# **Surgical Technique**

# Ex-PRESS Shunt for Choroidal Fluid Drainage in Uveal Effusion Syndrome Type 2 A Potentially Novel Technique

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Uveal effusion syndrome (UES) was reported by Schepens and Brockhurst in 1963,<sup>1</sup> and later described as a nanophthalmic disorder with a scleral abnormality.<sup>2</sup> Gass<sup>3</sup> described idiopathic UES and

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hypothesized that the cause was a congenital anomaly of the sclera and vortex veins that was related to aging, hormonal changes, or impairment of the permeability of the sclera. Using histological methods, Trelstad et al<sup>4</sup> reported the scleral abnor-

malities in UES. In addition, Forrester et al<sup>5</sup> observed migration of retinal pigment epithelial cells into the subretinal space.

For the treatment of UES, Brockhurst<sup>6</sup> described good surgical results with decompression of the vortex veins by scleral resection with a sclerotomy. Gass<sup>3</sup> reported an effective surgical procedure with a sclerectomy and sclerostomy without decompression of the vortex veins owing to the difficulty of isolating the vortex veins. Uyama et al<sup>7</sup> treated 19 eyes of 16 patients with UES by making a two-third-thickness scleral flap and performing a scleral excision to expose the underlying choroid.

The Ex-PRESS shunt (Alcon Laboratories) glaucoma implant is a small (2- to 3-mm-long and 0.4-mm-diameter tube) stainless steel nonvalved device that was designed to lower intraocular pressure by shunting aqueous humor from the anterior chamber into the subconjuntival space under a partial-thickness scleral flap space adjacent to the limbus.<sup>8</sup>

In the current study, we report the surgical outcomes of the use of an Ex-PRESS shunt device for choroidal drainage without vortex vein decompression for the management of 3 eyes with UES.

# Procedures

A retrospective medical record review was performed of 2 consecutive patients (3 eyes) with UES without nanophthalmos who underwent Ex-PRESS shunt implantation at the Clinica de Ojos de Maracaibo, Maracaibo, Venezuela, from March 2012 through March 2014 (**Table**). Institutional review board/ethics committee approval was obtained from Clinica de Ojos de Maracaibo and the patients signed a standard informed consent. This study adhered to the tenets of the Declaration of Helsinki for research involving human participants.

# Surgical Technique

After peribulbar anesthesia and standard asepsis and antisepsis in the operating room, a conjunctival incision was performed, bare sclera was exposed in the quadrant with more accumulation of fluid according to ultrasonography, and an oblique sclerotomy was performed with a 25-gauge needle 13 mm posterior to the limbus (to decrease the risk for infection), at which time, a small amount of suprachoroidal fluid was spontaneously drained. The Ex-PRESS shunt (model P-50) was then inserted obliquely in the sclerotomy to facilitate the continuous drainage of the suprachoroidal fluid (eFigure 1 in the Supplement). Balanced salt solution was injected into the anterior chamber with a 30-gauge needle to reconstitute intraocular pressure. The conjunctiva was closed with 7-0 polyglactin 910 suture and subconjunctival antibiotics were injected.

# **Report of Patients**

#### Patient 1

A woman in her 60s presented to the emergency department with a 15-day history of a "shadow" that affected the upper part of the visual field in her left eye. She had undergone cataract surgery in both eyes 3 years prior to presentation. Her best-corrected visual acuity (BCVA) without any refractive error was 20/20 OD and 20/200 OS. Fundus examination revealed choroidal detachment and shifting subretinal fluid underneath the exudative retinal detachment of the left eye. Ultrasonography was performed demonstrating thick sclera (1.5 mm), choroidal detachment, and exudative retinal detachment (eFigure 2 in the Supplement). The axial length was 22 mm in the right

Table. Demographic and Baseline Characteristics, Functional and Anatomic Results, and Complications in Eyes With Ex-PRESS Shunt for Choroidal Drainage in UES

		UFS Duration.	Previous Surgery	BCVA		CD and SRF		Total FU.	
Eye No./Sex/Age, y	Diagnosis	d		Baseline	Last FU	Baseline	Last FU	mo	Complications
1/F/65	UES	14	Cataract surgery	20/200	20/30	Yes	None	24	None
2/F/55	UES	21	None	20/400	20/40	Yes	None	24	None
3/F/55	UES	14	None	20/200	20/40	Yes	None	12	None
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Abbreviations: BCVA, best-corrected visual acuity; CD, choroidal detachment; F, female; FU, follow-up; SRF, subretinal fluid; UES, uveal effusion syndrome.

