

Contrast Sensitivity after Refractive Lens Exchange with A Multifocal Diffractive Aspheric Intraocular Lens

Teresa Ferrer-Blasco, PhD¹ • Santiago García-Lázaro, PhD² • César Albarrán-Diego, MSc³
Lurdes Belda-Salmerón, MSc⁴ • Robert Montés-Micó, PhD⁴

Abstract

Purpose: To evaluate distance and near contrast sensitivity (CS) under photopic and mesopic conditions before and after refractive lens exchange (RLE) and implantation of the aspheric AcrySof ReSTOR (SN6AD3 model) intraocular lens (IOL)

Methods: Sixty eyes of 30 consecutive patients after RLE underwent bilateral implantation with the aspheric AcrySof ReSTOR IOL. The patients sample was divided into myopic [n=15, mean spherical equivalent (SE) -6.87 ± 2.03 diopters (D)] and hyperopic (n=15, mean SE $+4.52 \pm 1.14$ D) groups. Monocular uncorrected visual acuity at distance and near (UCVA and UCNVA, respectively) and monocular best corrected visual acuity at distance and near (BCVA and BCNVA, respectively) were measured before and 6 months postoperatively. Monocular CS function was measured with the Stereo Optical FACT CS chart at distance and near, at three different luminance levels (85, 5 and 2.5 cd/m^2) before and after RLE. Postimplantation results at 6 months were compared with those found before surgery.

Results: Our results revealed that patients in both groups obtained good UCVA and BCVA after RLE at distance and near vision in relation to presurgery values. The safety and efficacy indexes were 1.06 and 0.87 for the myopic group and 1.02 and 0.86 for the hyperopic group at distance vision. At near vision, the safety and efficacy indexes were 1.24 and 1.16 for the myopic group and 1.15 and 1.13 for the hyperopic group. No statistically significant differences were found between the values of CS pre and post-RLE at distance and near, at any lighting condition and spatial frequency ($P > 0.002$).

Conclusion: RLE with aspheric AcrySof ReSTOR IOL in myopic and hyperopic population provided good visual function and yield good distance and near CS under photopic and mesopic conditions. Although mesopic CS is reduced at distance and near in relation to photopic CS, the performance is comparable to that obtained prior to surgery with the natural lens and better than published results of the spherical AcrySof ReSTOR model.

Keywords: Contrast Sensitivity, Multifocal Diffractive Aspheric Intraocular Lens

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1. Optics Department, University of Valencia, Spain

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Correspondence to: Robert Montés-Mico, PhD

Optics Department, Faculty of Physics, University of Valencia

C/ Dr. Moliner, 50. 46100. Burjassot (Valencia). SPAIN., Email: robert.montes@uv.es

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Introduction

Currently, refractive lens exchange (RLE) surgery in patient with high myopia or hyperopia and presbyopia is focused not only on restoring visual acuity (VA) at distance and near but also on providing the best visual quality to patients. Several studies showed satisfactory clinical results at distance vision after implantation of monofocal¹⁻⁵ and toric⁶ intraocular lenses (IOLs). In many cases these patients presented limited depth-of-focus because these IOLs cannot provide clear vision at near distance unless they are used in a monovision arrangement. Bifocal refractive,⁷⁻⁹ diffractive,¹⁰⁻¹³ spherical hybrid¹⁴⁻¹⁷ and aspheric hybrid¹⁸⁻²² IOLs generate two focal points along the optical axis to provide good monocular uncorrected distance and near visual acuity (UCVA and UCNVA, respectively) as well as functional intermediate vision. The bifocal IOLs have also been implanted in RLE with good results at distance and near vision.²³⁻³¹ Recent studies with the hybrid spherical AcrySof ReSTOR IOL (Alcon Laboratories, Inc.)^{25-27,30,31} in RLE surgery report satisfactory visual results. These hybrids IOLs combine refractive and diffractive optics to reduce the disadvantages of conventional refractive and diffractive IOLs in terms of contrast reduction.³²

Wavefront sensing technology recently has been applied in the design of new IOLs. The aspheric AcrySof ReSTOR (SN6AD3 model) was designed to produce negative spherical aberration (SA) to compensate the positive SA of the cornea.³³ This design aims to decrease unwanted visual phenomena associated with multifocal IOL performance and to increase the depth of focus, thereby improving image quality.¹⁸⁻²²

The contrast sensitivity (CS) function gives information on visual performance for a range of objective scales. The key question to be answered is whether this visual performance given by CS function in patients implanted with aspheric IOLs in RLE treatment is better or worse than the natural lens.

