·Clinical Research·

Combined pars plana vitrectomy and Baerveldt glaucoma implant placement for refractory glaucoma

Thalmon R. Campagnoli¹, Sung Soo Kim², William E. Smiddy¹, Steve J. Gedde¹, Donald L. Budenz³, Richard K. Parrish II¹, Paul F. Palmberg¹, William Feuer¹, Wei Shi¹

¹Department of Ophthalmology, Bascom Palmer Eye Institute, University of Miami Miller School of Medicine, Miami, FL 33136, USA

²Department of Ophthalmology, Gangnam Severance Hospital, Yonsei University College of Medicine, Seoul 135-720, South Korea

³Department of Ophthalmology, University of North Carolina, Chapel Hill, NC 27599, USA

Correspondence to: William E. Smiddy. Department of Ophthalmology, Bascom Palmer Eye Institute, University of Miami School of Medicine, 900 NW 17th St., Miami 33136, USA. wsmiddy@med.miami.edu

Received: 2015-01-27 Accepted: 2015-03-17

Abstract

• AIM: To evaluate outcomes of combined pars plana vitrectomy and Baerveldt glaucoma implant (PPV-BGI) placement for refractory glaucoma.

• METHODS: The medical records of 92 eyes (89 patients) that underwent PPV-BGI were retrospectively reviewed, including 43 eyes with neovascular glaucoma (NVG) and 49 eyes with other types of glaucoma (non-NVG).

• RESULTS: Outcome measures were visual acuity (VA), intraocular pressure (IOP), glaucoma medical therapy, complications, and success [VA >hand motions (HM), IOP \geq 6 mm Hg and \leq 21 mm Hg, no subsequent glaucoma surgery]. Cumulative success rates for the non-NVG group and NVG group were 79% and 40% at 1y, respectively (P=0.038). No difference in the rates of surgical success were found between pars plana and anterior chamber tube placement. Preoperative IOP (mean±SD) was 30.3±11.7 mm Hg in the Non-NVG group and 40.0±10.6 mm Hg in the NVG group, and IOP was reduced to 15±9.5 mm Hg in the non-NVG group and 15± 10.5 mm Hg in the NVG at 1y. Number of glaucoma medications (mean±SD) decreased from 2.7±1.3 in the non -NVG group and 2.8 ±1.3 in the NVG group preoperatively to 0.76±1.18 in the non-NVG group and 0.51±1.00 in the NVG group at 1y. Improvement in VA of \geq 2 Snellen lines was observed in 25 (27%) eyes, although only 33% of non-NVG eyes and 2.3% of NVG eyes maintained VA better than 20/200 at 1y. Nonclearing

vitreous hemorrhage was the most common postoperative complication occurring in 16 (17%) eyes, and postoperative suprachoroidal hemorrhages developed in 5 (5.4%) eyes.

• CONCLUSION: PPV-BGI is a viable surgical option for eyes with refractory glaucoma, but visual outcomes are frequently poor because of ocular comorbidities, especially in eyes with NVG. The location of tube placement does not influence surgical outcome and should be left to the discretion of the surgeon.

• **KEYWORDS:** glaucoma drainage device; pars plana vitrectomy; refractory glaucoma; neovascular glaucoma **DOI:10.3980/j.issn.2222–3959.2015.05.11**

Campagnoli TR, Kim SS, Smiddy WE, Gedde SJ, Budenz DL, Parrish RK II, Palmberg PF, Feuer W, Shi W. Combined pars plana vitrectomy and Baerveldt glaucoma implant placement for refractory glaucoma. *Int J Ophthalmol* 2015;8(5):916–921

INTRODUCTION

a laucoma drainage devices (GDDs) have traditionally been used in eyes at high risk for trabeculectomy failure, such as those with pediatric glaucoma, uveitic glaucoma, secondary glaucoma after penetrating keratoplasty, and neovascular glaucoma (NVG)^[1-8]. The GDD tube is usually inserted into the anterior chamber (AC), but it may be placed through the pars plana into the vitreous cavity if endothelial decompensation or AC corneal angle abnormalities are present ^[3,5,9-20]. Pars plana vitrectomy (PPV) is required to avoid vitreous occlusion of the tube ostium if placed into the vitreous cavity. A vitrectomy may also be required with tube insertion into the AC in the presence of vitreous prolapse, as is commonly seen in aphakia eyes. Glaucoma patients with coexisting posterior segment diseases may be considered for combined PPV and GDD implantation, such as patients with NVG who require both panretinal laser photocoagulation (PRP) and intraocular pressure (IOP) reduction ^[21-23]. Successful IOP reduction has been reported with the combination of PPV with implantation of various types of GDD's in case reports and small case series in eyes with complex diseases [2,9-18]. Studies of Baerveldt glaucoma implants (BGI) (Abbott Medical Optics,

Santa Ana, California, USA) combined with PPV and insertion into the vitreous cavity have not been widely published ^[11,14,20]. The purpose of this study is to report the outcomes of a large cohort of patients who underwent PPV and BGI placement (PPV-BGI), and to compare the outcomes between the NVG and non-NVG cases.

