparative review of the literature and learning from a 25-year experience. Neurochirurgie 2015;61:146-54.
- Kulkarni AV, Drake JM, Kestle JR, Mallucci CL, Sgouros S, Constantini S. Predicting who will benefit from endoscopic third ventriculostomy compared with shunt insertion in childhood hydrocephalus using the ETV Success Score. J Neurosurg Pediatr 2010;6:310-5.
- Labidi M, Lavoie P, Lapointe G, Obaid S, Weil AG, Bojanowski MW, et al. Predicting success of endoscopic third ventriculostomy: Validation of the ETV success score in a mixed population of adult and pediatric patients. J Neurosurg 2015;123:1447-55.
- Morgenstern PF, Osbun N, Schwartz TH, Greenfield JP, Tsiouris AJ, Souweidane MM. Pineal region tumors: An optimal approach for simultaneous endoscopic third ventriculostomy and biopsy. Neurosurg Focus 2011;30:E3.
- Morgenstern PF, Souweidane MM. Pineal region tumors: Simultaneous endoscopic third ventriculostomy and tumor biopsy. World Neurosurg 2013;79:S18.e9-13.
- Robinson S, Cohen AR. The role of neuroendoscopy in the treatment of pineal region tumors. Surg Neurol 1997;48:360-7.
- Samadian M, Maloumeh EN, Shiravand S, Ebrahimzadeh K, Sharifi G, Mousavinejad A, et al. Pineal region tumors: Long-term results of endoscopic third ventriculostomy and concurrent tumor biopsy with a single entry approach in a

- series of 64 cases. Clin Neurol Neurosurg 2019;184:105418.
- 10. Soleman J, Guzman R. Neurocognitive complications after ventricular neuroendoscopy: A systematic review. Behav Neurol 2020;2020:2536319.
- 11. Wasi MSI, Sharif SY, Gulzar F. Implication of image guidance

in endoscopic third ventriculostomy: Technical note. Surg Neurol Int 2020;11:87.

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