The purpose of this study was to show the visual quality by the CS function of patients implanted with the aspheric AcrySof ReSTOR SN6AD3 IOL, as a function of the illumination level under distance and near conditions, in 60 healthy, noncataractous eyes divided into

two groups (myopic and hyperopic) before and after RLE.

Methods

Study design

The present prospective interventional case series study was carried out on 60 consecutive eyes of 30 patients bilaterally implanted with of the aspheric AcrySof ReSTOR IOL. Inclusion criteria were age between 45 and 70 years and the motivation to no longer wear any form of spectacle or contact lens correction for distance and near. Exclusion criteria included ≥ 1 diopters (D) of corneal astigmatism, history of glaucoma or retinal detachment, corneal disease, previous corneal or intraocular surgery, abnormal iris, pupil deformation, macular degeneration or retinopathy, neuroophthalmic disease and history of prior ocular inflammation. The tenets of the Declaration of Helsinki were followed in this research. Informed consent was obtained from all patients after the nature and possible consequences of the study were explained. Approval from the Institutional Review Board was obtained.

Before the RLE procedure, patients underwent a complete ophthalmologic examination, including manifest and cycloplegic refraction, keratometry, slit-lamp biomicroscopy, Goldmann applanation tonometry, and binocular indirect ophthalmoscopy through dilated pupils. Axial length and anterior segment biometry were measured with the Zeiss Humphrey IOL Master biometer (Carl Zeiss Meditec, Inc., Dublin, CA). The SRK/T formula was used for the myopic patients. The Holladay II formula was used for IOL power calculation in hyperopic patients. The targeted refraction was emmetropia in all cases.

All surgeries in this study were performed by phacoemulsification with the Infiniti Vision System (Alcon, Fort Worth, TX) using topical anaesthesia and a clear corneal 2.2-3.2 mm incision. Phacoemulsification was followed by irrigation and aspiration of the cortex, and IOL implantation in the capsular bag. There were no complications in any of the cases.

IOL specifications

The aspheric AcrySof ReSTOR SN6AD3 pseudoaccommodative aspheric IOL uses

apodization, diffraction, and refraction. The apodized diffractive region is within the central 3.6 mm optic zone of the IOL. This area comprises 12 concentric steps of gradually decreasing (1.3 to 0.2 mm) height, creating multifocality from distance to near (2 foci). The refractive part of the optic surrounds the apodized diffractive region. This area directs light to a distant focal point for larger pupil diameters and is dedicated to vision at distance. The overall diameter of the IOL is 13.0 mm and the optic diameter, 6.0 mm. The IOL power used in this study varied from +10.00 to +30.00 D and incorporated a +4.00 D near addition (add). The IOL has an aspheric profile to correct positive SA of the cornea. The IOL material includes a blue light-filtering chromophore. It has been shown that the use of a blue-light filter is advisable because it prevents ultraviolet light alterations to the retina without disturbing CS and chromatic vision.^{34,35}

Visual performance measures

UCVA and UCNVA and monocular best corrected visual acuity at distance and near (BCVA and BCNVA, respectively) were measured before and after 6 months post-surgery. The measurements in distance vision were measured using 100% contrast Early Treatment Diabetic Retinopathy Study (ETDRS) charts (Optec 6500; Stereo Optical Co Inc, Chicago, Ill) under photopic conditions [85 candelas per square meter (cd/m^2)] at 4 m. The measurements in near vision were measured using the Logarithmic Visual Acuity Chart 2000 New ETDRS (Precision Vision, LaSalle, Ill) at 40 cm under photopic conditions ($85\text{cd}/\text{m}^2$). The safety index of the procedure (ratio mean BCVA postoperative / mean BCVA preoperative) and the efficacy index (mean UCVA postoperative / mean BCVA preoperative) were calculated at 6 month after surgery.

Monocular photopic and mesopic CS was measured with best distance correction using the Stereo Optical Functional Acuity Contrast Test (FACT Stereo Optical, Chicago, IL) in both groups before and 6 months after implantation. Absolute values of \log_{10} CS were obtained for each combination of patient, spatial frequency, and luminance; means and standard deviations were calculated. The levels of chart luminance were 85, 5 and 2.5

cd/m^2 , the first being photopic (i.e. the luminance recommended in the manufacturer's guidelines) and the others mesopic: room illumination was at similar levels. CS was measured first at the photopic level and then at the mesopic level. Patients were given 5 minutes to adapt to each level before testing. Pupil diameters were measured in each eye using a Colvard pupillometer (OASIS Medical Inc, Glendora, CA).

