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Psychophysical Impact and Optical and Morphological Characteristics of Symptomatic Non-Advanced Cataract.

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Psychophysical impact and optical and morphological characteristics of symptomatic non-advanced cataract

Abstract

Purpose To investigate whether psychophysical, morphological, and/or optical characteristics of symptomatic non-advanced cataract are complementary to, or more appropriate than, visual acuity (VA) for the purposes of recording visual data that reflect subjective visual difficulty in patients with cataract that exhibit relative sparing of high contrast acuity (0.4 logarithm of minimal angle of resolution (logMAR) scale or better). Methods Eighty-two patients with symptomatic non-advanced cataract and no other ocular pathology were asked to complete a validated questionnaire, and to perform a series of visual function assessments including: corrected distance VA (CDVA); photopic and mesopic contrast sensitivity; photopic and mesopic glare disability (GD); reading acuity and reading speed; stereoacuity; and retinal sensitivity. Optical and morphological characteristics of the cataract were evaluated by lens optical density and by the Lens Opacities Classification System III, respectively. Correlations between questionnaire score and each of these measures were calculated. Results Statistically significant negative correlations were observed between the Rasch-scaled questionnaire score and mesopic GD (at 3 and 6 cycles per degree (cpd); r = -0.396 (P < 0.01) and -0.451 (P < 0.05), respectively) and between the Rasch-scaled questionnaire score and photopic GD (at 3 and 6 cpd; r = -0.328 (P < 0.01) and -0.440(P < 0.01), respectively).

Conclusion Symptomatic non-advanced cataract, in the presence of good CDVA, is

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associated with measurable subjective visual difficulty, best reflected in a decrease in mesopic and photopic GD (at medium spatial frequencies). CDVA does not reflect the patient's visual dissatisfaction in such cases. *Eye* advance online publication, 10 June 2011; doi:10.1038/eye.2011.123

Keywords: psychophysical; symptomatic; non-advanced; cataract

Introduction

Age-related cataract is the most common eye disorder in the western world and, according to the World Health Organization, is one of the principal causes of blindness worldwide, therefore representing a major public-health issue.¹⁻⁴ Not surprisingly, cataract surgery by phacoemulsification and intraocular lens (IOL) implantation is the most commonly performed surgical procedure worldwide.^{3,5-7}

Valid evaluation of cataract and its visual consequences is indispensable for the purpose of investigating possible risk factors for cataract formation and for documenting progression of cataract in epidemiologic and cohort studies. Also, reliable and valid assessment of cataract and its visual sequelae is equally important for clinical practice, if the ophthalmologist is to accurately evaluate the impact of age-related cataract on patient's visual performance and experience, and therefore identify those who are likely to benefit from surgical intervention. This is especially true in an era of falling thresholds for cataract surgery.^{8,9}

Herman Snellen (1834–1908) developed a standardized test for visual acuity (VA) in

1862.¹⁰ This test is still widely used today and, in fact, represents the most frequently employed method of assessing visual function in cataract patients in clinical practice.¹¹ The greatest limitation of using VA measurement as a means of assessing visual function rests on the fact that acuity is a measure of the resolving power of the eye, but in terms of contrast sensitivity (CS) function it is defined as the highest spatial frequency that can be detected at 100% contrast.^{12,13} However, the real world contains few visual stimuli at 100% contrast, but is full of visual stimuli of contrast lower than 100%. Additionally, the well-lit environment of an ophthalmologist's office may not compare with the dimmer settings in which most individuals engage in daily activities. This, in turn, suggests that other (ie, non-acuity) measures of CS are better indicators of visual performance and experience for purposes of everyday life.14,15

The questionable ecological validity of VA charts hinders interpretation of patients' functional visual difficulty following the development of an ocular disease, such as age-related macular degeneration,¹⁶ cataract, glaucoma, and diabetic retinopathy.¹⁷

Consequently, it has been suggested that VA is not the most appropriate tool to investigate the impact of symptomatic non-advanced cataract on visual function and experience,¹¹ and complementary and/or alternative techniques that have been investigated include CS,¹⁸ glare disability (GD),¹⁹ reading performance,²⁰ and functional visual difficulty assessed by various questionnaires.²¹⁻²³ Morphological and optical characteristics of lens opacification are also important, and can be assessed by various grading systems²⁴ and Scheimpflug photography,^{25,26} respectively, and have the advantage of being independent of patient cooperation and ocular co-morbidity.

