focus on infants particularly likely to develop ROP. However, any study of the effects of β-blocker therapy must address the fragility of the patients to be tested and possible systemic and ocular adverse effects. Nevertheless, if topical β-blockers prove to be effective in preventing some cases of ROP, this opens the door for a more individualized approach to prevention of the disease, eg, using β-adrenergic receptor polymorphisms to guide ROP management.

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**Abbreviation:** ROP, retinopathy of prematurity.

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### Epithelial Downgrowth After Type 1 Boston Keratoprosthesis Manifesting as Tractional Retinal Detachment and Epiretinal Membrane

**Type 1 Boston keratoprosthesis (KPro) is a viable treatment option for corneal disease at high risk for graft failure with traditional penetrating keratoplasty. Postoperative complications of Boston KPro include retroprosthetic membrane, glaucoma, sterile vitritis, infectious endophthalmitis, corneal melt, extrusion, and retinal detachment.** To our knowledge, we report the first case of epithelial downgrowth (ED) of the posterior segment after Boston KPro placement.

**Report of a Case.** A 52-year-old man with a history of penetrating ocular injury to his right eye had open globe repair and cataract extraction in 1974, placement of a secondary anterior chamber intraocular lens in 1992, 2 failed penetrating keratoplasty procedures in 2004 and 2008, Baerveldt glaucoma tube implantation in 2005, astigmatic keratotomy in 2007, and, most recently, intraocular lens removal and type 1 Boston KPro placement in 2010. Histologic examination of the failed corneal graft excised at the time of KPro placement did not demonstrate ED. One week after KPro placement, visual acuity was 20/40 OD. Two months after KPro placement, he had pain and photophobia in the right eye.

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**Table. All Consented Patients With Type 1 ROP in at Least 1 Eye**

<table>
<thead>
<tr>
<th>Birth Weight, g</th>
<th>Race</th>
<th>Consented Patients With ROP, No. (N = 2320)</th>
<th>Patients Developing Type 1 ROP, No. (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;750</td>
<td>Non–African American</td>
<td>623</td>
<td>186 (29.9)</td>
</tr>
<tr>
<td></td>
<td>African American</td>
<td>315</td>
<td>36 (11.4)</td>
</tr>
<tr>
<td>750-999</td>
<td>Non–African American</td>
<td>726</td>
<td>140 (19.3)</td>
</tr>
<tr>
<td></td>
<td>African American</td>
<td>233</td>
<td>18 (7.7)</td>
</tr>
<tr>
<td>≥1000</td>
<td>Non–African American</td>
<td>331</td>
<td>25 (7.6)</td>
</tr>
<tr>
<td></td>
<td>African American</td>
<td>92</td>
<td>1 (1.1)</td>
</tr>
<tr>
<td>Total</td>
<td>Non–African American</td>
<td>1680</td>
<td>351 (20.9)</td>
</tr>
<tr>
<td></td>
<td>African American</td>
<td>640</td>
<td>55 (8.6)</td>
</tr>
</tbody>
</table>

**Abbreviation:** ROP, retinopathy of prematurity.
At his visit to us, visual acuity was 8/200 OD and 20/20 OS. No afferent pupillary defect was identified. Intraocular pressure was soft to palpation OD and 14 mm Hg OS. Slitlamp examination revealed an intraprosthetic membrane (Figure 1A). B-scan ultrasonography revealed anterior vitreous bands, tractional retinal detachment (TRD), and a small choroidal detachment (Figure 1B). Review of B-scan ultrasonographic images performed 2 months previously, before KPro placement, showed a normal posterior segment. A 23-gauge pars plana vitrectomy was performed to repair the TRD. Pars plana vitrectomy revealed a dense anterior vitreous band contiguous with a TRD and epiretinal membrane (ERM) extending across the macula. Extensive membrane peeling was performed, followed by fluid-air exchange, endolaser, and silicone oil injection. A large ERM specimen from the macula was sent for histopathologic evaluation. Microscopic evaluation of the ERM revealed mucosal epithelium containing goblet cells on hematoxylin-eosin, periodic acid–Schiff base, and Masson-Trichrome staining, consistent with ED (Figure 2). Immunohistochemical analysis with cytokeratin showed positive staining, revealing epithelial cells lining one side of the ERM (Figure 2). At the patient’s 2-month follow-up, slitlamp examination revealed recurrent intraprosthetic membrane and flat retina by indirect ophthalmoscopy and optical coherence tomography.