Data analysis

All examinations were performed preoperatively and 6 months after IOL implantation by one clinician who was unaware of the objective of the study. Correlation analysis was performed to assess the differences between pre and postsurgery outcomes at far and near and correlations in the CS measured at the 3 lighting conditions. Bonferroni correction was applied for multiple tests of correlation for the 5 frequencies ($P < 0.01/5$). A probability less than 0.2% ($P < 0.002$) was considered statistically significant.

Results

Sixty eyes of 30 patients had RLE with ReSTOR SN6AD3 IOL. Patients' demographics are shown in table 1. Eyes were divided into two groups, myopic ($n=15$, 10 females, 5 males) with an average preoperative mean spherical equivalent (SE) refraction of -6.87 ± 2.03 D (range, -4.25 to -11.00 D), and hyperopic ($n=15$, 9 females, 6 males) with an average preoperative SE of $+4.52 \pm 1.14$ D (range, 3 to 7 D). Mean patient age was 50.83 ± 4.08 years in the myopic group (range, 45 to 59 years) and 52.14 ± 5.14 years in the hyperopic group (range, 44 to 60 years). There was no statistically significant differences between-group in age ($P > 0.01$). Pupil diameter under mesopic conditions for the myopic group was 5.86 ± 0.69 mm and was 5.12 ± 0.63 mm in hyperopic group ($P < 0.01$). After surgery and IOL implantation, the pupils of all patients were round, without iris trauma, and showed good responsiveness to light. All eyes were available for examination at 6 months.

At 6 months after IOL implantation, the residual mean SE refractive error was -0.12 ± 0.40 D and -0.04 ± 0.46 in myopic and

hyperopic groups respectively. After surgery, none of the eyes required a secondary intervention. No potentially sight threatening complications such as persistent corneal edema, pupillary block, retinal detachment or endophthalmitis were observed during the postoperative period. In addition, no eye was in the need of Nd:YAG capsulotomy up to the postoperative last visit.

Table 2 shows the VA values before and after the IOL implantation for distance and near. After surgery, in myopic group the mean UCVA and BCVA were 0.08 ± 0.15 logMAR and 0.01 ± 0.02 logMAR, respectively. Statistically significant difference were found between pre and postsurgery BCVA results ($P < 0.01$). In hyperopic group the mean UCVA and BCVA were 0.05 ± 0.12 logMAR and 0.04 ± 0.10 logMAR, respectively. No statistically significant difference were found between pre and postsurgery results in this group ($P > 0.01$). In near vision, in the myopic group the mean UCNVA and BCNVA were 0.03 ± 0.05 logMAR and 0.01 ± 0.02 logMAR, respectively. In hyperopic group the mean UCVA and BCVA were 0.05 ± 0.08 logMAR and 0.04 ± 0.06 logMAR, respectively. In both groups we found statistically significant differences between pre and postsurgery BCNVA results ($P < 0.01$). In myopic group the safety index (ratio mean BCVA postoperative / mean BCVA preoperative) was 1.06 and 1.24 in distance and near vision respectively. The

efficacy index (mean UCVA postoperative / mean BCVA preoperative) was 0.87 and 1.16 in distance and near vision respectively. In hyperopic group the safety index was 1.02 and 1.15 in distance and near vision respectively. The efficacy index was 0.86 and 1.13 in distance and near vision respectively.

The mean values of $\log_{10}CS$ before and after RLE with the aspheric AcrySof ReSTOR IOL implantation are plotted as a series of CS functions in figure 1 for the myopic group and in figure 2 for the hyperopic group. Left column shows distance CS functions at the 3 luminance levels (85.5 and 2.5 cd/m^2) and right column shows the results of the measurements at near conditions. For comparison, mean measurements for myopic eyes implanted with spherical AcrySof ReSTOR SA60D3 IOL found by Blaylock et al²⁷ are included in figure 1. Similarly, mean measurements for hyperopic eyes implanted with the same IOL found by Ferrer-Blasco et al²⁸ are included in figure 2. To explore the statistical significance of differences between pre and post-RLE groups, t-tests were performed on the comparable data of the two groups (absolute $\log_{10}CS$ values) at each spatial frequency and illumination level. The results showed no statistically significant differences between the values pre and post-RLE at any distance, lighting condition and spatial frequency ($P > 0.002$).