The purpose of this study was to investigate whether psychophysical, morphological, and/or optical measures of cataract are complementary to, or perhaps more appropriate than, corrected distance VA (CDVA) for the purposes of recording visual data that reflect subjective visual difficulty experienced by patients with cataract that exhibit relative sparing of high contrast acuity (0.4 logarithm of minimal angle of resolution (logMAR) or better).

Materials and methods

Consecutive patients with symptomatic non-advanced cataract and no other ocular pathology, scheduled to undergo phacoemulsification cataract extraction and implantation of the Tecnis 1-Piece (T1P) IOL (Advanced Medical Optics Inc., Santa Ana, CA, USA) for visual improvement, were recruited from the clinics taking place at a high-volume surgical practice between July 2009 and February 2010. For the purposes of this study,

non-advanced cataract was defined as cataract presenting with CDVA better than or equal to 0.4, as measured in the clinic using the logMAR chart provided by Test Chart 2000^{PRO} (Thomson Software Solutions, Hatfield, UK) at a test distance of 4 m. Only one eye of each patient (the one with the worse CDVA) was recruited. The fellow eyes of 65 of the 82 study eyes were phakic, whereas 17 were pseudophakic. Ethics committee approval was secured from the Local Regional Ethics Committee and the research was conducted in accordance with the principles laid down in the Declaration of Helsinki. Written and informed consent for participation in the study was obtained from all recruited patients before recruitment in the study.

Exclusion criteria were as follows: any preoperative ocular co-morbidity, a history of ocular trauma, diabetes mellitus (according to self-reported medical history; screening for diabetes mellitus was not performed for the purpose of this study;), and previous intraocular surgery.

Data were collected 2–4 weeks before planned cataract surgery and included: questionnaire score; CDVA; CS; GD; reading acuity and reading speed; stereoacuity; retinal sensitivity; lens optical density (LOD); and Lens Opacities Classification System (LOCS) III score.

Subjective visual difficulty

Subjectively perceived visual difficulty in everyday life was evaluated using the Priquest questionnaire.²² The questionnaire was self-administered, but the examiner was available to answer patients' queries in relation to any of the questionnaire items. A copy of the items in the Priquest questionnaire is given in Table 1. Rasch analysis was applied to the questionnaire data, thus calibrating item difficulty and patient ability on the same scale.^{27,28}

Visual acuity

CDVA was measured monocularly and with the patient's best subjective refraction, using the logMAR chart provided by Test Chart 2000^{PRO} at a test distance of 4 m.

Contrast sensitivity

CS was measured using both a letter chart (Test Chart 2000^{PRO}) and the sine-wave grating contrast test system: 'Functional Acuity Contrast Test' (FACT) (Optec 6500 Vision Tester, Stereo Optical Co., Inc., Chicago, IL, USA).

Letter CS

At a constant room illuminance of 870 lx and with distance correction on, each patient was asked to identify

Table 1 Items in the Priquest questionnaire

Because of your vision, do you have difficulty with the following activities? (Response options: no difficulty, some difficulty, great difficulty, and extreme difficulty)

1. Reading newspaper print

- 2. Recognizing the faces of people you meet
- 3. Reading the prices of goods when you shop
- 4. Seeing to walk on uneven ground
- 5. Seeing to do needlework, etc
- 6. Reading TV text
- 7. Seeing to carry out a preferred activity/hobby
- 8. Do you feel that headlights, lamps, sunlight, or other lights dazzle you, reducing your vision?
- 9. Do you experience visual disturbances from differences
- (clarity, color, poor depth perception) between the two eyes?10. If you have a job, does your present vision cause any problems?
- 11. If you are a car driver, does your present vision cause any difficulties in driving?
- 12. If you look after yourself or care for someone at home, does your present vision cause any problems?

Table 2 Contrast levels tested with letter chart

Contrast (%)	CS
100	1.00
70.8	1.41
50.1	2.00
35.5	2.82
25.1	3.98
17.8	5.62
12.6	7.94
8.9	11.24
6.3	15.87
4.5	22.22
3.2	31.25
2.2	45.45
1.6	62.50
1.1	90.91
0.8	125.00
0.6	166.67

Abbreviation: CS, contrast sensitivity.

