A review of the surgical options for the correction of presbyopia

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ABSTRACT
Presbyopia is an age-related eye condition where one of the signs is the reduction in the amplitude of accommodation, resulting in the loss of ability to change the eye’s focus from far to near. It is the most common age-related ailments affecting everyone around their mid-40s. Methods for the correction of presbyopia include contact lens and spectacle options but the surgical correction of presbyopia still remains a significant challenge for refractive surgeons. Surgical strategies for dealing with presbyopia may be extracoronal (conical or scleral) or intraocular (removal and replacement of the crystalline lens or some type of treatment on the crystalline lens itself). There are however a number of limitations and considerations that have limited the widespread acceptance of surgical correction of presbyopia. Each surgical strategy presents its own unique set of advantages and disadvantages. For example, lens removal and replacement with an intracoronal lens may not be preferable in a young patient with presbyopia without a refractive error. Similarly treatment on the crystalline lens may not be a suitable choice for a patient with early signs of cataract. This article is a review of the options available and those that are in development stages and are likely to be available in the near future for the surgical correction of presbyopia.

INTRODUCTION
Presbyopia is an age-related reduction in the amplitude of accommodation and leads to the loss of ability in changing the eyes’ focus between far and near. The correction of presbyopia without resorting to spectacles and contact lenses (CLs) remains the Holy Grail for refractive surgeons as well as the billions of patients with presbyopia.

Numerous accommodative and pseudoaccommodative approaches to treat presbyopia surgically exist. Each has its own benefits and limitations, and may involve some degree of compromise between the distance and near visual acuities (VA). Accommodative approaches attempt to restore the true, dynamic and continuous range of the defocusing ability of the eye. Pseudoaccommodative approaches provide functional near vision from a variety of non-accommodative factors.

This review provides an overview of the options that are currently available for the surgical management of presbyopia.

PSEUDOACCOMMODATIVE APPROACHES

Corneal approaches

Excimer laser procedures

Monovision

Monovision with an excimer laser is a well-established technique that corrects one eye for distance vision (usually dominant eye) and the other eye for near vision, resulting in intentional anisometropia.1 This aim is to give functional near and distant VAs without the need for glasses. The mechanism that enables monovision to succeed is interocular blur suppression.

Studies have reported success rates ranging from 80–98%1–5 for monovision post laser vision correction (LVC), 91% for monovision after cataract surgery and 95% following clear lens extraction6 with good satisfaction. Surgically induced monovision is associated with a higher success rate than with CLs (91–98%), but it is unclear whether this is because it is harder to reverse the procedure or because of a multifocal corneal shape in LVC.

Limitations of monovision include compromising visual function, such as reduced low contrast VA and contrast sensitivity (CS), inability to incorporate an intermediate vision correction without compromising distance vision or near vision, reduced stereopsis, and small-angle esotropic shift.7–9

Multifocal corneal ablation
Multifocality achieved by excimer ablation sometimes known as presbyLasik, is interesting to refractive surgeons because it is familiar, seems less invasive than intraocular surgery and could theoretically be more controllable. However, this is against the conventional thinking for LVC where one usually attempts to minimise the higher order aberrations.

A variety of presbyopic LVC procedures exist.10–12 In peripheral presbyopic LVC, the peripheral cornea is ablated to create negative peripheral asphericity. Thus the central cornea is for distance vision and the mid-peripheral cornea for near vision (eg, Nidek Advanced Vision Excimer Laser; NIDEK, Gamagori, Japan).11 In central presbyopic LVC, the central area is ablated for near vision and the periphery for distance vision (eg, Supracor, Technolas Perfect Vision GmbH, München, Germany); and Pulsar (CustomVis, CV Laser, Perth, Australia).

Although optically the results are predictable and good, some patients find it difficult to adapt to the compromise and others are dissatisfied by the minor loss of distance VA.12,13

Presbyond Laser Blended Vision (Carl Zeiss Meditec, Jena, Germany) is an optimised laser treatment method attempting to improve on conventional monovision. The dominant eye is treated for distance vision to almost plano and the non-dominant eye is corrected to be slightly myopic for near vision to −1.5 D. This monovision treatment is enhanced by the use of a wavefront-optimised ablation profile to create a continuous refractive power gradient for the whole optical zone of the
cornea. Studies show that this treatment is a well-tolerated and effective procedure for treating patients with presbyopia.14–16

More recently, SCHWIND eye-tech-solutions (Kleinostheim, Germany) introduced its PresbyMAX software. This is a biaxial corneal modulation technique, based on the creation of a central hyperpositive area for near vision and leaving the peripheral cornea for far vision. Uthoff et al17 reported good distance and near visual outcomes in a 6-month follow-up study (table 1).

