Case Report

Diagnosis, Treatment, and Management Strategy of Meningioma during Pregnancy

Abstract

The most common benign tumor of the brain is meningiomas. Usually diagnosed between the ages of 40–60, they are more common in women. Studies have shown a strong relationship between hormones and malignancies. Although meningiomas are slow-growing tumors of the brain, pregnancy seems to induce its growth speed. Studies concerning meningiomas and hormone relationship may explain the reason why symptoms during pregnancy flare. More specifically, the estrogen and progesterone receptor may take an active role through signal transduction in inducing the growth of the tumor. Thus, the dilemma of pregnancy + meningioma arises. In this case, a 21-year-old pregnant woman with a giant meningioma diagnosed on the symptom of loss of sight is reported. Her pregnancy was terminated, and the tumor was excised. Her vision improved and the histopathological examination showed a progesterone receptor positive meningioma. It is a challenging decision to be made by the physician, the patient and the family when deciding if and when pregnancy should be terminated once an intracranial meningioma is diagnosed.

Keywords: Brain tumor, meningioma, pregnancy, termination, vision disorders

Introduction

Meningiomas are the most common benign and most common nonglial tumor of the brain. They are usually seen between the ages of 40–60 and are more common in postmenopausal women. 90% of meningiomas are benign, 6% are atypical, and 2% are malignant.[1] They orginate from arachnoid cap cells and may be diagnosed on symptoms or incidentally.[2] Infracranial malignancies are uncommon during pregnancy. According to Isla et al. during 12 years period, of 126,413 pregnant women 12 were diagnosed with intracranial tumors and only 2 of them were a meningioma.[3] However, it is believed that a strong relationship exists between malignancies in women and hormones.[4] Meningiomas are twice as common in women as men.[5] Meningiomas have been shown to grow faster during the luteal phase of the menstrual cycle and pregnancy.[6] Meningiomas have been shown to grow in size and become symptomatic during pregnancy. Among the reasons believed to flare the symptoms are water retention, engorgement of vessels, and the presence of sex hormone receptors on tumor cells leading to the growth of the tumor. Raised intracranial pressure may be misdiagnosed as hyperemesis gravidarium.[7] It is a difficult dilemma both for the physician and the patient in deciding whether the pregnancy should be terminated. Herein is described a case of a giant meningioma causing loss of sight during pregnancy.

Case Report

A 21-year-old female patient who had been suffering from a headache for about 2 years applied to an ophthalmologist with the loss of sight for the last 6 weeks. On her cranial magnetic resonance imaging (MRI), an intracranial tumor had been diagnosed and was referred to the neurosurgery clinic. Her cranial MRI showed a mass measuring approximately 45 mm *70 mm at the vertex level localized within the interhemispheric fissure, T1-weighted gadolinium-enhanced MRI images showed a characteristic dural tail sign, containing perifocal edema and compressing corpus callosum and the lateral ventricles [Figure 1]. There was no history of seizures and no significant neurological examination finding in the patient except a decrease in visual acuity. Due to the fact that the patient was in her reproductive age a β-human chorionic gonadotropin test was applied to an ophthalmologist with the loss of sight is reported. Her pregnancy was terminated, and the tumor was excised. Her vision improved and the histopathological examination showed a progesterone receptor positive meningioma. It is a challenging decision to be made by the physician, the patient and the family when deciding if and when pregnancy should be terminated once an intracranial meningioma is diagnosed.
analysis was ordered and found to be high (2598 mIU/ml N: 0–5.3). Being G2P1Y1 a pregnancy was possible thus she was consulted to the obstetrics and gynecology clinic. She was late to her last menstruation cycle for 5 weeks and 6 days, and her obstetric ultrasonography revealed a fundal localized gestational sac measuring approximately 6 * 8mm corresponding to a gestational age of 5 weeks. She was then evaluated multidisciplinary with a neurosurgeon, ophthalmologist, obstetrician, and anesthesiologist. With the consent of the family, her pregnancy was terminated, and the patient was transferred to our clinic.

On her careful examination, she was found to have bilateral homonymous superior quadrantanopia. She was then operated with bifrontal craniotomy and the mass was excised totally preserving the superior and inferior sagittal sinus. She was then extubated and taken to the neurosurgery intensive care unit for postoperative follow-up period. After 24 h the surgery her neurological examination was normal, and she was transferred to the ward and discharged with no postoperative complication on the fourth day. The pathological examination of the mass was reported as meningioma WHO Grade I.

On her first year follow-up, she had no symptoms, her sight had improved totally and the control cranial MRI showed no residue mass [Figure 2]. On the histological examination of the tumor, progesterone receptors were found to be positive leading to think that pre-existing asymptomatic meningiomas in women may flare during pregnancy due to increased hormone levels and become symptomatic. A year later, she became pregnant.

**Discussion**

Meningiomas which arise from arachnoid cap cells are usually found in the skull base and perivenous sinuses, where these cells are abundant. Even though most cases of meningioma are sporadic, people with neurofibromatosis-2 gene and exposure to ionizing radiation are at a higher risk. However, the role of sex hormones is yet unclear. Bernard was the first to diagnose a meningioma during pregnancy in 1898. The fact that intracranial meningiomas are twice as common in women as men, 10-fold increased risk of spinal meningioma in women than men, increase in the size of meningioma during pregnancy, commonly diagnosed with breast cancer in women and containing estrogen and progesterone receptors[4,10,11] leads to the conclusion that there may be a relationship with pregnancy and meningioma. These tumors tend to grow slowly however pregnancy seems to speed up this process creating symptoms.

