Case Report

Intraventricular ganglioglioma with unusual location and morphology–A case report

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Abstract

Ganglioglioma is one of the common tumour responsible for refractory temporal lobe epilepsy in children and young adults. Positioned in the group glioneural tumours, it is mostly of low grade histology. Here we outline a rare case of Intraventricular ganglioglioma at atypical location with atypical morphology, lacking its most classical symptom of epilepsy.

Introduction

Tumour of young adults and children, ganglioglioma constitutes 1% of all brain malignancies [1]. It is described as one of the most common malignancy associated with temporal lobe epilepsy along with DNET [2]. However intraventricular counterpart of ganglioglioma is less likely to be associated with seizures and is more common in females [2]. Despite being neoplasm of low grade histology, it is likely to show progression into anaplastic variants, hence early diagnosis and surgical resection is of utmost importance.

Case report

10 year old female child presented with history of chronic gradual onset headache for 1 year accompanied with occasional vomiting. There was progressive deterioration of vision for 1 year. No history of weakness in limbs or seizures was elicited.

MRI revealed well defined heterogeneously enhancing mass arising from the foramen of Munro (Figure 1). The mass was seen obstructing the 3rd ventricle causing bilateral lateral ventricular dilatation with periventricular ooze and corpus callosum thinning. The mass was hypointense to cortex on T1-weighted image (T1-WI) and intermediate signal intensity on T2-weighted image (T2-WI). Areas of blooming was seen within the mass on T2*-weighted image (T2*-WI) along with cystic changes. The mass was seen compressing the hypothalamus inferiorly and supra-optic recess superiorly. Preoperative diagnosis of central neurocytoma or intraventricular ependymoma was suggested.

Patient underwent right frontal craniotomy and decompression of intraventricular tumour. Intraoperatively the tumour was highly vascular and was fed by the choroidal arteries with adherence to surrounding structures. To avoid injury to surrounding vital structures only decompression of tumour was performed.

On follow up CT (Figure 2a) residual heterogeneous mass with haemorrhagic areas was seen at the foramen of Munro level with persisting obstructive hydrocephalus. Right VP shunt

was placed. Significant improvement in vision was recorded postoperatively.

Histopathology (Figure 2b) revealed tumour composed of neuronal and glia elements. The neuronal elements had dysplastic features with abnormal clustering and occasional binucleated and multinucleated cells. The ganglion cells were large having central to eccentric nuclei with abundant cytoplasm. Some of the cells had prominent nucleoli. The glial component showed dense fibrillary network and areas of cystic change. Few Rosenthal fibres and eosinophilic granular bodies were seen. There was mild vascular proliferation seen with perivascular cuffing of lymphocytes. Plenty of cyst macrophages along with chronic inflammatory cells composed of lymphocytes were also seen. Microcalcification was seen. The tumour showed marked capillary network resembling malformative angiomaticus component. No mitosis or necrosis seen.

Immunohistochemistry (IHC) revealed positive results for GFAP, synaptophysin, NF (indicates positive for ganglion cells), CD34 (indicates positive for endothelial cells) and Ki67 Labelling index were <1%. The final pathological diagnosis was Ganglioglioma.

No radio-chemotherapy was given as the tumour was of grade.

Discussion
Ganglioglioma is the most common neuronal-glial neoplasm of the CNS. Cyst with mural nodule has been described as its classical appearance (52%) on cross sectional imaging with temporal lobe as the most favoured site (38%-40%) [3]. However our patient lacked classical morphological appearance and site. Atypical appearances of ganglioglioma includes solid mass (43%) and cystic mass (5%). Atypical location of ganglioglioma include parietal lobe (30%), frontal lobe (18%), very rare location includes brainstem, cerebellum, spinal cord, pineal region, optic nerve optic chiasma and ventricles [4]. Tumor is most commonly seen in children and young adults [4].

On unenhanced CT 38% of ganglioglioma are hypodense, 32% have mixed attenuation, 15% Isodense, 15% hyperdense. Calcification is seen in 30% of cases. Approximately 16%-80% show enhancement on post contrast study. Ganglioglioma show variable MR features with most appearing hypointense to isointense to grey matter on T1 and hyperintense on T2. Rarely may they also appear hyperintense on T1. On T2 some areas of high signal are ideal for the diagnosis of ganglioglioma. On gadolinium enhanced images the lesion may show homogenous, ring like or no enhancement [1,2].

Hauck, et al., [5] and Goncalves, et al., [1], described the predilection for female gender in cases of intraventricular ganglioglioma in comparison to its cortical counterpart, which is also seen in our case. To our knowledge, only 3 cases of intraventricular Ganglioglioma with haemorrhage have been described in literature [6]. Goncalves, et al., [1], also described bizarre (eccentric and exuberant) pattern of calcification in these tumours.

Conclusion
Ganglioglioma is predominant cortical tumour, and intraventricular location is rare. Atypical clinical features depicted in our case is lack of seizures which is most common symptom of cortical Ganglioglioma. Female preponderance of intraventricular location is also of divergent behaviour. Hence differential of Ganglioglioma should be considered in young female with intraventricular mass.
References


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