While multifocal LVs represent a promising avenue for future presbyopic correction, outcome data is relatively sparse compared with other modalities.22–28

Conductive keratoplasty

Conductive keratoplasty (CK) is the successor of laser thermokeratoplasty. CK uses the application of low frequency radio waves to ‘shrink’ collagen fibrils within the mid-peripheral cornea. This causes a net steepening on the central cornea and thus increases the positive power of the eye. Radiofrequency energy is typically 0.6 W with a 0.6-s treatment time,25 delivered through a fine tip inserted into the peripheral corneal stroma in a ring pattern outside of the visual axis. Eight to 32 treatment spots are placed in up to three rings in the corneal periphery (6-mm, 7-mm and 8-mm optical zones) and striae form between the spots and create a band of tightening to steepen the cornea primarily to create monovision. Although this has shown to be a relatively safe technique and may present theoretical advantages over flap creation techniques (less invasive and no flap-related complications), long-term studies report high rate of regression and hence this is not a popular technique at present.18 26–28

Intrastromal femtosecond ring incisions

Although the primary application of femtosecond laser has been its use in the creation of Laser-Assisted in situ Keratomileusis (LASIK) flaps, its precision and safety features makes it a useful tool for many types of corneal refractive surgery, including intrastromal treatments. Typically, five concentric rings in the cornea stroma between 2 mm and 4 mm from the line of sight are created using a femtosecond laser. Studies with INTRACOR (Technolas Perfect Vision GmbH, München, Germany) have shown the technique to be efficient and safe.20 21

The main advantage of INTRACOR is that the corneal surface is not cut. The ring structure induces a localised biomechanical change in the tissue causing a slight central steepening of 1–2 dioptres (D). This steepening changes the spherical aberration (SA) and corneal asphericity, resulting in improvement in the near vision.29–32 To date, the results reported have shown an overall improvement of uncorrected near VA (UCNVA),19 29–32 However, some studies report no improvement in UCNVA at 1 month,19 30 reduced best distance corrected VA (BDCVA),19 31 and anterior corneal protrusion after hyperopic LASIK followed by INTRACOR.13 The treatment is usually performed in the non-dominant eye only. Further study is required on this treatment modality.

Conical inlays

Conical inlays (CIs) are intrastromal implants which are placed underneath a LASIK flap or into a femtosecond laser created corneal pocket. The pocket technique has a number of potential advantages: the majority of peripheral corneal nerves are preserved, allowing corneal sensitivity to be maintained, they are additive, do not remove tissue, preserve future options for presbyopic correction and may be used in pseudophakia and/or combined with IVC.34 In addition, they are all removable. The LASIK flap could be created with a microkeratome or with a femtosecond laser.

Complications reported with CI include hyperopic shift, haloes, a decrease in photopic and mesopic CS, corneal thinning and melting, broadened defocus curve and reduced simulated retinal blur in the implanted eye (Kamra Inlay, AcuFocus, Irvine, California, USA).22–35–37 With all inlay designs, centration is critical for proper performance, and a small displacement can make a clinically significant difference.38

At present, there are three types of corneal inlays:

- CIs that alter the index of refraction with a bifocal optic. The Flexixure MicroLens, (Presbia, Los Angeles, California, USA) and Icolens (Neoptics AG, Hunenberg, Switzerland) are currently in clinical trials although several studies have been presented. The Flexivue (precursor was the Invue) is the only CI using a refractive addition power.20
- The Icolens is a new CI and recently Baily has reported the 1-year visual outcomes (table 1).23
- The Raindrop Near Vision Inlay (ReVision Optics, Lake Forest, California, USA) is a CI that changes the corneal curvature. Garza et al24 reported good and stable results at 1 year.
- The Kamra CI relies on small-aperture optics to increase the depth of focus. Most of the published data demonstrates that monocural implantation of a small-aperture inlay results in sustained improvement in near vision and intermediate vision while maintaining good distance vision.22–35–37 However, the size, material and visibility of the Kamra CI can be a disadvantage compared with the other CIs.

Lenticular approaches

The ultimate goal of cataract extraction and clear lens extraction is to replace the crystalline lens with an intraocular lens (IOL) that simulates the original function of the crystalline lens and provides the patients with a full range of functional vision for all distances. Currently, the available IOLs can be grouped into accommodating (AIOls) or pseudoaccommodating IOLs (although the mechanism of action of some `accommodative lenses’ may be pseudoaccommodative in nature). With pseudoaccommodative multifocal IOLs (MIOLs), the patient has two or three points in focus but primarily perceives only the focused image of interest.39–40

Precise biometry, accurate IOL power calculation, good surgical technique as well as patient selection are crucial in achieving the best visual outcome and patient satisfaction.

Pseudophakic multifocal intraocular lens

Multifocal intraocular lenses are used following patients with cataract or in clear lens extraction and excellent clinical outcomes have been reported.42–44 However, patient dissatisfaction and secondary procedures, including IOL exchange, can also be significant.40 44 Some of the MIOLs are based on multifocal CL designs, however the visual results may differ between them. First, CLs and IOLs are placed in different locations in the eye which results in different plane corrections, and second, the CL moves during the blink versus the stability of the IOL. These differences could lead to different visual outcomes.