SUBJECTS AND METHODS

This study was approved by the Human Subjects Committee of the University of Miami Miller School of Medicine, and was conducted in accordance with the Declaration of Helsinki for research in human subjects. The medical records of all patients who underwent PPV-BGI at the Bascom Palmer Eye Institute between January 1, 1992 and May 31, 2007 by one vitreoretinal surgeon (Smiddy WE) were identified using Current Procedural Terminology (CPT) codes and retrospectively reviewed. Implantation of a BGI was performed by one of the other authors (Gedde SJ, Budenz DL, Parrish RK II and Palmberg PF). Demographic information, surgical indication, lens status, number of previous intraocular surgeries, preoperative AC angle status, presence of preoperative hyphema, presence of iris or angle neovascularization, adjunctive procedures performed during surgery, complications, pre- and post-operative best-corrected visual acuity (BCVA) and IOP, number of antiglaucoma medications, and post-operative interval until last follow-up examination were extracted from the medical record.

The surgical procedure consisted of a three-port, 20-gauge PPV performed in conjunction with the placement of a BGI in the superotemporal quadrant. Temporary restriction of aqueous flow through the implant was achieved with a tube ligation with a 7-0 polyglactin suture near the tube-plate junction. Fenestrations were created in the tube with a TG-130 needle just anterior to the polyglactin ligature with a needle for early IOP control according to the glaucoma surgeon's judgment. The tube was placed through the pars plana into the vitreous cavity either through a sclerotomy created by the 20-gauge microvitreoretinal blade or through a separate 23-gauge needle tract. It was the consensus of the glaucoma specialist that the Hoffman elbow modification of the BGI was not necessary and, hence, it was not used. A donor corneal or scleral patch graft was applied over the anterior portion of the tube to prevent erosion. Patients were monitored post-operatively by both the glaucoma and vitreoretinal surgeons.

Patients were categorized by diagnosis as NVG or non-NVG for the purpose of data analysis. The main outcome measure was success defined as BCVA \geq hand motions (HM), normal IOP (\geq 6 and \leq 21 mm Hg) with or without antiglaucoma medication, and no additional ocular surgery. Eyes that did not fail but required adjunctive glaucoma medications were classified as qualified successes, while those without

Table 1 Baseline characteristics of the study population						
Data analyzed	NVG	Non-NVG	Р			
No. of eyes	43	49				
No. of patients	41	48				
Age (a)						
Mean (SD)	65 (15)	62.51 (22.95)	0.55 ^a			
Median (range)	64 (36-92)	69 (5-90)				
Gender						
Male, <i>n</i> (%)	23 (53)	22 (45)	0.34 ^b			
Follow-up (mo)						
Mean (SD)	16 (15)	23 (21)	0.031^{a}			
Median (range)	11 (1-84)	15 (1-84)				
Lens status, n (%)			0.02 ^c			
Phakic	20 (47)	10 (20)				
Aphakia	2 (5)	5 (10)				
ACIOL	2 (5)	9 (18)				
PCIOL	19 (44)	25 (51)				
VA, logMAR						
Mean (SD)	2.23 (0.48)	1.51 (0.75)	$< 0.001^{a}$			
Median (range)	2.3 (0.7-3.0)	1.3 (0.18-3.0)				
IOP (mm Hg)						
Mean (SD)	40.0 (10.6)	30.3 (11.7)	$< 0.001^{a}$			
Median (range)	39 (17-60)	31 (6-55)				
Extent of angle closure by PAS, n (%)			0.24 ^c			
PAS (-) /open angle	13 (30)	15 (31)				
<180 degrees	13 (30)	8 (16)				
≥180 degrees	17 (40)	26 (53)				
Antiglaucoma medications, n mean (SD)	2.8 (1.3)	2.7 (1.3)	0.56 ^a			
Mean number previous intraocular surgeries, <i>n</i> mean (SD)		1.8 (1.4)	<0.001ª			

ACIOL: Anterior chamber IOL; IOP: Intraocular pressure; logMAR: Logarithm of minimal angle resolution; NVG: Neovascular glaucoma; PAS: Peripheral anterior synechia; PCIOL: Posterior chamber lens; VA: Visual acuity. ^aTwo-sample *t*-test; ^bFisher Exact test; ^cPearson Chi-square test.

glaucoma medications were categorized as complete successes. Hypotony was defined as IOP ≤ 5 mm Hg. Data from the clinic visit nearest to 12mo post-operatively (range, 7 to 18mo) were analyzed for 1y outcomes.

