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**QLAUKOMA MÜALİCƏSİNDƏ DRENAJ
CƏRRAHİYYƏSİNİN DÜNƏNİ, BU GÜNÜ VƏ SABAHİ
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XÜLASƏ

Hazırda qlaukoma geri dönməz korluq, əmək qabiliyyətinin itirilməsi və əlillik səbəbləri arasında lider mövqelərdən birini tutmağa davam edir və bununla da xüsusi sosial əhəmiyyət kəsb edir. Eyni zamanda, xəstələnmə hallarının artım tendensiyası qlobal xarakter daşıyır.

Medikamentoz terapiyasının bütün arsenalına və antiqlaukomatoz cərrahiyyə metodlarının təkmiləşdirilməsinə baxmayaraq, daha yüksək təhlükəsizlik profili və maksimal effektivliyi təmin edə bilən yeni texnologiyaların axtarışı, hazırlanması və tətbiqi oftalmocərrahlar arasında böyük maraq doğurmağa davam edir.

Biz 131 il dövr ərzində qlaukomanın müasir cərrahi müalicə metodları ilə bağlı mövcud elmi nəşrlərin təhlilini apardıq. Bu icmal kliniki praktikada qlaukoma zamanı geniş istifadə olunan drenaj cihazlarına həsr olunub. Xüsusilə, son onillikdə sürətlə inkişaf edən drenaj mikroinvaziv antiqlaukomatoz cərrahiyyə texnologiyasına, qlaukoma xəstəliyinin erkən və inkişaf etmiş mərhələlərində ənənəvi müalicəyə alternativ kimi xüsusi diqqət yetirilir. Məqalədə drenajların tətbiqinin tarixi, onların təsnifatı, modellərin xüsusiyyətləri və implantasiya üsulları araşdırılır.

Drenaj cihazlarından istifadə olunan cərrahi metodlarla bağlı müxtəlif aspektlərin öyrənilməsinə dair təqdim olunan klinik tədqiqat nəticələri bu metodların müasir qlaukoma müalicəsi konsepsiyalarına uyğunluğunu aydın şəkildə nümayiş etdirməklə yanaşı, onların tətbiq sahəsinin genişləndirilməsinin məqsədəuyğunluğunu göstərir. Cərrahi travmanı minimuma endirmək və davamlı hipotenziya effekti əldə etmək məqsədilə daha qabaqcıl, patogeneza baxımından əsaslandırılmış texnologiyaların axtarışının zəruriliyi heç bir şübhə doğurmur. Bununla belə, əminliklə demək olar ki, artıq bu gün drenaj cərrahiyyəsi müasir qlaukoma müalicə alqoritmində layiqli yer tutur.

Açar sözlər: *qlaukoma, qlaukoma drenaj cihazları, “Baerveldt” implant, “Ahmed valve”, “Molteno” implant, “PAUL” implant*

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DRAINAGE SURGERY IN GLAUCOMA TREATMENT: PAST, PRESENT AND FUTURE (LITERATURE REVIEW)

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SUMMARY

Currently, glaucoma continues to occupy one of the leading places among the causes of irreversible blindness, loss of ability to work and disability, thereby acquiring special social significance. At the same time, the trend of increasing incidence is global.

Despite the entire arsenal of drug therapy and the improvement of antiglaucoma surgery methods, the search, development and implementation of new technologies that can provide a higher safety profile combined with maximum efficiency continue to arouse increased interest of ophthalmic surgeons.

We analyzed relevant available publications for a 131-year period on the issue of modern methods of surgical treatment of glaucoma. This review is devoted to the topic of drainage devices widely used in clinical practice for glaucoma. Particular attention is paid to the actively developing technologies of drainage antiglaucoma surgery in the last decade as an alternative to traditional treatment for initial and advanced forms of glaucoma. The article considers the history of the use of drains, their classification, characteristics of models and methods of their implantation. The presented results of clinical studies on various aspects of the application of surgical methods using drainage devices not only clearly demonstrate their compliance with modern concepts in the treatment of glaucoma, but also dictate the expediency of expanding the indications for their use. There is no doubt that further research is needed for more advanced, pathogenetically substantiated technologies in order to minimize surgical trauma and achieve a stable hypotensive effect. However, we can confidently say that drainage surgery already today occupies a worthy place in the algorithm of modern glaucoma treatment.

Key words: *glaucoma, glaucoma drainage devices, Baerveldt implant, Ahmed valve, Molteno implant, PAUL implant*

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A comprehensive literature search was conducted in the PubMed, e-library, Scopus, Google Scholar databases using primary and secondary keywords: “glaucoma”, “glaucoma drainage devices”, “GDD”, “Baerveldt implant”, “Ahmed valve”, “Molteno implant”, “PAUL implant”. A total of 250 articles published between 1894 and 2025 were identified.

