Case Report

Multiple meningiomas in two male-to-female transsexual patients with hormone replacement therapy: A report of two cases and a brief literature review

Rahul Raj, Miikka Korja, Päivi Koroknay-Pál, Mika Niemelä

Department of Neurosurgery, University of Helsinki and Helsinki University Hospital, Helsinki, Finland

E-mail: *Rahul Raj - rahul.raj@hus.fi; Miikka Korja - miikka.korja@hus.fi; Päivi Koroknay-Pál - paini.koroknay-pal@hus.fi; Mika Niemelä - mika.niemela@hus.fi

*Corresponding author

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Abstract

**Background:** Exogenous sex hormones may play a role in meningioma development and growth. Thus, transsexual patients being on long-standing hormone replacement therapy (HRT) may be at particular risk for meningioma development and growth. Here we present two cases of two male-to-female transsexual patients taking HRT for an extended period of time with both patients requiring surgical treatment at our institution due to multiple growing meningiomas.

**Case Description:** The first patient was a 50-year-old genetic male (male-to-female transsexual) who presented 7 years after an extensive sex-change operation due to progressive bitemporal visual defects. The patient had been on HRT for approximately a decade. Radiological examinations showed a total of four meningiomas, one being a large suprasellar meningioma causing the symptoms. Three of the four meningiomas were operated on, but the patient’s vision could not be saved. Immunohistochemical (IH) analysis of the meningiomas showed both estrogen and progesterone receptor expression. The patient continued HRT and during follow-up regrowth of the meningiomas was noted. The second patient was a 48-year-old genetic male (male-to-female transsexual) who had been on HRT for two decades and also presented due to left-sided visual loss. Radiological examination showed four meningiomas, one being a left-sided sinus cavernous meningioma causing compression of the visual apparatus. This patient had a previous normal head computerized tomography scan dating back 10 years before his current presentation. Three of the four meningiomas were operated on with a slight improvement in visual acuity. IH analysis showed positive progesterone receptor expression but negative estrogen receptor expression.

**Conclusion:** Radiological screening may be justifiable for transsexual patients with a history of long-standing HRT and special focus should be paid to transsexual patients displaying new neurological symptoms or those already diagnosed with a meningioma.
INTRODUCTION

The role of sex hormones as a risk factor for meningioma development and growth has been a topic of debate. Several factors such as female gender, exposure to ionizing radiation, and familial cancer predisposition syndromes (e.g., neurofibromatosis type II) have been identified to be associated with an increased risk of meningiomas. A few previous case reports have described rapid development and growth of meningiomas in transgender individuals taking sex-hormone replacement therapy (HRT). On the other side of the spectrum, regression meningiomas after discontinuation of HRT have also been described.

Here we report two further cases of transgender patients with long-standing HRT developing several meningiomas and a brief overview of the literature on this topic.

CASE REPORTS

Case 1
A 43-year-old genetic male underwent a male-to-female sex-change operation. The patient had been taking HRT for several years before the sex-change operation. Following surgery, the patient continued HRT, consisting of estradiol cream (0.6 mg/g) twice a day and cyproterone acetate (Andocur®) 50 mg once a day.

This patient presented 7 years later with progressive bilateral visual compromise leading to complete left-sided vision loss. Clinical examination showed normal visual acuity on the right side (20/20). The left eye could only perceive light sensation, and clinical fundus examination revealed left-sided optic disk atrophy, with a dilated pupil and a limited direct pupillary light constriction reflex.

The patient was sent to neuroimaging and magnetic resonance imaging (MRI) showed a total of four meningiomas: (1) a suprasellar meningioma compressing the left optic nerve and chiasm causing the patient’s symptoms, (2) a small frontobasal midline meningioma at the level of cribriform plate, (3) a small frontal right-sided convexity meningioma, and another (4) large frontal right-sided convexity meningioma [Figure 1, upper row]. The patient was put on dexamethasone before the operation.