ETDRS letters in logMAR form, monocularly with the study eye, and at a distance of 4 m. The contrast of letters of predetermined size (and therefore predetermined spatial frequency) was reduced incrementally using the software's contrast adjustment function until the contrast threshold of the patient was reached (ie, the patient could read no more letters). Table 2 shows the increments of decreasing contrast tested and their corresponding CS. Six different letter sizes (and corresponding spatial frequencies) were tested, each having the following angular subtense in cycles per degree (cpd): 6/120 letters (1.2 cpd), 6/60 letters (2.4 cpd), 6/24 letters (6 cpd), 6/15 letters (9.6 cpd), 6/9.5 letters (15.16 cpd), and 6/6 letters (24 cpd).

Functional Acuity Contrast Test

At a room illuminance of 1.5 lx, monocularly with the study eye, and with distance correction, each patient was asked to look into the FACT. The patients were asked to look at five linear sine-wave grating charts of 1.5, 3, 6, 12, and 18 cpd, respectively. Each chart consisted of nine circular patches containing gratings of decreasing contrast, arranged in two rows (five patches above and four patches below). Patients were instructed to identify the orientation of the gratings by choosing 1 of 3 options: gratings tilted left (+15 degrees), gratings upright (0 degrees), or gratings tilted right (-15 degrees). Testing was performed under two different conditions: night (mesopic: 3 candelas (cd/m²)) and day (photopic: 85 cd/m^2), in that order.

Glare disability

The FACT was repeated in a similar manner as performed to test CS, under mesopic and photopic conditions, in that order, but this time with additional glare light (1 lx for night and 10 lx for day glare testing).

Reading performance

Reading speed and near VA were measured with an English version of the standardized Radner Reading charts, the reliability and reproducibility of which have been established, both for subjects with normal eyesight and for those with visual difficulty.²⁹

Stereoacuity

Stereoacuity was measured using the TNO stereo test (Lameris Instrumenten, Utrecht, Netherlands), an anaglyptic random dot test. The patients performed the quantitative part of the test (plates V–VII) and were awarded the stereoacuity of the smallest stereoacuity target that was correctly identified.

Retinal sensitivity

Retinal sensitivity was measured by performing microperimetry, with the Microperimeter MP 1 (Nidek Technologies Srl, Albignasego (PD), Italy). The patient's study eye was pharmacologically dilated with one drop of guttae Tropicamide BP 1% w/v minims (Chauvin Pharmaceuticals Ltd, Romford, UK) 20 min before the test. The central 10 degrees of fixation were examined. Retinal sensitivity was calculated for four areas: fixation (1 stimulus), within the central 5 degrees of fixation (average of 5 stimuli), between 5 and 10 degrees of 4

fixation (average of 8 stimuli), and within 10 degrees of fixation (average of 13 stimuli).

Lens optical density

LOD was measured using Scheimpflug images taken by the Pentacam Comprehensive Eye Scanner (software version 1.16; Oculus Inc., Wetzlar, Germany), the repeatability and validity of which have been established.³⁰ The study eye was pharmacologically dilated. Twenty-five images of the anterior segment of the eye were acquired. Density data were calculated by the software from the sum of the gray-scale values for the pixels along the axial length of the crystalline lens obtained from the cross-sectional image of the Scheimpflug camera. The density of the lens is standardized from 0 to 100. Therefore, 0 means the lens shows no clouding, 100 means the lens is completely opaque. The gray scale of the individual Scheimpflug images provides the basis for objective quantification. The 'densitometry along a line' part of the software was used to analyze the images and a mean LOD value was recorded directly from the visual axis line appearing in the Scheimpflug image.

LOCS III

Cataracts were categorized and graded clinically at the slit lamp by a single ophthalmologist using the LOCS III.²⁴Cataracts are graded in terms of nuclear opalescence/nuclear color (1–6), cortical opacity (1–5), and posterior subcapsular opacity (1–5).

Statistical analysis

Statistical analysis was performed using the software package PASW Statistics 18.0 (IBM Corp., Somers, NY, USA). Descriptive statistics were calculated for all measured variables, including demographic, ocular, refractive, psychophysical, cataract optical and morphological data, as well as data on subjective functional visual difficulty (questionnaire).