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eye and 22.5 mm in the left eye. She was diagnosed as having UES in the left eye. Despite treatment with steroids (oral prednisolone, 50 mg/d), the retinal findings and BCVA did not improve. Ex-PRESS shunt drainage surgery was performed in the inferotemporal quadrant (eFigure 3 in the Supplement and Video) without vortex vein decompression without complications. Two days postoperatively, a flat retina was seen on fundus examination. The retina remained attached during the 24-month follow-up, and an improved BCVA of 20/30 was achieved. We did not observe subretinal fluid or choroidal effusion by ultrasonography after surgery at 24 months (eFigure 4 in the Supplement).

## Patient 2

A woman in her 50s was referred owing to decreased visual acuity in her left eye for 3 weeks' duration. She had been wearing glasses for hyperopia since childhood. Her BCVA was 20/25 OD and 20/ 400 OS, and her spherical equivalents were +1 diopters (D) and +1.50 D in the left and right eyes, respectively. Fundus examination revealed uveal effusion involving the macula in the left eye. Axial length was 21 mm in the left eye and 22 mm in the right eye on A-scan ultrasonography. B-scan ultrasonography demonstrated thick sclera (1.6 mm), an exudative choroidal, and retinal detachment.

An Ex-PRESS shunt was implanted in the inferotemporal sclera quadrant in the left eye. One week postoperatively, a flat retina was seen on fundus examination and BCVA improved to 20/40. The retina remained attached during the 24 months of follow-up. Six months later, she presented with impaired visual acuity in her right eye. Her BCVA was 20/200 and the intraocular pressure was 16 mm Hg in the right eye and 18 mm Hg in the left eye. Fundus examination demonstrated supraciliary effusion and exudative retinal detachment. Ex-PRESS shunt implantation was performed in the inferotemporal quadrant. The subretinal fluid and the choroidal effusion disappeared 48 hours after surgery and BCVA improved to 20/30. The retina and choroid have remained attached during 12 months of follow-up.

# **Ex-PRESS Shunt for UES**

Uveal effusion syndrome is difficult to manage and often follows a relapsing course. Our 3 cases (eyes) can be classified as type 2 according to the classification by Uyama and colleagues<sup>7</sup> because they were not associated with nanophthalmos or hyperopia but had a thick sclera. Although Brockhurst<sup>6</sup> described good surgical results with decompression of vortex veins, other investigators<sup>3</sup> have questioned the usefulness of this technique because vortex veins are difficult to isolate. Uyama et al<sup>7</sup> reported that sclerectomy with a small sclerostomy under the scleral flap could be effective in both type 1 and type 2 UES because the abnormal sclera and increased transscleral outflow of intraocular fluid are thought to be the main causes of these disorders. However, this technique was not effective in type 3 UES, which develops in non-nanophthalmic eyes with normal eyeball size and normal scleral thickness.

We believe that isolation of the vortex veins is complicated and that vortex vein decompression is technically difficult to perform without complications such as vein rupture. Therefore, we elected to perform the Ex-PRESS shunt technique instead of full-thickness sclerotomy or scleral windows because it seemed likely to have a sufficient effect for resolving UES. We suspect that our technique may work by facilitating a constant drainage of uveal exudation from the suprachoroidal space to the subconjuntival space.

The actual surgical technique and implantation of the device are very simple. The tip of the Ex-PRESS shunt is not sharp and it is inserted obliquely so the chance to cause retinal damage is unlikely. In addition, considering the reported persistent and recurrence rates (23%-50%) for type 2 UES,<sup>7,9,10</sup> it seems that the cost may be worthwhile. Compared with other surgical procedures, our Ex-PRESS shunt technique could simplify the operative procedure, is less invasive, and reduces surgical time.

Limitations of our report include that we cannot know the longterm benefits and risks from 3 eyes and limited follow-up. At this point, we know that the procedure is feasible and safe for a limited time.

## ARTICLE INFORMATION

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