Effects of intraocular lens glistenings on visual function: a prospective study and presentation of a new glistenings grading methodology

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ABSTRACT

Objective To investigate the effect of intraocular lens (IOL) glistenings on visual performance and evaluate a new glistenings grading methodology.

Methods and Analysis Thirty-four patients (34 eyes) were recruited. Corrected distance visual acuity (CDVA), mesopic gap acuity (MGA), functional contrast sensitivity (FCS) and forward light scatter were measured (Advanced Vision and Optometric Tests, City Occupational, London, UK). The IOL centre was imaged and glistenings density graded by three observers using the Miyata scale and a new system. Inter-rater reliability, association between the two grading scales, and correlations between glistenings grades and visual performance parameters were evaluated.

Results The intraclass correlation coefficient between graders for the new grading system was 0.769 (95% Confidence Interval [CI] 0.636 to 0.868). There was a significant association between the Miyata scale and the new grading system for all graders (rs=0.533–0.895, p<0.001). There was no association between CDVA or MGA and glistenings grade (rs=−0.098, p=0.583 and rs=0.171, p=0.359, respectively). There was no association between FCS at mesopic light levels and glistenings grade (rs=−0.032, p=0.864), or the straylight parameter and glistenings grade (rs=0.021, p=0.916). No association was found between the integrated straylight parameter and glistenings grade (rs=0.078, p=0.701).

Conclusion The new glistenings grading scale was highly reproducible. In this cohort, glistenings in the same hydrophobic acrylic IOL after cataract surgery were not associated with changes in visual function, as assessed by a series of tests not previously used in glistenings research.

INTRODUCTION

Glistenings are vacuoles that can develop within intraocular lenses (IOLs) implanted as part of cataract surgery. They occur in all IOL materials but are mostly associated with hydrophobic acrylic IOLs.1–3 Glistenings form when water permeates through microchannels within the material to create small fluid-filled inclusions that are up to 30 µm in size.4–9

While most previous studies have demonstrated no significant effect of glistenings on vision,10–16 a few have found that high numbers of such vacuoles within IOLs can impair visual performance,5 9 especially high spatial frequency contrast sensitivity.10–15

Labuz et al16 published an in vitro model for predicting the straylight parameter from the density of glistenings. They found that ‘large numbers’ of glistenings were needed to cause straylight levels that might lead to

Key messages

What is already known about this subject?
► There is a limited amount of evidence that intraocular lens (IOL) glistenings may have an effect on visual function and in particular high spatial frequency contrast sensitivity.
► Previously described subjective grading scales of IOL glistenings in vivo vary in terms of defining the IOL area/volume under investigation, specifying imaging equipment settings and ambient illumination levels.

What are the new findings?
► We have described a new, highly reproducible grading methodology with optimum equipment settings specifically developed for the purpose of grading glistenings and with carefully controlled ambient illumination.
► In this cohort, glistenings in the same hydrophobic acrylic IOL after cataract surgery were not associated with visual function parameters, including visual acuity, positive and negative contrast sensitivity at 10 cycles per degree, and forward light scatter.

How might these results change the focus of research or clinical practice?
► The detailed method for evaluating IOLs in vivo and the new grading scale may help standardise IOL glistenings grading.
► The new finer, highly reproducible grading scale may be more suitable for comparing modern IOLs with small amounts of glistenings as well as IOLs with relatively large number of glistenings.
glare-related visual problems. However, in vivo the issue of accurately determining the numbers and density of IOL glistenings remains a challenge, especially with regard to eye/patient macromovements and micromovements during imaging, variation of light entering the eye with natural physiological variations and anomalies, and the limitations of current clinical diagnostic imaging technologies. As a result, it is difficult to test the precise relationship between glistenings density and visual performance in vivo.

