Advances in refractive corneal lenticule extraction

Matthias Fuest¹, Jodhir S. Mehta²,³,⁴,⁵*

Abstract:
Refractive errors are the leading cause of reversible visual impairment worldwide. In addition to the desired spectacle independence, refractive procedures can improve quality of life, working ability, and daily working performance. Refractive corneal lenticule extraction (RCLE) is a relatively new technique, dependent only on a femtosecond laser (FS). This leads to potential benefits over laser-assisted in situ keratomileusis (LASIK) including a quicker recovery of dry eye disease, a larger functional optical zone, and no flap-related complications. SMILE, available with the VisuMax FS (Carl Zeiss Meditec AG, Jena, Germany), is the most established RCLE application, offering visual and refractive outcomes comparable to LASIK. SmartSight (SCHWIND eye-tech-solutions GmbH, Kleinostheim, Germany) and CLEAR (Ziemer Ophthalmic Systems AG, Port, Switzerland) are two new RCLE applications that received Conformité Européenne (CE) approval in 2020. In this article, we review refractive and visual outcomes, advantages, and disadvantages of RCLE and also report on the latest advances in RCLE systems.

Keywords:
Cornea corneal surgery, laser refraction, ocular refractive surgical procedures

Introduction
Refractive errors are the leading cause of reversible visual impairment worldwide.[¹] Myopia is the most frequent form of refractive error.[²] The global prevalence of myopia is increasing, with the rate of increment being particularly alarming in many Asian countries.[²,³] Holden et al. predicted that the global prevalence of myopia will rise from 28% (2 billion people) in 2010 to 50% (5 billion people) in 2050.[³] The same study predicted that the global prevalence of high myopia will rise from 4% (227 million people) in 2010 to 10% (938 million) in 2050, making myopia a major public health issue.[³]

Corrective laser refractive surgery is one of the most commonly performed surgeries.[⁴] Refractive surgery has evolved over the decades. Advances in surface ablation techniques have seen a renaissance in its popularity.[⁵,⁶] Presbyopic treatment options have also expanded to include new ablation profiles, intracorneal implants, and phakic intracocular implants.[⁵] With the improved safety and efficacy of cataract surgery, a wider variety of intraocular lens implants, with modified optics, provide more options for refractive correction in carefully selected patients.[¹,⁴,⁵] In addition to the desired spectacle independence, refractive procedures have been reported to improve quality of life, working ability, and daily working performance.[⁷]

Refractive corneal lenticule extraction (RCLE) is a relatively new technique, dependent only on a femtosecond laser (FS), without the need for corneal flap creation.[⁸] Advances in this promising new technology will be discussed in this review.

Developments in Laser Refractive Surgery
The cornea is the most integral structure for refractive correction, being the most
accessible part of the eye and providing two-thirds of the total refractive power.\textsuperscript{[6,9]} Corneal ablation techniques can potentially treat most refractive errors (including myopia, hyperopia, astigmatism, and presbyopia) within a given range, by ablating corneal tissue into a specified shape, by the use of an excimer laser.\textsuperscript{[5,6,10]}

The evolution of keratorefractive surgery began with surface ablation techniques such as photorefractive keratectomy (PRK) that involved epithelial removal.\textsuperscript{[5]} More recently, excimer laser ablation has been used to remove the corneal epithelium directly (transepithelial PRK).\textsuperscript{[11]} The main advantage of transepithelial PRK is that the epithelial layer removal and stromal excimer ablation are performed sequentially, simplifying the treatment and saving time for patient and surgeon. Most reports have suggested that healing time and visual outcomes do not vary greatly among the different techniques of epithelium removal.\textsuperscript{[12]} In recent years, PRK (and in particular transepithelial PRK) has seen a revival in interest.\textsuperscript{[13]} Surface ablation has several potential advantages. Surface ablation has several potential advantages, e.g. the avoidance of flap-related complications and a greater residual stromal thickness (RST).\textsuperscript{[14]} Hence, it is useful in patients with thin corneas.\textsuperscript{[5]} While the refractive predictability of surface ablation is comparable with laser-assisted in situ keratomileusis (LASIK), myopic regression may be more common.\textsuperscript{[15]} Other disadvantages of PRK are prolonged pain, because of slow epithelial healing, and subepithelial scarring (haze), that can form as an unwanted healing response in the Bowman layer and anterior corneal stroma.\textsuperscript{[14]} Low-dose topical mitomycin-C (0.02%–0.04%) is used after excimer laser treatment to reduce haze formation.\textsuperscript{[17]}

