

DOI: 10.5455/2320-1770.ijrcog20140346

Case Report

Pregnant patient presenting with headache and medulloblastoma and hydrocephalous: a case report

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Received: 12 November 2013

Accepted: 9 December 2013

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ABSTRACT

Medulloblastoma, a primitive neuroectodermal tumor (PNET), is a second most brain tumor in children, accounting for about 20% of cases but is very rare in adults, 1% of all cancers in adults with only one third of cases occur in women and even less in pregnant women. It is primarily a cerebellar tumor. Symptoms include nausea, vomiting, loss of coordination, morning headache, seizures and personality changes. Treatment is surgical removal of tumor followed by radiotherapy. A grand-multipara presented in casualty with 30 weeks pregnancy & complains of headache, visual disturbances & disorientation. MRI findings were suggestive of medulloblastoma with hydrocephalus. Ventriculo-peritoneal shunting was done followed by termination of pregnancy. She delivered vaginally a live female child. After 15 days, her definitive neurosurgery was done. The uniqueness of this case is its rarity and symptoms of which are often masked by symptoms of pregnancy as syncope during pregnancy could be due to hypoglycemia, anxiety, orthostatic hypotension or cardiac causes.

Keywords: Medulloblastoma, Hydrocephalus, Pregnancy

INTRODUCTION

Medulloblastoma, a primitive neuroectodermal tumor (PNET), is a second most brain tumor in children, accounting for about 20% of cases but is very rare in adults (1% of all cancers in adults with only one third of cases occur in women and even less in pregnant women).¹ It is primarily a cerebellar tumor. On a hematoxylin and eosin stain, it appears as a small blue cell tumour. Symptoms include nausea, vomiting, loss of coordination, morning headache, seizures and personality changes. Medulloblastomas can metastasize to placenta during pregnancy, to spine and might recur in subsequent pregnancy. Treatment is surgical removal of tumor followed by radiotherapy.

CASE REPORT

A previously healthy 28 years old female G5P4 presented in Gynae OPD in Chirayu Medical College with 30 weeks pregnancy and complaints of headache, visual disturbances, vomiting dizziness and decreased mentation since 20 days. She was subjected to various investigations after neurosurgical opinion. Her MRI showed space occupying lesion in cerebellar region with hydrocephalous which was most likely medulloblastoma. Her USG showed alive 30 weeks pregnancy with mild oligohydraminos. Neurosurgeon planned urgent ventriculo-peritoneal shunt to relieve her compressive symptoms. After discussion with relatives and with their consent, explaining them fetal and maternal morbidity, patient underwent shunt surgery and termination of pregnancy so that definitive surgery could be planned thereafter. She delivered alive female child uneventfully

and child was discharged from NICU after 7days. After 15 days patient was posted for suboccipital craniotomy and surgical debulking. Patient is under close follow up and radiotherapy is planned.

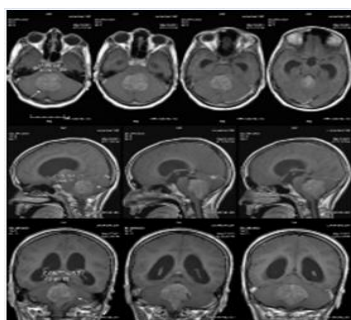


Figure 1: CT picture showing tumor in cerebellar region with hydrocephalus.

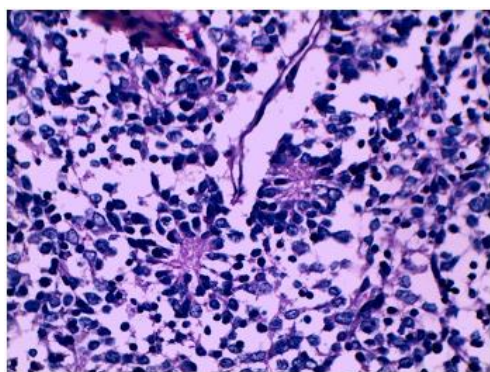


Figure 2: Histopathology picture showing blue cell tumor in rosettes.

DISCUSSION

Medulloblastoma is a cerebellar granule cell tumor, the most common cell type of cerebellum, and its robust proliferative capacity is thought to be regulated by sonic hedgehog pathway.² However, data suggests that cerebellar granule cells may be responsive to estrogen receptor stimulation. As such, pharmacologic blockade of estrogen receptors in cell cultures and xenograph models of medulloblastomas have shown to inhibit the migration and growth of medulloblastoma.³ These data suggest that a high levels of estrogen observed during pregnancy may be a contributing factor for this tumor in pregnancy.⁴ The uniqueness of this case is its rarity and symptoms are

often masked by symptoms of pregnancy as syncope during pregnancy could be due to hypoglycemia, anxiety, orthostatic hypotension or cardiac causes.⁵ In addition, the radiological report and gross features of tumor may be mimicked by other tumors as astrocytomas and meningiomas.^{6,7}

Funding: No funding sources

Conflict of interest: None declared

Ethical approval: Not required

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DOI: 10.5455/2320-1770.ijrcog20140346

Cite this article as: Sharma S, Tripathi A.

Pregnant patient presenting with headache and medulloblastoma and hydrocephalous: a case report. Int J Reprod Contracept Obstet Gynecol 2014;3:229-30.