Case Report

Adult Pilocytic Astrocytoma in the insula: Case report and review of the literature

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ABSTRACT

Introduction: Adult Pilocytic Astrocytomas (APA) are infrequent low grade tumors. While supratentorial APA is considered rare, insular APA is extremely rare.

Case presentation: We present a case of pure insular APA along with surgical outcomes. Tractography and functional MRI were obtained pre-operatively. The patient underwent neuro-navigation guided microsurgical resection with sub-cortical white matter mapping, utilizing intra-operative MRI guidance. The Sylvian fissure was opened to secure the M3 branches, and near total resection was achieved.

Clinical discussion: APA in the insula is a very rare presentation and is considered challenging. Its proximity to the middle cerebral and lenticulostriate arteries, motor areas, and language areas makes accessing and resecting the tumor challenging. A multidisciplinary approach by an experienced team is needed to plan the management of young adult patients and reach the best outcomes.

Conclusion: Implementing microsurgical techniques, modern imaging modalities and intraoperative mapping helps to achieve maximal safe resection without risking functions.

Introduction

Adult Pilocytic astrocytomas (APA) are rare tumors. They are classified as World Health Organization grade I gliomas [6]. PAs in general tend to be more prominent in children and adolescents. The incidence of PAs decreases significantly from 11.4% in children and adolescents to only 0.8% of CNS tumors in adults [7]. In addition, tumors location distribution seems to be related to the age of the patients. As it is more prominent to observe cerebellar presentations in pediatrics, in contrast to supratentorial location prominence in adults [10]. PAs present usually manifest with a space-occupying lesion affecting the posterior fossa as obstructive hydrocephalus with clinical pictures of headache, vomiting, nausea, and papilledema [9]. The excision of the tumor surgically is the treatment of choice aiming at tumor-free margins, with the least neurological insults. Without recurrence, further resection surgically is mainly performed. APA in the cerebral hemispheres is very rare in the literature, insular location is even extremely rare. Surgical resection of insular tumors is challenging due to their proximity to the middle cerebral artery (MCA), lenticulostriate arteries (LSA), motor areas, and language areas. Herein, we present a rare case of adult pilocytic astrocytoma presenting in the insula that have been managed at a cancer centre in Jordan and discuss it with the available literature. This case has been reported according to SCARE criteria for case reports [1].

Case report

A 23 year-old right-handed male patient presented to the ER by his parents with severe progressive headache associated with short-term memory loss and refractory seizures (KPS 60). The patient is known to have epilepsy since the age of 3 years and has a history of left-sided

Abbreviations: KPS, Karnofsky scale; MCA, middle cerebral artery; LSA, lenticulostriate arteries.
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insular brain tumor for four years. He was observed conservatively for four years through serial Magnetic Resonance Imaging (MRIs). On examination, the patient showed loss of tandem gait, pronator drift and weakness in the right side. CT scan was performed to exclude any active brain hemorrhage. Subsequent MRI revealed significant increase in the size of both cystic and solid parts of the tumor with midline shift and obstructive hydrocephalus (Fig. 1A, B). Awake craniotomy option was discussed with the patient but was uncooperative. Preoperative imaging assessment with Tractography MRI showed displacement of the corticospinal tract (Fig. 2) by the tumor. Functional MRI was performed to identify speech area (Fig. 2). Neuronavigation guided Surgical resection with cortical mapping and subcortical white matter mapping was performed by BM MD, MRCSI, FEBNS - a senior neurosurgical oncology and skull base surgery specialist practicing in a tertiary cancer centre-after opening the sylvain fissure to secure the M3 branches and corticotomy over the preplanned trajectory (Fig. 1, A). Motor cortex and corticospinal tract mapping using a bipolar probe as a stimulator was utilized (Fig. 3). Near total resection was safely achieved Figure (1C, D). Histopathological study revealed pilocytic astrocytoma with varying proportions of compacted bipolar cells and loose microcystic areas. Numerous Eosinophilic granular bodies are seen, as well as many entrapped ganglion cells. Rare mitotic figures are identified. The tumor cells are positive for GFAP, negative for CD34, PS3 and IDH-1. ATRX is lost in tumor cell nuclei. Ki67 index is around 4%. The postoperative KPS was 80 until the last follow up of 12 weeks.