SPSS (Version 17, SPSS Inc. Chicago, II, USA) was used for statistical analysis. Demographic data and preoperative data between the NVG group and non-NVG group were analyzed with a paired 2-sample *t*-test. IOP comparisons between two groups were analyzed with a paired 2-sample *t*-test. Rates of surgical success, surgical failure, postoperative hypotony, and visual acuity change were analyzed by Chi-square test. The incidence of complications was analyzed with the Fisher Exact test. Kaplan-Meier survival analysis was used to calculate success rates.

RESULTS

Baseline Characteristics Table 1 shows the baseline

Pars plana vitrectomy and Baerveldt glaucoma implant for refractory glaucoma

characteristics of the study population. Ninety two eyes from 89 patients were included in the study. The NVG group consisted of 43 eyes from 41 patients, and proliferative diabetic retinopathy (14 eyes, 33%) and central retinal vein occlusion (4 eyes, 9%) were the most common underlying etiologies. The non-NVG group consisted of 49 eyes of 48 patients, and the most common surgical indications were prior penetrating keratoplasty (8 eyes, 16%) and aphakia with vitreous prolapse (6 eyes, 12%). The median age (64y vs 69y), male gender (53% vs 45%), median follow-up interval (11mo vs 15mo), rate of \geq 50% angle closure (40% vs 53%), and number of preoperative glaucoma medications (2.8 vs 2.7) were not significantly different between the NVG group and non-NVG group, respectively. The non-NVG group had better preoperative vision (median logMAR 1.3 vs 2.3, P<0.001), lower median IOP (31 mm Hg vs 39 mm Hg, P < 0.001), and more previous intraocular surgeries (mean 1.8 vs 0.7, P < 0.001) in comparison to the NVG group.

Preoperatively there were a total of 55 pseudophakic, 7 aphakic, and 30 phakic eyes. Cataract extraction was performed during follow-up in the study in 8 (27%) phakic eyes. The NVG group was more commonly phakic (47%) than the non-NVG group (20%) at the time of BGI placement (P=0.023). Other ocular comorbidities included tractional retinal detachment (2 eyes; one due to PDR and one due to sickle cell retinopathy), retained lens material, aqueous misdirection, traumatic glaucoma, and aniridia (one each). Tube fenestrations were performed in 33 (67%) of non-NVG and in 40 (93%) of NVG eyes.

Surgical Outcomes Surgical outcomes are provided in Table 2. Surgical failure occurred in 34 (37%) of the 92 eyes with at least 1y follow-up, including 26 (60%) of the NVG group and 8 (16%) of the non-NVG group (P=0.038). Vision loss to NLP (16 eyes, 23%), IOP <6 or >21 mm Hg (6 eyes, 18%), phthisis or enucleation (14 eyes, 41%) and glaucoma reoperations (6 eyes, 18%) were the causes for treatment failure. Two eyes in the NLP category underwent enucleation due to a blind, painful eye. The reoperations included PPV due to vitreous hemorrhage (VH) or fibrovascular proliferation (2 eyes), BGI revision (1 eye), BGI replacement (2 eyes), and BGI removal (1 eye). Failure on treatment was evident within 6mo post-operatively in 20 (59%) of the 34 eyes. The cumulative probability of success at 1y using Kaplan-Meier survival analysis was worse in the NVG group than in the non-NVG group (40% vs 79%; P<0.001), a difference that persisted thereafter (Figure 1).