Correlations between the different psychophysical measures of visual function, as well as optical characteristics of the studied cataracts, and Rasch-scaled questionnaire score were investigated using Spearman's rank correlation. Tests were two-sided. A one-way analysis of variance test with a Games-Howell *post hoc* analysis was used to compare the Rasch-scaled questionnaire score between the different cataract morphology (LOCS III) subgroups, and also between the different stereoacuity subgroups. Levene's test was used to test for equal variances. In all analyses, a *P*-value <0.05 was considered statistically significant.

Results

Eighty-two patients (82 eyes) met the inclusion criteria and were recruited into this study. The demographic, ocular, refractive, psychophysical, cataract optical and morphological data, as well as data on subjective visual difficulty, are given in Table 3.

We found statistically significant negative correlations between Rash-scaled Priquest score and mesopic GD (measured at 3 and 6 cpd; r = -0.396 (P < 0.01) and -0.451 (P < 0.05), respectively) and between Rasch-scaled Priquest score and photopic GD (measured at 3 and 6 cpd; r = -0.328 (P < 0.01) and -0.440 (P < 0.01), respectively).

We also report statistically significant positive correlations between preoperative Rasch-scaled Priquest score and CS by letters (measured at 1.2 and 2.4 cpd; r = 0.235 (P < 0.05) and 0.272 (P < 0.05), respectively).

The questionnaire score was not related in a statistically significant way to any of the following: CDVA, photopic CS measured by letters (at any spatial frequency other than 1.2 and 2.4 cpd), mesopic or photopic CS measured by gratings (at any spatial frequency), mesopic GD (at any spatial frequency other than 3 and 6 cpd), photopic GD (at any spatial frequency other than 1.5 and 3 cpd), mean retinal sensitivity, reading acuity, and reading speed or LOD (P > 0.05 for all).

Questionnaire score was not related in a statistically significant way to LOCS III nuclear opalescence/color score subgroup (F(2,82) = 0.63, P > 0.05), LOCS III cortical score subgroup (F(3,82) = 0.62, P > 0.05), or LOCS III subcapsular score subgroups (F(2,80) = 0.82, P > 0.05; see Table 4).

Stereoacuity subgroups did not differ in a statistically significant way in terms of questionnaire score (F(5,78) = 6.30, P > 0.05; see Table 5).

Discussion

We carried out a study to evaluate a number of methods for assessing symptomatic non-advanced cataract and its visual consequences. The methods assessed included CDVA, photopic and mesopic CS (tested by letters (photopic only) and by gratings), photopic and mesopic GD (tested by gratings), reading performance (reading acuity and reading speed), stereoacuity, retinal sensitivity, LOD, and LOCS III. The purpose of this study was to investigate whether psychophysical, morphological, and/or optical measures of cataract are complementary to, or perhaps more appropriate than, CDVA for the purposes of documenting visual dissatisfaction attributable to cataract that exhibits relative sparing of high contrast acuity.

Table	3	Demogr	aphic,	ocular,	refrac	tive,	psychoph	ysical,
catarac	t o	ptical, ca	itaract 1	norphol	logical	and s	ubjective	visual
difficul	lty	data of	patient	s with	sympt	omatic	non-adv	vanced
catarac	t ar	nd no oth	ier ocula	ar patho	ology			

Table 3 (Continued)