Using a microkeratome to create a hinged flap, under which the corneal stroma was ablated by means of an excimer laser, LASIK was introduced in 1990.\textsuperscript{[18]} LASIK preserves the central corneal epithelium, which improves the comfort during the early postoperative period, allows for rapid visual recovery, and reduces the wound healing response.\textsuperscript{[10]} From 2004, FS was introduced to improve LASIK flap creation (FS-LASIK), with fewer flap-related complications (e.g., incomplete, buttonholed, free, or irregular flaps). The FS allowed customization of several flap parameters, for example, hinge position, flap diameter, hinge width, and flap thickness. This led to more precise and less variable flaps, and a lower incidence of epithelial ingrowth.\textsuperscript{[19,20]}

Laser refractive surgery is now widely recognized as safe and effective, yielding excellent results in patients with low-to-moderate amounts of refractive error.\textsuperscript{[9]} A review of 97 studies published since 2008 showed that up to 99.5% of patients who underwent LASIK met uncorrected distance visual acuity of better than 20/40 (considered spectacle independent), as many as 98.6% had refractive targets within ±1.0 diopter, and almost 98.8% were satisfied with their outcome.\textsuperscript{[21]} Moreover, complications that could lead to visual loss, such as corneal ectasia or infection, were very rare with laser eye surgery.\textsuperscript{[4]}

**Refractive Corneal Lenticule Extraction**

A precursor to modern RCLE was first described in 1996 using a picosecond laser to generate an intrastromal lenticule that was removed manually after lifting a flap in human donor eyes.\textsuperscript{[22]} However, significant manual dissection was required leading to an irregular surface. The switch to FS improved the precision, and studies were performed in rabbit eyes in 1998\textsuperscript{[23]} and in partially sighted eyes in 2003.\textsuperscript{[24]}

Following the introduction of the VisuMax FS [Carl Zeiss Meditec AG, Jena, Germany; Figure 1b], in 2007, intrastromal lenticule refractive correction was reintroduced, in a procedure called femtosecond lenticule extraction (FLEX). Sekundo et al. published the 6-month results of the first 10 fully seeing eyes treated in 2008\textsuperscript{[8]} and a larger series followed.\textsuperscript{[28]} The procedure still included the creation of a hinged flap, under which the stromal lenticule was extracted, both cut by FS
LASIK disrupts the corneal stroma, but visual recovery time was longer due to the lack of optimization in energy parameters and scan modes; further refinements led to much improved visual recovery times.[29]

Following the successful implementation of FLEX, the procedure was advanced to small incision lenticule extraction (SMILE or ReLex-SMILE). In SMILE, a dissector is passed through a small 2–5-mm incision to separate the lenticular interfaces and allow the lenticule to be removed, without the need to create a flap [Figure 2].[30] The SMILE procedure has gained popularity following the initial results of the first prospective trials.[30,31] In 2011, Zeiss received European conformity approval (Conformité Européenne, CE) for the SMILE application for corrections of myopia from −0.5 to −10.0 diopters (D) and astigmatism up to −5.0 D. SMILE was then approved by the US Food and Drug Administration (FDA) for the treatment of myopia in 2016. In 2018, astigmatism correction was added, so that the current SMILE FDA approval is for myopia of −1.0 to −10.0 D and astigmatism of −0.75 to −3.0 D with a maximum spherical equivalent (myopia plus half of the astigmatism) correction of no more than −10.0 D.[29]

Subsequently, other manufacturers have followed. The SCHWIND ATOS FS [SCHWIND eye-tech-solutions GmbH, Kleinostheim, Germany; Figure 1a] received CE approval for its SmartSight application in July 2020 and the FEMTO LDV Z8 platform [Ziemer Ophthalmic Systems AG, Port, Switzerland; Figure 1c] for its corneal lenticule extraction for advanced refractive (CLEAR) correction program in April 2020 [Table 1].[26,27]

In this review, we report on the latest advances in modern flap-free RCLE. When a particular RCLE application is discussed, we use the according abbreviations or acronyms [SmartSight, SMILE, or CLEAR; Table 1].