Complications of these MIOLs include reduction in quality of vision, especially loss of CS, dysphotopsia, and reduced intermediate vision and near vision.45

The discussion below is not an exhaustive list of the IOLs available or publications (it is beyond the scope of this article) but is representative for the common lenses used.
Table 1  Visual outcomes of presbyopia procedures

<table>
<thead>
<tr>
<th>Author</th>
<th>Procedure</th>
<th>Study design</th>
<th>Number of eyes</th>
<th>UCNVA</th>
<th>BDCNVA</th>
<th>Additional tests</th>
<th>Complications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Levinger et al⁴</td>
<td>Monovision induced by LASIK</td>
<td>1 year</td>
<td>38</td>
<td>0.06 logMAR (binocular)</td>
<td>Not available</td>
<td>CS reduced in mesopic condition. Near stereoacuity 57 s of arc 85.2% of satisfaction</td>
<td>Not reported</td>
</tr>
<tr>
<td>Greenbaum⁵</td>
<td>Monovision pseudophakia</td>
<td>1 year</td>
<td>120 cataract/20 CLE</td>
<td>0.0 logMAR or better (binocular) in 91% cataract and 95% CLE</td>
<td>Not available</td>
<td>91% acceptance cataract 95% acceptance CLE 1 dry eye 1 vitreous loss 1 iris atrophy 20% reported haloes and glare in cataract in CLE</td>
<td></td>
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<tr>
<td>Uthoff et al⁷</td>
<td>Multifocal corneal ablation</td>
<td>6 months</td>
<td>20 (emmetropic), 20 (hyperopic), 20 (myopic)</td>
<td>0.18 logRAD (emmetropic), 0.24 logRAD (hyperopic), 0.12 logRAD (myopic) binocular</td>
<td>Not available</td>
<td>CS significantly reduced in all groups 91% acceptance cataract 95% acceptance CLE</td>
<td></td>
</tr>
<tr>
<td>McDonald et al⁸</td>
<td>Conductive keratoplasty</td>
<td>6 months</td>
<td>143</td>
<td>0.18 logMAR or better in 77% (monocular)</td>
<td>Not available</td>
<td>76% reported very satisfied satisfied 1 dry eye 1 vitreous loss 1 iris atrophy</td>
<td></td>
</tr>
<tr>
<td>Menassa et al⁹</td>
<td>Intrastromal femtosecond Ring incision</td>
<td>18 months</td>
<td>25</td>
<td>0.2 logMAR (monocular)</td>
<td>Not available</td>
<td>Corneal steepening 0.90 D 36% reported rings around light sources</td>
<td></td>
</tr>
<tr>
<td>Limnopoulou et al¹⁰</td>
<td>Flexivue microlens inlay</td>
<td>1 year</td>
<td>47</td>
<td>0.14 logMAR (monocular), 0.13 logMAR (binocular)</td>
<td>Not available</td>
<td>HOA increased CS decreased 81.25% reported UCNA excellent</td>
<td>No surgical complications</td>
</tr>
<tr>
<td>Garza et al¹¹</td>
<td>The raindrop Inlay</td>
<td>1 year</td>
<td>20</td>
<td>&lt;0.1 logMAR (monocular and binocular)</td>
<td>Not available</td>
<td>Photopic CS no significant change 95% reported satisfied or very satisfied UCNA UCVA 100% satisfied or very satisfied UCDNA</td>
<td></td>
</tr>
<tr>
<td>Seyeddain et al¹²</td>
<td>The Kamra inlay</td>
<td>2 years prospective</td>
<td>24</td>
<td>0.1 logMAR (monocular)</td>
<td>Not available</td>
<td>0.1 logMAR UCVA 1 eye with epithelial ingrowth in the pocket 1 eye with epithelial iron deposit</td>
<td></td>
</tr>
<tr>
<td>Baily et al¹³</td>
<td>The Icolens corneal inlay</td>
<td>1 year</td>
<td>52</td>
<td>0.4 logMAR (monocular)</td>
<td>Not available</td>
<td>90% reported happy with the procedure 1.25–1.75 D increase in objective accommodation 0.18 logMAR or better in 95% UCNA</td>
<td>11 inlay explanted because minimal improvement UCNA No major complications</td>
</tr>
<tr>
<td>Hipsley (ASCRS 2011)</td>
<td>The LaserACE procedure</td>
<td>18 months</td>
<td>134</td>
<td>0.18 logMAR or better in 89%</td>
<td>Not available</td>
<td>Mean lines of improvement at near 2.3 monocular and 2.0 binocular</td>
<td></td>
</tr>
<tr>
<td>Berrow (2014)</td>
<td>PresVIEW scleral implant</td>
<td>3 months (ongoing 2 years FDA clinical trial)</td>
<td>28</td>
<td>Not available</td>
<td>0.3 logMAR 100% (monocular, binocular)</td>
<td>Mean lines of improvement at near 2.3 monocular and 2.0 binocular</td>
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</table>

Note: some visual acuities were converted to logMAR using the visual acuity conversion chart prepared by Jack T Holland.