cataract optical, catar difficulty data of pa	act morpholo tients with	ogical a sympto	and su matic	ibjective visual non-advanced	Variable	n (%)	Mean	± SD	Range
cataract and no other	ocular pathol	ogy			logGD				
Variable	n (%)	Mean	± SD	Range	(photopic conditions)				
					Frequency (cpd)				
Demographic					1.5	75 (91%)	1.34	4 0.20	0.85 to 2.00
Age (years)	82 (100%)	66.9	8.7	47 to 85	3	72 (88%)	1.52	7 0.20	1.00 to 2.06
0,0					6	47 (57%)	1.42	7 0.22	1.08 to 1.95
Condon					12	23 (28%)	1.1^{4}	4 0.14	0.90 to 1.34
Genaer	2 0 (24%)				18	13 (16%)	1.0	1 0.21	0.60 to 1.36
Male	29 (34%)				LogRAD	80 (98%)	0.33	3 0.21	0.1 to 1.1
Female	57 (66%)	_			LogRAD score	80 (98%)	0.34	4 0.21	0.1 to 1.1
Ocular					Reading speed (wpm)				
Laterality					Letter size				
Right eye	39 (45%)			_	12	80 (98%)	150	31	51 to 227
Left eve	47 (55%)			_	1.2	80 (08%)	144	21	51 to 227
	(111)				1.1	30(98%)	144	27	J1 to 221
P () (0.9	77 (94%)	149	3/	46 to 227
Kefractive					0.8	77 (94%)	149	36	16 to 221
SE study eye (D)	82 (100%)	+0.16	2.9	-16.5 to $+6.5$	0.7	77 (94%)	143	37	53 to 233
					0.6	72 (88%)	138	39	52 to 227
Psychophysical					0.5	71 (87%)	136	49	44 to 322
CDVA atudu ava	92(1000)	0.10	0.16	0.62 + 0.12	0.4	69 (84%)	129	42	47 to 210
CDVA study eye	82 (100%)	0.18	0.10	0.62 to -0.12	0.3	56 (68%)	115	38	58 to 200
CDVA fellow eye	82 (100%)	0.14	0.12	0.64 to -0.14	0.2	35 (43%)	100	39	43 to 183
LogCS by letters					Stereoacuity (s of arc)				
(photopic)					0	20 (23.8%))	_	_
Frequency (cpd)					480	5 (6%)			
1.2	79 (96%)	1.38	0.19	0.90 to 1.66	240	14(1670)	. —		
2.4	79 (96%)	1.31	0.25	0.45 to 1.66	120	14(10.7%)			
6	78 (95%)	1.00	0.31	0.15 to 1.49	120	30 (35.7%)			_
96	72 (88%)	0.78	0.20	0.15 to 1.20	60	13 (15.5%)) —		
9.0 15.16	72(0070)	0.70	0.29	0.15 to 1.20	30	2 (2.4%)			—
24	13 (16%)	0.34	0.27	0.15 to 0.60	15	0 (0%)	_	—	—
					Mean retinal sensitivity (dI	3)			
LogCS by gratings					Fixation	80 (98%)	14	5	0 to 20
(mesonic conditions)					Control 5 dogroos	80 (98%)	16	4	0 to 20
Frequency (cnd)					Central 5 degrees	80 (98%)	10	4	0 10 20
15	80 (0897)	1 44	0.22	0.85 ± 0.200	Between 5 and	80 (98%)	16	4	0 to 20
1.5	$\frac{30}{90}$	1.44	0.22	0.85 10 2.00	10 degrees				
3	77 (94%)	1.51	0.24	0.54 to 1.90	Central 10 degrees	80 (98%)	16	4	0 to 20
6	40 (49%)	1.31	0.20	1.08 to 1.65					
12	5 (6%)	0.99	0.12	0.90 to 1.18	Cataract optical $(n = 86)$				
18	2 (2%)	0.69	0.12	0.60 to 0.78	Average LOD	80 (98%)	12.4	1.7	9.8 to 18.2
					Maximum LOD	80 (98%)	39.8	10.6	16.9 to 72.5
LogCS by gratings					Minimum LOD	80 (98%)	5.5	1.1	2.4 to 12.7
Frequency (and)					Catavard 1 1 1 1				
1 E	PO (0907)	1.25	0.0.20	0.005 to 1.05	Cataract morphological (n	$l = \delta b$	a =		
1.5	80 (98%)	1.35	0.0.20	0.85 to 1.85	LOCS III nuclear scor	e 80 (98%)	2.5	0.7	1 to 4
3	80 (98%)	1.57	0.22	1.00 to 2.06	LOCS III cortical score	e 80 (98%)	2.1	1.0	1 to 4
6	57 (70%)	1.46	0.21	1.08 to 2.11	LOCS III subcapsular	80 (98%)	1.4	0.8	1 to 5
12	26 (32%)	1.14	0.17	0.90 to 1.48	score				
18	20 (24%)	0.93	0.21	0.60 to 1.23					
LogGD (mesopic conditions)					Subjective visual difficulty Rasch-scaled Priquest score	y (questionna 82 (100%)	aire) 2.15	5 0.36	(1.37 to 3.13)
Frequency (cpd)					Abbreviations: CDVA cor	rected-distand	e visu	al acuity.	cpd. cycles per
1.5	64 (78%)	1.31	0.23	0.85 to 2.00	degree: CS contrast sons	itivity: D di	iopter [.]	dB. deci	ibels: GD. olare
3	61 (74%)	1.45	0.22	1.00 to 1.90	disability LOCS long one	cities classific	ation of	vstem I	OD lens optical
6	23 (28%)	1.36	0.25	1.08 to 1.81	density: logRAD log	reading acu	itv. C	D etan	lard deviation
12	4 (5%)	1.04	0.16	0.90 to 1.18	SE spherical equivalent	UDVA 11	naided	-distance	visual acuity
18	3 (4%)	0 72	0.10	0.60 to 0.78	wind words per minute	, <i>CD 111,</i> U	macu	anstance	.iouur acuity,
10	5 (470)	0.72	0.10	0.00 10 0.70	wpu, worus per minute.				