Table 1. Demographic characteristics of participants

	Myopic group	Hyperopic group
Number of eyes	30	30
Age (years)	50.83 ± 4.08	52.14 ± 5.14
Range of age (years)	45-59	44-60
Gender (Male/Female)	20/10	19/11
IOL Power (D)	12.38 ± 1.97	27.23 ± 1.68
Range IOL power (D)	10-15.5	24-30
Axial length (mm)	25.93 ± 0.87	21.67 ± 0.74
Range axial length (mm)	23.79-27.39	20.31-23.23
Preoperative sphere (D)	-6.87 ± 2.03	4.52 ± 1.14
Range preoperative sphere (D)	4.25-11	3-7
Preoperative Cylinder (D)	0.55 ± 0.37	0.30 ± 0.31
Range preoperative cylinder (D)	0.-1.00	0-0.75
Preoperative keratometry (K)		
K1 (D)	44.29 ± 1.13	43.22 ± 1.50
K2 (D)	44.82 ± 1.25	43.83 ± 1.53
Postoperative mean pupil diameter (mm)		
85 cd/m^2	4.45 ± 0.82	3.53 ± 0.84
5 cd/m^2	5.86 ± 0.69	5.12 ± 0.63
2.5 cd/m^2	6.01 ± 0.58	5.81 ± 0.61
Range preoperative keratometry (K)	42.5-46	40.5-47
IOL: Intraocular lens		
D: Dioptres		
Means \pm standard deviation		

Table 2. Monocular visual acuity results for distance and near vision. Mean and standard deviation logMAR (logarithm of the minimum angle of resolution) values before and after 6 months post-AcrySof ReSTOR SN6AD3IOL implantation

	Myopic group			Hyperopic group		
	Before	After	P	Before	After	p
Distance (4m)						
UCVA		0.08±0.15			0.11±0.14	
BCVA	0.03±0.05	0.01±0.02	<0.01	0.05±0.12	0.04±0.10	0.30
Near (40 cm)						
UCNVA		0.03±0.05			0.05±0.08	
BCNVA	0.10±0.01	0.01±0.02	<0.01	0.10±0.01	0.04±0.06	<0.01

UCVA: Uncorrected logMAR distance visual acuity
 BCVA: Best distance-corrected logMAR visual acuity
 UCNVA: Uncorrected near logMAR visual acuity
 BCNVA: Best distance-corrected near logMAR visual acuity