This case presented with a bilateral homonymous superior quadrantanopia. Even though there was no direct compression of the optic chiasm, which would normally explain this defect, there are cases in the literature where parasagittal and interhemispheric meningiomas cause disturbance of the visual field.[12] The lesion in this case with a massive size compressed the ventricles superiorly causing a downward shift along with increased intracranial pressure findings such as a headache causing a probable compression of the optic chiasm or optic nerve.

80% of women and 34% of men with meningiomas were found to have progesterone receptors.[13] These receptors have been shown to take an active role in the growth of the tumor.[14] However unlike the ras system these receptors do not use a direct transcription pathway but a unique sequence activated by the binding of progesterone to its receptor.[15] Unfortunately in vitro studies have shown these receptors to disappear after 2 or 3 passages. Also, the use of anti-estrogen drugs in studies has not yielded promising results. At the same time, meningiomas have been shown to contain platelet derived growth factor,[16] vascular endothelial growth factor (VEGF),[17] glucocorticoid,[18] and epidermal growth factor[19] receptors. Aside from its location, size, and venous invasion, the presence of VEGF receptors determines the length of the perifocal edema.[20] Prolactin receptors have also been found to be positive in meningiomas and closely associated with the growth rate.[21,22] Rich in prolactin receptors, meningiomas are more likely to grow during and after pregnancy causing physiological pituitary hyperplasia elevating diaphragm sellae compressing the optic nerve and chiasm causing loss of vision.

![Figure 1: Preoperative T1-weighted gadolinium enhanced magnetic resonance imaging, axial view (a) and coronal view (b) showing a large interhemispheric meningioma compressing the lateral ventricles](image1)

![Figure 2: A year after the surgery, T1-weighted gadolinium enhanced magnetic resonance imaging axial view (a) and coronal view (b) showing no remnant mass](image2)
Although radiotherapy is a treatment modality, surgery is usually preferred for the treatment of meningiomas. This decision is based on symptoms, age, radiological features, postoperative morbidity, patient preference and when a definite diagnosis is necessary. For this patient, we used the Cleveland Clinic Comorbidity, Localization, Age, Size, Symptoms algorithm. The meningioma in our patient measured 45 mm × 70 mm, contained perifocal edema and was causing visual symptoms. There is no algorithm yet present for handling meningiomas during pregnancy. After the termination of the pregnancy, surgical excision of the tumor yielded positive results. Unlike the 18 meningiomas operated during pregnancy by Kanaan et al., this patient’s visual symptoms improved completely after 1-year.

If a meningioma is diagnosed during pregnancy the patient herself and the age of the fetus must be evaluated first. An operation decision should be decided upon multidisciplinary taking into consideration of first the patient, second the fetus and last the decision of the family. A surgical excision without the termination of the fetus must be handled carefully, taking into consideration of intraoperative blood loss, hypotension, hypovolemia, and hypoxia. Corticosteroids may be safely used during per operative and per partum period. 2–4 mg dexamethasone every 6 h is usually the preferred choice. Since mannitol is able to pass through the maternal fetal barrier it should only be preferred under urgent conditions. Monotherapy should be the preferred therapy for seizures. It should be kept in mind that complications of seizures are more serious than the side effects of antiepileptics. Women with history of seizures should take folic acid vitamin-K1 at the early stages of pregnancy to prevent the risk of neural tube defect.

A meningioma diagnosed during pregnancy causes the surgeon to ponder whether to perform surgery immediately or wait for the end of the pregnancy. If an emergent situation is at hand, such as wide spread edema, midline shift, change in consciousness, paresis, acute hydrocephalus, or acute total neurological deficit such as loss of vision, a dual surgery along with the excision of the tumor is usually the preferred choice. Since mannitol is able to pass through the maternal fetal barrier it should only be preferred under urgent conditions. Monotherapy should be the preferred therapy for seizures. It should be kept in mind that complications of seizures are more serious than the side effects of antiepileptics. Women with history of seizures should take folic acid vitamin-K1 at the early stages of pregnancy to prevent the risk of neural tube defect.

Conclusion

Studies have shown the growth rate of meningiomas during pregnancy accelerates. However even though a strong relationship between hormones and meningiomas exists, further research must show a definite effect of hormones and hormone therapy for the treatment of these tumors. Theoretically meningiomas in women arise from differentiated arachnoid cap cells sensitive to female sex hormones. Studies up to date need to remind the surgeon that women with meningioma during their fertility period must be evaluated for pregnancy and if pregnancy is the case the patient must be evaluated multidisciplinary. If the patient suffers from increased intracranial pressure symptoms, a risk of herniation or neurological deficit, then the termination of the fetus must definitely be evaluated. Losing precious time at this critical stage in deciding whether termination should be performed may alter the benign course of meningioma causing prolonged or permanent neurological deficits. When neurological symptoms arise and if an obstetrician would not recommend otherwise, termination of the pregnancy would be more beneficial to the patient. In addition, the presence of headache, vomiting and seizure during pregnancy should not be diagnosed as hyperemesis gravidarum without a detailed neurological examination. If there is any suspicion of an intracranial tumor, abnormal fundoscopic examination, visual deficit, focal seizures or lateralizing finding a cranial MRI must be ordered.

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References