Meningiomas 1, 2, and 4 (suprasellar, frontobasal midline, frontal right-sided convexity) could all be surgically accessed in one single session with adequate results (Simpson grade III for suprasellar and frontobasal meningiomas and Simpson grade I for convexity meningioma). The histological diagnosis in all these meningiomas revealed World Health Organization (WHO) grade I tumors with low proliferative index (MIB-1 <5%). Further immunohistochemical (IHC) profiling revealed a high level of progesterone receptor expression and lower level expression of estrogen receptors.

Immediately after surgery, the limited function that had been present in the left eye remained unchanged, but unfortunately the visual acuity of the right eye acutely deteriorated (to light perception only). Postoperative head computed tomography (CT) showed no intracranial hemorrhage. The patient was taken back to the operating room for exploration. Intraoperative finding showed acute swelling of the right optic nerve, and a right-sided optic canal decompression and optic nerve sheath fenestration were performed. Unfortunately, vision could not be restored in the right eye and the patient became legally blind since both eyes suffered from compromised vision. After close inspection of anesthesiology reports, no hypotensive insult before or during the first or second operation was noted.

Postoperative MRI showed complete removal of (1) suprasellar meningioma, (2) frontobasal midline meningioma, and (4) frontal right-sided convexity meningioma [Figure 1, lower row]. Clinical examination did not show any improvement in visual function and the patient developed bilaterally dilated pupils without any pupillary reaction to light. This was corroborated by bilateral optic atrophy and bilateral blindness. The remainder of the postoperative course was uneventful, and HRT was continued.

Key Words: Estrogen, hormone replacement therapy, meningioma, progesterone, transsexual

Figure 1: Upper row shows the pre-operative magnetic resonance imaging (MRI) showing four meningiomas: (1) a suprasellar meningioma, (2) a small frontobasal midline meningioma in front of the suprasellar meningioma, (3) a right-sided midline front convexity meningioma, and a (4) large right-sided frontal convexity meningioma. Lower row shows the 1-week postoperative MRI where meningiomas (1), (2), and (4) have been totally removed. Numbers 1–4 refer to meningiomas as mentioned in the main text.
The patient was followed with sequential imaging, and 1 year after surgery a repeat MRI showed no regrowth of resected meningiomas but slight growth of small frontal right‑sided midline convexity meningioma (3) that had been left untouched [Figure 2, upper row]. HRT was continued unchanged.

At 2 years, postoperative MRI showed regrowth of a (5) new suprasellar meningioma and further growth of unoperated frontal right‑sided midline convexity meningioma (3). The patient was managed expectantly and HRT was continued unchanged [Figure 2, middle row].

Four years after operation, another MRI showed significant growth of (5) suprasellar meningioma and (3) of the previous unoperated right‑sided midline frontal convexity meningioma [Figure 2, lower row]. The patient’s neurological status has not changed, and the patient did not develop any new deficits. This prompted a discussion with the patient regarding the safety of his HRT, but the patient remained determined to continue HRT. Thus, due to progressive growth of meningiomas, a second operation has been scheduled.

Case 2
This is a 48‑year old male‑to‑female transsexual (genetic male), who had been taking HRT for 21 years before neurosurgical presentation. The HRT consisted of estradiol cream (1 mg/g) three times a day and cyproterone acetate (Andocur) 50 mg once a day.

The patient presented with progressive left‑sided vision loss. Clinical examination revealed near‑normal visual acuity in the right eye (20/20) but extremely compromised visual acuity in the left eye (20/250) with left‑sided optic atrophy on fundus examination. A brain MRI was obtained that revealed four meningiomas: (1) in the left‑sided sinus cavernous, (2) over the left‑sided frontal convexity, (3) over the left‑sided temporal convexity, and (4) right‑sided tentorial [Figure 3, upper row].
Moreover, the patient had undergone a contrast-enhanced head CT 10 years before the current presentation due to transient vertigo and the respective CT was assessed as completely normal.

Three of the meningiomas (sinus cavernous, left-sided frontal, and temporal) were operated on and the left optic nerve was decompressed in the same surgical sitting at about 5 months after symptom had started. Surgery was successful and achieved Simpson grade I for convexity meningiomas and Simpson grade IV for sinus cavernous meningioma. Postoperative MRI showed complete removal of convexity meningiomas and a reduction in the size of sinus cavernous meningioma [Figure 3, lower row]. Histopathological diagnosis revealed all meningiomas to be of WHO grade I (MIB-1 <5% in all). IH analysis showed positive staining for progesterone receptor expression but was negative for staining of estrogen receptor expression. Immediately after surgery, the patient experienced subjective visual improvement. The remainder of the postoperative course was uneventful.

