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Title: Threshold limit of postoperative astigmatism for patient satisfaction after clear lens extraction and multifocal intraocular lens implantation

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Corresponding Author: Professor jonathan E Moore, FRCOphth, PhD

Corresponding Author's Institution: Mater Hospital

First Author: Jonathan E Moore

Order of Authors: Jonathan E Moore; McNeely Richard; Eric Pazo; Andrew Spence; Zack Millar; Olivier Richoz; Andrew Nesbit; Tara Moore

Abstract: PURPOSE: To determine the degree of tolerance towards different magnitudes of residual refractive and corneal astigmatism, and angles of corneal astigmatism following implantation of an asymmetric multifocal intraocular lens (MIOL).

SETTING/VENUE: Cathedral Eye Clinic, Belfast, Northern Ireland, UK

DESIGN: Retrospective comparative case series.

METHODS: The study enrolled 117 patients (234 eyes) with a mean age of 64 years undergoing clear lens extraction (CLE) and implantation of a Lentis Mplus LS-312 MF30 MIOL. Refraction, uncorrected (UDVA) and best-corrected (CDVA) distance visual acuities, uncorrected near (UNVA) and intermediate (UIVA) visual acuities, and a quality of vision (QOV) questionnaire were evaluated. Groups were categorised based upon the magnitude of residual refractive astigmatism and secondly on postoperative corneal astigmatism. The groups were $\leq 0.50D$, and $>0.50D$ of residual refractive astigmatism and $\leq 0.50D$, $0.51D$ to $0.75D$, $0.76D$ to $1.00D$, $>1.00D$ of corneal astigmatism.

RESULTS: There was a significant difference in UDVA ($P=0.003$), refractive sphere ($P=0.001$) and defocus equivalent ($P<0.001$) between residual refractive astigmatism groups, but no difference in QOV ($P=0.28$). The same was found for corneal astigmatism with UDVA ($P<0.001$) and QOV ($P=0.16$). The angle of corneal astigmatism in relation to inferior nasal placement of the near segment did not statistically affect visual and subjective outcomes.

CONCLUSIONS: The Lentis Mplus MIOL appears to subjectively tolerate well residual refractive and corneal astigmatism despite a statistically significant difference in UDVA with higher magnitudes of residual astigmatism. The angle of residual corneal astigmatism in relation to

MIOL placement does not have a significant effect on objective or subjective outcomes.

Title:

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2 Threshold limit of postoperative astigmatism for patient satisfaction after clear lens
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4 extraction and multifocal intraocular lens implantation
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10 **Short Running Head:**

11 Threshold limit of MIOL to residual astigmatism
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17 **Authors:**

18
19 Mr Richard N McNeely BSc^{1,2}, Dr Eric Pazo MD^{1,2}, Mr Andrew Spence BSc¹, Mr
20
21 Zack Millar¹, Dr Olivier Richoz MD PhD¹, Dr Andrew Nesbit BA, PhD², Professor
22
23 Tara CB Moore PhD² and Professor Jonathan E Moore PhD FRCOphth ^{1,2}
24
25
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27

28
29 1 Cathedral Eye Clinic, Belfast Campus, University of Ulster, Belfast, UK.
30

31
32 2 Biomedical Sciences Research Institute, University of Ulster, Coleraine, UK.
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54 Name and address of author to receive reprint requests:
55

56 **Professor Jonathan E Moore, Cathedral Eye Clinic, 89-91 Academy Street,**
57
58 **Belfast, County Antrim, BT1 2 LS. jmoorecathedral@gmail.com**
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Abstract

PURPOSE: To determine the degree of tolerance towards different magnitudes of residual refractive and corneal astigmatism, and angles of corneal astigmatism following implantation of an asymmetric multifocal intraocular lens (MIOL).

SETTING/VENUE: Cathedral Eye Clinic, Belfast, Northern Ireland, UK

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METHODS: The study enrolled 117 patients (234 eyes) with a mean age of 64 years undergoing clear lens extraction (CLE) and implantation of a Lentis Mplus LS-312 MF30 MIOL. Refraction, uncorrected (UDVA) and best-corrected (CDVA) distance visual acuities, uncorrected near (UNVA) and intermediate (UIVA) visual acuities, and a quality of vision (QOV) questionnaire were evaluated. Groups were categorised based upon the magnitude of residual refractive astigmatism and secondly on postoperative corneal astigmatism. The groups were $\leq 0.50D$, and $>0.50D$ of residual refractive astigmatism and $\leq 0.50D$, $0.51D$ to $0.75D$, $0.76D$ to $1.00D$, $>1.00D$ of corneal astigmatism.

RESULTS: There was a significant difference in UDVA ($P=0.003$), refractive sphere ($P=0.001$) and defocus equivalent ($P=<0.001$) between residual refractive astigmatism groups, but no difference in QOV ($P=0.28$). The same was found for corneal astigmatism with UDVA ($P=<0.001$) and QOV ($P=0.16$). The angle of corneal astigmatism in relation to inferior nasal placement of the near segment did not statistically affect visual and subjective outcomes.

CONCLUSIONS: The Lentis Mplus MIOL appears to subjectively tolerate well residual refractive and corneal astigmatism despite a statistically significant difference in UDVA with higher magnitudes of residual astigmatism. The angle of residual corneal astigmatism in relation to MIOL placement does not have a significant effect on objective or subjective outcomes.

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Introduction

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5 Multifocal intraocular lens (MIOL) technology is often utilised in modern cataract
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7 extraction surgery and refractive procedures, providing excellent levels of visual
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9 performance at a range of distances¹⁻⁴ and spectacle independence.^{5,6} However not
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11 all individuals are content postoperatively. Reported problems include reduced
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13 contrast, glare, haloes and/or starbursts around lights.^{1,7} Patients may also report
14
15 substandard near or intermediate vision which affects their ability to see clearly at
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17 different working distances.^{8,9} It is important to consider what produces these
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19 negative side effects and reduces overall patient satisfaction. One factor that
20
21 appears to affect postoperative vision and quality of life is uncorrected astigmatism.¹⁰
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23 Various MIOL studies have found patient complaints of blurred vision, which in the
24
25 majority of cases has been caused by ametropia and / or astigmatism, a well-
26
27 recognised cause of patient dissatisfaction with symmetric MIOLs.^{11,12} The effect of
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29 astigmatism upon uncorrected distance visual acuity has been shown to have a
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31 greater effect in symmetric MIOLs compared to monofocal IOLs.¹³
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41 Rotationally asymmetric MIOLs have since been introduced which differ from
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43 rotationally symmetric MIOLs, as they only possess two sections, a surface-
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45 embedded near section and a larger distance section, providing two different foci
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47 opposed to concentric rings in previous MIOL design. With only two transition zones,
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49 rotationally asymmetric MIOLs provide excellent visual results and reduce
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51 dysphotopsic side effects¹⁴ and improve contrast sensitivity.¹⁵
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1 To our knowledge the effect of uncorrected residual astigmatism on this new
2 asymmetric MIOL design has not been investigated. Therefore, this study sought to
3 determine and quantify, where possible, the effect of residual refractive astigmatism
4 upon subjective QOV following the implantation of an asymmetric MIOL. The effect
5 of postoperative corneal astigmatism and the relationship between the MIOL position
6 compared to the angle of the steepest corneal meridian, upon subjective QOV was
7 also measured. The aim of this study is to determine the degree of tolerance patients
8 have to different levels of residual astigmatism and to the angle of this astigmatism
9 with relation to the position of the asymmetric MIOL, before there is a significant
10 deleterious impact upon QOV.
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27 **Patients and Methods**

