Multiple Intracranial Meningiomas - A Case Report

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INTRODUCTION

Meningiomas are benign tumors, accounting for 13-20% of all primary central nervous systems tumors. Multiple meningiomas are rare and the term is used for more than one meningioma occurring in several intracranial locations without signs of neurofibromatosis. The prognosis of multiple intracranial meningiomas is similar to benign solitary meningioma. We present a rare case of multiple intracranial meningiomas in a 45 years old female patient.

Keywords: Meningiomas, Multiple, Intracranial.

CASE REPORT

A 45 years old female patient presented with headache, seizures and upper and lower limb weakness. MRI brain revealed multiple meningiomas along right parietal parasagittal region attached to convexity of dura. The lesions were excised and sent for histopathological examination.

Histopathological Examination

We received 5 globoid grey white firm tissue pieces with attached dura. The largest tissue piece of the tumor was 7.5 cm in diameter. Cut section of all tumor pieces were grey white and homogenous (Fig. 1, 2).

Microscopic examination of all tumor deposits showed neoplastic spindle shaped cells arranged in whorled pattern. Neoplastic cells showed elongated, flat nuclei without atypia & scanty cytoplasm. The tumor showed numerous psammoma bodies. All tumor deposits showed similar microscopic grade. Considering the histopathological features, the diagnosis was given as multiple meningiomas WHO grade I (Fig. 3, 4).

DISCUSSION

A widely accepted hypothesis for multiple meningiomas is spread of clones of neoplastic cells through cerebrospinal fluid as well as hematogenous spread [3]. Different etiological factors were considered in the development of meningiomas, one of which is deletion of chromosome 22 in patients with type II neurofibromatosis. The second etiological factor is hormone. Higher frequency of meningiomas is seen in female patients. Progesterone receptors are found in 80% meningiomas [4]. In our case, the patient was premenopausal female. The most common location of multiple meningiomas are supratentorial convexity and parasagittal area, posterior fossa being the rare site [5]. Similar location was found that is parietal parasagittal region attached to convexity of dura in our patient.

Histopathological examination of multiple meningiomas reveals similar features as in solitary type, however simultaneous occurrence of different grade of malignancies is seen in 1/3rd of patients [6]. The common histological types in multiple meningiomas are- psammomatous, fibroblastic, meningoeipithelial and transitional types [7]. In our case the microscopy revealed a psammomatous meningiomas grade I in all the lesions. Clinically differential diagnosis of multiple meningiomas includes intracranial metastases. Radiological examination helps in detecting the location of these tumors. MRI is more helpful than CT to detect the exact site of tumors. Surgery is the treatment of choice for the multiple meningiomas. Surgical excision is
done in the case of symptomatic meningiomas and asymptomatic meningiomas more than 3 cm in size [8]. The prognosis of multiple meningiomas is same as for solitary one.

**CONCLUSION**

Multiple meningiomas are rare and are commonly observed in female patients. Clinically they can mimic intracranial metastasis and histopathological examination helps in making differential diagnosis.

**REFERENCES**

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