MYOPIC

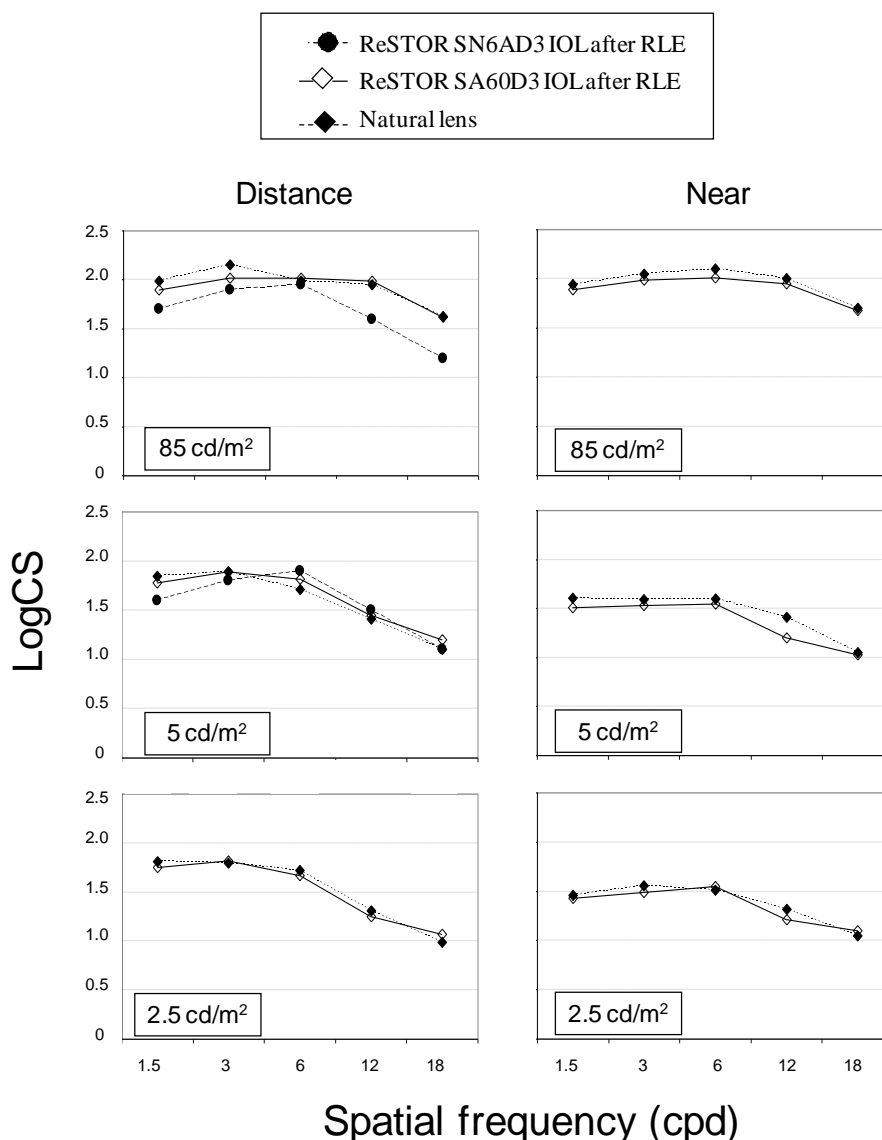


Figure 1. Contrast sensitivity functions before (natural lens) and after refractive lens exchange with the aspheric AcrySof ReSTOR implantation in myopic group as a function of the distance and FACT chart luminance (85, 5 and 2.5 cd/m²). For comparison, results for spherical AcrySof ReSTOR IOL from Blaylock et al²⁷ are included. CS: Contrast sensitivity, cpd: cycles per degree

HYPEROPIC

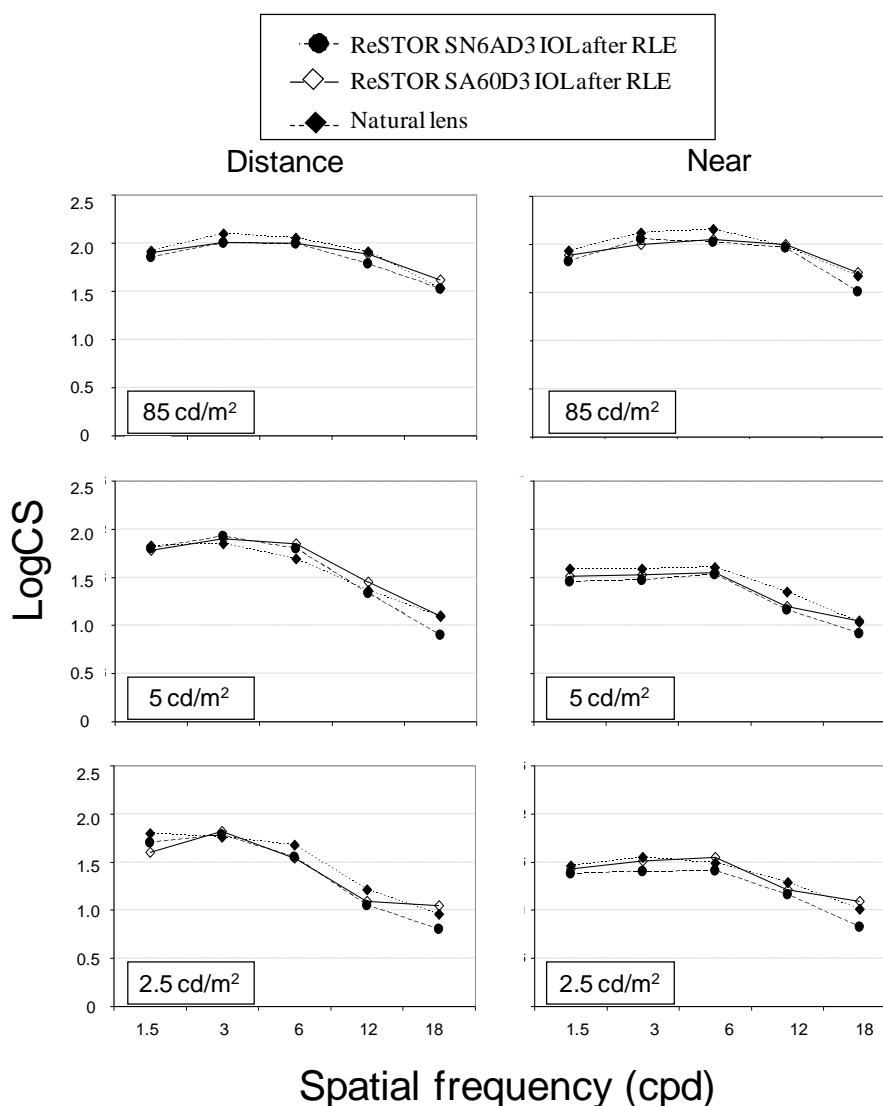


Figure 2. Contrast sensitivity functions before (natural lens) and after refractive lens exchange with the aspheric AcrySof ReSTOR implantation in hyperopic group as a function of the distance and FACT chart luminance (85, 5 and 2.5 cd/m^2). For comparison, results for spherical AcrySof ReSTOR IOL from Ferrer-Blasco et al²⁸ are included. CS: Contrast sensitivity, cpd: cycles per degree