LOCS III classification	Questionnaire score		F	P-value	
	Mean	± SD			
LOCS III nuclear opales	cence/color	score			
2	0.56	0.27			
3	0.51	0.29			
4	0.62	0.21	(2,82) = 0.63	>0.05	
LOCS III cortical score					
1	0.55	0.31			
2	0.53	0.26			
3	0.52	0.27			
4	0.68	0.24	(3,82) = 0.62	> 0.05	
LOCS III subcapsular s	core				
1	0.56	0.27			
2	0.62	0.27			
3	0.44	0.33	(2,80) = 0.82	> 0.05	

Table 4	Variation in subjective visual difficulty (questionnaire
score) in	different LOCS III classification subgroups

Abbreviations: *F*, degrees of freedom; LOCS III, Lens Opacities Classification System III; SD, standard deviation.

 Table 5
 Variation in subjective visual difficulty (questionnaire score) in different stereoacuity subgroups

Stereoacuity (s of arc) ^a	Question	aire score	F	P-value	
	Mean	$\pm SD$			
0	0.65	0.29			
480	0.57	0.36			
240	0.55	0.18			
120	0.48	0.28			
60	0.48	0.29	(5,78) = 6.30	> 0.05	

Abbreviations: *F*, degrees of freedom; SD, standard deviation.

^aThe stereoacuity of 30 s of arc subgroup contained only two patients and was therefore excluded from the above analysis due to its small size.

There was no statistically significant correlation between CDVA of the study eye and subjective visual difficulty in our study. This finding is consistent with the authors' suspicion that, in the context of symptomatic non-advanced cataract, high contrast VA, in isolation, is not an appropriate psychophysical test to evaluate the visual impact of the disease process, and the importance of this finding rests on the fact that CDVA is still the most commonly used psychophysical test in the assessment of cataract and its visual sequelae.

However, there was a statistically significant negative correlation between subjective visual difficulty (as measured by the Rasch-scaled questionnaire score) and mesopic GD (measured at 3 and 6 cpd) and photopic GD (measured at 3 and 6 cpd). This is an important clinical finding because it suggests that, for cataracts with little or only marginal impact on high contrast VA, mesopic and photopic GD measured at medium spatial frequencies is a more valid psychophysical tool than CDVA, and one that better reflects subjective visual difficulty attributable to the lens opacity.