Titles, summaries, and keywords were pre-screened for relevance to the topic, resulting in 187 articles being selected for full-text assessment. The remaining 120 relevant publications were included in the final review and cited. They consisted of 56 randomized controlled trials (RCTs), 12 retrospective meta-analyses, and a number of additional observational and experimental studies, as well as national protocols and guidelines, which provide a comprehensive overview of the Glaucoma Drainage Device (GDD) theoretical justifications and clinical implementation.

Purpose

This review was conducted to highlight the current situation and trends in glaucoma drainage surgery, to evaluate the efficacy, safety and sustainability of various GDDs, to inform about innovative technologies and development trends in this segment of glaucoma treatment. Data of 12 major meta-analyses were presented to qualify the practical effectiveness of GDD as the very important and significant part of glaucoma management.

Glaucoma, as a chronic progressive neurodegenerative disorder, has been one of the leading causes of irreversible blindness worldwide for many decades [1]. In 2010, 4,5 million people were diagnosed with bilateral blindness due to glaucoma and by 2020, this number had increased to 5.9 million [2, 3]. According to the World Health Organization, various types of glaucoma have caused blindness to approximately 8 million people globally [4]. The global prevalence of glaucoma is estimated at 3.54%, with the

number of cases projected to increase from 64.3 million in 2013 to 111,8 million in 2040 [5, 6].

Their irreversible consequences of glaucoma highlight its significance in both medical and social contexts. In Russia, glaucoma is the leading cause of primary disability due to eye disease, being the cause of 31% of all disability cases, depending on the region [7]. In the United States glaucoma affects about 2% of adults over 40 with incidence rates increasing significantly in older population [8, 9]. In Azerbaijan, the burden of glaucoma averages $14.2 \pm 1.3\%$ of all eye and adnexa diseases, recorded between 2010 and 2019, with a prevalence rate of $14.5 \pm 18.7\%$ [10].

Early detection of glaucoma is the cornerstone of combating this serious disease, which has irreversible consequences and a high disability rate. Aggressive proactive screenings and prioritizing the etiopathogenetic features of glaucoma when choosing treatment strategies are crucial for its effectiveness and play a decisive role in reducing its impact. However, according to some studies, even with an adequate treatment, about 10% of patients still lose their vision [11, 12].

Drug therapy remains relevant and is recommended by various protocols as a first-choice or starting treatment. Considering the serious expansion of pathogenetically targeted antiglaucoma medicines and their increased effectiveness, the therapeutic approach has become a vector-forming in the arsenal of modern ophthalmologists [13, 14, 15, 16]. Prostaglandin analogues, beta-blockers, rho kinase inhibitors (ROCK), diuretics, carbonic anhydrase inhibitors, cholinergic agonists, alpha agonists represent the main groups of medications using for the treatment of several types of glaucoma [17, 18, 19, 20].

However, some authors argue that the shift toward an active observational therapeutic strategy has led to negative consequences, resulted in deterioration in surgical outcomes [21]. Thus, a large-scale study on the clinical and epidemiological

characteristics of glaucoma in the CIS countries and Georgia revealed that surgical treatment was performed in 19.3% of cases at the initial stage of the disease, and in 37.2% and 37.4% of cases at advanced and severe stages, respectively [22]. Considering the late diagnostic of glaucoma, surgical treatment remains the primary method for achieving long-term reduction of intraocular pressure (IOP) to target levels necessary to maintain visual function [23, 24, 25].

Another important risk factor in glaucoma progression is the IOP fluctuations and controlling it is crucial for glaucoma treatment due to destructive effect on the optic nerve. [26]. However, the clinical polymorphism of glaucoma doesn't allow us to consider any of the many actual surgical techniques as universal and most effective.

Currently, the only reliable way to stabilize the glaucoma process is to reduce the level of ophthalmotonus to target IOP in the range of 12-21 mmHg, which is confirmed by the results of several large multicenter studies [27, 28]. Thus, based on the main indicator – targeting low IOP, the criteria for qualifying the stabilization of glaucoma process are the absence of additional instillations, need for re-surgery and availability of vision-threatening complications [29].

The first-choice surgery – trabeculectomy, remains the “gold standard” in glaucoma filtration surgery, but obtained results are temporary, unpredictable and unstable due to fibrosis and scarring. Almost 60% of filtration blebs cause to function within 15 years, often a much shorter period, and subsequent treatment is limited to cyclodestructive procedures [30, 31].

This is why the revolutionary idea of implanting drainage systems in glaucoma treatment has become increasingly popular in recent years and is considered as a full-fledged alternative to trabeculectomy and cyclodestruction [32]. Modern models of implants made from intact materials, as well as the latest surgical techniques based on innovational approaches, have brought to

significant increase of number of drainage surgeries even in the early stages of glaucoma [33, 34].

Drainage surgery in the treatment of glaucoma has become much more relevant in most cases. Implantation of the Glaucoma Drainage Device (GDD) is the main treatment tactic for drug-uncontrolled glaucoma and the most common penetrating surgery for glaucoma in the world. Its advantages include a long-lasting hypotensive effect and a relatively controlled IOP level. Originally intended for the treatment of secondary glaucoma and ineffective trabeculectomy, it is now becoming the first-choice surgery type for refractory glaucoma [35, 36, 37, 38].