Tube Location The tube was placed into the AC in 48 (52%) eyes (including 20 pseudophakic, 4 aphakic, and 24 phakic eyes) and into the vitreous cavity in 44 (48%) eyes (including 35 pseudophakic, 3 aphakic, and 6 phakic eyes). The tube was more frequently placed into the AC in the NVG

Table 2 Surgical outcomes					
Data analyzed	NVG <i>n</i> =43	Non-NVG <i>n</i> =49	Р		
Tube location, <i>n</i> (%)			<0.001 ^a		
AC	35 (81)	13 (27)			
РР	8 (19)	36 (73)			
Visual Outcome at final visit, n (%)			0.007 ^a		
Improved	10 (23)	27 (55)			
Stable	13 (30)	10 (20)			
Worse	20 (47)	12 (24)			
NLP ≥20/200	14 (33) 1 (2.3)	2 (4) 16 (33)	<0.001 ^b		
Mean IOP at 1a, mm Hg (SD)	15 (10.5)	15 (9.5)	0.425		
Treatment outcomes at final visit, n (%)			<0.001 ^a		
Complete success	8 (19)	26 (53)			
Qualified success	9 (21)	15 (31)			
Failure	26 (60)	8 (16)			
Mean number of glaucoma medications at $1a, n$ (SD)	0.51 (1.00)	0.76 (1.18)	0.343		

AC: Anterior chamber; IOP: Intraocular pressure; NLP: No light perception; NVG: Neovascular glaucoma; PP: Pars plana. ^a Analyzed by Pearson Chi-square test; ^bFisher Exact test.

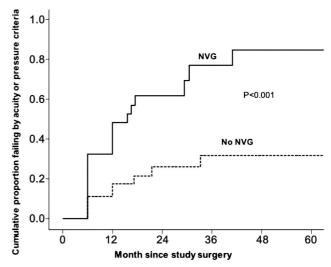


Figure 1 Kaplan-Meier plot of cumulative proportion of eyes failing visual acuity or intraocular pressure criteria for successful outcome.

group (81%) than in the non-NVG group (27%) (P<0.001), and in phakic (80%) compared to aphakic or pseudophakic (38%) eyes.

Mean IOP at one-year follow-up visit was 15 mm Hg for both NVG and Non-NVG groups (P=0.425) and the mean number of antiglaucoma medications was reduced from a mean of 2.8 in the NVG group and 2.7 in the non-NVG preoperatively to 0.51 and 0.76, respectively (P=0.343). Paired *t*-test for postoperative compared to preoperative.

Visual Results Visual results were better in the non-NVG group (Table 2). After one-year follow-up, 16 (37%) of the NVG eyes presented visual acuity ranging between HM and

counting fingers and only one eye (2%, VA = 20/70)presented visual acuity $\ge 20/200$, while 16 (33%) of 42 non-NVG eyes achieved Snellen visual acuity $\ge 20/200$ (*P*=0.001). Visual acuity was stable or improved in 75% of non-NVG eyes (*vs* 53% in the NVG group).

Complications Table 3 shows complications and associated surgical procedures during the first year of follow-up. The complication, only intraoperative а suprachoroidal hemorrhage, occurred in a non-NVG patient after a keratoplasty. Postoperative complications penetrating occurred in 28 (65%) NVG and 24 (49%) non-NVG eyes. The most common complications that were present at or beyond 1wk post-operatively in NVG patients included non-clearing VH and/or hyphema in 16 eyes (37%). In contrast, none of the non-NVG eyes had a VH by the first post-operative week. The persistence of VH/hyphema likely indicated continuing hemorrhage due to progressive underlying disease. Postoperative VH was an important prognostic factor in the NVG group; 88% of NVG patients with non-clearing VH ultimately were surgical failures. Postoperative suprachoroidal hemorrhage requiring surgical drainage within 2wk post-operatively occurred in 5 (10%) non-NVG eyes, two of which developed phthisis. Suprachoroidal hemorrhage did not occur in any NVG cases. Post-operative retinal detachment (3 cases in each group, 7% of total) or retinal breaks without retinal detachment (1 eye in the non-NVG group, 1% of total) also occurred (Tables 3, 4). DISCUSSION

This study has demonstrated that PPV combined with GDD can offer respectable results, considering it is a treatment option for the most advanced cases, and specifically that eves with non-NVG have a better prognosis. Potential advantages of this procedure in selected patients with uncontrolled IOP includes simultaneous management of glaucoma and posterior segment disorders and the option of pars plana placement of the drainage tube for patients with anterior segment comorbidities ^[12,14,16,20]. Previous studies of combined procedures have been limited to small cases of various types of GDD's^[2,9-18], offered limited comparison outcomes of NVG to other indications, used broader criteria for success (e.g. inclusion of light perception final visual acuity as success), or lacked details of visual acuity results [14,16,17,20]. The present study included a large number of patients who underwent a single type of GDD, and outcomes are subdivided into eyes with NVG and those with other glaucoma diagnoses.