Discussion

Pilocytic astrocytoma is very rare in adults, with only scarce data available on the tumor characteristics, clinical course and prognostic factors. To our knowledge, this is the first report describing pilocytic astrocytoma in the insula for an adult patient. The presentation of the tumor usually mimics the manifestation caused by a space-occupying lesion affecting the posterior fossa such as obstructive hydrocephalus with a clinical picture of headache, seizures, vomiting, nausea, and papilledema.

The tumors location in adults goes in another direction when compared with the pediatric population, as its more prominent in the posterior fossa. Ye et al. [13], Ishkanian et al. [5] and Bond et al. [2] all have published studies that analyzed the patients data and found that the tumor distribution was infratentorial in more than 50% of the cases, which is similar to pediatric PA.

Theeler et al. [11] have analyzed the data of 127 adult patients with PA and found a predominance of supratentorial tumors in 53% of the cases. However, 13% of their cohort were reported as ventricular tumors, without further specifications. Boschetti et al. [3] showed the same with 73.9% supratentorial distribution in their adult APA cohort.

For all PA’s, the excision of the tumor surgically is the treatment of choice aiming at tumor-free margins, with the least neurological insults. With recurrence, further resection surgically is usually performed.

Cyrinea et al. [4] investigated lesions in the eloquent and

![Fig. 1. Brain MRI with contrast; A) Axial preopertaive, B) Coronal preopertaive, C) Axial post pertaive, D) Coronal postopertaive views.](image-url)
deep-seated areas like the optochiasmatic area, diencephalon, and brainstem. These locations are significant challenges in the face of surgical interference while biopsy and subtotal resection may be the only and best treatment choice. In all, it raises the suspicion that a more limited resection extent, due to constrained surgical accessibility, could be related to the elevated progress rates and worse prognosis attributed to specific sites. Though the resection extent usually has a connection to the survival rate in most of the studies assessing these factors, a resection extent appraisal can be complicated and cause a biased investigation.

The extent of resection (EOR) is a main prognostic factor in Overall Survival OS, malignant transformation, and progression-free survival (PFS). Yahanda et al. [12] investigated 284 subjects undergoing grade I gliomas treatment. They had 248 tumor resections; the higher is the EOR, the more improved is the five-year OS with $p = 0.013$ and the five-year PFS with $p < 0.0001$.

Resection of the tumor enhanced the life quality for patients manifesting with intractable epilepsy. Regarding the evidence presented, trying to maximize the tumor resection and minimize the neurological effect should be aimed at insular glioma intervention. The challenge that every neurosurgeon encounters is to control both seizures and lesions while preserving most of the functions. Collected experience in resecting these tumors established significant flexibility in the region. Recognition of social cognition, cognitive functions, and language intraoperatively of the right non-dominant hemisphere (RnDh) utilizing intraoperative brain mapping (ioBM) could be of utmost importance in enhancing the resection extent of low-grade gliomas and positively affect the neuropsychological and clinical result at six months [8] which might permit the majority of patients to reimburse overtime after aggressive resections of insular gliomas given that motor locations recognized by stimulation and essential language are maintained in surgery. Implementing microsurgical techniques, modern imaging modalities and intraoperative mapping helps to achieve maximal safe resection without risking functions.

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**Consent for publication**

Written informed consent was obtained from the patient for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

**Author contribution**

Baha’eddin A. Muhsen MD1: Performed the intervention, supervised and reviewed the manuscript.

Ansam Ghzawi MS2: Manuscript writing, final revision.

Hasan Hashem MD5: Editing and reviewing the manuscript. Maher Elayyan1, Bayan Maraqa MD3 and Mahmoud Al Masri4: finalizing the review process.

**Research registration (for case reports detailing a new surgical technique or new equipment/technology)**

N/A.

**Guarantor**

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Appendix A. Supplementary data
Supplementary data to this article can be found online at https://doi.org/10.1016/j.amsu.2021.102300.

Fig. 3. Motor mapping in the lower limbs.

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