Previously published in vivo glistenings grading systems have typically used estimating or counting methodologies with a single examiner, differing reported slit-lamp (SL) magnifications (16×, 17 21 25×, 18 40×), 18 ordinal scale ranges ([0 to 2], 11 18 [0 to 3], 2 17 [0 to 3+], 21 [0 to +4], 20 [trace to 3+], 19 [trace to 4]), 5 22 and different highest grade cut-offs ([>50], 21 [40 ‘per field’], 2 18 [150 per mm²], 7 [200 per mm²]). Previous studies that have used subjective evaluation and grading of glistenings at the SL or from SL-derivered photographs have not defined the region of the IOL being studied, specified illumination levels or image capture parameters. 2 5 11 17 19 21 22 Moreover, correlation between glistenings grading at the SL and that of IOL images has been found to be only moderate. 22

To try and more accurately describe the relationship between IOL glistenings and visual performance, in this study we developed a new, precisely defined, subjective, finer, zonal grading system based on high-quality colour digital SL images taken with a described, fixed protocol and using three graders. Agreement of glistenings grading between the graders was analysed and the new scale compared with a previously described and published grading system in widespread usage. 7 To investigate the association between glistenings and visual performance, we employed a range of tests at photopic and mesopic light levels, including logarithm of the minimum angle of resolution (logMAR) visual acuity, straylight measurements, gap acuity and functional contrast sensitivity (FCS).

MATERIALS AND METHODS

Thirty-four patients were recruited between September 2017 and February 2018. All had undergone uncomplicated cataract surgery with phacoemulsification and were implanted with monofocal, spherical, hydrophobic acrylic IOLs (Alcon AcrySof SA60AT). Given the novel methodologies for visual testing employed in this investigation, it was designed as a pilot study to help generate results that could power a future main study. Inclusion criteria were age between 18 and 100 with corrected distance visual acuity (CDVA) in the study eye equal to or better than 0.2 logMAR. Prior to study entry, all subjects underwent an ophthalmic examination, including mydriatic funduscopy and, where appropriate, macular optical coherence tomography scans. Exclusion criteria included signs of dry eye, corneal opacities, failure of pupillary miosis beyond 5.0 mm, posterior capsule opacification, glaucoma, age-related macular degeneration, retinal vascular disorders, previous retinal detachments, neuro-ophthalmological conditions, inherited retinal disorders or pathology, previous strabismus surgery or history of amblyopia, previous transient ischaemic attack, cerebrovascular attack, or other vaso-occlusive disease or epilepsy.

Assessment of glistenings

Following pupillary mydriasis, with tropicamide 1% and phenylephrine 2.5%, central images of the IOLs were taken with a 5-megapixel digital camera (Topcon DC-4, Topcon, Tokyo, Japan) mounted on an SL (Topcon SL-701, Topcon). To obtain the best images of IOL glistenings, prior to starting this study, one author (NS) systematically evaluated different SL and digital camera settings to optimise quality of images.

The same mesopic conditions in the examination room were maintained, with only the assessor’s screen on minimal brightness turned away from the patient (ambient illuminance on the SL table did not exceed 0.5 lux). A vertical slit beam of 10.0 mm by 2.0 mm at an angle of 40° and 16× objective magnification was used with the SL set to maximum brightness to illuminate the centre of the IOL within the pupil. For the Topcon DC-4 camera, an ISO of 800 was used with a shutter speed of 1/30 s, a sharpness of ‘+32’ (default), a denoising of ‘0’ (default), a contrast of 50 (default) and the ‘auto-brightness’ setting at ‘off’.

The best of five colour digital SL images from each study eye (one per patient) was selected for analysis. Each was processed by fitting the pupil, identifying its centre and overlaying a 5.0 mm by 1.0 mm grid divided into 1.0 mm² areas (figure 1).

Three ophthalmologists (NS, NM and DPSO’ B) assessed and graded the digital images independently and within each of the five defined 1.0 mm² grid squares (Z+2, Z+1, Z0, Z−1, Z−2) by counting the number of glistenings they could identify within each separate area. Graders disregarded artefacts such as anterior or posterior IOL surface particulates (eg, pigment), IOL scratches and scuff marks (eg, from the IOL loading device or forceps), posterior capsule irregularities, and vitreous floats. To reduce assessor bias, image strips were randomised and presented to the graders for evaluation in PowerPoint (Microsoft, Washington, USA) by a fellow researcher not involved in grading (CH).

Grades of glistening density were then assigned to each 1.0 mm² area according to the following 8-point ordinal scale; grade 0=no visible glistenings, grade 1=1–10 per mm², grade 2=11–20 per mm², grade 3=21–30 per mm², grade 4=31–40 per mm², grade 5=41–50 per mm², grade 6=51–60 per mm² and grade 7 >61 glistenings per mm².