Comment. Epithelial downgrowth of the posterior segment is rare but can manifest within months after eye surgery. McDonnell et al reported ED occurring 3 months after ruptured globe repair, lensectomy, and pars plana vitrectomy, wherein TRD and ERM were found to consist of nonkeratinized, stratified squamous epithelium. Our case demonstrates a similar rapid progression of ED into the posterior segment after surgery. B-scan ultrasonography documented a normal-appearing posterior segment before KPro placement. Two months after KPro placement, B-scan ultrasonography showed the large TRD due to ED.

Epithelial downgrowth after KPro placement has been a subject of historical interest. Girard observed that ED only occurred in early KPro models previous to 1972. The Boston KPro was approved by the US Food and Drug Ad-
Bilateral Uveal Effusion and Angle-Closure Glaucoma Associated With Bupropion Use

Bupropion hydrochloride, an aminoketone antidepressant, is a dopamine reuptake inhibitor with norepinephrine and nicotinic acetylcholine receptor antagonist actions. We report the first case to our knowledge of uveal effusion and bilateral angle-closure glaucoma associated with bupropion use.

Report of a Case. A 40-year-old healthy white woman with a history of depression had bilateral blurry vision starting the morning of her visit. Her ocular history was significant only for myopia. She reported excellent vision with −6.00–diopter sphere (DS) contact lenses prior to her visit. Her only medications were ibuprofen, last used 1 month prior, and bupropion hydrochloride, 100 mg 3 times a day, which she started 2 weeks prior. Ten years earlier, she had taken an uncertain dose of bupropion for an unknown duration without incident.

At her initial visit, visual acuity was 20/200 OD and 20/400 OS while wearing −6.00–DS contact lenses. Intraocular pressure was 35 mm Hg OU, both pupils reacted to light, and slitlamp examination revealed mild corneal edema and shallow anterior chambers bilaterally (Figure 1A and B). Gonioscopy revealed appositional angle closure bilaterally (Figure 1C and D). Fundus examination showed healthy nerves with a cup-disc ratio of 0.2 OU. A diagnosis of bilateral angle-closure glaucoma was made. Treatment was started in the emergency room with 1 dose of each of the following: pilocarpine hydrochloride, 1%, eye drops; timolol maleate, 2%/dorzolamide hydrochloride, 0.5%, eye drops; brimonidine tartrate, 0.15%, eye drops; latanoprost, 0.005%, eye drops; and oral acetazolamide, 500 mg.

The next day in the Glaucoma Service, intraocular pressure was 22 mm Hg OU. Ultrasound biomicroscopy showed bilateral choroidal effusions causing shallow angles (Figure 1E and F), and B-scan ultrasonography showed diffuse 360° of suprachoroidal hypoechoogenicity consistent with uveal effusions (Figure 1G and H). Autorefration demonstrated a myopic shift to −16.00 DS OU, supporting a diagnosis of bilateral angle-closure glaucoma with myopic shift secondary to uveal effusions. Bupropion, acetazolamide, and pilocarpine were discontinued, and treatments with prednisolone acetate and cyclopentolate hydrochloride eye drops were started.

Two days later, her visual acuity improved to 20/70 OD and 20/100 OS with contact lenses, her intraocular pressures normalized, and her angles were open. All treatments with eye drops were stopped. At 1 week, her examination findings normalized (Figure 2A-D) and repeat ultrasound biomicroscopy (Figure 2E and F) and B-scan ultrasonography (Figure 2G and H) showed complete resolution of the uveal effusions. One month later, her visual acuity was 20/20 OU while wearing −6.00–DS contact lenses. She started treatment with escitalopram oxalate for depression. Nine months later, her examination findings remained stable without effusions.

Comment. Drug-induced uveal effusions with resultant bilateral angle-closure glaucoma and myopic shift are uncommon but have been reported with a variety of medications, most notably sulfon-based medications such as topiramate. The mechanism for drug-induced uveal effusions is unclear. Some cases appear to be dose dependent as lower doses of the inciting medication may not