

Intraocular lens power calculation in short eyes

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Abstract

Purpose To compare the accuracy of the Hoffer Q and SRK-T formulae in eyes below 22 mm in axial length, using biometry measured with partial coherence interferometry (PCI), without a customised ACD constant.

Methods Data were retrospectively and prospectively collected by identifying eyes of axial length below 22 mm in the records of the intraocular lens (IOL) master and in preoperative notes. Biometry was performed using PCI and IOL power was calculated using both SRK-T and Hoffer Q formulae. Refractive outcome was measured and the accuracy of the two formulae compared.

Results Forty-one eyes of 41 patients were identified with an axial length <22 mm. Axial lengths ranged from 21.96 to 20.29 with a mean of 21.51 mm, and IOL power ranging from 23 dioptres (D) to 29 D. The Hoffer Q formula showed a mean prediction error of 0.61 D (SD 0.80) compared with the SRK-T, which showed a mean prediction error of 0.87 D (SD 0.829). A paired *t*-test found that the Hoffer Q was significantly more accurate than the SRK-T formula ($P < 0.001$).

Conclusions Hoffer Q was found to be more accurate than the SRK-T formula in this series of eyes <22 mm axial length when customised ACD constants are not used. Royal College of Ophthalmologists guidelines may need to be adjusted in accordance with these findings. This study underlines the importance of monitoring outcomes, and suggests different customisations are needed for different formulae, with a higher correction if the SRK-T formula is used for short eyes.

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Introduction

Intraocular lens (IOL) power calculation accuracy has been of great importance in cataract surgery for many years but with an increasing demand for low postoperative refractive error and increasingly high patient expectations, the need for accurate power calculation is paramount. In the past two decades, there have been advances in both surgical technique and in the accuracy of biometric measurements. The routine use of phacoemulsification and sutureless incisions combined with continuous curvilinear capsulorhexis has improved the predictability of refractive outcome.¹ In addition, optical biometry with partial coherence interferometry (PCI) has been shown to yield significantly better IOL power prediction, and therefore refractive outcome in cataract surgery, than ultrasound biometry.² Improvements in measurement and surgical technique have raised a debate about the best IOL calculation formula to obtain the most precise prediction of refractive outcome.

The 2001 Royal College of Ophthalmologists (RCOphth) Cataract Surgery Guidelines issued recommendations on biometry and IOL calculation according to the axial length of the eye,³ shown in Table 1. These guidelines were based on Hoffer's recommendations,⁴ and did not discuss A constant customisation, which had been recommended by Hoffer. The National Biometry audit, published in 2004, found poor adherence to these guidelines, with only 36% of departments using two or more formulae, and only 4% using the Hoffer Q formula in eyes with axial lengths below 22 mm.⁵ The current 2004 RCOphth Guidelines advise use of Hoffer-Q⁴ or SRK-T⁶ formulae for eyes <22 mm length, as well as use and checking of the appropriate ACD constants.⁷

Hoffer's original series⁴ included 36 eyes below 22 mm in axial length, and suggested the Hoffer Q may be more accurate than the SRK-T (not statistically significant, $P = 0.83$, probably

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due to the relatively small size of the series). Hoffer also investigated the accuracy of the Hoffer Q against the SRK-T again in 2000⁸ (in addition to the Holladay 1 and Holladay 2 formulae) and again the Hoffer Q appeared more accurate than the SRK-T, but with only 10 eyes in the series below 22 mm in axial length, no statistical significance was tested.

The purpose of this study was to compare the accuracy of the Hoffer Q and SRK-T formulae in eyes below 22 mm in axial length, using biometry measured with PCI, without a customised ACD constant, to ascertain whether one formula was more accurate than the other.

Materials and methods

Data were collected retrospectively and prospectively by identifying eyes of patients undergoing routine cataract surgery with an axial length below 22 mm from the biometry database and in preoperative notes. Biometry was performed using PCI (IOLMaster, Zeiss Humphrey Systems) and IOL power was calculated using the SRK-T formula and the manufacturer’s A constant. In addition, the Hoffer Q formula was calculated with the same biometric data for each eye.