Previously, GDD was used when fistulizing surgeries were considered to be low-effective or impossible for technical reasons. These included patients with excessive postoperative scarring of the conjunctiva or its pronounced pathology, active neovascularization of the iris or anterior chamber angle. However, subsequent observations have demonstrated more positive results compared with trabeculectomy in medically uncontrolled glaucoma with a lower risk of filtration impairment [39, 40]. These devices have been shown to be successful in controlling IOP in eyes that have previously undergone trabeculectomy, as well as in uveitic, neovascular, pediatric, and progressive glaucoma [41, 42].

Thus, neovascular glaucoma as the main type of refractory glaucoma is pathogenetically associated with severe ischemic eye diseases, such as proliferative diabetic retinopathy and ischemic occlusion of the central retinal vein. Panretinal photocoagulation and intravitreal anti-VEGF injections are used in the treatment of neovascular glaucoma to reduce ischemia and angiogenic factors, but they are not sufficient for sustainable IOP control. Currently, the main surgical treatment for neovascular glaucoma is the implantation of antiglaucomatous drainage devices, and in cases of pars plana vitrectomy, it is the only effective option [43, 44, 45, 46, 47].

Main indications for GDD implantation:

- Neovascular glaucoma
- Traumatic glaucoma
- Uveitic glaucoma
- Pseudoexfoliative glaucoma
- Pigmentary glaucoma
- Pediatric glaucoma
- Iridocorneal endothelial syndrome
- Post-penetrating keratoplasty glaucoma
- Post-vitreoretinal surgery glaucoma
- Aphakic/Pseudophakic glaucoma
- Failed trabeculectomies
- Sturge-Weber's syndrome

The main criteria to qualify the GDD:

- IOP control
 1. *Target range 6-18 mmHg*
 2. *IOP reduction from the baseline $\geq 20-30\%$*
 3. *Sustained IOP control without significant fluctuations*
- Medication reduction
- Stability of the visual field and optic nerve
- Maintenance or improvement of Visual Acuity (VA)
- Endothelial cell density
- Safety profile
 1. *Minimal or manageable complications: hypotony, tube exposure, endophthalmitis, corneal decompensation*
 2. *No device-related severe adverse events*
- Surgical success rate
 1. *Complete success (target IOP achieved without medications or reoperation)*
 2. *Qualified success: (target IOP achieved with or without medications, but no reoperation)*
 3. *Failure: (IOP > defined upper limit, IOP < 5 mmHg on ≥ 2 visits, vision loss from complications, or device removal)*

The idea of using drainage to reduce intraocular pressure dates back over 100 years. In 1886, French ophthalmologist Wecker

proposed using a gold wire as a drainage device. However, trauma of the tissues caused by its movement led to the abandonment of this revolutionary idea [48]. The use of setons for aqueous humor drainage began in 1906 when silk threads were implanted in anterior chamber via paracentesis [49]. Zorab used horsehair to drain the anterior chamber during the treatment of hypopyon and, a year later, to enhance aqueous humor outflow in glaucoma surgery [50]. However, the postoperative period was often complicated by active formation of connective tissue. Bick experimented with magnesium wire and tantalum foil, but these attempts failed due to poor material fixation, leading to frequent extrusion and ocular infections [51]. One study described the use of platinum drains. Kuljaca et al. implanted a Teflon drain in 12 patients, achieving a positive hypotensive effect in 75% of cases. However, Teflon induced significant connective tissue formation and this is limiting its clinical use [52].

Various materials, including vanadium steel, nickel-titanium alloy, stainless steel, and even gold, were tested as alternatives. Analysis of these studies revealed that the overall effectiveness of drainage devices directly depends on the material used. The material's characteristics determine the inflammatory and reparative response of surrounding tissues. While this response is a protective function of connective tissue, it can lead to scar tissue formation, obstructing outflow pathways and reducing the hypotensive effect. Minimizing this risk requires materials with high biocompatibility and low toxicity [53].

The first antiglaucomatous drainage device, the Molteno Implant (**Figure 1**), was introduced over 50 years ago by Professor Anthony Molteno (1938–2023). His pioneering GDD work started in University of Cape Town, South Africa, and then prolonged at the University of Otago, Dunedin, New Zealand.



Figure 1. *Graphic image of Molteno Implant.*

This valveless device allowed aqueous humor to flow from the anterior chamber through a silicone tube to an end plate, where it was absorbed in the sub-Tenon's space. Fibrous capsule formation around the plate was expected within 4-6 weeks postoperatively, regulating the outflow of aqueous humor [54, 55]. Molteno developed several modifications of the device, including a pediatric version. In 1981, he introduced a double-plate implant but later returned to single-plate designs with larger (175 or 230 mm²) and more flexible configurations. Nowadays, the Molteno drainage device line is widely used worldwide and is manufactured by Innovative Ophthalmic Products Inc. (Costa Mesa, CA, USA) and Molteno Ophthalmic Limited (Dunedin, New Zealand). Newest models of Molteno GDD are: Molteno3 Single Plate (Molteno3 S), Molteno3 Double Plate (Molteno3 D) and Molteno3 Mini. Improvements include lower profile design, flexible biocompatible silicone tube, sutureless plate fixation system and integrated ripcord ligature for flow control [56 - 59].



