

Baerveldt implant for secondary glaucoma due to iris melanoma

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Background: Proton beam therapy (PBT) is effective in the treatment of iris melanoma. Reported complications after PBT are radiation-induced cataract and raised intraocular pressure (IOP). Filtering glaucoma surgery has generally been avoided because of fears of seeding.

Case report: A 37-year-old man presented with a self-discovered, pigmented lesion on his right iris. Four years later, the pigmented lesion was diagnosed as an iris melanoma, because of documented growth. The patient was treated with PBT but developed secondary glaucoma one month later. The IOP could not be controlled despite maximal medical therapy and selective laser trabeculoplasty (SLT). Finally, Baerveldt implant surgery was performed, resulting in an IOP lowering to 10 mmHg and stabilization of the glaucomatous visual field loss.

Conclusion: Our case demonstrates that Baerveldt implant surgery is a reasonable therapy for glaucoma following successful radiotherapy of iris melanoma.

Keywords: iris melanoma, proton beam therapy, secondary glaucoma, Baerveldt implant surgery

Introduction

Iris melanoma comprises 3% to 10% of all uveal melanomas and is the most common primary malignancy of the iris.^{1,2} It tends to have a ten-year metastasis rate of 3% to 6%.¹⁻³ Possible therapies for iris melanoma include iridectomy, iridocyclectomy, plaque brachytherapy, proton beam therapy (PBT), and enucleation. PBT is generally selected if the tumor is too extensive for surgical excision or if such surgery is rejected because the surgical iris coloboma is expected to cause unacceptable photophobia or cosmetic deficit. The main complications after PBT are radiation-induced cataract and raised IOP. Trabecular scarring may play a role in the latter.⁴ Secondary glaucoma occurs in 7% of eyes with untreated iris melanoma and in 30% of patients with microscopically-confirmed iris melanoma, occurring mostly because of tumor infiltration of the trabecular meshwork with outflow obstruction.^{5,6} Traditionally, filtering glaucoma surgery has been avoided in patients with iris melanoma because of fears that such aqueous drainage might encourage subconjunctival or intraorbital tumor seeding.^{2,3} Nevertheless, insights into the biology of uveal melanomas and outcome studies after PBT encouraged us to treat a patient with a Baerveldt tube implantation because of rapidly deteriorating vision and uncontrollable glaucoma.

Case report

A 37-year-old man discovered a pigmented iris lesion in his right eye. The best-corrected visual acuity (BCVA) was 6/6. Ophthalmologic examination showed a

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pigmented iris lesion between the 5.30 and 6.30 o'clock meridians with a basal diameter of 3.5 mm and a thickness of 1.4 mm. Gonioscopy showed pigment deposition in the inferior and nasal parts of the iridocorneal angle. The IOP was 25 mmHg. Ultrasonography showed no involvement of the ciliary body. The left eye was normal. Initially, the tumor showed no growth; however, four years after presentation, the iris lesion had changed to a diffuse tumor, extending from the 4.30 to 6.30 o'clock meridians with seeding onto the iris surface from the 3.30 to 7.00 o'clock meridians. The pupil also became oval (Figure 1a). Gonioscopy showed pigment deposition in the iridocorneal angle between the 1.00 and 10.00 o'clock meridians. The BCVA decreased to 6/7.5. The patient refused a biopsy for histological examination. Because of documented growth, the tumor was diagnosed as an iris melanoma.

The entire anterior segment was treated with PBT (53.1 Gy, administered in four fractions over four days). One month later (Figures 1b and 1c) the patient developed secondary glaucoma with IOP levels fluctuating between 20 and 43 mmHg, despite maximal medical therapy. The optic disc showed normal cupping.

SLT was performed, placing 25 shots with a total energy of 15 mJ in the nasal quadrant. The IOP decreased slightly but only transiently. Further treatment of the temporal quadrant did not lead to a reduction of IOP. Meanwhile, the optic disc cupping had become pathological and glaucomatous visual field loss had progressed from mild loss to very extensive loss within two months.