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31 This study enrolled patients undergoing clear lens extraction (CLE) with bilateral
32 implantation of Lentis Mplus MIOLs. Patients were first divided into two groups
33 depending on the magnitude of residual refractive astigmatism found following
34 subjective refraction. The two groups were $\leq 0.50D$ and $>0.50D$. Patients were then
35 categorised based on the magnitude of postoperative corneal astigmatism into 4
36 groups. The groups were $\leq 0.50D$, $0.51D$ to $0.75D$, $0.76D$ to $1.00D$, $>1.00D$. The
37 postoperative corneal astigmatism was measured using the NIDEK OPD-Scan II
38 ARK-10000 aberrometer. The magnitude of corneal astigmatism was defined as the
39 difference between the steep and flat corneal meridians. Patients with corneal
40 astigmatism $> 0.50 D$ were then categorised by the angle of the steepest corneal
41 meridian in relation to the position of the MIOL (Figure 1). Preoperatively at the slit-
42 lamp the horizontal and vertical axis were marked at the limbus. The MIOL was
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1 implanted with the near segment positioned inferiorly and nasally deviated half way
2 between the vertical and horizontal 'limbal marks'. Therefore, the long axis of the
3
4 MIOI was positioned around 135 degrees in the right eye and 45 degrees in the left
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6 eye. Patients with the steepest meridian between 0 to 22.5 degrees, 158 to 180
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8 degrees, and 68 to 112.5 degrees in either the right or left eye were considered to
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10 obliquely cross the vertical axis. Patients with the steepest meridian at an angle of 23
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12 to 67.5 degrees in the right eye and an axis of 113 to 157.5 degrees in the left eye
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14 were categorised together because they both crossed perpendicularly to the vertical
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16 axis. Patients with an axis of 113 to 157.5 degrees in the right eye and 23 to 67.5
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18 degrees in the left were grouped together because the steepest meridian ran parallel
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20 to the vertical axis in the respective eyes.
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29 Informed consent was obtained from all patients. The research adhered to the tenets
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31 of the Declaration of Helsinki and was approved by the local research ethics
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33 committee. The patients were advised of the possible risks of the operation and the
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35 possible need for further corneal laser refractive surgery.
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41 **Patient assessment**

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45 Full ophthalmological assessment was performed preoperatively and
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47 postoperatively. The examination included a medical history, autorefractometry (OPD-
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49 Scan II ARK-10000, Nidek), subjective refraction (RT-5100 Auto Phoropter Head
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51 Nidek), uncorrected (UDVA) and corrected (CDVA) distance visual acuities, defocus
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53 equivalent based on subjective refraction, uncorrected near (UNVA) and uncorrected
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55 intermediate (UIVA) visual acuities. These results were evaluated with logMAR
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1 charts (6m) and with Radner reading charts in M notation (40cm and 70cm).
2 Biometry was carried out preoperatively with the IOLMaster (Carl Zeiss Meditec AG).
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4 Pupil size, corneal topography, angle kappa and wavefront examinations (OPD-Scan
5 II ARK-10000, Nidek) were performed. The OPD-Station Software was utilised to
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7 report higher order aberrations (HOAs) across a 6mm pupil up to the sixth radial
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9 order.¹⁶ It has been found that aberrations above the sixth order have extremely
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11 small impact on the overall aberration.¹⁷ Slit-lamp microscopy, tonometry, dilated
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13 funduscopy, and OCT of the retina were completed. Each patient was assessed
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15 within 6 weeks and then 6 months postoperatively. The position of the vertical axis of
16
17 the MIOL was assessed postoperatively to confirm an axis of 135 degrees in the
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19 right eye and 45 degrees in the left eye.
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29 A quality of vision (QOV) questionnaire was completed preoperatively and at the
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31 second postoperative assessment. This assessed how annoyed the patients were by
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33 the questioned symptoms. Patients responded either not at all (0), a little (1), quite
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35 (2) or very (3). The patient also rated their vision out of 10; 0 representing the worst
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37 and 10 the best.
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43 **Intraocular lenses**

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48 The Lentis Mplus is a rotationally asymmetric MIOL consisting of an aspheric
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50 distance vision zone and a separate sector shaped near vision zone. It has a
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52 refractive design with seamless transition between the two sections of the MIOL. The
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54 Lentis Mplus MIOL is available with either a +1.50D, +2.00D or +3.00D near
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segment addition. In this study each patient received bilateral implantation of the Lentis Mplus LS-312 MF30 (+3.00D addition).

Surgical technique

The same experienced surgeon performed all operations with standard on-axis clear corneal phacoemulsification surgery. An incision of 2.75mm was used to reduce postoperative corneal astigmatism and the incision was made on the steepest meridian to evade the introduction of oblique astigmatism. A 5.00mm capsulorhexis and implantation of the MIOL into the capsular bag was performed. The vertical axis (near segment) was positioned inferiorly with slight nasal deviation in each eye. The refractive aim was emmetropia.

Statistical analysis

Descriptive and statistical analysis was performed using SPSS for Windows (Statistical Package for the Social Sciences, Version 22, Chicago, Illinois, USA) and Excel (Microsoft; Redmond, Washington, USA). The Kolmogorov-Smirnov test was used to assess normality. Independent t tests and one way Anova were utilised for parametric data with a post hoc Tukey test to compare the results among groups. The Kruskal-Wallis test was used to compare nonparametric data and the Wilcoxon signed rank-test was used to compare nonparametric data among groups. A paired sample t test was also used to test the significance between preoperative and postoperative HOAs. For all statistical analysis, the level of significance was $P < 0.05$.

Results

Demographics

This study included 117 patients (234 eyes) with a mean age of 64 years, ranging from 44 to 87 years (Table 1).

The magnitude of residual refractive astigmatism

Figure 2a displays the linear regression analysis between the residual refractive astigmatism and the UDVA, where a weak correlation was found ($R^2=0.12$). Figure 2b shows a slightly stronger association between UDVA and the defocus equivalent ($R^2=0.23$). The patients were then divided into two groups. There were 216 eyes with $\leq 0.50D$ and 18 eyes with $>0.50D$ postoperative refractive cylinder. Table 2 displays the objective results for the different groups. The group with residual refractive astigmatism $\leq 0.50D$ achieved better UDVA than those with $>0.50D$ ($P=0.003$, independent t test). There was also a greater magnitude of defocus equivalent in the $>0.50D$ group ($P=<0.001$, independent t test). Comparison of the individual symptom responses is highlighted in Figure 3 and the QOV scores in Figure 4. There was no significant difference in individual responses or QOV scores ($P=0.28$, independent t test).

