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Awake craniotomy for fenestration of motor cortex neuroglial cyst: A case report

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Case Report

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ABSTRACT

Background: Neuroglial cysts (NCs) are uncommon benign cysts covered by an epithelial layer, accounting for <1% of all intracranial cysts. The optimal management approach for these cysts remains a subject of debate. Given their rarity, management principles used for arachnoid cysts can be applied to NCs.

Case Description: We present a case of a 35-year-old male without prior medical history, who presented to the neurosurgery clinic with complaints of absence seizures. A neurological examination revealed subtle weakness in the left upper limb. Brain magnetic resonance imaging demonstrated a large cystic lesion in the posterior frontal and anterior parietal lobes of the right hemisphere. The patient underwent an awake craniotomy, during which a cystoventricular fenestration was performed on the motor cortex cyst. Histopathological examination confirmed the diagnosis of NC. At the 4-month follow-up, the patient experienced complete recovery, with normal strength (5/5) in all limbs and absence of seizure remission.

Conclusion: This case highlights the successful use of awake craniotomy for the fenestration of an NC in the motor cortex. Given the rarity of NCs, there is no consensus on the optimal treatment strategy. However, in this particular case, the patient achieved complete recovery without any new neurological deficits following the procedure.

Keywords: Awake craniotomy, Cyst fenestration, Glioependymal cyst, Neuroepithelial cyst, Neuroglial cyst

INTRODUCTION

Neuroglial cysts (NCs), also referred to as glioependymal cysts or neuroepithelial cysts, are benign cystic formations characterized by the presence of an epithelial lining. These cysts are considered rare, accounting for <1% of all intracranial cysts. They can emerge at various locations within the neuraxis.^[2,3] The purpose of this paper is to present a unique case involving a neuroglial cyst located in the motor cortex, which was successfully managed through an awake craniotomy procedure and cystoventricular fenestration. Notably, to the best of our knowledge, there have been no previously published reports detailing the utilization of awake craniotomy for the treatment of NCs occurring in eloquent cortical regions. Hence, this study aims to provide valuable insights by sharing our experience in managing this particular case, serving as a reference for future research and clinical practice.

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DESCRIPTION OF THE CASE

A 35-year-old male with no known medical illnesses presented to the neurosurgery clinic with a 2-month history of absence seizures with a frequency of 4–7 times a day. He had weakness in his left-hand grip, with a power examination of 4 ± 5 . A satellite sign in the left upper limb was observed on the forearm-rolling test. Other than the left upper limb weakness, his remaining neurological examination was unremarkable.

Brain magnetic resonance imaging (MRI) with contrast showed a large nonenhancing right cystic intra-axial lesion involving the posterior frontal and anterior parietal lobes measuring $6.7 \times 6.1 \times 5.3$ cm AP × TS × CC (anteroposteriorly, transversely, and coronal). [Figures 1a-c] The patient was planned for awake craniotomy and cyst fenestration.

Surgical technique

Phase one (asleep)

Operative preparation and nerve block

After general anesthesia, scalp sterilization commenced. A nerve block was used on the right side on the supraorbital, supratrochlear, zygomaticotemporal, auriculotemporal, and greater occipital and lesser occipital nerves. The supraorbital, supratrochlear, zygomaticotemporal, and auriculotemporal were also blocked on the left side. Local anesthetic was also applied on the sites of Mayfield pins.

Position and incision

The patient was positioned supine, and his head was flexed and tilted slightly to the left. A three-point Mayfield skull clamp was placed. The incision was marked to be straight, extending from the midline to 2 cm just above the ear pinna. Intra-operative images are illustrated in Figures 2a-c.

Exposure

After retracting the pericranium, two burr holes were placed on the incision's superior and inferior poles, dissecting the dura, and a craniotomy was completed. The dura was opened in a curvilinear fashion with the base posteriorly. The brain was bulging, and furosemide bolus was given. The first phase of the operation had finished.

Phase two (awake)

The patient was roused from sleep for the second phase of the operation, and a neurological assessment was conducted. His

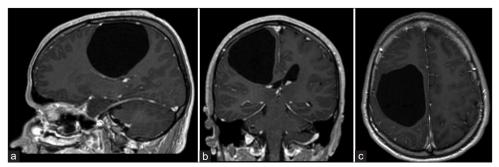


Figure 1: Preoperative MRI: (a) sagittal, (b) coronal, and (c) axial/transverse.

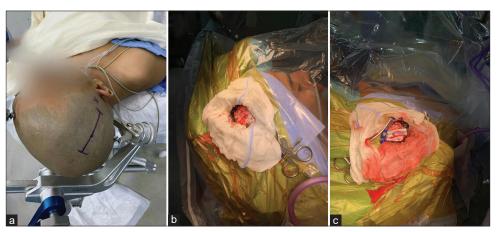


Figure 2: Intraoperative images of (a) the position and incision marking, (b) after minicraniotomy, and (c) the location of corticectomy.

results were consistent with the preoperative baseline, and a series of tests were conducted to evaluate the language, the left upper limb, and the left lower limb. These tests were repeated throughout the phase with no abnormalities detected.