Since the patient was keen to preserve the remaining vision in this eye, despite advice about a possible risk of extraocular spread, we finally placed a Baerveldt tube in the anterior chamber (Figure 1d and Figure 2). The IOP decreased

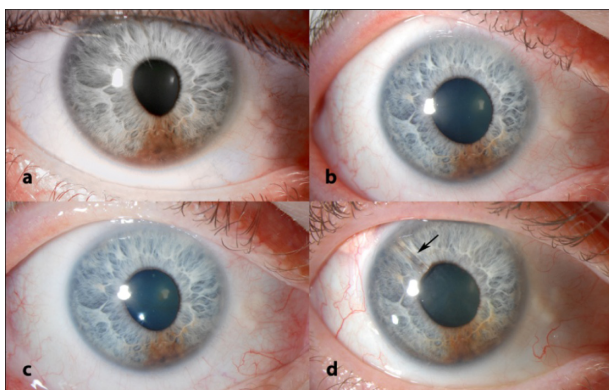


Figure 1 Slit lamp figures. Figure 1a shows the iris lesion when diagnosed as iris melanoma. Figure 1b shows the lesion after PBT. Figure 1c shows the lesion after PBT. Figure 1d shows the Baerveldt tube in the temporal/superior quadrant.

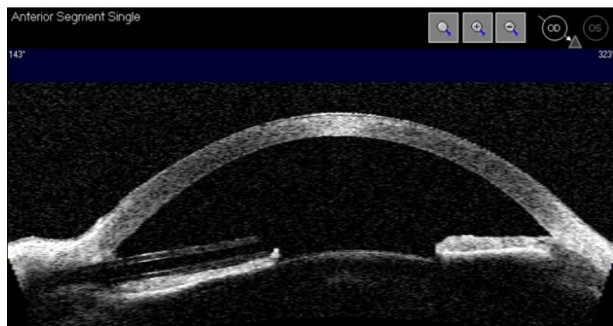


Figure 2 A Visante OCT scan showing the Baerveldt tube in the anterior chamber to the left.

to 13 mmHg. After one year of follow-up, the IOP was stable at 10 mmHg with the additional use of dorzolamide and timolol. The visual field showed no further deterioration. The BCVA was 6/15. The patient declined systemic screening for metastasis; however, he remains under intensive ophthalmic surveillance.

Discussion

Secondary glaucoma after PBT for iris melanoma can be difficult to control with medical therapy.⁷ We report successful lowering of the IOP in such a case by using a Baerveldt glaucoma implant.

To our knowledge this approach has not been reported previously. As a rule, drainage surgery is avoided after treatment of iris melanoma, probably because of concerns that tumor cells might seed through the drainage fistula and metastasize to other parts of the body. Therefore, we initially attempted to lower the IOP by SLT. Insights from genetic studies on uveal melanomas suggest, however, that these tumors metastasize almost exclusively if they show loss of chromosome 3⁸ or class II gene expression profile.⁹ There is growing evidence that metastasis starts at a very early stage, before the patient even presents to the ophthalmologist.¹⁰ For these reasons, there are now considerable doubts that glaucoma drainage surgery would enhance risks of metastasis spread to the rest of the body by providing an exit route from the eye.

Another concern is that the iris melanoma can recur and seed through the tube into the subconjunctival and orbital tissues. We consider these risks to be small, firstly, because local tumor recurrence is rare after PBT and, secondly, because the tumor was located far from the internal opening of the tube (Figure 1d). In any case, the patient is being monitored closely so that appropriate treatment can be administered without delay in case of re-growth of the tumor.

Further studies with more patients and long-term follow-up are indicated to evaluate the safety and efficacy of the Baerveldt glaucoma implant for the treatment of secondary glaucoma after PBT for iris melanoma.

Disclosures

The authors report no conflicts of interest in this work.

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