The magnitude of residual corneal astigmatism

1 The different groups were $\leq 0.50\text{D}$ (n=99), 0.51D to 0.75D (n=53), 0.76D to 1.00D
2 (n=41) and $>1.00\text{D}$ (n=41) of corneal astigmatism. The mean corneal astigmatism
3 found postoperatively was $-0.63 \pm 0.36\text{ D}$. Patients with $\leq 0.50\text{D}$ achieved UDVA of -
4 $0.07 \pm 0.08\text{ logMAR}$ which was significantly better than those with 0.76 D to 1.00 D
5 $(0 \pm 0.08\text{ logMAR})$ and $>1.00\text{D}$ $(0 \pm 0.11\text{ logMAR})$, outlined in Table 3. There was no
6 significant difference in individual symptoms and overall QOV scores between the
7 groups.
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10 Residual corneal astigmatism axis in relation to the MIOL

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19 There were 135 eyes with corneal astigmatism $>0.50\text{D}$ and 99 of these were in the
20 oblique group, 22 in the perpendicular group and 14 in the parallel group. There was
21 no significant difference in objective outcomes between eyes with varying angles of
22 corneal astigmatism (Table 4). There were 60 patients with the steepest corneal
23 meridian crossing the vertical axis of the MIOL in each eye, 13 patients with steepest
24 corneal meridian crossing perpendicular to the vertical axis in each eye and 6
25 patients with a parallel relationship to the MIOL in each eye. There was no significant
26 difference in individual questioned symptoms and overall QOV scores between the
27 groups (Figure 5).
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48 **Discussion**

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53 It is common to have varying levels of residual astigmatism following lens
54 replacement surgery and its magnitude can be difficult to predict due to multiple
55 dependent factors.¹⁸ Residual refractive astigmatism affects visual acuity^{13,19} and is
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1 one of the main causes of blurred vision^{11,12} following implantation of both monofocal
2 IOLs and MIOLs. However, it is not known how residual astigmatism affects
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4 asymmetric MIOLs. Therefore, this study sought to determine what effect residual
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6 refractive astigmatism and corneal astigmatism had upon QOV after implantation of
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8 an asymmetric Lentis Mplus MIOL.
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12 The current study demonstrates there is a significant difference in UDVA between
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14 patients who have ≤ 0.50 D compared to those with > 0.50 D of residual refractive
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16 astigmatism. It was also observed that the defocus equivalent was greater in the
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18 >0.50 D group confirming that the defocus equivalent was not a significant
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20 cofounding factor to the effect of increasing cylinder upon UDVA (Table 2). There
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22 was no significant difference in either UNVA or UIVA between the two residual
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24 refractive astigmatism groups. With traditional symmetric MIOLs it has been found
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26 that the distance visual acuity is significantly affected by uncorrected astigmatism.
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28 Hayashi et al¹⁹ found that with 0.50 D, 1.00D, 1.50D and 2.00D of simulated
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30 astigmatism distance visual acuity was significantly reduced at each magnitude of
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32 astigmatism with both a +3.00 D and +4.00 D addition symmetric MIOL (AcrySof
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34 ReSTOR). The same was found for monofocal IOLs, however the reduction in
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36 distance visual acuity was worse with the MIOLs. There was no significant difference
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38 between the MIOLs and monofocal IOLs in distance visual acuity up to 1.00 D and
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40 with astigmatism of 1.50 D and greater the distance and intermediate vision was
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42 better with monofocal IOLs. The near visual acuity was significantly better with the
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44 MIOL group up to 1.00 D of astigmatism. This study suggests that the MIOLs are
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46 useful up to 1.00 D of astigmatism. Hayashi et al¹³ found that with symmetric MIOLs
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48 (AMO Array) that the fraction of treated eyes able to achieve 0.16logMAR decreased
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1 for every 0.50D of astigmatism. Of the 30 eyes in their study, 24 achieved 20/29
2 (0.16 logMAR) and 20/50 (0.40 logMAR) with no astigmatism, 21 eyes with 0.50 D,
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4 13 eyes with 1.00 D, 8 eyes with 1.50 D and no eyes achieved this level of acuity
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6 with 2.00 D and 2.50 D of astigmatism. This study also compared these results to a
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8 monofocal IOL where they found that the MIOL group achieved a significantly worse
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10 distance visual acuity with 0.50 D, 1.00 D and 1.50 D of astigmatism, however the
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12 near visual acuity was better in the MIOL group. With higher levels of astigmatism,
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14 2.00 D and 2.50 D of astigmatism, the MIOL group showed significantly worse
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16 distance and intermediate visual acuity than the monofocal group. This study
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18 concludes that MIOLs are more affected by residual astigmatism than monofocal
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20 IOLs. These studies highlight that residual astigmatism has an effect on visual acuity
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22 at all distances however it appears the multifocality of MIOLs is not affected until
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24 astigmatism is greater than 1.00D. It appears that monofocal IOLs are better at
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26 distance and intermediate when astigmatism is greater than 1.00 D. In our study
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28 there were only 18 eyes with astigmatism of 0.75 D and greater because our
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30 refractive aim was emmetropia. This is one shortcoming of our study allowing only
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32 two separate groups of differing magnitudes of residual refractive astigmatism. It
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34 would be beneficial to have more groups to allow more detailed analysis between
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36 different magnitudes of astigmatism determining the exact level that begins to affect
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38 visual performance. However, it is clear from our study that there is a statistically
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40 significant difference in UDVA between the groups. Therefore, in this study it would
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42 appear that the objective UDVA is affected by increasing levels of astigmatism in a
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44 similar fashion to symmetric MIOLs,¹³ however the >0.50D of residual refractive
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46 astigmatism group still displays an excellent level of unaided visual acuity. As
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48 discussed in the study by Hayashi et al¹³ they report 21 eyes out of 30 (70%)
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1 achieved distance visual acuity of 0.16 logMAR for distance and 0.40 logMAR for
2 near with 1.00D and 1.50 D of astigmatism. In our study 14 patients out of 18 (77%)
3 displayed distance and near visual acuity of this level with astigmatism of 0.75 D and
4 above. However, unlike previous studies of symmetric MIOLs, residual refractive
5 astigmatism does not negatively affect UNVA and UIVA.¹⁹ Further analysis of
6 objective findings with asymmetric MIOLs is required and direct comparison to both
7 traditional symmetric MIOLs and monofocal IOLs would be beneficial.
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19 Objective outcomes only give an indication of visual performance therefore we
20 sought to determine how residual refractive astigmatism affected subjective
21 outcomes through the use of a QOV questionnaire. Analysis of the individual
22 symptoms investigated through the questionnaire in this study for each of the two
23 refractive astigmatism groups are shown in Figure 3. There was no significant
24 difference between the two groups for each of the questioned symptoms. In a study
25 by de Vries et al¹¹ of 76 eyes implanted with diffractive MIOLs in which 64.5% of
26 patients complained of blurred vision due to ametropia or astigmatism. The overall
27 mean refractive cylinder was -0.95D. In another study of 43 eyes with diffractive
28 MIOLs 25% had residual astigmatism of $\geq 0.75D$, and patients that complained of
29 blurred vision had mean astigmatism of 1.55D compared to the other patients where
30 the mean astigmatism was 0.53D.¹² A previous study found that patients with MIOLs
31 and astigmatism $>1.00D$ reported significantly bigger haloes compared to individuals
32 with astigmatism of $<1.00D$.²⁰ This contrasts with the findings from our study where
33 there was no significant difference in subjective visual complaints, such as glare and
34 dysphotopsias, between the groups. However our results with this asymmetric MIOL
35 were in broad agreement with those of Dick et al²⁰ which found that monofocal IOLs
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did not show a significant difference in frequency of reported haloes and glare between patients with <1.00D and >1.00D of astigmatism. An important aspect of the questionnaire used is that the patient is asked to report their overall QOV out of 10. This provides an overall indication of how satisfied an individual is with their QOV. In this current study each group achieved excellent QOV scores with no significant difference between the groups. Although there is a significant difference in UDVA with higher residual refractive astigmatism, similar to symmetric MIOLs this is not reflected in individual symptom responses or the overall QOV scores. As there was only one case with higher levels of residual defocus equivalent >1.50 D it is difficult to know absolute patient tolerances, however no one in this cohort required further laser refractive surgery despite its free availability. It would appear that asymmetric MIOLs are subjectively more tolerant to higher degrees of astigmatism.