In addition, the central three zones for the new scale were also graded. This was undertaken to see if it provided greater reproducibility. In addition, a 3 mm pupil is the average pupil size under photopic conditions for most individuals over the age of 70. 23

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Figure 1  Slit-lamp image of pupillary centre (slit-lamp beam width and height setting: 2.0 mm and 10.0 mm, respectively) with an overlay of a 5.0 mm by 1.0 mm grid, which was divided into 1.0 mm by 1.0 mm areas, with the central area bisecting the pupillary centre.

In addition, all three graders assessed each digital image as a whole with the previously described Miyata scale. Agreement of glistenings grading between the examiners and association between the two scales were evaluated.

Visual testing
Photopic visual acuity was assessed using an Early Treatment Diabetic Retinopathy Study backlit chart at 4 m (Precision Vision, Illinois, USA).

Advanced Vision and Optometric Tests (AVOT, City Occupational, London, UK) were used under mesopic conditions where ambient illuminance on the display surface did not exceed 0.3 lux and background display screen luminance was 1 Cd m⁻²:
1. Gap acuity using a Landolt ring for both positive and negative contrast at a viewing distance of 3 m. The orientation of the Landolt ring was restricted to four locations and the subject pressed one of the four buttons available on a bespoke keypad to indicate the location of the gap.
2. FCS using a Landolt ring (as for mesopic gap acuity) with a 3’ gap for both positive and negative contrast at a viewing distance of 3 m. A gap size of 3’ (Landolt ring size 15’ or 10 cycles per degree) was employed to avoid eyestrain and to minimise the effects of microfluctuations of accommodation. FCS is a measure of contrast sensitivity that has been found to be relevant in occupational environments.
3. Forward light scatter test. The forward light scatter test implemented in the AVOT system used a flicker cancellation method. It employed a single ring of fixed size designed to produce specified luminance levels in the plane of the pupil from a viewing distance of 70 cm. The participant used a single button on a bespoke keypad to neutralise the flicker that appeared on the screen. Both ‘low threshold’ and ‘high threshold’ were evaluated. The software calculated the straylight parameter and integrated straylight parameter.

The association between glistenings grade and all the tested visual performance parameters described above were tested.

Statistical methods
One eye was randomly selected for each participant. Data organisation and descriptive statistics were handled with Excel 2013 (Microsoft Corporation, WA, USA), with further statistical analyses performed with Minitab V.14 (Minitab, Pennsylvania, USA).

The subjective glistenings grade was calculated as the median of all five zones for each grader since the data were skewed and the grades were on an ordinal scale. The overall grade used in the analysis of association with visual performance parameters was taken as the median of all three graders. Inter-rater reliability was assessed using the intraclass correlation coefficient. Agreement between grades for the median of the central three
zones and the median of all five zones was assessed using weighted kappa. Association between the new grading system and the Miyata scale was assessed with a χ² test for independence. Associations between glistenings grade and visual performance parameters were measured using Spearman’s rho. The significance level was taken to be 5% in all cases.

**RESULTS**

**Patient demographics**

The mean subject age was 72.9±5.1 years (Standard Deviation [SD], range 63–85 years). Eleven subjects (32%) were female and 23 (68%) were male. Study participants had undergone cataract surgery between 5 and 66 months previously (median 14 months), with 50% of the participants having had surgery 9.75–14.25 months previously. Random selection resulted in 24 (71%) right and 10 (29%) left eyes.

**Glistenings grades**

Of the 34 IOLs evaluated with the new grading scale, 20 (59%) had grade 1 glistenings, 8 (23%) grade 2, 3 (9%) grade 3, 1 (3%) grade 4 and 2 (6%) grade 5 when considering all five zones.

With regard to the central three zones, 17 (50%) had grade 1, 9 (26%) grade 2, 5 (15%) grade 3, 1 (3%) grade 4, 1 (3%) grade 5 and 1 (3%) grade 7 glistenings.

Using the median grade for all three graders with the Miyata scale, of the 34 eyes, 1 (3%) was grade 0, 27 (79%) grade 1, 9 (26%) grade 2, 5 (15%) grade 3, 1 (3%) grade 4 and 2 (6%) grade 5 when considering all five zones.