BDCNVA, best distance corrected near visual acuity; CLE, clear lens extraction; CS, contrast sensitivity; FDA, Food and Drug Administration; HOA, high order aberration; LogMAR, log minimum angle of resolution; logRAD, logarithm of the reading acuity determination; UCDVA, uncorrected distance visual acuity; UCNA, uncorrected intermediate visual acuity; UCIVA, uncorrected near visual acuity.

Reference:
Refractive MIOLs

Refractive MIOLs have the incorporation of two different powers integrated into two or more typically circular refractive zones. Due to each lens zone having a different effective refractive power, the image quality can depend on the pupillary response to light and the accommodation reflex.47

The ReZoom (Abbott Medical Optics (AMO), Irvine, California, USA) is a refractive MIOL (the original model being called the ARRAY) and the Food and Drug Administration (FDA) approved it in 2005. It is a three-piece MIOL and has five refractive optical zones; zones 1, 3 and 5 are adjusted for far vision, while zones 2 and 4 are adjusted for near vision. This design gives good distance vision and good intermediate-range vision although the reading performance is variable.48 Disadvantages of this lens, like many MIOLs, include dysphotopsia.49

The M-flex MIOL (Rayner IOLs Limited, Hove, UK) is based on a multitoned refractive aspherical optical technology, with either 4 or 3 annular zones (depending on IOL base power) providing +3.0 D or +4.0 D of additional refractive power at the IOL plane (equivalent to +2.25 D or +3.0 D at the spectacle plane). Cezon et al50 reported good visual performance and high rate of spectacle independence at 1 year.

Refractive MIOLs appear to be associated with more photic phenomena compared with diffractive MIOLs.51 Photic phenomena are among the most frequent reasons for patient dissatisfaction following implantation of MIOLs.52

Diffractive MIOLs

These are based on the principle of diffraction, whereby light slows down and changes direction when it encounters an obstacle.53 These lenses use microscopic steps (diffractive zones) across the lens surface. As light encounters these steps, it is directed towards the distant and near focal points (the amount of light is directly related to the step height as a proportion of wavelength). Diffractive MIOLs can be subdivided into apodised (gradual reduction in diffractive step heights from centre to periphery) or non-apodised (uniform height): both categories are wavelength. Diffractive MIOLs can be subdivided into apodised (gradual reduction in diffractive step heights from centre to periphery) or non-apodised (uniform height): both categories are designed to reduce the severity of night haloes compared with refractive MIOLs.47 Examples include the ReSTOR (Alcon Lab, Fort Worth, Texas, USA) (apodised) and Tecnis Multifocal (Abbott Medical Optics, Santa Ana, California, USA) and AT LISA 809 IOL (Carl Zeiss Meditec, Hennigsdorf, Germany) (both non-apodised). Most studies report good and stable distance vision and near vision, leading to low spectacle dependence and high patient satisfaction.54–56 Although these designs have good visual outcomes, their weakest points can be their inability to provide good levels of vision at an intermediate distance and loss of CS.

Aiming to improve intermediate vision, trifocal MIOL designs were introduced in the market: AT LISA trio 839MP novel design (Carl Zeiss Meditec, Hennigsdorf, Germany), FineVision (PhysIOL SA, Liège, Belgium) and MIOL-Record trifocal IOL (Reper NN, Nizhny Novgorod, Russia). Results reported so far of these lenses show a significant improvement in uncorrected VA at all distances. The trifocal designs may be the emerging refractive MIOLs.47 Examples include the ReSTOR (Alcon Lab, designed to reduce the severity of night haloes compared with periphery) or non-apodised (uniform height): both categories are wavelength. Diffractive MIOLs can be subdivided into apodised (gradual reduction in diffractive step heights from centre to periphery) or non-apodised (uniform height): both categories are designed to reduce the severity of night haloes compared with refractive MIOLs.47 Examples include the ReSTOR (Alcon Lab, Fort Worth, Texas, USA) (apodised) and Tecnis Multifocal (Abbott Medical Optics, Santa Ana, California, USA) and AT LISA 809 IOL (Carl Zeiss Meditec, Hennigsdorf, Germany) (both non-apodised). Most studies report good and stable distance vision and near vision, leading to low spectacle dependence and high patient satisfaction.54–56 Although these designs have good visual outcomes, their weakest points can be their inability to provide good levels of vision at an intermediate distance and loss of CS.