Glaucoma type was a major factor affecting surgical success in this study, with NVG eyes fairing substantially worse than non-NVG eyes. The Kaplan-Meier cumulative success one year post-operatively (40% in NVG eyes and 79% in the non-NVG eyes) in the current study was similar to previously reported success rates of 43% to 78% for BGI which lumped all glaucoma categories ^[24-26]. Our study design did not permit

 Table 3 Complications and associated surgical procedures during 1y follow-up

Ly follow-up	NVG	Non-NVG
Complications	n=43	<i>n</i> =49
Choroidal detachment	3	6
Retinal breaks with RD	3	3
Retinal breaks without RD	0	1
Unclearing VH ^a	16	0
Endophthalmitis	2	0
Flat AC	1	0
Tube blockage by vitreous	0	2
IOL dislocation	0	3
Exposed tube	0	1
Corneal endothelial failure	2	4
Enucleated	2	0
Phthisis	12	2
Repeated surgery	3	3
Cataract progression	1	0
(18 phakic eyes) VA LP or NLP	8	0
IOP <6 or >21 mm Hg	3	3
Total (number of eyes)	56 (28)	28 (24)

IOL: Intraocular lens; IOP: Intraocular pressure; NLP: No light perception; NVG: Neovascular glaucoma; RD: Retinal detachment; SCH: Suprachoroidal hemorrhage; VH: Vitreous hemorrhage; VA: Visual acuity; LP: Light perception. ^aUnclearing VH: Vitreous hemorrhage not cleared till 1mo post-operatively.

 Table 4 Postoperative complications on first postoperative day

Complications	NVG <i>n</i> =43	Non-NVG n=49
Hypotony	3	13
Hyphema	26	7
VH	25	10
SCH	0	5
Total (number of eyes)	54 (26)	35 (20)

SCH: Suprachoroidal hemorrhage; VH: Vitreous hemorrhage.

a controlled evaluation of combined PPV/BGI in regards to BGI placement only or placement by location, but a reasonable inference from these data is that the tube site may be effectively decided by the surgeon based upon constraints in an individual eye. The more frequent pars plana placement in non-NVG eyes (73%) was probably due to associated anterior segment pathology that was more chronic or reflected previous surgical attention, such as vitreous prolapse commensurate with aphakia and pseudophakia.

The poor visual outcome of NVG patients in the current series is consistent with those reported previously, and is likely attributable to their poorer baseline visual acuity and natural history. A previous study reporting 56% success and 28% qualified success among 50 NVG eyes considered eyes with LP visual acuity as successes, and did not offer a detailed visual acuity analysis ^[20]. Still they acknowledged "serious complications" in 10% of eyes, consistent with the findings of the current study which offers a sober prognosis among NVG eyes. A recent retrospective study evaluated the

Pars plana vitrectomy and Baerveldt glaucoma implant for refractory glaucoma

performance of PPV-BGI in 89 NVG eyes and found a cumulative success rate of 67% in 73 eyes (82% of total) that completed follow-up (6mo post-operatively), however, the authors also considered eyes with LP visual acuity as successes cases ^[27]. Fifty-five eyes (62%) in the aforementioned study presented improved or stable visual acuity, a result that somewhat resembles ours, once 53% of the NVG eyes in this series presented stable or improved visual acuity and 8 (19%) had light perception at 1-year follow up visit. Also of note is that fifty-one eyes (58% of total) in this recent case series received intravitreal anti-VEGF injection perioperatively, in contrast to only 2 eyes (2%) in our study.

Progression of NVG implies increasing retinal ischemia and increased production of vascular endothelial growth factor (VEGF)^[28,29]. The principal weakness of the current study is that the interval during which these cases were studied predated our more regular use of anti-VEGF therapy for eves undergoing combined PPV-BGI. Studies using off-label intravitreal injection of bevacizumab (IVB) have reported short-term efficacy and safety in NVG treatment, in association with or not with other treatment modalities^[30-35]. IVB has also been reported as an effective adjuvant for surgical treatment of NVG [36-48]. Blocking anti-VEGF activity to interrupt the vicious cycle of NVG aggravation is probably helpful in improving the surgical outcome of NVG, but a prospective randomized trial, or at least a larger, case-control study specifically aiming to define the best setting (s) for its surgical adjunctive use would likely be more definitive, but at this point probably impractical. Any retrospective study is subject to the biases of case ascertainment and individual surgeon preferences; these were at least mitigated by this being a consecutive case series, there being one vitreoretinal surgeon (but several glaucoma surgeons), and little variation in the retinal portion of the procedure.