Surgical technique

Using neuronavigation to pinpoint the thinnest and most superficial part of the cyst, the surgical team performed coagulation of the pia and a corticectomy. A ventricular cannula was inserted into the cyst, and 10 cc of clear fluid resembling cerebrospinal fluid (CSF) was released. The cyst deflated, and a second corticectomy was done to inspect the cyst wall. A thin, whitish membrane was observed, and a biopsy sample was taken. Multiple septations were found, which were fenestrated and connected. The cyst was fenestrated to the ventricle.

Phase three (asleep) and closing

The operation moved to the third "asleep" phase of the awake craniotomy, and the patient was sedated again. Hemostasis was achieved; then, the dura was approximated primarily

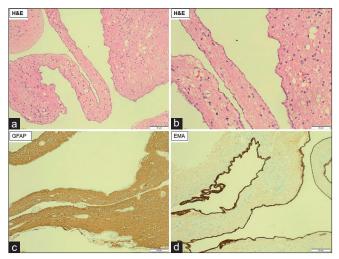


Figure 3: (a-d) Histopathology slides.

by silk, followed by fibrin sealant and a hemostatic agent. Microplates and screws fixed the bone, and bone dust was used to plug the burr holes. Pericranium was approximated, and subcutaneous tissue was sutured. The skin was closed by staplers.

Postoperative course

Histopathology specimen showed nonciliated cuboidal to simple squamous (nonkeratinizing) epithelial surrounded lining by gliosis and calcification. Immunohistochemistry study showed cytokeratins (CK)+, epithelial membrane antigen (EMA) focal weak+, S100+, glial fibrillary acidic protein (GFAP)-. The tissue diagnosis is a neuroepithelial cyst. [Figures 3a-d] Postoperative day 2, his weakness started improving with the upper and lower limb power examination of 4+/5. He was discharged on levetiracetam in stable condition.

In a clinic follow-up appointment 4 months after the operation, the patient made a full recovery with a power of 5/5 in all limbs, no episodes of absence seizure since the surgery. Follow-up MRI showed interval reduction in the cystic lesion, measuring $3.7 \times 3 \times 3.5$ cm compared to the preoperative $6.7 \times 6.1 \times 5.3$ cm [Figures 4a-c].

DISCUSSION

NCs are a rare type of intracranial cysts, comprising <1% of all cases.^[1-3] There are no specific symptoms for NCs, and they are often found incidentally; however, the clinical presentation is dependent on the location and size of the cyst and may include headaches, motor weakness, visual disturbance, or dizziness.^[2,3]

MRI is the imaging of choice for NCs, with features similar to most intracranial benign cystic lesions, which include smooth and rounded borders, a nonenhancing wall, and minimal mass effect even in large lesions. MRIs show NCs to be isointense to CSF in all sequences.^[1-3]

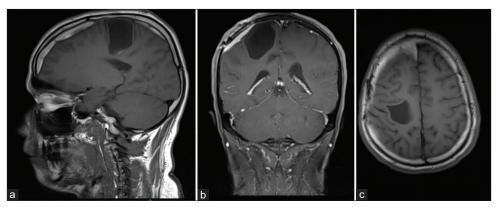


Figure 4: Postoperative MRI: (a) sagittal, (b) coronal, and (c) axial/transverse.

The surgical principles used to treat arachnoid cysts can be applied to the management of NCs and other intracranial benign cysts. As this is a rare condition, a consensus on the treatment strategy has not been established. The most common procedure implemented in the literature was cystoventricular fenestration, and none reported awake craniotomy.^[3] The surgery of choice for NCs depends on the location of the cyst. As in our case, the cyst was located in a part of the eloquent cortex, the motor cortex, which indicated awake craniotomy. Given the lack of consensus regarding the management of intracranial NCs, this case report details the results and outcomes of employing awake craniotomy and cystoventricular fenestration as a treatment modality for such lesions.

CONCLUSION

This paper presents a case of a young adult with subtle motor weakness in his left upper limb and absence of seizure who made a full recovery after an awake craniotomy. This case highlights the importance of considering awake craniotomy as a treatment option for NCs located in the eloquent cortex. Further, research is needed to establish a consensus on the treatment strategy for NCs.

Declaration of patient consent

Institutional Review Board (IRB) permission obtained for the study.

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Conflicts of interest

There are no conflicts of interest.

Use of artificial intelligence (AI)-assisted technology for manuscript preparation

The author(s) confirms that there was no use of artificial intelligence (AI)-assisted technology for assisting in the writing or editing of the manuscript and no images were manipulated using AI.

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