Figure 2. *Graphic image of Krupin Valve*

In 1979, Professor Theodore Krupin from Northwestern University Medical School (Chicago, USA) introduced the first valved antiglaucomatous drainage device, where the valve provided resistance to aqueous outflow. The valve automatically opened at an IOP of 11-16 mmHg and closed at 1-3 mmHg. Krupin Valve played a significant role in the evolution of GDD and in common glaucoma surgery (**Figure 2**). It rarely used today due to more effective and predictable alternatives [60, 61, 62].

In early 1980s, Dr. Stephen S. Shocket conceived and developed the idea of using a tube connected to a 360-degree encircling band as a drainage for glaucoma treatment with initial cases dating to around 1983-1985. The ACTSEB (Anterior Chamber Tube Shunt to an Encircling Band) remained a niche technique used in some unpredictable situations, but it has historical value in the development of latest GDDs as non-valved and biologically modulated procedure. Later, in association with Dr Ahmed was created the Shockmed valve, which combined the ACTSEB procedure with Ahmed Valve implantation [63, 64, 65].

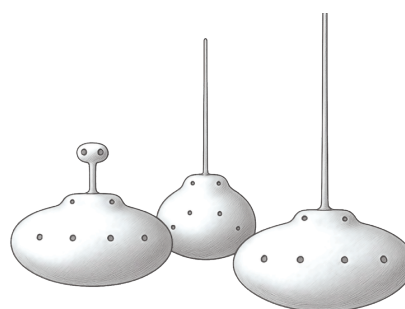


Figure 3. *Graphic image of Baerveldt® Glaucoma Implant*

Professor George Baerveldt supported the concept of non-restrictive drainage and, in 1990, (FDA approved in 1995) introduced his model of the GDD in California, USA, which quickly gained global popularity. The American Glaucoma Society awarded him the Innovator Award for his extraordinary

contributions to glaucoma research and surgical innovations [66 – 70]. The new drainage device used softer silicone, but the plate size was significantly larger (250 and 350 mm²). Over the years, numerous modifications of the Baerveldt implant (**Figure 3**), including pediatric versions, have been developed. But generally, it stays relatively unchanged due to well-established performance. These are manufactured by Johnson & Johnson Vision Care, Inc., and distributing by Advanced Medical Optics, Inc. (Santa Ana, CA, USA) [71, 72, 73].



Figure 4. *Graphic image of Ahmed® Glaucoma Valve*

In 1993, Abdul Mateen Ahmed introduced the Ahmed® Glaucoma Valve, a valved device with self-regulating silicone membranes that open automatically when IOP exceeds 8-10 mmHg. The device has undergone numerous modifications, significantly improving its effectiveness and ease of use (**Figure 4**). These modifications have focused on simplifying implantation techniques, enhancing IOP control, and reducing acute postoperative hypotony [74, 75, 76]. Pressure-sensitive venturi valve opens when IOP exceeds 8-10 mmHg. This GDD, as well as its newest versions, are manufactured by New World Medical Inc., USA.

It should be noted that despite the large numbers of GDD models from different manufacturers, the implantation of the device has common similar technical principles and treatment regulations. The superotemporal quadrant is the gold-standard site for GDD implantation. It provides the easiest surgical

access, avoids superior oblique tendon, minimizes risk to long posterior ciliary nerves and provides a large surface area for bleb formation. If superotemporal placement is not available due to scarring, conjunctival fibrosis or previous surgery, GDD can be implanted in different locations. Thus, superonasal quadrant is more difficult for exposure due to being near superior oblique tendon and nasal structures, inferonasal – more technically challenged and higher risk of tube-corneal touch because of inferiorly shallower AC, inferotemporal – harder exposure and higher risk of plate extrusion and patient discomfort. GDD's plate should be sutured 8-10 mm posterior to the limbus on sclera and must be placed under Tenon's capsule and conjunctiva to reduce the risk of exposure.

The tube of GDD can be positioned in several anatomical sites depending on the patient's ocular status and surgical goals. Anterior chamber is most common position, when tube passes through peripheral cornea into AC. It is the easier technique with predictable results, suitable for standard eyes with adequate chamber depth. Creates the risks of endothelial cells loss and corneal decompensation. Ciliary sulcus or Posterior Chamber position is useful in pseudophakic eyes, in eyes with shallow AC and high risk of corneal damage. It's more difficult technically and can create the iris contact and pigment dispersion.

After vitrectomy, the GDD plate is placed as usual on sclera, but the tube is directed into the pars plana, 3-3.5 mm posterior to limbus, and covered with a patch graft. Complete vitrectomy is essential to avoid vitreous blocking the tube.