Any surgical case series could have longer follow-up examination intervals, but the nature of the tertiary practice setting of the current study dictated these constraints; the rapid declaration of clinical course in these sorts of cases probably establishes the long term outcome early post-operatively.

The current study, although retrospective and non-randomized, shows encouraging results with the performance of PPV-BGI (especially for non-NVG eyes), offering the glaucoma specialist a viable option in selected cases of refractory glaucoma. The best suited eyes for pars plana insertion of the GDD are those with narrow or flat angles, corneal decompensation, or those with florid rubeosis iridis. The apparently disappointing results with NVG in our series suggest the need to optimize the treatment approach for those eyes, possibly through earlier PPV-BGI intervention and/or use of anti-VEGF agents.

ACKNOWLEDGEMENTS

Conflicts of Interest: Campagnoli TR, None; Kim SS, None; Smiddy WE, None; Gedde SJ, None; Budenz DL, None; Parrish RK II, None; Palmberg PF, None; Feuer W, None; Shi W, None.

REFERENCES

1 Chow K, Mora J. Practice preferences for glaucoma drainage device implantation and cyclodestruction in Australia and New Zealand. *J. Glaucoma* 2012;21(3):199-205

2 Lloyd MA, Heuer DK, Baerveldt G, Minckler DS, Martone JF, Lean JS, Liggett PE. Combined Molteno implantation and pars plana vitrectomy for neovascular glaucomas. *Ophthalmology* 1991;98(9):1401–1405

3 Lloyd MA, Sedlak T, Heuer DK, Minckler DS, Baerveldt G, Lee MB, Martone JF. Clinical experience with the single-plate Molteno implant in complicated glaucomas. Update of a pilot study. *Ophthalmology* 1992;99 (5):679-687

4 Mermoud A, Salmon JF, Alexander P, Straker C, Murray ADN. Molteno tube implantation for neovascular glaucoma. Long-term results and factors influencing the outcome. *Ophthalmology* 1993;100(6):897-902

5 Knape RM, Szymarek TN, Tuli SS, Driebe WT, Sherwood MB, Smith MF. Five-year outcomes of eyes with glaucoma drainage device and penetrating keratoplasty. *J Glaucoma* 2012;21(9):608-614

6 Lloyd MA, Baerveldt G, Heuer DK, Minckler DS, Martone JF. Initial clinical experience with the Baerveldt implant in complicated glaucomas. *Ophthalmology* 1994;101(4):640–650

7 Ceballos EM, Parrish RK 2nd, Schiffman JC. Outcome of Baerveldt glaucoma drainage implants for the treatment of uveitic glaucoma. *Ophthalmology* 2002;109(12):2256-2260

8 Budenz DL, Gedde SJ, Brandt JD, Kira D, Feuer W, Larson E. Baerveldt glaucoma implant in the management of refractory childhood glaucomas. *Ophthalmology* 2004;111(12):2204-2210

9 Azuara-Blanco A, Katz LJ, Gandham SB, Spaeth GL. Pars plana tube insertion of aqueous shunt with vitrectomy in malignant glaucoma. *Arch Ophthalmol* 1998;116(6):808-810

10 Chalam KV, Gandham S, Gupta S, Tripathi BJ, Tripathi RC. Pars plana modified Baerveldt implant versus neodymium:YAG cyclophotocoagulation in the management of neovascular glaucoma. *Ophthalmic Surg Lasers* 2002; 33(5):383–393

11 Faghihi H, Hajizadeh F, Mohammadi SF, Kadkhoda A, Peyman GA, Riazi-Esfahani M. Pars plana Ahmed valve implant and vitrectomy in the management of neovascular glaucoma. *Ophthalmic Surg Lasers Imaging* 2007;38(4):292-300

12 Gandham SB, Costa VP, Katz LJ, Wilson RP, Sivalingam A, Belmont J, Smith M. Aqueous tube-shunt implantation and pars plana vitrectomy in eyes with refractory glaucoma. *Am J Ophthalmol* 1993;116(2):189–195

13 Kaynak S, Tekin NF, Durak I, Berk AT, Saatci AO, Soylev MF. Pars plana vitrectomy with pars plana tube implantation in eyes with intractable glaucoma. *Br J Ophthalmol* 1998;82(12):1377-1382

