A RARE CASE OF SPACE OCCUPYING LESION OF BRAINSTEM IN AN ELDERLY MALE PATIENT

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INTRODUCTION

Brainstem tumours are defined as lesions which occur in the midbrain, the pons, or the medulla oblongata. The tumour’s extension is considered focal when it occupies <50% of the axial brainstem diameter, and the extension is considered diffuse when the lesion is poorly demarcated and is >50% of the brainstem diameter [1–3]. Brainstem glioma is the most frequent tumour of the region. A clear bimodal age distribution supports the distinction between brainstem gliomas in children and adults. In contrast with the paediatric population in which brainstem gliomas represent up to 20% of brain tumours and exhibit a rather homogeneous and unfavourable course, adult brainstem gliomas are rare (accounting for only 1%–2% of adult brain gliomas) and heterogeneous with varying radiological patterns and variable prognosis[4,5]. Histologically, adult brainstem gliomas can have an astrocytic, oligodendrogial, or mixed appearance, with astrocytic tumours further characterized as either pilocytic or diffusely infiltrative.

CASE REPORT

An elderly male patient aged 65 presented to us with history of swaying towards left side of the body since 1 month with normal higher mental functions and neurological examination suggestive of cerebellar ataxia. MRI Brain plus contrast was suggestive of an irregular, ill-defined heterogeneous enhancing lesion with few necrotic areas within and few foci of blooming on FFE (Fast Field Echo imaging technique) with significant perilesional oedema involving right thalamus and brainstem showing Choline peak on MR Spectroscopy.

KEYWORDS: Neurological examination; Cerebellar ataxia; MRI brain; FFE.

ABSTRACT

An elderly male patient aged 65 presented to us with history of swaying towards left side of the body since 1 month with normal higher mental functions and neurological examination suggestive of cerebellar ataxia. MRI Brain plus contrast was suggestive of an irregular, ill-defined heterogeneous enhancing lesion with few necrotic areas within and few foci of blooming on FFE (Fast Field Echo imaging technique) with significant perilesional oedema involving right thalamus and brainstem showing Choline peak on MR Spectroscopy.

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DISCUSSION

On evaluating for the space occupying lesion of the brain by neuroimaging and other relevant investigations to rule out other systemic involvement and infective pathology, differential diagnosis considered were 1) High grade glioma 2) Multicentric glioma 3) Glioma with metastasis. Glioma with metastasis was ruled out as the history, clinical examination and investigations revealed no evidence of tumour elsewhere in the body. High grade gliomas are malignant, often rapidly progressive brain tumours that are divided into anaplastic gliomas and glioblastoma based on their histologic and molecular features. Multicentric gliomas are widely separated lesions whose simultaneous presence cannot be attributed to dissemination through commissural pathways, CSF channels, and blood or by local extension through satellite formation [6]. As neuroimaging in this case does not show any evidence of disseminated lesions, high grade glioma can be considered as our probable diagnosis. Conventional radiotherapy is the standard treatment for diffuse intrinsic low-grade brainstem gliomas in adults (the median survival is 5 years)[6].

CONCLUSION

Differential diagnosis considered were High grade glioma, multicentric glioma and Glioma with Metastasis and we speculate that this patient had high grade glioma. Despite significant advances in neuroradiology techniques, a purely radiological classification remains imperfect in the absence of a histological diagnosis. A biopsy may often be reasonably avoided in the diffuse non-enhancing forms, obtaining histological proof seems necessary in many contrast-enhanced brainstem lesions because of the wide variety of differential diagnosis in adults.

REFERENCES


Table 1. Shows various investigations done and their results

<table>
<thead>
<tr>
<th>INVESTIGATIONS</th>
<th>RESULT</th>
</tr>
</thead>
<tbody>
<tr>
<td>HIV Rapid</td>
<td>Negative</td>
</tr>
<tr>
<td>Blood Glucose</td>
<td>159 mg/Dl</td>
</tr>
<tr>
<td>Serum Creatinine</td>
<td>1.0 mg/dL</td>
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<tr>
<td>Serum Sodium</td>
<td>135 mmol/L</td>
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<tr>
<td>Serum Potassium</td>
<td>3.8 mmol/L</td>
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<tr>
<td>Total Leucocyte Count</td>
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<td>Hemoglobin</td>
<td>14.0 gm%</td>
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<tr>
<td>RBC</td>
<td>4.74 millions/cmm</td>
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<tr>
<td>ESR</td>
<td>20</td>
</tr>
<tr>
<td>Platelet</td>
<td>2.69 lakhs/cmm</td>
</tr>
</tbody>
</table>

Fig 1. Irregular, ill-defined lesion appearing isotense on T2/ FLAIR

Fig 2. Heterogenous enhancement with few non-enhancing necrotic areas within