Figure 5. *Graphic image of Ahmed ClearPath®*

Pars plana tube placement is preferable after corneal transplantation, when shallow anterior chamber, extensive peripheral anterior synechiae, pediatric eyes with corneal risk and in patients undergoing combined PPV+GDD implantation [77-81].

Today, antiglaucomatous drainage devices come in various sizes, materials, and designs, with or without valves to regulate IOP. The non-valved **Ahmed ClearPath®** was developed as part of Ahmed family of implants and has the pre-threaded tube design to reduce the error in tube positioning, rounded posterior plate to minimize the conjunctival stress and low-profile plate to avoid the erosion risk (**Figure 5**). It requires the intraoperative ligature, stenting and fenestrations to manage the flow [82, 83]. Newest ClearPath® ST offers a smaller profile tube of 127 µm and a 457 µm outer diameter. This option allows surgeons for more flexibility to accommodate different patient needs and surgical preferences.



Figure 6. Graphic image of Aurolab Aqueous Drainage Implant

The non-valved **Aurolab Aqueous Drainage Implant** is a GDD developed by Aurolab, the manufacturing division of Aravind Eye Care System, India (**Figure 6**). It was modeled to offer a low-cost alternative for managing refractory glaucoma. It is made from medical-grade silicone, with the plate size – 350 mm² and ligature and intraluminal stenting to prevent postoperative hypotony. Aurolab creates a subconjunctival filtering bleb by draining aqueous humor through a silicone tube into episcleral plate, which allows for gradual absorption into surrounding tissue. The mechanism is very similar to Baerveldt GDD, but due to significantly

lower cost is suitable for the large-scale use in public health systems [84-87].

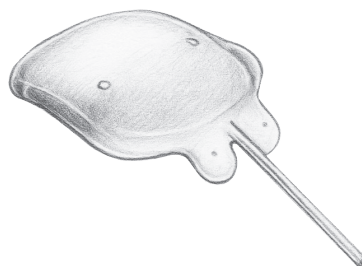


Figure 7. Graphic image of Susanna GDD

The **Susanna Glaucoma Drainage Device** was developed by Dr. Remo Susanna, Brazil, designed as a non-valved implant intended as a low-cost alternative for managing the refractory glaucoma (**Figure 7**). It follows up the successful design principles of other large-plate GDDs, but its thinner plate with fenestrations enhances stability, making it less mobile comparing with other GDDs. Some authors presented 86.4% success rate with very few severe complications (4.5%), another publication states that at 6 months successful rate were 73% for neovascular and 86% for refractory glaucoma [88 - 90].

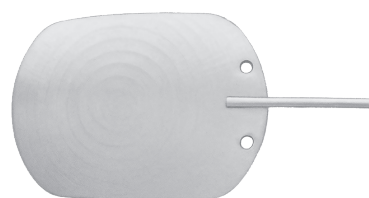


Figure 8. Graphic image of Virna GDD

Virna GDD is the newest non-valved implant designed to lower the IOP. This device was developed by Rohto Corp., Indonesia, to cover the local need for glaucoma drainage implants (**Figure 8**). Virna designed for use in advanced or uncontrolled glaucoma cases [91, 92].

The **Paul® Glaucoma Implant** is a newer generation non-valved glaucoma drainage implant designed to lower the IOP in patients with refractory or advanced glaucoma (**Figure 9**). It was developed by Advanced Ophthalmic

Innovations, France, in collaboration with Dr. Paul Chew and designed to improve the safety and efficacy while minimizing complications associated with the earlier devices.

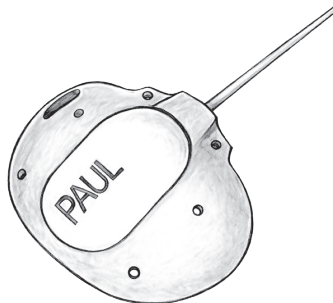


Figure 9. Graphic image of Paul® Glaucoma Implant

Main advantages are thinner and more flexible tube to decrease corneal endothelial cells loss, smaller inner diameter (0.127 mm) to lower the risk of postoperative hypotony, lower-profile plate made from biocompatible medical-grade silicone to minimize the conjunctival erosion. In accordance with some clinical studies IOP reduction after Paul GDD implantation is 50-60%, medication burden reduced by 2-3 medications in average and lower complications rate [93 - 98].

It should be noted that pediatric versions of drainage devices have recently begun to be actively used in the treatment of glaucoma [83]. The most recognized GDD for implantation in children are Ahmed FP8, Molteno, Baerveldt and Paul usually with some modifications. The choice depends on the child's age, eye size, glaucoma type and surgeon preference. Here are more risks for encapsulation and fibrosis due to stronger healing response in children. Tubes may need revisions or repositioning as the eye grows [99 - 102].