There was some disparity between the refractive astigmatism and the corneal astigmatism found postoperatively. This may be due to in part to measurement error or the MIOL internally affecting the total refractive astigmatism present. Accurate measurement of this however can be somewhat difficult with the OPD scan due to the inability of Zernike polynomials to adequately decompose wavefront aberrations from both the distance and juxtaposed near add. Characteristically this can be interpreted in the OPD as coma rather than simply an increase in spherical power in the near add.²¹ This current study only found a significant increase in trefoil aberrations when comparing preoperative to postoperative which again may be inaccurate and due to the aforementioned problems. We therefore assessed the isolated effect of corneal astigmatism upon objective outcomes and patient satisfaction. The mean corneal astigmatism found in this study was -0.63 ± 0.36 D

1 and is comparable to other studies where Elkady et al²² found the mean astigmatism
2 to be $-0.63 \pm 0.62D$ after microincision cataract surgery. Our study showed that the
3 level of UDVA was significantly worse than the $\leq 0.50D$ group when astigmatism was
4 $0.76D$ to 1.00 ($P=<0.001$) and $>1.00D$ ($P=<0.001$) as outlined in table 3, however
5 again UDVA remained at an excellent clinical level. Similar to the previously
6 assessed refractive astigmatism this difference in UDVA did not appear to affect the
7 patient satisfaction as no significant difference was found between the postoperative
8 corneal groups. Likewise with residual refractive astigmatism there was no significant
9 difference in reports of blurred vision and haloes as previously found with increasing
10 levels of induced astigmatism.^{11,12,20}

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26 We also sought to determine the effect of the axis of the corneal astigmatism in
27 relation to the vertical axis of the asymmetric MIOL on postoperative objective and
28 subjective outcomes. To our knowledge this is the first study to determine the effect
29 of uncorrected astigmatism on asymmetric MIOLs and therefore the first study to
30 determine the effect of the axis on postoperative outcomes. In this current study we
31 found that the relation between the vertical axis of the MIOL and the corneal
32 astigmatism did not have a significant effect on objective or subjective outcomes.

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36 In summary this study highlights that asymmetric MIOLs are significantly affected by
37 residual astigmatism for UDVA however UNVA and UIVA appears to be unaffected.
38 Additionally, an increasing magnitude of residual astigmatism does not significantly
39 affect subjective outcomes. Also when implanting an asymmetric MIOL inferiorly with
40 nasal displacement the angle of the steepest corneal meridian in relation to this
41 placement does not affect either objective outcomes or overall subjective QOV. This

gives the clinician a better understanding of the effects of residual astigmatism after
CLE on patient satisfaction and will help the surgeon decide on clinical management
of residual astigmatism.

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What was known:

- Astigmatism affects unaided visual acuity in both symmetric MIOLs and monofocal IOLs. The greater the magnitude of residual astigmatism the worse the objective outcomes.
- Uncorrected residual astigmatism causes subjective reports of haloes and blurred vision in symmetric MIOLs.

What this study added:

- This rotationally asymmetric MIOL is not subjectively affected by increasing magnitudes of residual refractive or corneal astigmatism.
- The angle of the residual corneal astigmatism in relation to the position of the MIOL does not affect objective outcomes and overall patient satisfaction.

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Legends

Figure 1. Diagram displaying the relationship between the position of the MIOL and the various categorised angles of residual astigmatism. The vertical axis of the MIOL (reading segment inferiorly and nasally displaced) is represented by the line at 135 degrees in the right eye and 45 degrees in the left eye. *Top row:* Angles from 0 to 22.5 degrees and 68 to 112.5 in both the right and left eye cross the vertical axis of the MIOL obliquely and are categorised together in the oblique group. *Middle row.* Angles from 23 to 67.5 in the right eye and 113 to 157.5 in the left eye cross the MIOL perpendicularly and are categorised together in the perpendicular group. *Bottom row.* Angles from 113 to 157.5 in the right eye and 23 to 67.5 in the left eye cross the MIOL perpendicularly and are categorised together in the parallel group.

Figure 2a. Assessment of the relationship between the residual refractive astigmatism and UDVA, 6 months postoperatively. Linear regression analysis demonstrates a weak relationship between residual refractive astigmatism and UDVA of 234 eyes. (UDVA = uncorrected distance visual acuity; D = dioptres)

Figure 2b. Linear regression analysis demonstrates a weak relationship between the defocus equivalent and UDVA of 234 eyes at the 6 month postoperative assessment. (UDVA = uncorrected distance visual acuity; D = dioptres)

Figure 3. Histogram showing the mean overall individual symptom scores for different magnitudes of residual refractive astigmatism at the 6 months postoperative assessment for 234 eyes. The x-axis indicates the questioned symptom against the

1 average responses (0 = not at all, 1 = a little, 2 = quite, 3 = very) on the y-axis for
2 each of the four categorised groups of residual refractive astigmatism. A higher
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5 average score indicates the patient is more affected by the symptom.
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10 **Figure 4.** Histogram showing the mean overall QOV scores for different magnitudes
11 of residual refractive astigmatism at the 6 months postoperative assessment for 234
12 eyes. The QOV is rated out of 10, with 0 being the worst and 10 being the best.
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14 There was no significant difference between groups with varying magnitudes of
15 residual refractive astigmatism, $P=0.28$. (QOV = quality of vision; D = dioptres)
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24 **Figure 5.** Histogram showing the mean overall QOV scores for different angles of
25 residual corneal astigmatism in relation to the position of the MIOL. Eyes had corneal
26 astigmatism of greater than 0.50 D at the 6 months postoperative assessment. The
27 QOV is rated out of 10, with 0 being the worst and 10 being the best. There was no
28 significant difference between groups with varying angles of residual corneal
29 astigmatism, $P=0.43$. (QOV = quality of vision)
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Synopsis file

This study provides vital information on the effect of residual astigmatism on the Lentis Mplus MIOL where it was found that increasing magnitudes did not significantly affect postoperative patient satisfaction.