**Reliability and reproducibility of the new grading system**

The intraclass correlation coefficient, for the median of the grade for all five zones, was 0.769 (95% CI 0.636 to 0.868), indicating a ‘good’ level of agreement between graders. The intraclass correlation coefficient for the median of the grade of the central three zones was slightly higher at 0.813 (95% CI 0.696 to 0.895). Intraobserver agreement between the grades for all five zones versus the central three zones for the three graders was assessed using quadratically weighted kappa. The values of 0.948, 0.875 and 0.854, respectively, demonstrated ‘very good’ levels of agreement. The association between the Miyata and the new grades (median of the five zones) for all three graders was statistically significant (p≤0.003).

**Association between subjective assessment of glistenings and visual function logMAR distance acuity**

For the 34 study eyes, the mean logMAR uncorrected distance acuity was 0.06±0.12 (SD) (median 0.01, range −0.1 to 0.5). The mean CDVA was −0.04±0.07 (SD) (median −0.06, range −0.2 to 0.18).

There was no association between glistenings grade and high-contrast CDVA when the median grade for the five zones was considered (r_s=−0.098, p=0.583) nor when the central three zones were considered (r_s=−0.045, p=0.80).

**Gap acuity (minutes of arc)**

At positive target contrast, there was no association between gap acuity and grade of glistenings when the median grade for the five zones was considered (r=0.171, p=0.359) nor when median grade for the central three zones was considered (r=0.137, p=0.461). Similarly, at negative target contrast, there was no association between gap acuity and grade of glistenings when the median grade for the five zones was considered (r=0.009, p=0.962) nor when the median grade for the central three zones was considered (r=0.007, p=0.968). There was no statistically significant difference (p=0.691) between positive and negative mesopic gap acuity. Three of 34 (9%) subjects could not complete the mesopic gap acuity test.

**Straylight**

There was no association between the straylight parameter, k, and the median grade of glistenings for neither the five zones (r_s=0.091, p=0.916) nor for the central three zones (r_s=0.050, p=0.803). Similarly, there was no association between the integrated straylight parameter, k’, and the median grade of glistenings for neither the five zones (r=0.078, p=0.701) nor the central three zones (r=0.088, p=0.663). Seven out of 34 (21%) subjects could not complete the light scatter test.

**DISCUSSION**

The results of the present study add to the evidence that glistenings, unless possibly present in extremely large amounts, have a minimal effect on visual performance in vivo. Our results add new knowledge because they confirm the results of some previous studies,10–16 but used novel visual assessment methods and a novel grading system using three graders and strictly controlled imaging and ambient illuminance parameters.

Our results demonstrate good agreement between graders and a high level of correlation with an existing scale.7 Our grading system offers benefits over that described by Miyata7 as it has precisely defined imaging and ambient illuminance parameters, as well as definition of the region of the IOL being analyzed, which can be reproduced by other clinical investigators. Indeed, we find the Miyata system was somewhat confusing as it originally described the grading scale in three dimensions29 or per mm³ but later1 referenced the scale to a two-dimensional image or per mm². This might explain why the high number of glistenings in the highest Miyata grade does not seem to correspond to the number of glistenings observed in the actual reference images.7,29

Glistenings are small fluid-filled inclusions, up to 30 µm in size.4–9 Detection of glistenings is dependent on their luminance contrast and not size. Therefore, provided there is enough light reflected they will be seen, although some appear fainter than others. Rather than size, one...
of the main difficulties is to resolve multiple glistenings which are close together. To help overcome this, our digital image strips were divided into five separate 1 mm² areas, centred on the pupil and presented at full screen size on a computer monitor for grading. Three graders performed the grading, allowing us to evaluate both the interobserver and intraobserver reliability of our method, which was very high. Such accuracy of a grading system based on counting is not possible, with observers counting at the SL; this is due to factors such as patient discomfort and movement due to bright light and lack of reference points, and the large numbers of glistenings requiring counting in some cases.

As discussed above, unlike previous studies that used digital images and counting methodologies, we systematically analysed and optimised the parameters of a commercially available SL-based digital imaging system under controlled conditions to reduce artefacts and maximise the quality of glistenings images taken in vivo. In addition, we identified, imaged and graded the same central pupillary area of each IOL and divided this area for analysis into five 1 mm² areas in which the numbers of glistenings could be reliably identified and counted. Finally, to provide a further level of consistency, we excluded any eyes and patients for analysis with ocular and/or neurological comorbidities that might affect visual performance and all participants had the same design of IOL. Using our system, we analysed both the grades from all five 1 mm² zones and the central three zones. Using the central three zones may be more relevant to the photopic pupil size of those aged 70 or above, but have been used in other vision research. FCS has been shown to be a more sensitive indicator of changes in the quality of the retinal image caused by small residual refractive errors, higher order aberrations and/or scattered light. As the FCS test provides a measure of contrast sensitivity that has been shown to be relevant in occupational environments, it seems appropriate for evaluating how IOL glistenings can affect image changes caused by increased scattered light and aberrations and for evaluating the ‘real-life’ visual effects of IOL glistenings.