Non-valved devices include:

- Molteno (IOP Inc., USA, and Molteno Ophthalmic Limited, New Zealand),
- Baerveldt (Advanced Medical Optics, Inc., USA),
- Eagle Vision implants (Eagle Vision, Inc., USA),
- Ahmed ClearPath (New World Medical,

Inc., USA)

- Aurolab Aqueous Drainage Implant (Aurolab, Aravind Eye Care System)
- Paul Glaucoma Implant (PGI, Advanced Ophthalmic Innovations, Singapore).

Valved devices include:

- Krupin Eye Valve with Disk (Hood Laboratories, Pembroke, MA),
- Ahmed Glaucoma Valve (AGV, New World Medical, USA),
- Schockmed valve (a modified Ahmed valve using the Shocket's ACTSEB technique)



Figure 10. Graphic image of Eye Watch System

Some newly designed GDDs can be adjusted post-operatively to reduce the incidents of hypotony and hypertension and thus creates a 3rd group: GDD with adjustable valve. These devices include the eyeWatch (eW, Rheon Medical, Switzerland) (**Figure 10**). The eyePlate has a deformable silicone tube that can undergo targeted compression to alter its cross-sectional area and thereby change fluidic resistance. Post-operatively IOP may be changed non-invasively by moving the position of an internal magnetic rotor with an external control unit – the eyeWatch Pen. A pilot study announces a complete success rate of 40%. Adjustable GDDs may reduce the need for intra-operative measures, such as tube ligation, and enable better postoperative IOP control. Some clinics used the part of EyeWatch system – eyePlate-300 and eyePlate-200 – as a stand-alone GDD [103, 104, 105].



Figure 11. *Graphic image of GORE Glaucoma Drainage Implant*

The **GORE Glaucoma Drainage Implant** (GORE GDI) is an investigational non-valved glaucoma drainage device developed by W.L.Gore & Associates, Inc., USA, (www.gore.com) using expanded polytetrafluoroethylene (ePTFE), a biocompatible, microporous material to prevent the excessive scarring (**Figure 11**). This device demonstrates an improved tissue integration than silicone- or propylene-based GDDs and allow to expect potentially lower rates of encapsulation [106, 107]. Considering modern trends in development of drainage surgery, it should be noted that many scientists in different medical centers and countries work on the creation of new GDD. It is noteworthy that these developments do not modify current devices, but carry fundamentally new ideas and methods.



Figure 12. *Graphic image of VisiPlate*

VisiPlate is a next-generation anti-glaucomatouse device classified as a hybrid – minimally invasive GDD developed by Avisi Technologies, USA. It is an ultra-thin ($\approx 20 \mu\text{m}$), extremely low-profile, multi-layer nanostructure made of titanium. It is biocompatible and designed to reduce the inflammation and scarring (**Figure 12**).

The device made from 400 nm aluminum oxide plate coated with a 2- μm layer of parylene-C and consists of multiple hexagons with intervening channels to provide slow outflow. Thus, prevent hypotony and create comfortable for the patient less-elevated bleb [108].

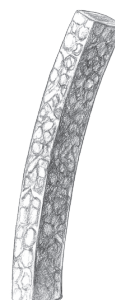


Figure 13. *Graphic image of Alloflo Uveo*

IANTREK, White Plains, USA, and Dr. T. Ianchulev with colleagues introduced new **Alloflo Uveo** uveoscleral dual bio-spacers micro interventional biostent-reinforced cyclodialysis technique to enhance supraciliary aqueous drainage (**Figure 13**). Biostent comprised decellularised sclera allograft tissue microtrephined into a polymer tubular implant [109, 110].



Figure 14. *Graphic image of Calybreye® TGT® Surgical*

The **Calybreye® Titratable Glaucoma Therapy® (TGT) Surgical System's**, Myra Vision Inc., USA, is engineered to put control in the hands of ophthalmologists through unprecedented adjustable outflow (**Figure 14**). The Calibreye Shunt's 10.0×1.6 mm overall dimensions and very low profile (0.30 mm thick) intended to provide posterior bleb

formation to increase aqueous drainage. has 3 distinct outflow channels: the standard is open in all settings, medium and large channels are valve controlled. In accordance with press release of Myra Vision, they secured FDA Approval to initiate US IDE Study in Glaucoma. (www.myravision.com)

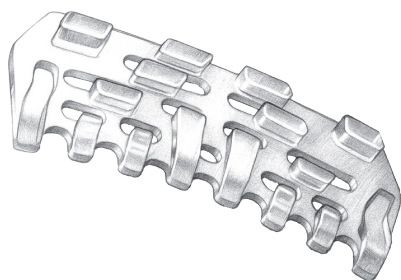


Figure 15. *Graphic image of Intercil® Uveal Spacer*

The **Intercil® Uveal Spacer**, Ciliatech SAS, France, is the first ever Cilioclinal Interpositioning Device (CID), belongs to a new category of glaucoma procedure known as the Cilioclinal Interpositioning. Intercil® doesn't enter the anterior chamber, is ab-externo implanted and bleb frees (**Figure 15**). Ciliatech engineered it to create the localized gap between the ciliary muscle and the sclera. This spacing decreases the resistance to the passage of aqueous humour into this area and will lead to a physiological reduction of IOP by re-establish the natural uveo-scleral flow to the choroidal circulation. Trapezoidal shaped implant ($7 \times 3.4 \times 0.6$ mm) is a non-resorbable one-piece device made of hydrophilic acrylic and characterized by high biocompatibility. It designed with grooves and corrugations to maximize the outflow from the anterior chamber to the suprachoroidal space [111]. Ciliatech are currently working with key regulatory bodies around the world in order to obtain formal licenses in preparation for formal commercialization. Intercil® received CE mark approval in August 2025 (www.cilia.tech).