It was recommended to the patient to taper down the HRT, but long-term follow-up is still lacking.

**DISCUSSION**

Here we report our observations of long-term use of HRT and the growth of multiple meningiomas in two man-to-female transsexual patients. In the first patient, growth of several meningiomas was noticed (over a 4-year follow-up following removal) and it seems probable that this is related by the use of exogenous HRT. Since the patient had not undergone any brain imaging before their clinical presentation, we cannot demonstrate the causal role of HRT and risk of meningioma development. However, given the receptor expression profile of the pathological specimen, this explanation appears intuitive. The link between exogenous HRT and meningioma is further supported in the second case, by the fact that the patient had undergone a contrast-enhanced head CT scan 10 years before the discovery of his meningiomas, and that CT was read out as completely normal. Thus, it seems likely that the exogenous HRT had a role in the development and growth of these patients’ multiple meningiomas.

Epidemiological studies have revealed that females are more likely to develop meningiomas in the second half of life, and due to the fact that many meningiomas may express progesterone and estrogen receptors, it has been hypothesized that both endogenous and exogenous sex hormones may increase the risk for meningioma development and growth.\(^{2,6,7,11,18}\) A study showed that up to 88% of all meningiomas express progesterone receptors and about 40% of tumors express estrogen receptors, with no differences in gender of the afflicted patients.\(^{12}\) The first patient’s pathological specimen was positive with regard to both estrogen and progesterone receptor expression, whereas the second patient’s pathological specimen was positive only with regard to progesterone receptor expression. Thus, it seems that meningioma sex-hormone receptor expression per se does not account alone for the noted sex differences in meningioma risk. Rather, it might be the continuously raised levels of sex hormones that directly affect the target tissue and induce meningioma growth. This is supported by the fact that in the experimental setting antiprogesterone treatment reduces meningioma volume regardless of progesterone-receptor expression status.\(^{16}\)

Previous studies have shown an association between HRT and risk of meningioma occurrence in post-menopausal women, especially when taking estradiol supplements only.\(^{2,3,11}\) Androgen deprivation therapy in male patients with prostate cancer has also been linked to an increased development of meningiomas.\(^{13,14}\) Although we cannot show definite proof, it seems plausible that there is an association between HRT and risk of meningiomas\(^{15,17}\) and it appears to naturally fit this concept that meningioma risk would be increased in male-to-female transsexual patients with long-standing HRT. The real incidence of meningiomas in male-to-female transsexuals remains, however, unknown. This scenario might be an under- or overdiagnosed problem. A previous study from the Netherlands found that only 3 of 2810 male-to-female transsexual patients were diagnosed with meningiomas.\(^{17}\) The authors further estimated the incidence of meningiomas in male-to-female transsexuals to be 126/100,000 (which appears to be higher than expected when compared with the observed incidences.
The use of exogenous sex hormones may play a role in the development and growth of meningiomas. The risk may be dose-dependent, which would suggest that risk is especially high in male-to-female transsexuals taking high doses of exogenous sex hormones over long periods. Radiological screening may be justifiable for transsexual patients with a history of long-standing high-dose exogenous sex-hormone therapy and special focus should be paid to transsexual patients displaying new neurological symptoms or those already diagnosed with meningiomas. Still, further case-control matched epidemiological studies are necessary to determine the real risk of meningioma in transsexual persons.

**CONCLUSION**

The use of exogenous sex hormones may play a role in the development and growth of meningiomas. The risk may be dose-dependent, which would suggest that risk is especially high in male-to-female transsexuals taking high doses of exogenous sex hormones over long periods. Radiological screening may be justifiable for transsexual patients with a history of long-standing high-dose exogenous sex-hormone therapy and special focus should be paid to transsexual patients displaying new neurological symptoms or those already diagnosed with a meningioma.

**Declaration of patient consent**

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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**Conflicts of interest**

There are no conflicts of interest.

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