

Removal of Densiron-68 with a 23-gauge transconjunctival vitrectomy system

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Abstract

Purpose To report a new approach for removal of Densiron-68 via pars plana with a 23-gauge transconjunctival sutureless vitrectomy system (TSVS).

Methods Prospective, interventional case series. Ten eyes (4 phakic, 5 pseudophakic, 1 aphakic) of 10 patients underwent Densiron-68 (1480 mPa viscosity and 1.06 g/ml specific gravity) removal via pars plana with a suction pressure of 600-mmHg vacuum through a short 23-gauge silicon cannula.

Results Densiron-68 was completely removed from all eyes. Retinal reattachment was achieved in all cases. The intraocular pressure was 20.9 (SD 3.5) mmHg at baseline, 12.2 (SD 4) mmHg at day 1 postoperatively, and 13.6 (SD 2.9), 15.4 (SD 2.5), and 16 (SD 1.8) mmHg after 1 week, 1 month, and 3 months, respectively. Five eyes needed suture of at least one sclerotomy. Postoperative hypotony (≤ 8 mmHg) was seen in 2 out of 10 eyes (20%). No additional postoperative procedure was necessary.

Conclusions Active removal of Densiron-68 with a 23-gauge short cannula is a simple, innovative, and safe technique that can help reduce surgical trauma.

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Introduction

Densiron-68 (Fluoron, Neu-Ulm, Germany) is an intraocular tamponade used for inferior retinal tears owing to its specific gravity being

greater than water. Over the past few years it has gained increasing popularity, thanks to more and more encouraging anatomical and functional results.^{1,2} The drawback of the use of Densiron-68 may be related to the difficulty of its removal. The gravity acting on the bubble, the force of adhesion between Densiron-68 and the retina, and its viscosity may cause special problems during its removal.³ We report a 23-gauge transconjunctival technique for active removal of Densiron-68 through a short 23-gauge cannula.

Patients and methods

Ten consecutive eyes underwent active removal of Densiron-68 via pars plana, using a 23-gauge transconjunctival sutureless vitrectomy system (TSVS). Densiron-68 was removed 102 (SD 48) days after the initial surgery for RRDs with inferior retinal breaks (between 4 and 8 clock hours). The duration of critical time period for removal of oil differs from case to case. We removed Densiron-68 according to the retina status and clinically significant amount of emulsification.

Complete ophthalmic examinations, including best-corrected visual acuity (BCVA), intraocular pressure (IOP), and indirect ophthalmoscopy, were scheduled at baseline and at 1 day, 1 week, 1 and 3 months after the removal of Densiron-68.

Surgical technique

The surgical procedure was based on a 23-gauge TSVS using a one-step system (Alcon Laboratories Inc., Fort Worth, TX, USA). The conjunctiva was displaced and 30°-angled incisions were made with a combined 23-gauge blade-trocar system to obtain tunnels parallel to the corneoscleral limbus. The IOP was

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maintained at 40 mmHg using the vented gas forced infusion of the ACCURUS system. Densiron-68 (1480 mPa viscosity) was removed with suction of 600-mmHg vacuum through a short 23-gauge silicon cannula (PolyTip® VFI Cannula, 0.6 mm(23 G) × 7.0 mm, MedOne Surgical, Sarasota, FL, USA) (Figure 1).

The main endpoints of our study were rate of complete removal of Densiron-68 and rate of anatomical success with retinal reattachment. We also analyzed the changes in IOP and the occurrence of complications.

Results

Ten eyes (4 phakic, 5 pseudophakic, and 1 aphakic) of 10 patients (5 males and 5 females) with a mean age of 58.5

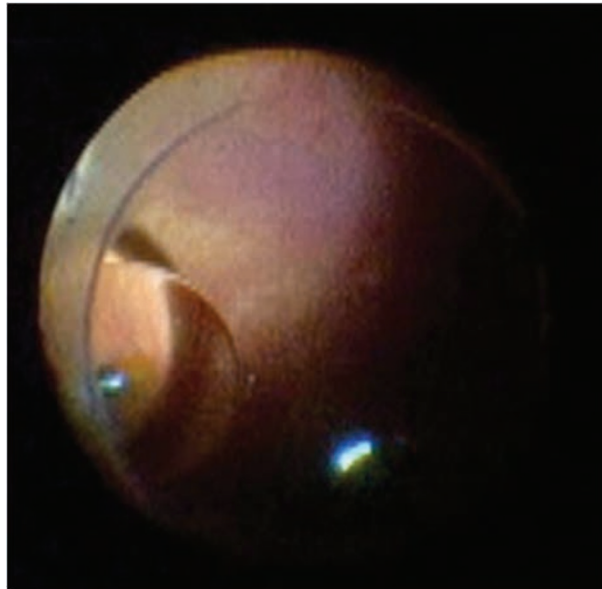


Figure 1 Densiron-68 bubble levitated by the short 23-gauge cannula in the middle of the vitreous cavity.