Figure 1
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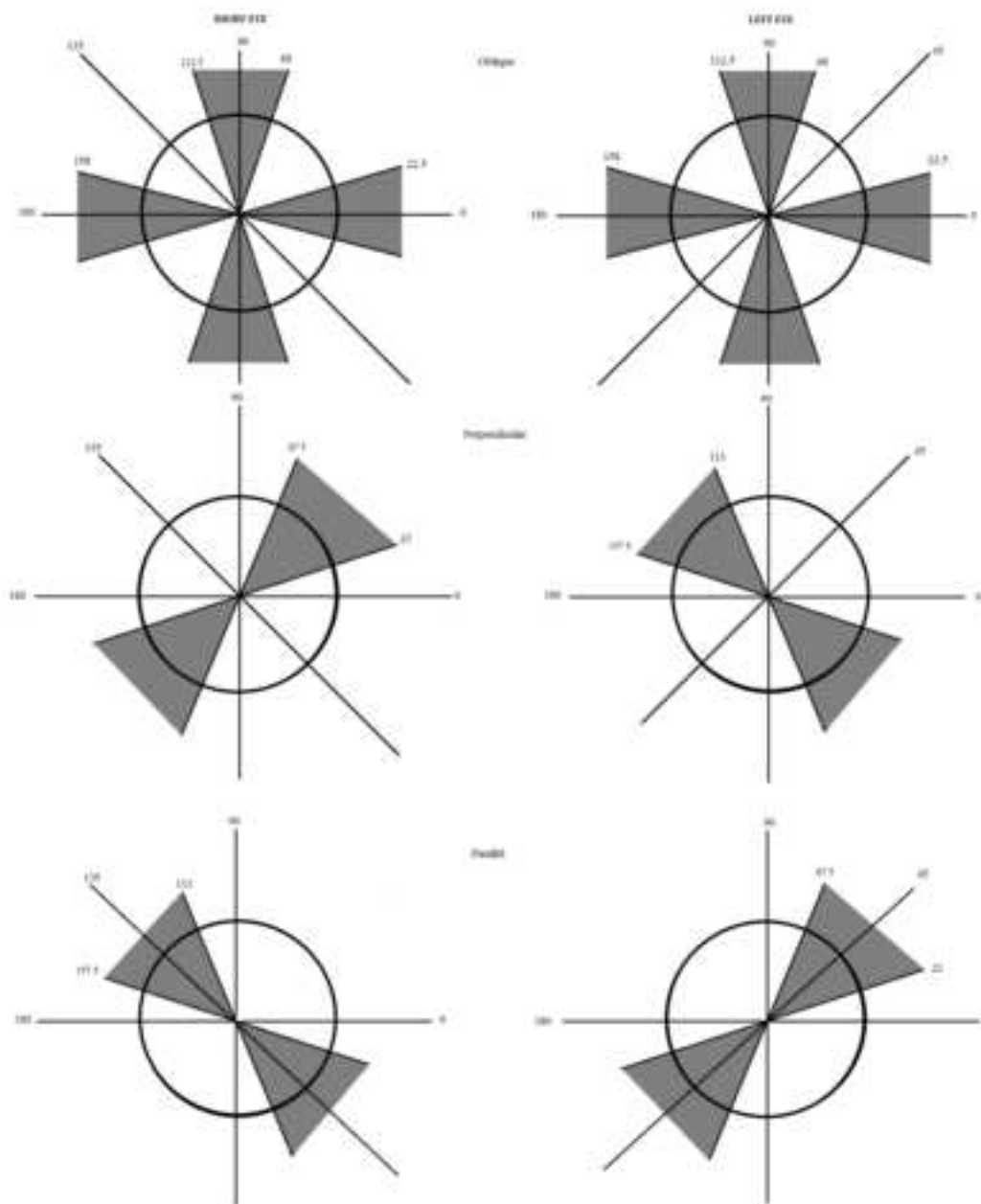


Figure 2a

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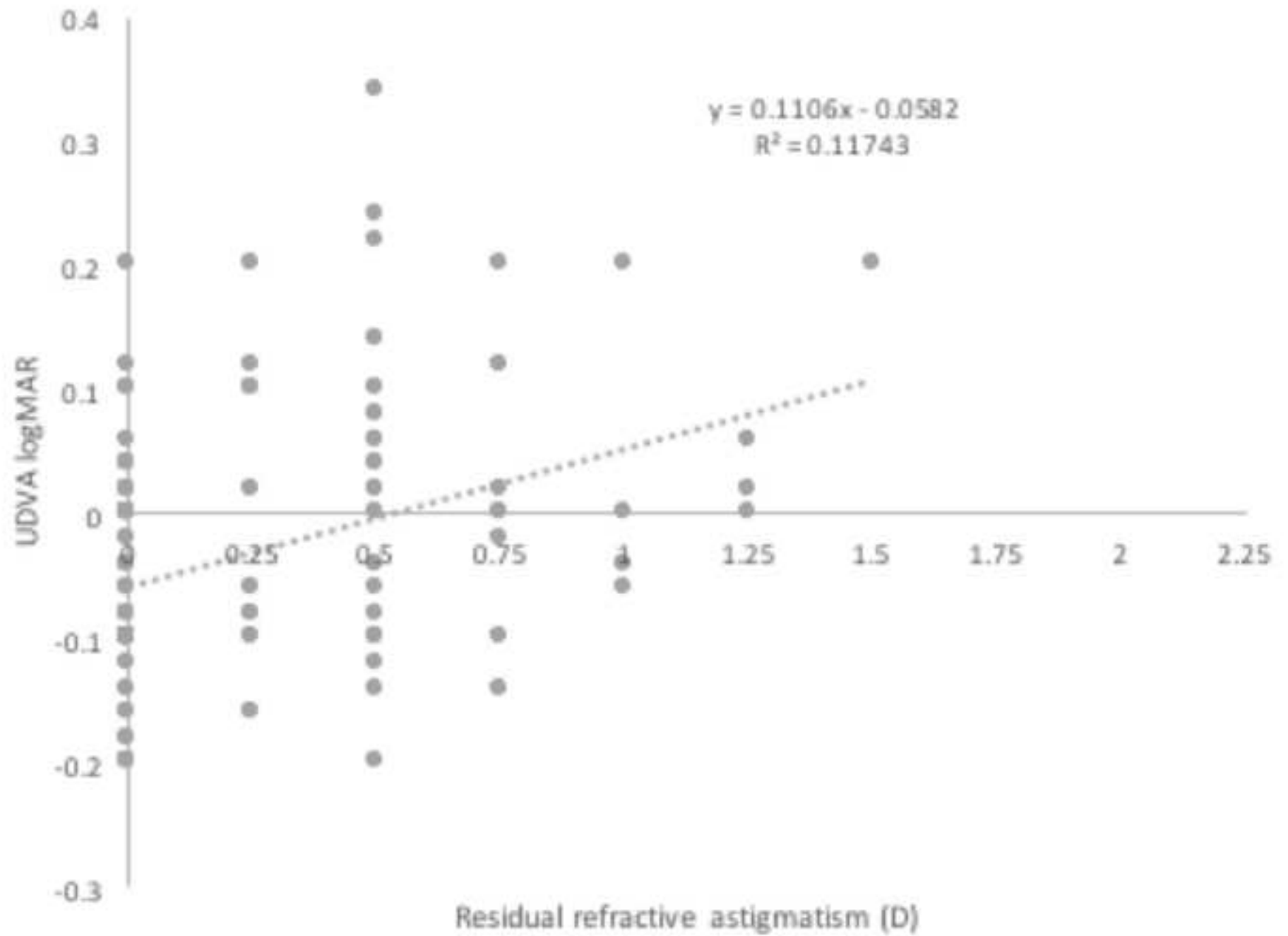