Discussion

The removal of the crystalline lens and replacement with a pseudophakic lens for the purposes of reducing or eliminating refractive errors has been labelled with many titles, including clear lensectomy, clear lens phacoemulsification, clear lens replacement, clear lens exchange, presbyopic lens exchange, and RLE. It is a simple, effective technique to correct high ametropia. Several studies report good safety, efficacy,

predictability and stability in RLE with monofocal¹⁻⁵ and toric IOLs.⁶ Others studies evaluating the clinical, functional, and quality-of-life outcomes after implantation of the multifocal spherical IOLs^{23-28,30-31} show that these lenses improves high level of distance vision with additional benefit of increased range of near vision without additional correction in non-cataractous eyes. Blaylock et al²⁷ and Ferrer-Blasco et al²⁸ evaluated the

CS under photopic and mesopic conditions before and after implantation of the spherical AcrySof ReSTOR IOL. These authors found postoperatively a reduction in mesopic CS in relation to photopic CS but with a performance comparable to the CS preoperative. This study is the first one assessing CS after RLE and implantation of a multifocal IOL with an aspheric design.

Our results revealed that aspheric AcrySof ReSTOR IOL provided good and comparable safety and efficacy indexes in both groups. The safety and efficacy indexes were 1.06 and 0.87 for the myopic group and 1.02 and 0.86 for the hyperopic groups at distance vision. These values agree with those found previously by Fernández-Vega et al²⁵ in 112 patients after RLE with bilateral spherical AcrySof ReSTOR IOL. There was no statistically significant difference between pre and postoperative BCVA in hyperopic group ($P>0.01$), however we found statistically significant difference in the myopic group ($P<0.01$; Table 2). Magnification and minimization of the retinal image in myopic and hyperopic patients, respectively, may play a significant role in this difference. At near vision, the safety and efficacy indexes were 1.24 and 1.16 for the myopic group and 1.15 and 1.13 for the hyperopic groups. The patients obtained an UCVA of 0.03 logMAR and 0.05 logMAR that improved to 0.01 logMAR and 0.04 logMAR in myopic and hyperopic groups with the best distance correction. Statistically significant differences were found between BCVA pre and postsurgery ($P<0.01$).

One of the most important assessments for successful RLE with multifocal IOLs use involves precise preoperative measurement of axial length and accurate IOL power calculation. Keratometry and anterior chamber depth and axial length measurement should be performed as accurately as possible. In patients with high ametropia, axial length is difficult to measure, and then, power selection should be performed carefully. We chose the SRK/T formula for myopic patients because it has been shown to be accurate in eyes with a greater axial length.³⁹ The Holladay II formula was used for hyperopic patients, considering that for short eyes (about 22.0 mm) it performs better than the SRK/T formula.⁴⁰

If we focus now on CS we must first point out that the light distribution between distance and near foci depends on pupil diameter and varies from approximately 40% to 90% of the light to the distance focus for this IOL.³² Pupil diameter postoperative in myopic group under photopic conditions was 4.45 ± 0.82 mm, so the percentage of light for the distance/near focus was about 80/20. In the mesopic conditions pupil diameter was 5.86 ± 0.69 mm, about 90/10 of the light goes to distance/near focus. In hyperopic group the pupil diameter measured postoperative was 3.53 ± 0.84 mm and 5.12 ± 0.63 mm in photopic and mesopic conditions respectively, so the percentage of the light for the distance/near focus was about 70/30 and 85/15.