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Table 2  Visual results of intraocular lenses (IOLs)

<table>
<thead>
<tr>
<th>Author</th>
<th>IOL</th>
<th>Design</th>
<th>Study design</th>
<th>Number of eyes</th>
<th>UCNVA</th>
<th>BDCNVA</th>
<th>Additional tests</th>
<th>Complications</th>
</tr>
</thead>
<tbody>
<tr>
<td>Forte and Ursoleo</td>
<td>ReZoom</td>
<td>Refractive</td>
<td>2 years</td>
<td>55</td>
<td>0.10 logMAR (monocular)</td>
<td>Not available</td>
<td>▶ UCVA 0.07 logMAR (monocular).</td>
<td>7% patients reported moderate glare</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>▶ 5% patients reported moderate halo</td>
<td>Not reported</td>
</tr>
<tr>
<td>Cezon et al</td>
<td>Rayner M-flex</td>
<td>Refractive</td>
<td>1 year</td>
<td>32</td>
<td>0.28 logMAR (monocular)</td>
<td>0.28 logMAR (monocular)</td>
<td>▶ UCVA 0.15 logMAR (monocular).</td>
<td>▶ 2 eyes implant replacements</td>
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<td></td>
<td></td>
<td></td>
<td>▶ CIVA 0.15 logMAR (monocular).</td>
<td>▶ 2 eyes with cystoid macular oedema</td>
</tr>
<tr>
<td>Kohnen et al</td>
<td>AcrySof ReSTOR MA60D3</td>
<td>Diffractive</td>
<td>6 months</td>
<td>127</td>
<td>0.14 logMAR in 66.9% (binocular)</td>
<td>0.14 logMAR in 71.2% (binocular)</td>
<td>▶ 84.6% spectacle independence for near vision</td>
<td>▶ 1 eye flat macular oedema</td>
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<td></td>
<td></td>
<td>▶ 8.5% patients reported severe glare</td>
<td>▶ 1 eye macular oedema with fibrous reaction</td>
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<td></td>
<td></td>
<td>▶ 4.2% patients reported severe halo</td>
<td>▶ 1 cystic maculopathy</td>
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<td></td>
<td>▶ 2 eyes implant replacements</td>
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<td></td>
<td>▶ 2 eyes with cystoid macular oedema</td>
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<td></td>
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<td>▶ 1 eye flat macular oedema</td>
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<td>▶ 1 eye macular oedema with fibrous reaction</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>▶ 1 cystic maculopathy</td>
<td></td>
</tr>
<tr>
<td>Packer et al</td>
<td>Tecnis multifocal</td>
<td>Diffractive</td>
<td>1 year</td>
<td>244</td>
<td>0.20 logMAR (monocular)</td>
<td>0.18 logMAR (monocular)</td>
<td>▶ 95.5% spectacle independence for distance vision</td>
<td>▶ 10.3% patients reported moderate glare</td>
</tr>
<tr>
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<td></td>
<td></td>
<td>▶ 86.6% spectacle independence for near vision</td>
<td>▶ 2.6% patients reported severe glare</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>▶ 94.6% satisfied</td>
<td>▶ 84.8% spectacle independence</td>
</tr>
<tr>
<td>Mojís et al</td>
<td>AT Lisa tri 839MP</td>
<td>Diffractive trifocal</td>
<td>6 months</td>
<td>60</td>
<td>0.20 logMAR (monocular)</td>
<td>0.17 logMAR (monocular)</td>
<td>▶ UCVA 0.08 logMAR (monocular)</td>
<td>▶ UCVA 0.08 logMAR (monocular)</td>
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<td>▶ DCIVA 0.08 logMAR (monocular)</td>
<td>▶ UCVA 0.19 logMAR (monocular)</td>
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<td></td>
<td></td>
<td>▶ NAVQ Rasch scores satisfaction at near 15.9 logits</td>
<td>▶ Not reported</td>
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<tr>
<td>Sheppard et al</td>
<td>Finevision trifocal</td>
<td>Diffractive trifocal</td>
<td>2 months</td>
<td>30</td>
<td>Not available</td>
<td>Not available</td>
<td>▶ UCVA, DCVA 0.2 logMAR, Scotopic</td>
<td>▶ Not surgical complications</td>
</tr>
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<td></td>
<td>▶ CS 0.2 log unit below standard values at all spatial frequencies</td>
<td>▶ 94% patients reported spectacle freedom</td>
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<td></td>
<td>▶ 25% patients reported haloes</td>
<td>▶ 25% patients reported haloes</td>
</tr>
<tr>
<td>Voskresenskaya et al</td>
<td>MIOL-Record trifocal</td>
<td>Diffractive trifocal</td>
<td>6 months</td>
<td>36</td>
<td>0.10 logMAR (monocular)</td>
<td>0.10 logMAR (monocular)</td>
<td>▶ UCVA, DCVA 0.2 logMAR, Scotopic</td>
<td>▶ Not surgical complications</td>
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<td></td>
<td>▶ CS 0.2 log unit below standard values at all spatial frequencies</td>
<td>▶ 94% patients reported spectacle freedom</td>
</tr>
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<td>Shah (2014)</td>
<td>Lentis Mplus X LS-313</td>
<td>Rotationally asymmetrical</td>
<td>3 months</td>
<td>34</td>
<td>0.18 logMAR (monocular)</td>
<td>0.15 logMAR (monocular)</td>
<td>▶ UCVA, DCVA 0.2 logMAR, Scotopic</td>
<td>▶ Not surgical complications</td>
</tr>
<tr>
<td>Venter et al</td>
<td>SBL-3</td>
<td>Rotationally asymmetrical</td>
<td>3 months</td>
<td>106</td>
<td>0.12 logMAR (monocular); 0.08 logMAR (binocular);</td>
<td>0.11 logMAR (monocular); 0.08 logMAR (binocular);</td>
<td>▶ UCVA 0.16 logMAR (monocular) and 0.13 logMAR (binocular)</td>
<td>▶ UCVA 0.16 logMAR (monocular) and 0.13 logMAR (binocular)</td>
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<td>▶ DCIVA 0.15 logMAR (monocular) and 0.1 logMAR (binocular)</td>
<td>▶ UCVA 0.15 logMAR (monocular) and 0.1 logMAR (binocular)</td>
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<td></td>
<td>▶ 94.4% satisfied or very satisfied</td>
<td>▶ 94.4% satisfied or very satisfied</td>
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<td>▶ 86.8% had no difficulty at all or little difficulty performing tasks that require good close-up vision</td>
<td>▶ 86.8% had no difficulty at all or little difficulty performing tasks that require good close-up vision</td>
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<td>▶ Efficacy ratio of 80%</td>
<td>▶ Efficacy ratio of 80%</td>
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<td>▶ Safety ratio of 94%</td>
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<td></td>
<td>▶ Slight pupil ovalisation in 10% eyes</td>
<td>▶ Slight pupil ovalisation in 10% eyes</td>
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Continued
Table 2  Continued