Market Scope (www.market-scope.com) recently published the 2025 Glaucoma Surgical Device Market Report, which provides an analysis based on six years of

global sales data and evaluates emerging trends for future projections. According to the report, microstents represent the largest share of utilization (44.8%), followed by canal surgery devices (23.5%), laser-based procedures (12.3%), tube shunts (10.9%), and subconjunctival shunts (8.4%). This information is extremely important for understanding the prevalence of certain devices and, accordingly, surgical techniques. It impartially demonstrates the preferences of surgeons, based on the results of their clinical practice and highlights the further development (**Table 1**).

Narrative review of meta-analysis

There is a vast number of studies dedicated to the GDDs, their advantages and disadvantages, implantation techniques, efficacy and complications. In this review we would like to present the results of latest large multicenter meta-analysis of GDDs. Thus, typically refers to comprehensive research studies that combine data from multiple clinical centers and trials to evaluate the common information in managing of glaucoma. This improves the accuracy of conclusions about treatment efficacy, safety and risk factors.

The meta-analysis performed by Serhan H.A. and co-authors reviewed 8 studies (474 eyes) for comparison between GDD (Ahmed valve and Baerveldt) implanted and trabeculectomy underwent eyes condition. It revealed no significant difference in IOP reduction, but GDDs were associated with higher rates of complications: cystoids macular edema (15% vs 4%, $P < 0.001$), need for revision surgery (11% vs 6%, $P = 0.04$), and uveitic flare (5% vs 0%, $P = 0.001$). However, trabeculectomy had a higher risk of cataract progression (7% vs 1%, $P < 0.001$) [112].

3 studies comprised by Dr. Raja S. and team, a total of 176 individuals with refractory glaucoma, with 107 patients received the AADI and 69 patients received the BGI. This meta-analysis, using a random-effects

Table 1. Main characteristics of GDD

Type	GDD name	Manufacturer	Dimensions	Inner Diameter	Outer Diameter	Material	IOP reduction	Flow control
Valved	Ahmed® Glaucoma Valve FP7/FP8, S2	New World Medical, Inc., USA	184 mm ² / 96 mm ² , 184 mm ²	0.305 mm	0.635 mm	Flexible silicone/Rigid polypropylene	40-50%	Venturi-based valve mechanism
	Krupin Valve with Disk	Alcon Inc., Switzerland/ USA	183 mm ²	0.30 mm	0.64 mm	Medical –grade silicon	40-55%	Pressure-sensitive valve
Adjustable valve	EyeWatch® System with EyePlate	Rheon Medical, Switzerland	250 mm ²	0.30 mm	0.64 mm	Titanium housing/silicon plate	40-55%	External magnetically adjustable valve
Non-valved	Molteno3® Glaucoma Drainage Device S150/D270	Molteno Ophthalmic Ltd, NZ/IOP inc., USA	150 mm ² 270 mm ²	0.305 mm	0.635 mm	Medical-grade polypropylene or silicone	55-60%	Ligatura/ Ripcord/ venting slits
	Baerveldt® Glaucoma Implant	Johnson & Johnson Surgical Vision	250 mm ² 350 m ² / 0.84mm	0.3 mm	0.63 mm	Medical-grade silicone	40-60%	Ligatura/ ripcord/ staged opening
	Ahmed ClearPath®	New World Medical, Inc., USA	250 m ² 350 mm ² / 1.9mm	0.305 mm	0.635 mm	Medical-grade silicon	45-50%	Ligatura/ intraluminal stenting
	Paul® Glaucoma Implant	Advanced Ophthalmic Innovations, Switzerland	342 mm ²	0.127 mm	0.467 mm	Silicone	50-60%	Ligatura/ venting slits
	Aurolab Aqueous Drainage Implant	Aurolab, Aravind Eye Care System, India	350 mm ²	0.3 mm	0.63 mm	Medical-grade silicon	50-65%	Ligatura/ intraluminal stenting
	Susanna Glaucoma Drainage Implant	Kinner Silicone Rubber Industria Comercio Ltda., Brazil	200 mm ² /0.5 mm	230 µm	530 µm	Medical grade, soft silicon	55-65%	Ligation/ fenestration
	Virna® Glaucoma Drainage Device	Virna S.p.A., Italy	350 mm ²	0.30 mm	0.60 mm	Medical-grade silicon	60-70%	Ligation/ stenting/ fenestration
	GORE® Excluder® Glaucoma Drainage Implant	W.L. Gore & Associates, USA	200-230 mm ²	0.30 mm	0.60 mm	Expanded Polytetrafluoro-ethylene (ePTFE, Gore-Tex®)	35-50%	Ligation/ stenting/ fenestration
Non-Valved Hyb	VisiPlate® Glaucoma Drainage Device	Avisi Technologies, USA	96 mm ²	0.07-0.1 mm	0.25-0.3 mm	Composition of alumina and Parylene-C	30-40%	Gradual flow control due to very small inner lumen