14 Luttrull JK, Avery RL. Pars plana implant and vitrectomy for treatment of neovascular glaucoma. *Retina*1995;15(5):379-387

15 Scott IU, Alexandrakis G, Flynn HW, Jr., Smiddy WE, Murray TG, Schiffman J, Gedde SJ, Budenz DL, Fantes F, Parrish RK. Combined pars plana vitrectomy and glaucoma drainage implant placement for refractory glaucoma. *Am J Ophthalmol* 2000;129(3):334–341

16 Smiddy WE, Rubsamen PE, Grajewski A. Vitrectomy for pars plana placement of a glaucoma seton. *Ophthalmic Surg* 1994;25(8):532-535

17 Varma R, Heuer DK, Lundy DC, Baerveldt G, Lee PP, Minckler DS.

Pars plana Baerveldt tube insertion with vitrectomy in glaucomas associated with pseudophakia and aphakia. *Am J Ophthalmol* 1995;119(4):401–407

18 Malone PE, Herndon LW, Muir KW, Jaffe GJ. Combined fluocinolone acetonide intravitreal insertion and glaucoma drainage device placement for chronic uveitis and glaucoma. *Am.J Ophthalmol* 2010;149(5):800–806.e1

19 Tello C, Espana EM, Mora R, Dorairaj S, Liebmann JM, Ritch R. Baerveldt glaucoma implant insertion in the posterior chamber sulcus. *Br J Ophthalmol* 2007;91(6):739-742

20 Luttrull JK, Avery RL, Baerveldt G, Easley KA. Initial experience with pneumatically stented Baerveldt implant modified for pars plana insertion for complicated glaucoma. *Ophthalmology* 2000;107(1):143–150

21 Bartz-Schmidt KU, Thumann G, Psichias A, Krieglstein GK, Heimann K. Pars plana vitrectomy, endolaser coagulation of the retina and the ciliary body combined with silicone oil endotamponade in the treatment of uncontrolled neovascular glaucoma. *Graefe's Arch Clin Exp Ophthalmol* 1999:237(12):969–975

22 Kono T, Shiga S, Takesue Y, Sakamoto T. Long-term results of pars plana vitrectomy combined with filtering surgery for neovascular glaucoma. *Ophthalmic Surg Lasers Imaging* 2005;36(3):211-216

23 Lima VC, de Moraes CG, Gentile RC, Sidoti PA, Prata TS, Liebmann JM, Will DV, Tello C, Rosen RB. Combined Baerveldt glaucoma implant and scleral buckling surgery for patients with retinal detachment and coexisting glaucoma. *J Glaucoma* 2013;22(4):294–300

24 Hodkin MJ, Goldblatt WS, Burgoyne CF, Ball SF, Insler MS. Early clinical experience with the Baerveldt implant in complicated glaucomas. *Am J Ophthalmol* 1995;120(1):32-40

25 Siegner SW, Netland PA, Urban RC Jr, Williams AS, Richards DW, Latina MA, Brandt JD. Clinical experience with the Baerveldt glaucoma drainage implant. *Ophthalmology* 1995;102(9):1298-1307

26 Krishna R, Godfrey DG, Budenz DL, Escalona-Camaano E, Gedde SJ, Greenfield DS, Feuer W, Scott IU. Intermediate-term outcomes of 350-mm (2) Baerveldt glaucoma implants. *Ophthalmology* 2001;108(3):621-626

27 Kolomeyer AM, Seery CW, Emami-Naeimi P, Zarbin MA, Fechtner RD, Bhagat N. Combined pars plana vitrectomy and pars plana baerveldt tube placement in eyes with neovascular glaucoma. *Retina* 2015;35(1): 17–28

28 Aiello LP, Avery RL, Arrigg PG, Keyt BA, Jampel HD, Shah ST, Pasquale LR,Thieme H, Iwamoto MA, Park JE, Nguyen HV, Aiello LM, Ferrara N, King GL. Vascular endothelial growth factor in ocular fluid of patients with diabetic retinopathy and other retinal disorders. *N Engl J Med* 1994;331(22):1480–1487

29 Tripathi RC, Li J, Tripathi BJ, Chalam KV, Adamis AP. Increased level of vascular endothelial growth factor in aqueous humor of patients with neovascular glaucoma. *Ophthalmology* 1998;105(2):232-237

30 Avery RL. Regression of retinal and iris neovascularization after intravitreal bevacizumab (Avastin) treatment. *Retina* 2006;26(3):352-354