<table>
<thead>
<tr>
<th>Author</th>
<th>IOL</th>
<th>Design</th>
<th>Study design</th>
<th>Number of eyes</th>
<th>UCNVA</th>
<th>BDCNVA</th>
<th>Additional tests</th>
<th>Complications</th>
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<tbody>
<tr>
<td>Alio and Mulet</td>
<td>The AMO multifocalphakic IOL</td>
<td>Refractive</td>
<td>1 year</td>
<td>34</td>
<td>0.20 logMAR</td>
<td>Not available</td>
<td>UCIVA 0.00 (binocular)</td>
<td>▶ Mean endothelium cell loss less than 5%</td>
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<td></td>
<td>prototype</td>
<td></td>
<td></td>
<td></td>
<td>(binocular)</td>
<td></td>
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<td>▶ 4 IOL explantations because of dissatisfaction</td>
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<tr>
<td>Cumming et al</td>
<td>Crystalex AT-4S</td>
<td>Accommodative</td>
<td>1 year FDA clinical trial</td>
<td>263 eyes</td>
<td>0.20 logMAR or better in 93.5% (binocular)</td>
<td>0.20 logMAR or better in 83.9% (binocular)</td>
<td>DCIVA 0.1 in 95% eyes (binocular)</td>
<td>▶ Endophthalmitis 1 eye</td>
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<td></td>
<td>(binocular)</td>
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<td>▶ 12 eyes IOL dislocation</td>
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<td>▶ 2 eyes retinal detachment</td>
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<td></td>
<td></td>
<td></td>
<td>▶ 1 eye iridectomy</td>
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<td></td>
<td></td>
<td></td>
<td>▶ 1 eye with persistent corneal oedema</td>
</tr>
<tr>
<td>Sanders and Sanders</td>
<td>Tetraflex</td>
<td>Accommodative</td>
<td>1 year FDA clinical study</td>
<td>255</td>
<td>0.4 logMAR in 77%</td>
<td>0.4 logMAR or better in 67%</td>
<td>90% could read ≥ 80 wpm at the 0.2 logMAR print size</td>
<td>Malpositioning of S IOLS</td>
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<td>Mastropasqua et al</td>
<td>1CU Human Optics</td>
<td>Accommodative</td>
<td>2 years</td>
<td>14</td>
<td>Not available</td>
<td>0.2 logMAR at 6 months 0.48 logMAR at 2 years</td>
<td>AA 1.9 D at 6 months and 0.30 D at 2 years</td>
<td>Anterior and posterior opacification in 100% of cases</td>
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<tr>
<td>Alió (2009)</td>
<td>The NuLens</td>
<td>Accommodative</td>
<td>1 year</td>
<td>10 (Cataract and atrophic macular degeneration)</td>
<td>Increase of 3.8 Jaeger rows (6 months)</td>
<td>Not available</td>
<td>Cross-section measurement of IOL of 0.09 mm (equivalent to 10 D)</td>
<td>▶ 1 posterior synechia inducing IOL tilt</td>
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<td>▶ 1 capsulorhexis edge capture by the haptic endplate inducing high myopia</td>
</tr>
</tbody>
</table>

Note: some visual acuities were converted to logMAR using the visual acuity conversion chart prepared by Jack T Holland.