Figure 2b
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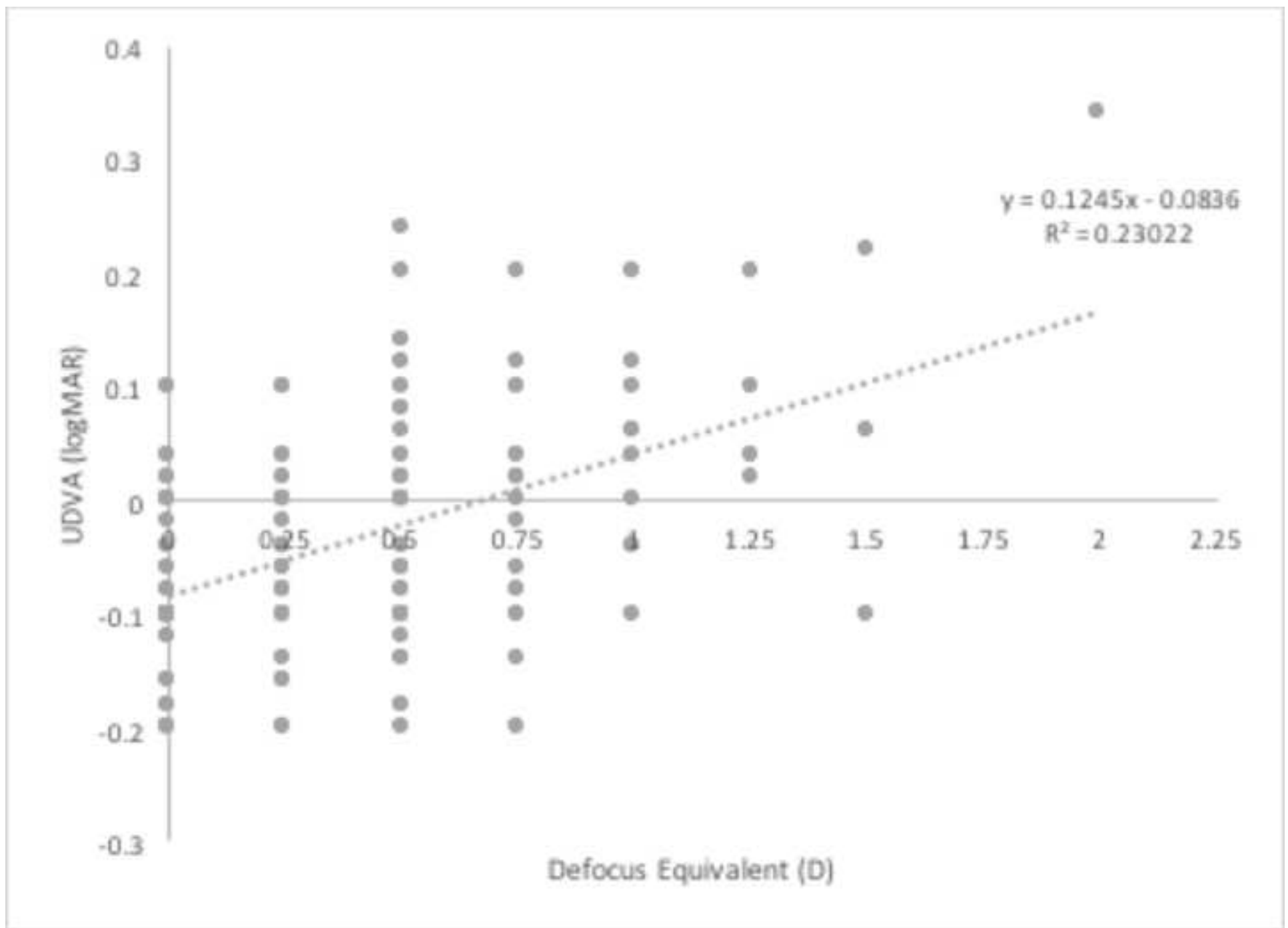


Figure 3
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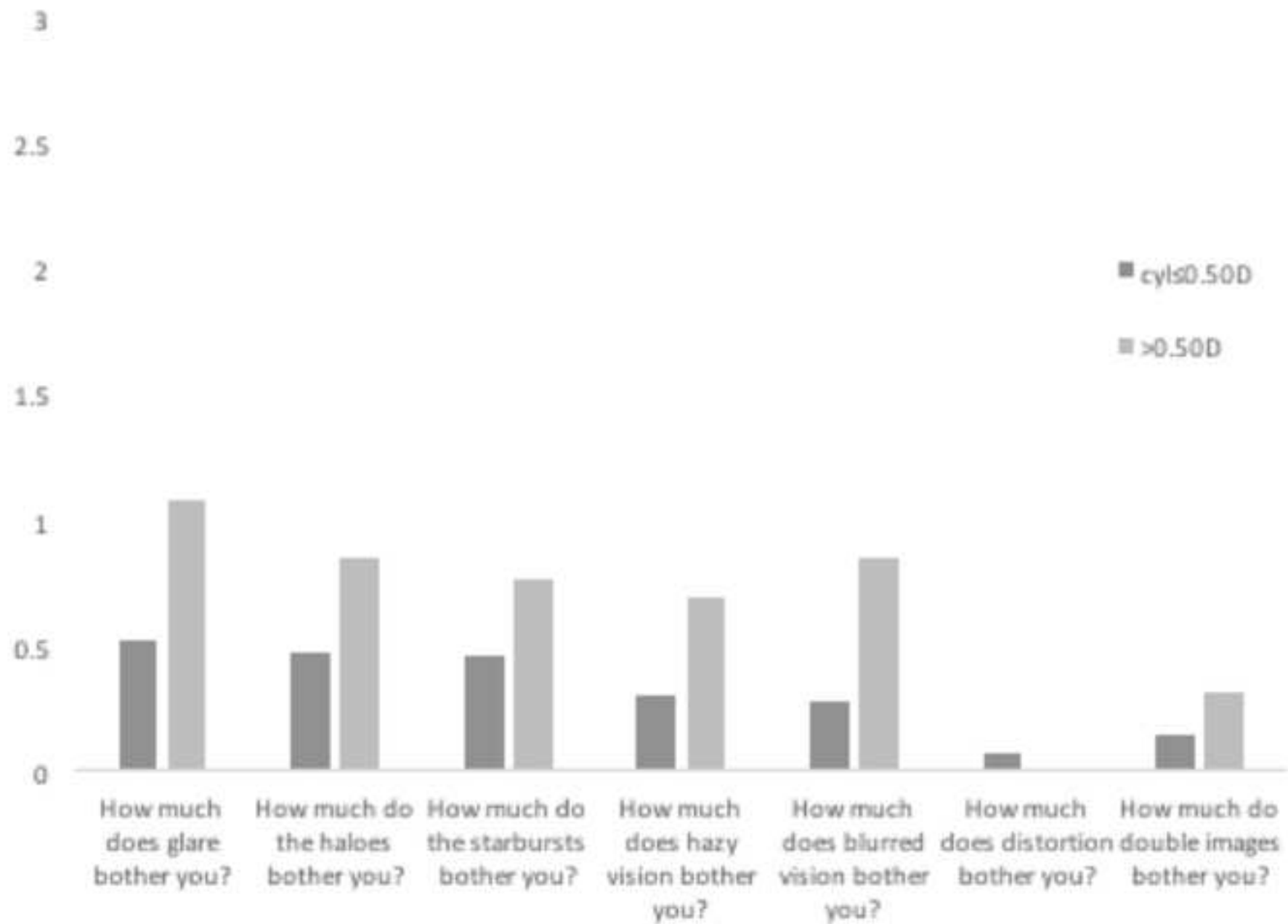