Advantages of Refractive Corneal Lenticule Extraction

With laser refractive surgery already achieving excellent clinical visual outcomes, it is often difficult to demonstrate that newer procedures are superior to the established techniques. A recent meta-analysis revealed a similar safety, efficacy, and predictability of SMILE compared to FS-LASIK in the correction of myopia.[32] Similarly, Ang et al. recently published their results of a contralateral randomized clinical trial, in 70 patients, performing FS-LASIK in one eye and SMILE in the other. They found excellent, comparable outcomes in terms of refractive predictability, efficacy, and safety for both procedures at 3 months and also after 1 year of follow-up.[33] Vision-related quality of life has also been found to be comparable between SMILE and LASIK.[33-38] However, potential benefits of RCLE over LASIK are a quicker recovery of dry eye disease (DED), a larger functional optical zone, and no flap-related complications.[36]

DED is one of the most common side effects after laser refractive surgery.[37] LASIK disrupts the corneal nerves during the creation of the corneal flap and during the stromal ablation. RCLE is a flapless procedure with a smaller corneal incision that results in less damage to the anterior corneal innervation, making it theoretically less prone to DED.[37,38] Both RCLE and LASIK induce a transient worsening in dry eye parameters, but there is evidence showing that RCLE has a less negative impact on ocular surface parameters and allows for an earlier recovery.[37,39] Analyzing ocular surface changes in 15 patients post SMILE and 32 post FS-LASIK, Gao et al. found lower fluorescein staining scores after 1 week, longer tear breakup times after 1 and 3 months, higher central corneal sensitivity at 1 week, 1 and 3 months as well as lower levels and a faster decrease of interleukin-6 and nerve growth factor in tears following SMILE compared to FS-LASIK.[40]

Corneal inflammation may play a role in postoperative DED after SMILE and FS-LASIK, but more evidence is needed to support this hypothesis. Studies have also shown faster corneal nerve regeneration[41] and improvement of corneal sensitivity[42] following SMILE, supporting the evidence for less DED.

It was previously shown that the strength of the corneal stroma decreases from anterior to posterior.[43] By preservation of a stronger anterior stromal lamella, RCLE may offer a theoretical biomechanical advantage over LASIK. Even though this has been shown using theoretical[44] and laboratory experiments,[45] three

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randomized controlled trials found no statistically significant differences between SMILE or flap-based procedures concerning corneal hysteresis or corneal resistance factor, as measured with the Ocular Response Analyzer (Reichert Inc., Depew, USA). However, the authors speculated that the current commercially available diagnostical technology might not be optimal for detecting such differences in biomechanical stability.

Using the Pentacam (Oculus Optikgeräte GmbH, Wetzlar, Germany), it has been demonstrated that the postoperative “effective/functional optical zone” is significantly larger in SMILE than that in FS-LASIK based on the same diameter of optical treatment zone. This could potentially reduce pupil-related photopsia, particularly in patients with large mesopic pupils.

### Table 1: Overview of current femtosecond lasers (FS) that offer Conformité Européenne (CE) marked modern refractive corneal lenticule extraction (RCLE) applications²⁵-²⁷

| FSL                        | ATOS SCHWIND                  | VisuMax Zeiss                  | Z8 Ziemer                      |
|---------------------------|-------------------------------|-------------------------------|-------------------------------|
| RCLE program              | SmartSight                    | SMILE                         | CLEAR                         |
| Possible refractive cuts  | Flap/lenticule                | Flap/lenticule                | Flap/lenticule/arcuate incisions |
| Weight (kg)               | <275 kg                       | 870                           | 215                           |
| Approval                  | CE                            | FDA/CE                        | CE                            |
| Treatment range (diopters, D) |                  |                               |                               |
| Sphere                    | -0.5 to -12.0                 | -0.5 to -10.0                 | -0.5 to -10.0                 |
| Cylinder                  | 0 to -6.0                     | 0 to -5.0                     | 0 to -5.0                     |
| SEQ                       | -0.5 to -14                   | -0.5 to -12.5                 | -0.5 to -12.5                 |
| Repetition rate           | Up to 4 Mhz                   | 500 kHz                       | Up to 20 Mhz                  |
| Energy per pulse (nJ)     | 75-135                        | 110-150                       | <<100                         |
| Laser/patient interface   | Machine-fixed                 | Machine-fixed                 | Handheld                      |
| Contact glass on suction system | Curved (20 mm)             | Curved (22 mm)                | Flat                          |
| iOCT                      | No                            | No                            | Yes                           |
| Automatic detection of pupil | Yes                           | No                            | Yes                           |
| Pupil central offsetting  | Yes                           | No                            | Yes                           |
| Cyclotorsion compensation | Yes                           | No                            | Yes                           |
| Centration                | Eye tracking guided (semi-automated) centration | Manual Eye tracking guided (semi-automated) centration | |
| Real-time video recording | Yes                           | Yes                           | No - schematic graphic        |
| Postsuction lenticule adjustment | Lateral electronic adjustment | Nil                           | Lateral electronic adjustment |
| Laser pattern             | Arc segments-centrifugal/centrifugal | Centripetal/centrifugal     | Spiral raster                 |
| Lenticule shape           | No side cut                   | 10 µm side cut                | No side cut                   |
| Number of treatments performed | 700+                          | 3.5 million+                  | 400+                          |
| Incisions (mm)            | 1, 2-5                        | 1-3, 2-5                      | 1-2, 1.5-4                    |
| Advantages                | Recentering after docking     | Established technique         | Recentering after docking     |
|                          | Eye tracking with pupil recognition and cyclotorsion compensation | Many treated cases            | Eye tracking with pupil recognition and cyclotorsion compensation |
|                          | Mobile device                 | Low IOPs during suction       | Mobile device                 |
|                          | High repetition rate          |                                 | Small footprint               |
|                          | Lower pulse energy            |                                 | High repetition rate          |
|                          |                                 |                                 | Low pulse energy              |
| Disadvantages             | New procedure lacking clinical experience | No recentering after docking | New procedure lacking clinical experience |
|                          |                                 | No eye tracking                |                               |