31 Davidorf FH, Mouser JG, Derick RJ. Rapid improvement of rubeosis iridis from a single Bevacizumab (Avastin) injection. *Retina* 2006;26(3): 354–356

32 Mason JO 3rd, Albert MA Jr, Mays A, Vail R. Regression of neovascular iris vessels by intravitreal injection of bevacizumab. *Retina* 2006;26(7): 839–841

33 Oshima Y, Sakaguchi H, Gomi F, Tano Y. Regression of iris neovascularization after intravitreal injection of Bevacizumab in patients with proliferative diabetic retinopathy. *Am J Ophthalmol* 2006;142 (1): 155-158

34 Iliev ME, Domig D, Wolf-Schnurrbursch U, Wolf S, Sarra GM. Intravitreal Bevacizumab (Avastin) in the treatment of neovascular glaucoma. *Am J Ophthalmol* 2006;142(6):1054-1056

35 Ehlers JP, Spirn MJ, Lam A, Sivalingam A, Samuel MA, Tasman W. Combination intravitreal bevacizumab/panretinal photocoagulation versus panretinal photocoagulation alone in the treatment of neovascular glaucoma. *Retina* 2008;28(5):696–702

36 Cornish KS, Ramamurthi S, Saidkasimova S, Ramaesh K. Intravitreal bevacizumab and augmented trabeculectomy for neovascular glaucoma in young diabetic patients. *Eye (Lond)* 2009;23(4):979–981

37 Kahook MY. Bleb morphology and vascularity after trabeculectomy with intravitreal ranibizumab: a pilot study. *Am J Ophthalmol* 2010;150 (3): 399-403.e1

38 Grewal DS, Jain R, Kumar H, Grewal SP. Evaluation of subconjunctival bevacizumab as an adjunct to trabeculectomy: a pilot study. *Ophthalmology* 2008;115(12): 2141–2145

39 Beutel J, Peters S, Luke M, Aisenbrey S, Szurman P, Spitzer MS, Yoeruek E; the Bevacizumab Study Group, Grisanti S. Bevacizumab as adjuvant for neovascular glaucoma. *Acta Ophthalmol* 2010;88(1):103–109 40 Miki A, Oshima Y, Otori Y, Kamei M, Tano Y. Efficacy of intravitreal bevacizumab as adjunctive treatment with pars plana vitrectomy, endolaser photocoagulation, and trabeculectomy for neovascular glaucoma. *Br J Ophthalmol* 2008;92(10):1431–1433

41 Zhang HT, Yang YX, Xu YY, Yang RM, Wang BJ, Hu JX. Intravitreal bevacizumab and Ahmed glaucoma valve implantation in patients with neovascular glaucoma. *Int J Ophthalmol* 2014;7(5):837-842

42 Sevim MS, Buttanri IB, Kugu S, Serin D, Sevim S. Effect of intravitreal bevacizumab injection before Ahmed glaucoma valve implantation in neovascular glaucoma. *Ophthalmologica* 2013;229(2):94–100

43 Mahdy RA, Nada WM, Fawzy KM, Alnashar HY, Almosalamy SM. Efficacy of intravitreal bevacizumab with panretinal photocoagulation followed by Ahmed valve implantation in neovascular glaucoma. *J Glaucoma*2013;22(9):768-772

44 Marey HM, Ellakwa AF. Intravitreal bevacizumab with or without mitomycin C trabeculectomy in the treatment of neovascular glaucoma. *Clin Ophthalmol* 2011;5:841-845

45 Alkawas AA, Shahien EA, Hussein AM. Management of neovascular glaucoma with panretinal photocoagulation, intravitreal bevacizumab, and subsequent trabeculectomy with mitomycin C. *J Glaucoma* 2010;19 (9): 622-626

46 Takihara Y, Inatani M, Kawaji T, Fukushima M, Iwao K, Iwao M, Tanihara H. Combined intravitreal bevacizumab and trabeculectomy with mitomycin C versus trabeculectomy with mitomycin C alone for neovascular glaucoma. *J Glaucoma* 2011;20(3):196–201

47 Zhou MW, Wang W, Huang WB, Chen SD, Li XY, Gao XB, Zhang XL. Adjunctive with versus without intravitreal bevacizumab injection before Ahmed glaucoma valve implantation in the treatment of neovascular glaucoma. *Chin Med J (Engl)* 2013;126(8):1412–1417

48 Mahdy RA. Adjunctive use of bevacizumab versus mitomycin C with Ahmed valve implantation in treatment of pediatric glaucoma. *J Glaucoma* 2011;20(7):458-463