model, demonstrated a statistically borderline significant reduction in postoperative IOP in patients with the AADI compared to the BGI group at 3 months (MD: -2.74; 95% CI [-5.47; -0.01]; $p = 0.05$; $I^2 = 0\%$), while at 6 months,

the decrease was not statistically significant (MD: -3.77; 95% CI [-8.42; 0.88]; $p = 0.11$; $I^2 = 68\%$). By the 12 months follow-up, no significant difference in postoperative IOPs was observed between the 2 groups (MD:

-1.19; 95% CI [-3.90; 1.53]; $p = 0.39$; $I^2 = 0\%$). Surgical success rate in 2 groups had a statistically nonsignificant difference and similar overall complication incidence rate [113].

The meta-analysis performed by Raja A. et al. earlier indicates that the Molteno implants demonstrate greater success in sustaining low IOP, however the Ahmed valve exhibits fewer overall complications [114].

Dr. Hong M. included 825 eyes (820 patients) from 6 studies to his meta-analysis comparing the results of Aurolab Aqueous Drainage Implant and Ahmed Glaucoma Valve implantations. Weighted mean difference of the IOP reduction between those implants were 0.58 (95% CI: 0.07 - 1.09) at 3 months, 0.44 (95% CI: 0.11 - 0.77) at 6 months, 2.20 (95% CI: 0.63 - 3.77) at 12 months, and 3.24 (95% CI: 1.73 - 4.75) at the follow-up endpoint; of the reduction in antiglaucoma medication were 0.87 (95% CI: 0.61 - 1.13) at 6 months, 1.04 (95% CI: 0.66-1.42) at 12 months and 0.93 (95% CI: 0.52 - 1.34) at the follow-up endpoint. The pooled odds ratio comparing AADI and AGV were 3.64 (95% CI: 2.44 - 5.45) for the complete success rate and 1.72 (95% CI: 1.24 - 2.39) for qualified success rate [115].

18 studies (946 eyes) follow-up the Paul Glaucoma Implant (PGI) implantation were analysed by Dr. Carla M. and colleagues. The PGI demonstrated mean IOP reduction ranged from 14,8 mmHg to 19,1 mmHg, depending on follow-up duration and patient characteristics. Complete success rates ranged from 38.4% to 75%, while qualified success rates were consistently high, reaching up to 93.2% [116].

32 studies (1221 eyes, 885 patients) were included in the study hold by Dr. Stallworth J. and colleagues. The findings show that Ahmed and Baerveldt shunts substantially reduced IOP for at least 24 months of childhood glaucoma, with similar findings among device types and glaucoma etiologies [117].

Dr. Ramdas W.D. and co-authors from Erasmus Medical Center, Netherlands,

performed the meta-analysis to compare their own results with the 24 studies from PubMed database. It showed no significant differences between patients with and without uveitis in the IOP reduction (44.9% vs 42.8% respectively), postoperative weighted mean difference of - 17.8 mmHg and 2.2 lower number of IOP-lowering medications in uveitic glaucoma compared to -13.2 mmHg and 3.5 in current study, respectively [118].

13 cohort studies (562 eyes) were included in meta-analysis performed by Dr. Pan S.Y. and his team. The effects of trabeculectomy with or without antifibrotic agents, valved or non-valved GDDs and medical treatment were compared. According to their results, valved GDD was more effective in controlling IOP and preserving the corneal grafts. UTherefore, they assume the valved GDD as the preferred surgical approach for the treatment of post-PKP glaucoma [119].

One of the latest reviews Dr. Garg A. stated, that the future of glaucoma surgery lies in balancing the safety and efficacy of traditional GDDs with the minimally invasive nature of MIGS. While GDDs still crucial for advanced cases, MIGS are expected to expand their reach. Innovations in materials, smart technologies, and personalized treatment approaches will shape the future landscape, offering patients safer, more effective options to manage their glaucoma [120].

Conclusion

Summarizing, we can estimate the GDDs as a pathogenetically justified and clinically essential modality in the modern management of glaucoma. The active development of MIGS as another modern method of glaucoma treatment in no way diminishes the role of GDD, forming and expanding a specific niche for each of them. An analysis of current trends indicates that the future modernization of GDDs will focus on three key domains: enhancing safety, improving efficacy, and maximizing patient comfort.

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