iOCT=Intraoperative OCT, OCT=Optical coherence tomography, IOP=Intraocular pressure, SEQ=Spherical equivalent, FDA=US Food and Drug Administration, CE=Conformite Européenne, FSL=Femtosecond laser

### Refractive Corneal Lenticule Extraction

**Complications and Management**

Despite the fact that RCLE has been shown to be a safe technique, it is not free of complications and one of the disadvantages of the procedure has been a longer learning curve for the surgeon, when compared with conventional FS-LASIK procedures. With the introduction of FS-created flaps, and improved excimer laser ablation profiles, FS-LASIK has demonstrated a remarkable consistency among surgeons of different experiences. However, RCLE still requires the manual removal of the lenticule following laser creation [Figure 2].

Patient head and eye movement can lead to suction loss, a common and feared complication of SMILE,
with an incidence of 0.5%–4.4%. Suction loss is more commonly observed after SMILE, because of the lower suction pressure and longer duration compared to FS-LASIK flap creation. Several factors can predispose a patient to suction loss, for example, the longer duration of suction to create the posterior refractive and the anterior nonrefractive planes in the stroma. Ocular factors include a small palpebral aperture, loose corneal epithelium, excessive reflex tearing, and poor fixation. Individual factors include patient anxiety and inability to follow instructions. However, Reinstein et al. proposed a clear management protocol and decision tree for this event. Options range from continuing with SMILE or converting to LASIK to complete the treatment on the same day, to aborting the surgery and repeating SMILE or another laser refractive surgery at a later date. In the vast majority of cases, outcomes are excellent with no significant differences in vision or refraction compared to uncomplicated cases.

Identifying the lenticule edge at the beginning of the surgery is crucial to ensure lamellar separation in the correct plane and to prevent lenticule mis-dissection. The surgeon can for instance make mistakes while performing the dissection and inadvertently dissect the posterior surface first. This would then compact the lenticule into the anterior cap, making anterior plane dissection more difficult and subsequent lenticule extraction more challenging. However, with advances in technology and improvements in surgical technique and instruments, it has been demonstrated that the majority of cases, outcomes are excellent with no significant differences in vision or refraction compared to uncomplicated cases.

As RCLE has gained popularity over the last decade, the treatment has continuously been optimized. Initially, the main issue was the delayed visual recovery relative to LASIK. However, detailed research into nomograms, energy levels, and spot spacing has significantly improved visual recovery, without compromising the ease of lenticule separation.