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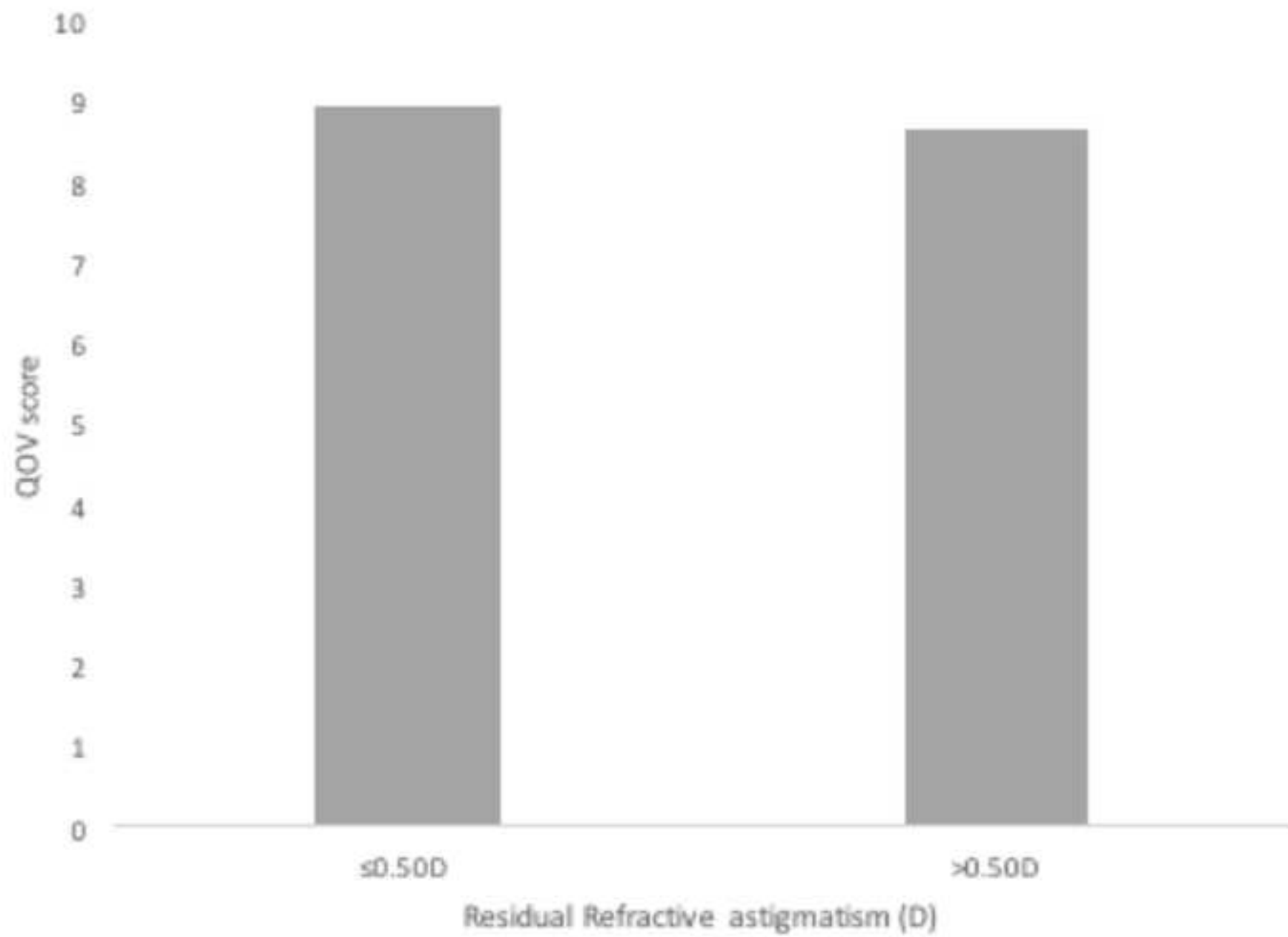


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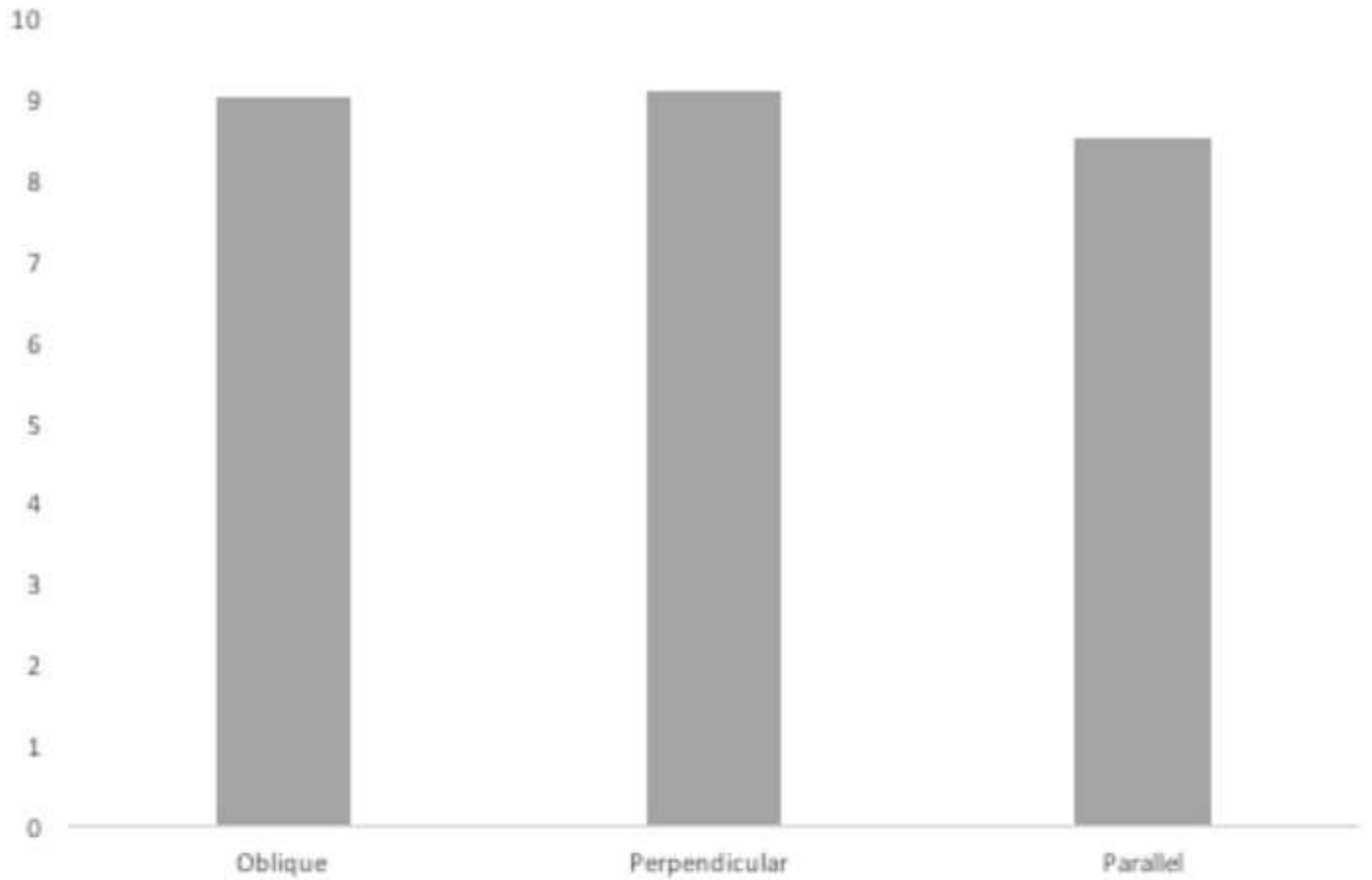