The data also suggest that a value of mesopic GD at 3 cpd of < 29 (corresponding to inability to correctly identify the fourth gratings patch presented during the test), a value of mesopic GD at 6 cpd of <23(corresponding to inability to correctly identify the third gratings patch presented during the test), a value of photopic GD at 3 cpd of <40 (corresponding to inability to correctly identify the fifth gratings patch presented during the test), and a value of photopic GD at 6 cpd of <33 (corresponding to inability to correctly identify the fourth gratings patch presented during the test) are good indicators of subjective visual difficulty caused by cataract that is relatively sparing of high contrast acuity. In other words, if a patient with lenticular opacity and good high contrast CDVA complains of visual difficulty in the absence of ocular co-morbidity, and yet cannot correctly identify the presented gratings patches during GD testing, at the aforementioned levels of CS, then the lenticular opacity is indeed likely to be the cause of his/ her subjective visual difficulty. Of note, it is perhaps unsurprising that symptomatic non-advanced cataracts are best judged by a test which represents a function of light scatter, reduced retinal illumination and blur effect of cataract, such as mesopic and photopic GD at medium spatial frequencies. It is well known that developing cataracts cause increased intraocular forward light scatter, which causes visual difficulty due to loss of retinal image contrast, and this type of visual difficulty is called GD.^{19,31} It has also been previously reported that the amount of scatter can be evaluated indirectly by measuring GD,³² that GD offers a discriminative and valid measure of visual assessment in cataract,³³ including cataract with good high contrast VA in the presence of functional complaints,^{34,35} that GD improves after cataract surgery,³⁶ and that this improvement is independent of improvement in high contrast acuity.37 We suggest that testing mesopic and photopic GD at a medium spatial frequencies will help assess symptomatic non-advanced cataracts. The clinical test we suggest is both user- and clinic-friendly requires no more patient cooperation than a standard VA test, and consumes a considerably shorter amount of time ($\sim 3 \min$).

Interestingly, there was no statistically significant correlation between questionnaire score and average LOD. This finding suggests that LOD is unlikely to represent an objective surrogate or alternative to psychophysical testing in an eye with cataract that has little impact on high contrast VA.

Also, there were no statistically significant differences in terms of questionnaire score between different LOCS III nuclear opalescence/color score subgroups, cortical score subgroups or subcapsular score subgroups, indicating that this grading system of cataract morphology (LOCS III) is unlikely to represent a valid means of documenting the lens opacity in a way that reflects the patients' symptoms attributable to the cataract, in cases where there is relative sparing of high contrast acuity.

We report a lack of correlation between subjective visual difficulty and retinal sensitivity, indicating that this test is also unlikely to represent a useful means of recording visual data that reflect subjective visual difficulty in patients with symptomatic non-advanced cataract.

Stereoacuity, in our study, was not found to be indicative of overall subjective visual difficulty attributable to the lens opacity. However, it should be borne in mind that the TNO stereotest, used here, requires excellent fusion and is also known to be adversely affected by increasing age in normal individuals who report no difficulty in judging distances in everyday tasks.³⁸ In other words, our findings may indicate that stereoacuity is a psychophysical test which is indeed affected by symptomatic non-advanced cataract (reflected in the observation that the vast majority of patients (over 82%) had stereoacuity of 120s of arc or less, and over 23% of patients had unmeasurable levels of stereoacuity), but may not be a good reflection of subjective visual difficulty in the presence of good CDVA in the fellow eye.

Conclusion

In conclusion, this study demonstrates that symptomatic non-advanced cataract, in the presence of good high contrast CDVA and in the absence of coexisting ocular pathology, is associated with subjective functional visual difficulty (measurable by a validated questionnaire), which is best reflected in a decrease in mesopic and photopic GD measured at medium spatial frequencies. High contrast VA does not reflect the patient's visual dissatisfaction in such cases. Ophthalmologists should consider testing mesopic and/or photopic GD at a medium spatial frequency to quantify the impact of symptomatic non-advanced cataract in those patients who report functional visual difficulty, not reflected in their high contrast VA test result. This clinic- and userfriendly test requires no more patient cooperation than a standard high contrast VA test, and takes <3 min to conduct. The collection of such preoperative data will represent a valid record of adversely affected vision where standard high contrast VA may be essentially normal or near normal, and will inform professional practice by assisting ophthalmologists and their patients

in deciding the best time for cataract surgery, an issue which is becoming increasingly important in an era of falling thresholds for this procedure.

Summary

What was known before

 It has been suggested that visual acuity is not the most appropriate tool to investigate the impact of symptomatic non-advanced cataract on visual function and experience, and complimentary and/or alternative techniques that have been investigated include contrast sensitivity, glare disability, reading performance, and functional visual difficulty assessed by various questionnaires.

What this study adds

• Our research shows that symptomatic non-advanced cataracts are associated with subjective visual difficulty (measurable by a validated questionnaire), best reflected in a decrease in mesopic and photopic contrast sensitivity under conditions of glare, at medium spatial frequencies.

Conflict of interest

The authors declare no conflict of interest.

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