Postoperative complications after SMILE are similar to those after LASIK. Diffuse lamellar keratitis (DLK) after SMILE has been reported to occur with an incidence of 0.45%. It usually resembled the appearance following LASIK but may rarely present as a sterile multifocal inflammatory keratitis. However, topical steroid therapy resolved all reported cases of DLK, and in all cases, there were no sequelae and no adverse effect on refractive or visual outcome. Epithelial ingrowth after SMILE has been described in cases following incisional tears of the smile pocket or inadvertent implantation of loose epithelium following lenticule removal. It can be treated by meticulous irrigation of the pocket, manual scraping using a blunt spatula, and forceps removal of epithelial strands. Particularly, eccentric epithelial nests have also been successfully treated with a neodymium-doped yttrium aluminum garnet (Nd:YAG). In contrast to LASIK, which can be retreated by a flap re-lift, SMILE enhancement is more complicated. In cases of primary under/overcorrection or myopic regression following RCLE, different techniques have been applied. One of the most popular choices is surface ablation, which preserves the flap-free approach of the primary procedure and a higher RST. However, the aspect of pain and a slow visual recovery might render it less appealing, particularly if it needs to be done bilaterally. The SMILE cap can be converted into a FS-LASIK flap for secondary excimer laser application (CIRCLE). Thin-flap LASIK, with a new flap anterior to the SMILE interface, can also be created. These options offer a faster visual recovery, however, at the price of flap creation. Each enhancement method usually requires adjusted nomograms and generates specific tissue responses with a different impact on corneal biomechanics dependent on the previous SMILE parameters, especially the cap thickness.

SMILE is now a mature and established procedure that provides patients with safe and effective outcomes with current reports demonstrating that the visual and refractive results are similar to LASIK. However, it is still a first-generation procedure, and as we have seen LASIK mature over the years, through multiple iterations driven in large by improvements in technology, we expect to see the same with SMILE.
Characteristics of the Different Refractive Corneal Lenticule Extraction Systems

CLEAR is a new CE marked proprietary RCLE procedure for myopic correction of sphere −0.5 D to −10 D and cylinder 0 D to −5 D [Table 1]. CLEAR is an optional software upgrade on the FEMTO LDV Z8 platform. The platform itself can also be used for cataract surgery (i.e., femtosecond laser-assisted cataract surgery), corneal transplantation, pterygium surgery, LASIK flap creation, and presbyopic correction (pockets for inlays). The Z8 is based on a low-energy (<<100 nJ) high-frequency (up to 20 MHz) concept, where the miniaturized scanning optic, integrated into the handpiece and its high numerical aperture, creates highly focused laser pulses. Advantages of the low-energy concept are the decreased stromal gas generation and an accurate laser focus. The high-frequency repetition rate leads to overlapping pulses. The laser pulses are guided from the laser source through an articulated moveable arm to a handpiece that is adaptable in position and height with a very close working distance to the eye. In addition, the FEMTO LDV Z8 offers a wide range of centration options and the options of recentering the treatment area after having performed the docking, which is not possible with the current VisuMax Laser System.

Recently, Izquierdo et al. reported on their initial experience with CLEAR. They marked the visual axis on the slit lamp before CLEAR. If the surgeon was not satisfied with the centration after docking, the Z8 allowed recentering of the treatment area, including correction of cyclotorsion by adjustment on the touchscreen monitor, without having to release the suction. This ensured accurate centration on the visual axis and adjusted cyclotorsion, which is especially important in patients with cylindrical corrections. In 5 eyes of 5 patients treated, two small incisions of 3.0 mm width were cut separately at 35° and 145° positions, with an entrance angle of 90°. Each incision allowed for the posterior and anterior surfaces of the lenticule to be delineated directly and independently aiding one of the most difficult and important steps of the lenticule extraction technique. In CLEAR, the surgeon can adjust the distance between the incisions and the angle position for greater comfort and can also decide if 1 or both incisions were created and used. In all the cases reported by Izquierdo et al., the lenticule had a diameter of 6.5 mm without adding any additional edge thickness. During the cutting of the lenticule, the applied vacuum level to the eye was 700 mbar, the same as habitually used for LASIK flaps with the Z8. A complete dissection and removal of the lenticule was achieved in all cases without any intraoperative complications, and at postoperative day 1, all patients had a clear cornea. In the postoperative period, 1 patient presented with a mild stromal haze in the interface that resolved with 2 weeks of topical corticosteroid treatment 4 times a day. The refractive and visual outcomes were excellent and comparable to established SMILE and LASIK procedures. However, follow-up was only 1 month. Izquierdo et al. speculated that the guiding incisions and tunnels leading to the anterior or posterior interface have the potential to shorten the learning curve for surgeons and lower the complication rate of RCLE, which needs to be verified in future larger case series.

Another ability of the Z8 not mentioned by the previous authors is the inbuilt intraoperative OCT (iOCT) [Figure 3]. This feature may also be used following lenticule creation. This could be useful in more complicated cases, for example, CLEAR in posttransplant cases.