There are limitations to our study. We evaluated only the number of glistenings, while their size and surface portion might also affect visual function. The glistenings grades in our study participants had a narrow IQR probably because the time post cataract surgery was closely grouped around a median of 14 months and due to manufacturing changes introduced in the past few years in the IOL implanted in these patients to limit glistenings. Our results, however, included patients with what would have been described by a previous grading system as mild to severe levels of glistenings. Indeed, if we applied the grading scale as described by Christiansen et al. to our cohort, then 24% of our cases (8 out of 34 eyes) would have ‘severe glistenings’ (in these eyes at least one of the five squares was graded as our grade ‘5’ by at least one grader). We believe that our scale by virtue of its eight grading steps allows more detailed grading of the highest glistenings densities than other grading systems, which tend to group such changes together. Moreover, by allowing finer stratification of glistenings densities that may not have been isolated by other grading scales, it may be more suitable for comparing modern IOLs with fewer glistenings.

It is difficult to compare our grading system with the semiautomated counting methods, which are largely operator-dependent and may detect image artefacts as well as true glistenings. It has been postulated that Scheimpflug images might be used to detect glistenings in vivo. However, it appears that this method is not suitable for such evaluation as it cannot distinguish between light scatter due to glistenings and that from other optical changes, such as aqueous–IOL interface, posterior capsule or debris on the IOL surface. Indeed, Biwer et al found that a Scheimpflug device did not provide images of required resolution to perform automated counting of separate glistenings. This has also been our experience with a Scheimpflug device (Pentacam HR; Oculus Optikgeräte, Wetzlar, Germany).

Colin et al. reported that approximately a third of their AcrySof IOL cohort had no glistenings, a third had grade 1 glistenings and about a third had grade 2 glistenings (the most severe grade in their system). Similarly, Christiansen et al. found that 65% of their AcrySof patients had ‘trace’ glistenings, while grades ‘1+’ and grade ‘2+’ accounted for 25% and grades ‘3+’ and ‘4+’ accounted for 5% each. Our results, where, for the central three zones, 50% had grade 1, 41% had grades 2–3 and 9% had grade 4 or above, appear to show a similar range of IOL glistenings.

Three of the 34 (9%) of our subjects could not complete the FCS test, and 7 of 34 (21%) could not complete the light scatter test. The AVOT tests we used are essentially psychophysical tests requiring high levels of cognitive functioning. The limitation of the FCS test is its length of time if a participant is not able to resolve the 3’ Landolt ring gap accurately, and the light scatter test is limited because it cannot test participants with very low flicker sensitivity. These limitations of test methods leading to some incomplete data are also found in previously reported studies, using other assessment methodologies. Colin and Orignac reported that in only 53 of 97 cases (54%) could valid measurements be produced when they measured intraocular light scatter in their cohort of patients with glistenings, with the C-Quant test (Oculus Optikgeräte, Wetzlar, Germany).

In common with many previous studies, we have demonstrated no significant effect of glistenings on visual
function as measured by a series of novel tests under strictly controlled conditions. In order to standardise glistenings grading in vivo, we developed a new protocol that uses three graders and that images IOL with optimised digital camera parameters under controlled ambient illumination, which may be a useful tool for future research.

**Contributors**

NS: conception, design, collecting data, analysis and interpretation of data, drafting the article, revising the article. DPSO'B: conception, design, analysis and interpretation of data, drafting the article, revising the article. NM: conception, design, data analysis, drafting the article, revising the article. CCH: conception, design, analysis and interpretation of data, drafting the article, revising the article.

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**Competing interests**
DPSO'B reports grants and personal fees from Alcon, outside the submitted work. CCH reports grants from Alcon, outside the submitted work.

**Patient consent for publication**
Not required.

**Ethics approval**
This study was approved by London - Bloomsbury Research Ethics Committee. This research conformed to the tenets of the Declaration of Helsinki.

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