years (SD 8.4) underwent active removal of Densiron-68 through a 23-gauge TSVS. No intraoperative complications were reported. Suture placement was not necessary in any of the sclerotomies in 5 out of 10 eyes. Three eyes needed suture of 1/3 and two eyes of 2/3 of the sclerotomies. Densiron-68 was completely removed from all eyes, with no clinically significant residual oil in the vitreous cavity seen at follow-up examinations (Table 1). Multiple small droplets of Densiron-68 may remain in the posterior segment after removal of the main body of silicon oil. These residual droplets can be easily removed with a 23-gauge flute needle with passive aspiration.

The time taken to remove similar volumes of Densiron-68 was approximately the same with a short 23-gauge cannula as it was with a short 20-gauge cannula.

Following tamponade removal, retinal reattachment was achieved in all cases (100%). Mean BCVA improved from 0.70 (SD 0.51) to 0.48 (SD 0.9) logMAR.

The IOP was 20.9 (SD 3.5) mmHg at baseline, 12.2 (SD 4) mmHg at 1 day, and 13.6 (SD 2.9), 15.4 (SD 2.5), and 16 (SD 1.8) mmHg at 1 week, 1 and 3 months, respectively (Table 1).

Anterior chamber shallowing occurred in 1 out of 10 eyes (10%). Localized bleb formation was not seen. Three out of five phakic patients developed cataract during the follow-up. Postoperative hypotony (≤ 8 mmHg) was seen in 2 out of 10 eyes (20%), but resolved spontaneously within 1 week.

Discussion

The conventional way of removing heavy silicon oil with a 0.90 mm(20 G) × 28 mm cannula reaching the lowest part of the vitreous cavity is associated with risk of complications mainly related to the larger scleral openings (entry site tears) and to the high suction

Table 1 Demographics and outcomes of study patients

Patient no.	Age	Sex	Eye	Lens status	Intraocular pressure					Sclerotomy needed suture
					Baseline	1 day	1 week	1 month	3 month	
1	53	Male	Right	Phakic	20	14	14	14	16	2/3
2	62	Male	Right	Phakic	16	12	10	12	12	0
3	70	Female	Left	Pseudophakic	22	10	10	12	16	0
4	55	Male	Left	Phakic	25	12	16	16	18	1/3
5	70	Male	Right	Pseudophakic	18	8	14	16	16	0
6	45	Female	Left	Pseudophakic	26	22	20	20	18	2/3
7	48	Female	Right	Pseudophakic	20	10	14	14	14	0
8	58	Female	Right	Aphakic	16	8	12	16	16	1/3
9	60	Male	Right	Pseudophakic	22	12	12	16	16	1/3
10	64	Female	Right	Phakic	24	14	14	18	18	0

pressure applied close to the retina through a relatively large bore opening.^{3,4} If the needle is momentarily emptied of the viscous oil, the eye might suddenly collapse and the needle tip could damage the retina.

Stappler *et al*⁵ have recently proved that the removal of Densiron-68 is possible using a short 20-gauge steel needle. A combination of forces, such as the interfacial tension between the cannula and the oil, as well as the phenomenon of tubeless siphoning, makes it feasible and reduces the risk of iatrogenic damage.

We report that it is also possible to remove Densiron-68 using a 23-gauge cannula of only 7.0 mm in length using 600 mmHg of suction pressure. Densiron-68 is made of polymers and behaves as non-Newtonian fluids. When such non-Newtonian fluids are forced through a narrow opening, the three-dimensional structure of the molecules unfolds, thereby storing up energy in the process.⁵ The key point of this technique is to not lose the contact between the tip of the short cannula and the oil bubble. The only problem is related to the postoperative leaking of sclerotomies. After removal of silicon oil, the vitreous body base is mostly contracted and compressed; hence the remaining peripheral vitreous cannot seal the sclerotomies as happens for macular surgery, when a broad rim of peripheral vitreous is not removed.⁶ To prevent such postoperative hypotony, we prefer to put a single transconjunctival suture on the sclerotomy in case of leakage after the removal of the trocar.

In conclusion, active removal of Densiron-68 with a 23-gauge short cannula is a simple, innovative, and safe technique, which can help reduce surgical trauma. It can be performed in phakic, pseudophakic, and aphakic eyes. Controlled studies with larger series are warranted.

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