Performing experimental CLEAR on enucleated porcine eye balls during a medical exhibition in China, Wang et al. found longer time on suction peak pressure, total laser application, and total surgery time spent during CLEAR compared to SMILE. However, the CLEAR group included OCT scanning and offsetting before performing the laser procedure, which took more time for these crucial steps. The anterior and posterior lenticule surfaces analyzed by scanning electron microscopy were smoother in CLEAR compared to SMILE. This atraumatic, smooth cutting of high-frequency low-energy FS has been shown before in rabbit corneas by Riau et al. using the FEMTO LDV Z6 predecessor model. By placing laser spots directly adjacent to each other, the Z6 and Z8 are believed to create a smooth stromal interface.

Further comparative studies are necessary to evaluate whether these differences in lenticule surface roughness created by different laser platforms could result in different clinical performances with regard to postoperative refraction, optic quality, and tissue response.

Figure 3: Intraoperative optical coherence tomography (iOCT) can be used in CLEAR on the FEMTO LDV Z8 platform (Ziemer Ophthalmic Systems AG, Port, Switzerland)
Taiwan J Ophthalmol

The FEMTO LDV Z8 uses a flat contact glass on its docking system, which is directly in contact with the cornea during laser applications. Previous studies have proven that the flat suction cone in FEMTO LDV models induced a higher intraocular pressure (IOP) than the curved suction cone of the VisuMax. [78,79] Although extremely uncommon, [80] the higher IOP theoretically increases the risk of ocular vessel occlusions or visual field loss. However, these complications have not been reported following FS-LASIK with the Z8 at the same suction pressure. [81,82] An advantage of the high suction is the reduction in the risk of suction loss, during the procedure. However, more clinical cases are required to validate the latter comment.

SmartSight is the new RCLE application available with the SCHWIND ATOS FS that received CE approval in 2020. [26,83] The relatively small design and “lightweight” of the ATOS laser, similar to the Z8, is supposed to allow mobility of the device.

For SmartSight, the ATOS offers eye tracking, featuring pupil recognition, and cyclotorsion compensation to allow precise centering. Eye tracking, the centering options (pupil, vertex, and user-defined offset), and the cyclotorsion compensation are known to be particularly helpful in the correction of astigmatism. [26]

The ATOS has a curved patient interface, which can be used for SmartSight and for making flaps in FS-LASIK. The curved geometry aims to reduce the pressure on the eye and patient discomfort during the contacting process. A sophisticated contact glass design allows the cutting of large flaps and RCLE treatment diameters up to 9.6 mm. [26]

The ATOS FS uses a high repetition rate up to 4 MHz to allow a quick cutting and has a high numerical aperture for optimized resolution. The energy per pulse of 75–135 nJ is lower than SMILE but higher than in CLEAR [Table 1].

According to the manufacturer, like in CLEAR, SmartSight does not use any side cuts and consequently does not have a minimal thickness as in SMILE. The lenticule tapers toward the periphery following a refractive progressive true transition zone, in an attempt to reduce epithelial remodeling and refractive regression. [26,83]

Pradhan and Mosquera described no serious complications during the treatment of their first 185 SmartSight patients. They did witness a slight tendency for overcorrection and improvement of uncorrected vision took a little longer than with LASIK, with 70% of patients reaching 6/9, decimal 0.6 by day 1, improving to >70% in 6/6 by 1 week. [83] However, this has been described for RCLE previously and is not considered a limitation. [84] A greater number of clinical cases will allow for further nomogram refinement to improve outcomes.

ZEISS is currently developing a new generation FS named VISUMAX 800. [25] It incorporates a significantly faster laser source, which is intended to bring the time needed for a complete lenticule creation down to approximately 10 s. The new device will feature computer-assisted centration and cyclotorsion compensation. As compared to the current version of the VisuMax, the VISUMAX 800 will have a significantly smaller footprint as well. CE certification and commercialization are anticipated for the second half of 2021. [25]

Alcon (Geneva, Switzerland) and Johnson & Johnson (New Brunswick, USA) are also working on RCLE programs for their laser systems. However, further details were not published at the time of composing this review (March 2021).

To summarize, 2020 has seen the approval of two new devices for RCLE. Future studies will have to investigate whether capabilities of the new machines, for example, eye tracking, can further improve the visual outcomes already achieved, and make them more universally consistent.

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Conflicts of interest
The authors declare that there are no conflicts of interests in this paper.

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