Correlation between central corneal thickness, applanation tonometry, and direct intracameral IOP readings

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Abstract

Background—Several authors reported incorrect high intraocular pressure (IOP) values in eyes with a thick cornea using applanation tonometry. This hypothesis was checked by comparing applanation tonometry with direct intracameral manometry.

Methods—73 patients, scheduled for intraocular surgery, were enrolled. Immediately before surgery, the following were registered: (i) central corneal thickness (CCT), (ii) applanatory IOP (Perkins/Tonopen), and (iii) intracameral IOP.

Results—The difference between applanatory and intracamerally measured IOP was completely independent of CCT ($y = -3.43 + 3.8x$; where $y$ is the difference between applanatory and intracamerally measured IOP (mm Hg) and $x$ is CCT (mm); $r^2 = 0.002$; $p = 0.72$).

Conclusions—There is no systematic error of applanation tonometry with increasing CCT. Therefore it is inadequate to recalculate IOP based on regression formula of applanatory IOP versus CCT.

When Goldmann and Schmidt presented their applanation tonometer in 1957, they stressed certain possible sources of error. Central corneal thickness (CCT) was one of them. Later on some groups reported thicker corneas in patients with higher intraocular pressure (IOP) compared with thinner corneas in patients with lower IOP.

Varying corneal rigidity was claimed to be responsible for false applanatory IOP readings with different CCT. Some authors recommended a recalculation of the IOP depending on the CCT with a correction factor ranging from 1 to 6.8 mm Hg per 0.1 mm CCT. The intention of this study was to reevaluate this assumption by correlating the CCT with applanation tonometry and intraocular IOP readings.

Patients and methods

A total of 73 patients (73 eyes) were prospectively enrolled. All patients gave their consent before their inclusion in the study. They were scheduled for intraocular surgery for glaucoma ($n = 31$) or retinal diseases ($n = 42$).

The mean age was 40.7 years, ranging from 13 to 88 years. Patients with more than 1.5 D astigmatism or corneal abnormalities were excluded.

After retrobulbar or general anaesthesia measurements were performed in the theatre with the patients supine. While one investigator (NF) measured CCT and applanatory IOP, a second investigator, the surgeon, performed intracameral IOP measurements. Four surgeons participated in this study.

An eyelid retractor was placed and the CCT was determined with an ultrasonic pachymeter (Pachette, DGH-Technology Inc). The IOP was measured first with the Perkins tonometer followed by the Tonopen (Mentor). The eyelid retractor was removed and the patient was prepared for surgery.

Every operation started by placing the eyelid retractor. Then the intracameral measurement was performed immediately. We used a special device developed by the University of Düsseldorf. It consists of an invasive blood pressure monitor as used in intensive care (Cardiocap II; Datatex Engstrom) connected to a specially designed transducer (Monitoring Kit; Abbott) and a steel cannula. This cannula was placed in the anterior chamber for about 10 seconds until the readings on the monitor were stable (Fig 1). Thereafter, the cannula was removed and surgical treatment begun. The agreement of methods was validated calculating the mean difference (md) and the standard deviation of the differences (SD) between applanatory and intracamerally measured IOP.

Results

The CCT values ranged from 0.448 to 0.713 mm (mean 0.58 (SD 0.054) mm).

The applanatory IOP readings varied from 8 to 32 mm Hg (mean 17.5 (6.5) mm Hg) using the Perkins tonometer, and from 7 to 38 mm Hg (mean 18.7 (7.2) mm Hg) using the Tonopen.
There was a strong correlation between these two methods ($r^2=0.866$; $p<0.0001$). The mean of the two methods was calculated and used as applanatory IOP (IOPappl) for further statistics.

Intraocular IOP readings varied from 8 to 37 mm Hg (mean 19.5 (6.5) mm Hg). Comparing IOPappl and intracameral readings, the regression formula is $y=2.91+0.78x$ (where $y$ is IOPappl (mm Hg) and $x$ is the IOP measured by cannula (mm Hg); $r^2=0.56$; $p<0.0001$). In four patients the difference varied by more than 10 mm Hg.

Calculating the methodical agreement between applanation and intracameral measurement, the mean difference was 1.2 mm Hg and the standard deviation of the differences was 4.6 mm Hg. To determine the correlation, we calculated the mean difference and the standard deviation of the differences between IOPappl and intracameral IOP. Accordingly, 95% of these differences ranged within plus or minus 4.6 mm Hg of the mean. This indicates a moderate agreement between the methods used. There is no doubt that applanation tonometry is the clinical gold standard; however, we believe intracameral measured IOP values reflect the “true” IOP more accurately.

Goldmann was aware that corneal rigidity has to be considered if IOP is measured by applanation. Although he claimed CCT altered the measurement, he expected no clinically relevant misreadings. However, several authors described a positive correlation and recommended a recalculation of IOP depending on CCT.

Intraocular readings are independent of CCT. If applanation tonometry gives artificially high values in patients with thick corneas and artificially low values in patients with thin corneas, intraocular and applanatory readings should diverge as CCT rises.

In our study, the difference between intracameral and applanation tonometry did not increase with increasing CCT. Based on these findings it is concluded that applanatory readings are presumably not methodically influenced by CCT.

As the 95% confidence bands show (Fig 2), a minimal positive as well as a minimal negative correlation is unlikely. But it cannot be excluded completely. The statistical uncertainty, however, is of no relevance in clinical practice.

As can be seen from the same figure IOP values of thicker as well as of thinner corneas fluctuate widely above and below the regression line. Obviously, applanatory IOP readings can be too high as well as too low, independent of corneal thickness. Therefore, one cannot predict whether the “true” IOP value is higher or lower than a given applanatory IOP reading.

From this result we conclude that the use of any global recalculation formula is unsuitable to find the true IOP in clinical practice.

Furthermore, there is no need to combine the measurement of applanatory IOP and pachymetry. In cases of doubt an intracameral IOP measurement is recommended.

Proprietary interest: Nil.


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