AA, amplitude of accommodation; BDCNVA, best distance corrected near visual acuity; CIVA, corrected intermediate visual acuity; CS, contrast sensitivity; DCIVA, distance corrected intermediate visual acuity; LogMAR, log minimum angle of resolution; MIOL, multifocal intraocular lens; UCIVA, uncorrected intermediate visual acuity; NAVQ, Near Assessment of Vision Questionnaire; piOL, phakic intraocular lens; UCNVA, uncorrected near visual acuity.
axial movement of the optic resulting from action of the ciliary muscle. A hinge between the optic and haptics allows the lens to move forward as the eye focuses on near objects and backward as the eye focuses on distant objects, thereby increasing the dioptrical power of the pseudophakic eye.

It has been reported that an IOL optical shift of 1.0 mm can offer about 1.0 D of accommodation in a single-optic IOL and between 2.5 D and 3.0 D in an IOL with two-lens optics. In addition, the amount of accommodative result depends on several factors, such as the position of the optics in the capsular bag, the posterior chamber and the refractive power of the IOL. The Crystalens HD (Bausch & Lomb, Rochester, New York, USA) the Tetraflex HD (Lenstec, St Petersburg, Florida, USA) and the 1CU Human Optics (Human Optics AG, Erlangen, Germany) are examples of single-optic AIOLs and have all been extensively used. Visual performance reported with these AIOLs is promising, however capsule opacification and loss of accommodative ability with time are often present.

The single-optic passive-shift IOLs are considered pseudoadcommodative because they have limited accommodative ability, as their anterior movement is insufficient to provide functionally significant amplitudes of accommodation. Hence, dual-optic devices were developed such as the Synchrony IOL (AMO, Irvine, California, USA), and the Sarfarazi IOL (Shenasa Medical, Carlsbad, California, USA). The configuration of these devices with a high positively powered mobile anterior optic, connected to a stationary negatively powered posterior optic, is designed to increase the potential accommodative amplitude. Published results of both IOLs are limited but have shown positive results in small cohorts. They may be relatively difficult to handle, technically.

There are some IOLs, that change their shape or curvature with accommodative effort, in different stages of development. The FluidVision lens (PowerVision, Belmont, California, USA) drives fluid of a polymer-matched refractive index from the IOL’s soft haptics through channels to a fluid-driven internal activator. One-year follow-up showed that the base IOL powers were accurate and stable, VAs were good, and patients showed more than 5.00 D of accommodation on average (American Society of Cataract and Refractive Surgery (ASCRS) 2011). The NuLens (NuLens, Herzliya Pituah, Israel), a sulcus-based accommodating IOL, is still under development although it has been implanted in 10 eyes with cataract and atrophic macular degeneration showing at 1 year, reporting this IOL may result in up to 10 D of accommodation and the Superior Accommodating IOL (Human Optics AG, Erlangen, Germany) is designed to mimic the behaviour of natural lens and it is under development.

The lens filling techniques have been under investigation for years. It consists of replacing the lens with a soft gel that would allow modifying the shape for accommodation. The Medennium SmartLens IOL (Medennium, Irvine, California, USA) is a ‘smart’ hydrophobic acrylic material with unique thermodynamic property. When implanted into the capsular bag, the body’s temperature causes the material to transform into a gel-like polymer and take the shape of the natural lens. To the knowledge of the authors no data has been published yet. It should be noted that objective measurement of the accommodative capability of AIOLs is extremely difficult to obtain.

The femtosecond laser seeks to restore the flexibility that has been lost by making precise incision patterns within the lens without opening the capsule.

Preclinical studies have been performed in human cadaver and animal lense which have demonstrated safety, increased lens flexibility and no production of cataract. A feasibility study with the LENSAR (LENSAR, Orlando, Florida, USA) in 80 subjects with cataract showed that a third showed an improvement in objective accommodation measured with the Grand Seiko WR-3100 K autofractor (Grand Seiko, Fukuyama, Japan) and over 50% showed an improvement in subjective accommodation with the push-down method. Over 40% also showed an increase in the best distance-corrected near VA (results presented at ASCRS 2014).

Currently, there is another femtosecond laser-based therapy study for the treatment of presbyopia in Germany (The Human Eye study Cologne/Rostock). This clinical study is being conducted at two sites: University of Rostock University Eye Hospital, Rostock, Germany and Augenklinik am Neumarkt, Köln, Germany. Fifteen eyes in each site (n=30 eyes) have been recruited. However, to the knowledge of the authors, no data has been reported.