Patients underwent routine clear-corneal phacoemulsification and Allergan MA60 foldable IOL insertion into the capsular bag. Following uneventful surgery, patients were reviewed in the eye clinic 2–3 weeks

following surgery. Objective measurement of the refractive status of the eye was obtained with the autorefractor (Canon AR-2, Canon, Tokyo) and this was converted into a spherical equivalent value, which was taken as the refractive outcome. The predictive errors were calculated for both formulas—the predictive error was defined as the difference in the predicted value from the biometry for each formula and the spherical equivalent value of the postoperative objective refraction. The accuracy of the two methods was compared using a paired *t*-test.

Results

Patients undergoing routine cataract surgery at our hospital between April 2004 and March 2005 were identified. Forty-six patients undergoing cataract surgery with axial length <22 mm were identified and of these five were excluded: the notes/biometry/postoperative autorefractation were incomplete for three patients, one patient’s biometry related to a lens exchange, and one patient developed postoperative endophthalmitis. In total, 41 eyes of 41 patients were included. Axial lengths ranged from 20.29 to 21.96 mm, with a mean of 21.51 mm, and the IOL power inserted ranged from 23 dioptres (D) to 29 D (mean 25.73 D, median 26 D). The planned refraction aimed for was determined by individual surgeons, based on the preoperative biometry (using the SRK-T formula and manufacturer’s A constant).

Comparison of actual outcomes with predicted outcomes using PCI biometry and IOL calculation formulae, without any correction/A constant customisation, resulted in a more hypermetropic outcome than predicted for both formulae. Figure 1 shows the distribution of the predictive errors in this series using the SRK-T formula. The Hoffer Q formula’s mean prediction error of 0.61 D was lower than the mean prediction error of the SRK-T formula (0.87 D), shown in Table 2. Table 2 also shows the ranges for the two formulae, with the Hoffer Q having a higher proportion of patients achieving +/–1 D of the predicted outcome than the SRK-T (63 vs 54%). A paired *t*-test found that the Hoffer Q was significantly more accurate than the SRK-T formula (*P* < 0.001).

If a customised correction factor for each formula was applied (+0.61 D for Hoffer Q and +0.87 D for SRK-T added to the manufacturer’s A constant), then the two formulae resulted in similar outcomes, with 80% of subjects within 1 D of the predicted value for the SRK-T and 83% for the Hoffer Q, as shown in Table 3.

Discussion

To our knowledge this series is the first to show the Hoffer Q formula is statistically significantly more

Table 1 Royal College guidelines on formula usage according to axial length interval

Axial length (mm)	RCOphth 2001 Guideline	RCOphth 2004 Guideline
<22	Hoffer Q	Hoffer Q or SRK-T
22.0–24.5	Average Hoffer Q Holladay and SRK-T	Hoffer Q, Haigis or SRK-T
24.6–26.0	Holladay	SRK-T
>26.0	SRK-T	SRK-T

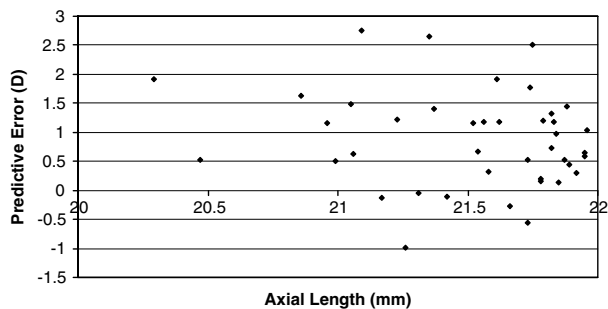


Figure 1 Predictive error (in dioptric spherical equivalent) in a series of 41 short eyes (<22 mm axial length), using biometry obtained by PCI and the SRK-T formula.