Analysing the results for distance, under photopic conditions, performance was very similar in both groups when compare the results of CS functions before and after surgery. There is some loss in photopic retinal image contrast with the aspheric AcrySof ReSTOR IOL although this difference was not significant ($P>0.002$). Due to diffractive nature of the IOL, for small pupils a maximum of 40% of the total light passing through the pupil contributes to either the distance or near image, and the remainder of the light decreases retinal image contrast.³² The light energy focused on the distance focus was about 80% and 70% in myopic and hyperopic groups respectively and these values are enough to achieve a CS comparable with the natural lens, both with appropriate distance correction. A recent study with another diffractive IOL design has shown that a 65% of light energy for the distance focus is also acceptable for achieving good photopic distance CS.¹⁸

Under low lighting conditions, there is a little reduction in CS function results obtained with multifocal IOLs compared with the results with the natural lens at 5 cd/m^2 and 2.5 cd/m^2 , particularly at higher spatial frequencies in both groups. This trend agrees with classic data for the effect of luminance level on CS.³⁶ At these mesopic levels, pupil diameters are substantially larger, and it seems reasonable to attribute the observed reduction in mesopic CS at higher spatial frequencies to the additional blur introduced by the larger diameter, out-of-focus zones of the multifocal IOL.

At near, the photopic and mesopic CS functions were similar pre and post-RLE in both groups. These results agree with those found previously by Ferrer-Blasco et al.²⁸ In comparison with distance vision the reduction of CS under dim conditions correlated with the reduction of light energy concentrated at the near focus. In photopic the percentage of the light for the distance focus were 80% and 70% and in mesopic conditions approximately the 20% and 30% of the light contributes to near image in myopic and hyperopic groups respectively. These percentages leading to noticeably worse the results of CS at near. Increasing pupil diameter implies that more light is distributed for the distance foci and then better CS is expected for distance CS functions compared to near CS function at mesopic levels. One should consider that the loss in retinal image contrast has little effect on acuity as measured with high-contrast letters, since contrast can be reduced to quite low levels before acuity is affected.³⁶

For comparison in myopic group we included in figure 1 the mean values obtained for the spherical AcrySof ReSTOR as found by Blaylock et al.²⁷ in 19 myopic patients (mean preoperative SE -3.89 ± 2.14 D). We should point out that these CS results were measured at 50 foot-candles in photopic level and 3 foot-candles in mesopic level with the SIFIMAV Vision Tester (Designs For Vision; Sydney, Australia). Caution should be exercised when comparing results, among different studies. The differences in conditions and measure systems between studies may play a role in the differences found but it is obvious that the results of spherical AcrySof ReSTOR return to be lower.

In hyperopic group we had included in figure 2 the results obtained by Ferrer-Blasco et al.²⁸ in 50 hyperopic patients (mean preoperative SE 2.18 ± 1.17) at the same conditions and with the same test in patients implanted with the spherical AcrySof ReSTOR. At higher spatial frequencies, the CS results of the spherical AcrySof ReSTOR IOL were lower compared with CS results of the aspheric IOL and natural lens. Differences between both IOLs become more evident when lighting conditions are reduced and pupil diameter is increased. This is an expected

result considering the reduction of ocular SA for large pupil diameters when an aspheric IOL is implanted, compared with that found in eyes with a spherical IOL.³³ Maxwell et al.²⁹ compared the optical performance in RLE with 6 presbyopia-correcting IOLs of different designs (3 spherical models: Crystalens AT-50SE, AcrySof ReSTOR SA60D3 and ReZoom NXG1; 3 aspheric models: AcrySof ReSTOR SN6AD3, Acri.Lisa 366D, and Tecnis ZM900) found that the aspheric AcrySof ReSTOR SN6AD3 IOL showed superior optical properties, demonstrated in modulation transfer function and the United States Air Force 1951 Resolution Target in optical bench testing.

If we now compare the postsurgery CS function results found in both groups, similar outcomes were reported (Figure 3). Statistically significant differences were found related to pupil size between both groups ($P < 0.01$), the pupil size in myopic group was slightly higher compared to that found in the hyperopic group (Table 1). According to the light distribution between distance and near foci, in this group a higher percentage of the light contributes to distance image but we need also to consider that there is an increasing effect of higher order optical aberrations especially in hybrid IOLs when the pupil increases. The balance between the advantage and disadvantage of the large pupils (light distribution versus optical aberrations) could be the reason of finding similar results in both groups.

In general, the results found in both groups were similar in distance and near vision at the different lighting conditions (Figure 3) and pre and postsurgery (Figures 1 and 2) without statistically significant differences ($P > 0.002$). Previous literature about CS with a multifocal IOL found a reduction compared to that obtained with a monofocal IOL.⁸ Our results were similar possibly because in these patients there was already a high degree of contrast-reducing scatter previous surgery (natural eye), giving a reduced CS function as compared with normal young eyes. Replacing the scattering crystalline with an IOL would remove this scatter, even if the diffractive IOL itself reduced the retinal image contrast by its simultaneous bifocal properties.