Scleral modification

Extraocular approaches have been developed based on Schachar’s theory. This model states that accommodation results from an increase of zonular traction at the lens equator to increase the lens diameter, therefore, presbyopia occurs as a result of increased lens growth causing a reduction in the space between the lens and the ciliary body (circumlenticular space), such that upon contraction the zonules can no longer exert their effect on the lens due to a loss of tension. MRI studies have shown that the circumlenticular space decreases with age as a result of the inward movement of the ciliary muscle ring that occurs with advancing age and an increase of the lens thickness. However, goniovideography, infrared photography and MRI studies have shown that the lens decreases in diameter and surface area with accommodation. Despite the controversy of this theory, Schachar postulated that expanding the dimensions of the overlying scleral wall by pulling the ciliary muscle away from the equatorial edge of the lens, would reverse the process of presbyopia and increase accommodative amplitude. LaserACE (Ace Vision Group, Silver Lake, Ohio, USA) and VisAbility Implant System surgery Scleral Implants (Refocus Group, Dallas, Texas, USA) were originally developed on the basis of this theory but the actual mechanism of action is still under investigation.

Laser-assisted presbyopia reversal

Laser assisted presbyopia reversal aims to restore dynamic accommodation increasing pliability in the sclera and net forces of the ciliary muscles on the lens facilitating accommodation. The postulated mechanism of action of laser assisted presbyopia reversal is to decrease ocular rigidity. The procedure is performed using a handheld fibre-optic handpiece that delivers pulses of an erbium:YAG laser ablating a diamond matrix pattern of nine laser spots into each oblique quadrant of the sclera. These are presumed to decrease the distance between the ora serrata and the scleral spur, restore the anatomical relationships of the system and free the ciliary muscle to contract normally. The spots delivered in a diamond matrix pattern of nine laser spots into each oblique quadrant. The results so far (in 134 eyes of 67 patients after 18 months follow-up) are promising. Hipsey reported restoration of 1.25–1.75 D of objective accommodation, which remained stable through 18 months in initial results (2011 ASCRS meeting) (table 1).
Scleral expansion bands
Scleral expansion band surgery for the treatment for presbyopia is based on the model of accommodation theorised by Schackar.

Scleral expansion band surgery has therefore been used for this purpose, but previous studies have demonstrated mixed results and have demonstrated limited success with temporary improvement in amplitude of accommodation.

Most recently, Refocus group has developed The VisAbility Implant System scleral implants. The technique consists of implanting four prostheses (size of a grain of rice) within elongated pockets in the sclera. The prostheses are thought to exert traction on the sclera in the region overlying the ciliary body which expands the sclera and the underlying ciliary body: thus restoring the effective working distance of the ciliary muscle and increasing the amplitude of accommodation. The actual surgical technique has evolved markedly from the initial use of manual diamond blade to the current use of disposable scleratome improving considerably the accuracy of tunnel creation. Furthermore, the original implant was a one-piece device, which was pushed into place and was difficult to thread through the tunnel. This one-single piece had a tendency to slip out of the tunnel over the long term resulting in a return or regression of patients’ preoperative near vision. Nowadays, the implant is a two-piece locking implant that prevents the implanted device from slipping out. Currently, Refocus group is conducting a FDA clinical trial (table 1).

CONCLUSION
There have been significant developments in surgery for presbyopia over the last decade achieving relatively good outcomes but each modality has its own advantages and disadvantages and sometimes compromises. However, to properly compare interventions it is necessary to encourage researchers to report best distance-corrected near VA rather just UCNA to minimise any confounding effect of myopia and astigmatism on results.

Other options for the management of presbyopia should not be forgotten, for example, it has been suggested that the use of miotics to increase depth of focus could help those suffering from presbyopia, and this would represent a type of reversible treatment. However there is little published evidence with this form of treatment, although 200 emmetropic eyes have been reported as having been treated in South America.

In the next few years it is likely that the introduction of different IOLs will be seen, as well as the development of new pharmacological treatments and technologies to provide patients with better visual outcomes and then possibly restoration of true accommodation to the presbyopic eye will be seen.

Contributors
All three authors were involved in the planning, writing and reviewing of this manuscript.

Competing interests
SAN and SS have received unrestricted grants or been involved in commercial studies for Bausch and Lomb, Lenstec, Abbot Medical Optics, Oculentis GMBH, FineVision, Topcon Europe Medical BV, Refocus Group and LENSSAR. SS is a consultant to Lenstec, Oculentis GMBH, Topcon Europe Medical BV, Refocus Group and LENSSAR and is medical director and equity holder for CustomVis.

Provenance and peer review
Not commissioned; externally peer reviewed.

REFERENCES
Review