Table 2 Predictive errors and predictive ranges for the SRK-T and Hoffer Q formulae

	SRK-T	Hoffer Q
Mean (SD) error	0.87 (0.829)	0.61 (0.80)
Mean absolute (SD) error	0.98 (0.701)	0.78 (0.654)
±0.25 D of predicted value	17%	27%
±0.5 D of predicted value	24%	44%
±1.0 D of predicted value	54%	63%

Table 3 Predictive ranges for the SRK-T and Hoffer Q formulae with corrective factor adjustment.

Formula and 'Correction Factor'	SRK-T + 0.87 D	Hoffer Q + 0.61 D
±0.25 D of predicted value	12%	15%
±0.5 D of predicted value	49%	44%
±1 D of predicted value	80%	83%

accurate than the SRK-T, and is also the largest series investigating biometry accuracy in eyes below 22 mm in axial length. Our results also indicate that the Hoffer Q is more accurate than the SRK-T when customised ACD constants are not used, which was the case for 53% of departments in the UK in the second National Biometry Audit.⁹ The SRK-T formula was developed to increase accuracy in high myopia⁶ and the Hoffer Q formula for use in extreme hyperopia,⁴ and this study confirms the previous impression that the Hoffer Q is more accurate than the SRK-T for this group of patients. Although current Royal College guidelines allow the SRK-T to be used for all axial lengths, these results highlight the need to audit the refractive outcomes of myopes, hypermetropes, and emmetropes separately.

Use of the SRK-T formula tended to result in a more hypermetropic refraction than the Hoffer Q. However, if a different correction factor was applied (0.61 D for the Hoffer Q and 0.87 for the SRK-T), then the results of the two formulae are comparable (Table 3). These results emphasise the importance of individual surgeons and departments to closely audit their results, and to adopt customised A constants depending on the formula used and previous outcomes. Based on our results, a greater correction than normal is required in hyperopic eyes to achieve emmetropia, particularly if the SRK-T formula is used.

Any measurement error in the axial length of a short eye will have a larger effect on the final refraction, as a given measurement error is a much larger percentage of the axial length in a short eye and hence this could be an important source of error. This is of particular importance in applanation ultrasound biometry, as this

method is known to have the disadvantage of potentially depressing the cornea during use leading to a falsely short axial length measurement. The use of the PCI in this study should have minimised biometry errors as this method has a higher resolution than standard ultrasound transducers,^{10,11} is highly reproducible¹² and avoids any contact with the eye.

The prospective study of 1817 eyes of all axial lengths by Murphy *et al*¹³ achieved a final refraction within 1 D of the biometry-predicted value in 72.3% of subjects, and may be considered a benchmark for current cataract practice and a useful audit standard. However, Murphy *et al* did not state the proportion of eyes below 22 mm or the biometry prediction error for this subgroup, and therefore comparison between the current study and their data is not valid.

A potential source of error in this study was the use of the autorefractor as the method of measuring the spherical equivalent for postoperative outcome. Other studies have found there is no significant difference in the accuracy of the autorefractor and subjective refraction in assessment of the final achieved refractive outcome for LASIK¹⁴ and cataract surgery,¹⁵ indicating that the autorefractor is an acceptable outcome measure of final refractive status. The advantage of use of the autorefractor measure to be able to include all patients with axial length <22 mm outweighed the disadvantage of trying to rely on later (possibly more accurate) refraction from optometrists, with the likelihood of incomplete data collection and possible ascertainment bias.

In conclusion, the Hoffer Q was found to be more accurate than the SRK-T formula with statistical significance in this series of eyes <22 mm axial length. To our knowledge, this has not been reported previously, and this is the largest series specifically reporting the refractive outcomes of short eyes in cataract surgery. RCOphth Guidelines may need to be adjusted in accordance with these findings; failing this, surgeons may need to factor a different (higher) customisation constant for eyes of small axial length, depending on the formula used, and to audit their own results in this group of patients.

Acknowledgements

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