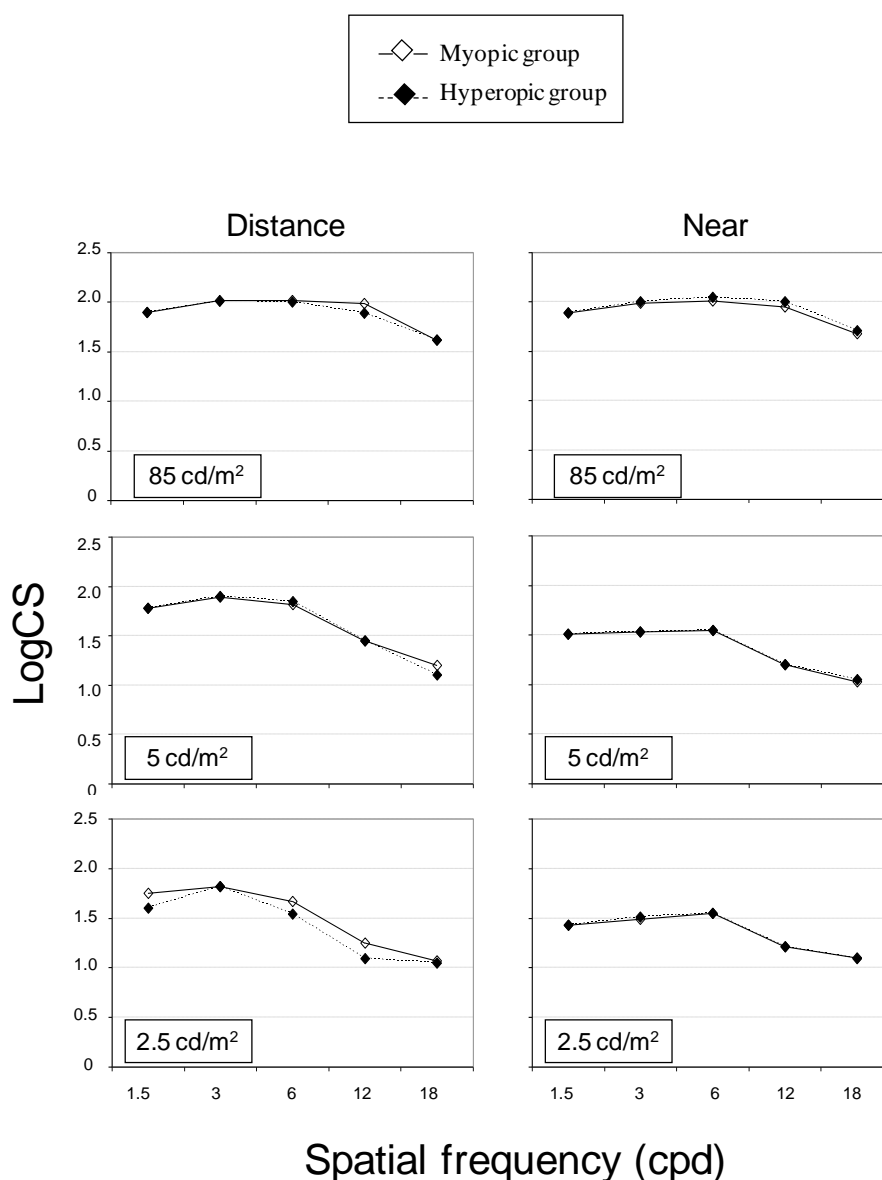


Figure 3. Contrast sensitivity functions after refractive lens exchange with the aspheric AcrySof ReSTOR implantation in myopic and hyperopic group as a function of the distance and FACT chart luminance (85, 5 and 2.5 cd/m²). CS: Contrast sensitivity, cpd: cycles per degree

Conclusion

In conclusion RLE with aspheric AcrySof ReSTOR in presbyopic population provided good visual function and yield good distance and near CS under photopic and mesopic conditions in myopic and hyperopic patients. Although mesopic CS is reduced at distance and near in relation to that found under photopic conditions, the performance is comparable to that obtained prior to surgery with the natural lens and better than the results obtained in patients implanted with

spherical AcrySof ReSTOR. Further studies are needed to assess the stability, photic phenomena, such as starbursts and halos, patient satisfaction, the role of pupil size after this multifocal IOL implantation and intermediate VA for comparison with others IOLs profiles.

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