Table 1 Demographics

Demographics	
Patients (n)	117
Male, n (%)	48 (41)
Female, n (%)	69 (59)
Age (y)	
Mean \pm SD*	64 \pm 8.31
Median	64
Range	44, 87
Sphere (D†)	
Mean \pm SD	0.41 \pm 3.83
Median	1.50
Range	-16.50, 5.75
Cylinder (D)	
Mean \pm SD	-0.56 \pm 0.56
Median	-0.50
Range	-2.25, 0
LogMAR CDVA††	
Mean \pm SD	0.01 \pm 0.12
Median	0
Range	-0.20, 0.60

**SD = standard deviation; †D = dioptres; ††CDVA= corrected distance visual acuity*

Table 2

Between group comparisons of objective outcomes with different magnitudes of residual refractive astigmatism at the second postoperative assessment (6 months).

	≤0.50 D* (n=216)	>0.50D (n=18)	P Value
LogMAR			
UDVA†			
Mean ± SD††	-0.05 ± 0.09	0.02 ± 0.10	0.003
Range	-0.20, 0.34	-0.14, 0.20	
Sphere (D)			
Mean ± SD	0.08 ± 0.38	0.51 ± 0.47	0.001
Range	-1.50, 1.50	-0.25, 0.75	
Defocus equivalent (D)			
Mean ± SD	0.31 ± 0.34	0.83 ± 0.24	<0.001
Range	0, 2.00	0.50, 1.25	
UNVA§ (M notation)			
Mean ± SD	0.60 ± 0.17	0.64 ± 0.23	0.35
Range	0.32, 1.60	0.40, 1.25	
UIVA (M notation)			
Mean ± SD	0.96 ± 0.24	1.10 ± 0.38	0.15
Range	0.50, 2.00	0.80, 2.00	

**D = dioptres; †UDVA = uncorrected distance visual acuity; ††SD = standard deviation; §UNVA = uncorrected near visual acuity ; ||UIVA = uncorrected intermediate visual acuity*

Table 3

Between group comparisons of objective outcomes with different magnitudes of corneal astigmatism at the second postoperative assessment (6 months).

	≤0.50 D* (n=99)	0.51D to 0.75D (n=53)	0.76D to 1.00 (n=41)	>1.00D (n=41)	P Value
LogMAR					
UDVA†					
Mean ±			0 ± 0.08	0 ± 0.11	
SD††	-0.07 ± 0.08	-0.05 ± 0.09			<0.001
Range	-0.20, 0.20	-0.20, 0.24	-0.16, 0.20	-0.20, 0.34	
Sphere					
(D)					
Mean ±			0.16 ± 0.42	0.06 ± 0.45	
SD	0.10 ± 0.34	0.13 ± 0.46			0.72
Range	-0.50, 1.25	-1.00, 1.50	-0.50, 1.25	-1.50, 1.00	
Defocus					
equivalent					
(D)					
Mean ±	0.28 ± 0.30	0.37 ± 0.43	0.39 ± 0.33	0.43 ± 0.43	0.11
SD					
Range	0, 1.25	0, 1.50	0, 1.25	0, 2.00	
UNVA§ (M					
notation)					
Mean ±			0.65 ± 0.24	0.58 ± 1.26	
SD	0.58 ± 0.16	0.62 ± 0.20			0.14
Range	0.40, 1.25	0.32, 1.25	0.40, 1.60	0.40, 0.80	
UIVA (M					
notation)					
Mean ±			0.99 ± 0.20	1.04 ± 0.38	
SD	0.95 ± 0.20	0.97 ± 0.28			0.45
Range	0.80, 1.60	0.63, 1.60	0.80, 1.25	0.50, 2.00	

**D = dioptres; †UDVA = uncorrected distance visual acuity; ††SD = standard deviation ; §UNVA = uncorrected near visual acuity ; || UIVA = uncorrected intermediate visual acuity*

Table 4

Between group comparisons of objective outcomes for different angles of corneal astigmatism greater than 0.50 D at the second postoperative assessment (6 months or greater).

	Oblique axis groups (n=99)	perpendicular (n=22)	Parallel (n=14)	P Value
UDVA* (LogMAR)				
Mean ± SD†	-0.03 ± 0.10	0.01 ± 0.10	0 ± 0.08	0.23
Range	-0.20, 0.34	-0.10, 0.22	-0.10, 0.20	
Sphere (D††)				
Mean ± SD	0.12 ± 0.42	0.08 ± 0.61	0.16 ± 0.33	0.87
Range	-1.50, 1.25	-1.00, 1.50	0, 1.25	
Cylinder (D)				
Mean ± SD	-0.22 ± 0.35	-0.18 ± 0.22	-0.18 ± 0.42	0.80
Range	-1.25, 0	-0.50, 0	-1.50, 0	
UNVA§ (M notation)				
Mean ± SD	0.62 ± 0.21	0.62 ± 0.15	0.57 ± 0.10	0.65
Range	0.40, 1.60	0.32, 1.00	0.40, 0.80	
UIVA (M notation)				
Mean ± SD	1.02 ± 0.32	0.96 ± 0.22	0.94 ± 0.21	0.60
Range	0.50, 2.00	0.63, 1.25	0.80, 1.25	

*UDVA = uncorrected distance visual acuity; †SD = standard deviation; ††D = dioptres; §UNVA = uncorrected near visual acuity; ||UIVA = uncorrected intermediate visual acuity