

Comparative Study of Intraocular Pressure Measurements with Airpuff, iCare and Goldmann Applanation Tonometers

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ABSTRACT

Purpose: To compare the difference in intraocular pressure (IOP) measurements by Airpuff, iCare and Goldmann Applanation Tonometers (GAT).

Study Design: Comparative analytical study.

Place and Duration of Study: Department of Ophthalmology, Mohi-ud-Din Teaching Hospital, Mirpur Azad Kashmir, from June 2020 to August 2020.

Methods: Twenty-five patients (50 eyes) were included in this study. IOP was measured in each eye firstly by Airpuff tonometry, then by iCare tonometry and lastly by Goldmann applanation tonometer. Three consecutive readings were taken in each eye. If there was a difference of 2 mm Hg or more among the readings, measurement was repeated. Once we got three readings, their average was taken and analyzed. Comparison of IOP readings between these tonometers was done.

Results: Mean IOP was 15.84 ± 2.736 mm Hg with Airpuff Tonometer, 14.48 ± 2.435 mm Hg with iCare Tonopen and 14.74 ± 2.489 mm Hg with Goldman Applanation Tonometer. The difference between the mean Airpuff and Goldman Applanation Tonometer readings was 1.10 mm Hg which was not statistically significant (p-value = 0.083). The difference between the mean Goldman Applanation Tonometer and iCare Tonopen readings was 0.26 mm Hg which is also not statistically significant (p-value = 0.867). But, the difference between the mean iCare Tonopen and Airpuff Tonometer readings was -1.36 mm Hg which was statistically significant (p-value = 0.02).

Conclusion: It is concluded that IOP readings taken by iCare Tonopen and Airpuff Tonometer are comparable to those taken by Goldman Applanation Tonometer and iCare Tonopen underestimates the IOP when compared with Airpuff Tonometer.

Key Words: Airpuff Tonometer, Glaucoma, Goldmann Applanation Tonometer, iCare Tonometer, Intraocular Pressure.

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INTRODUCTION

Intraocular pressure (IOP) is the aqueous pressure determined by the equilibrium between the amount of aqueous humor produced, by the ciliary body, and drained through the trabecular meshwork.¹ Tonometry is the method by which IOP is measured and its

accurate measurement is necessary in evaluation of patients at risk of glaucoma.² An accurate IOP measurement is mandatory component of ophthalmological examination in clinical practice.

The range of IOP in the general population is about 11–21 mm Hg.³ In some cases, glaucomatous changes are noticed even with IOP values less than 21 mm Hg i.e. Normal tension glaucoma (NTG).⁴ While in other cases with IOP more than 21 mm Hg, no glaucomatous changes are detected i.e. Ocular hypertension.⁵ There are some factors, which influence IOP value, one of most important such factor is central corneal thickness (CCT).⁶

Glaucoma can be classified in different ways.⁷ There are many methods to measure IOP. In Airpuff tonometry, the central corneal surface is flattened by a jet of air and time needed to do so is directly proportional to IOP. As it is done without topical anesthesia and there is no contact made with the eye, it is ideal for community screening. For more accurate readings, at least three readings are taken to get an average.⁸ iCare tonopen is based on rebound tonometry in which there is a thin wire with attached 1.8 mm plastic ball, when probe decelerates upon corneal contact, the magnitude of deceleration is directly proportional to IOP.⁹ No anesthesia is needed in this procedure as well, thus it is also helpful for community screening. While in Goldmann applanation tonometry (GAT), Imbert–Fick principle is applied according to which the pressure (P) inside a dry thin-walled sphere is equal to the force (F) needed to flatten its surface divided by the area (A) which is flattened (i.e. $P = F/A$). This requires topical anesthesia and there is corneal contact made. For this reason disinfection between patients is needed thus it is not suitable for mass screening.¹⁰ Disposable tonometer prisms and caps can also be used to address the need of repeated disinfection between patients.

The objective of this study was to compare the IOP measurements by Airpuff tonometer, iCare tonometer and Goldmann Applanation Tonometer (GAT) and to evaluate the accuracy and reliability of the three IOP measurement methods.

METHODS

This study was carried out at the Department of Ophthalmology, Mohi-ud-Din Teaching Hospital, Mirpur Azad Kashmir. Study period was from 1st June 2020 to 31st August 2020. Ethical approval was

obtained from the Ethical Committee of the hospital. Informed written consent was obtained from all patients and detailed counseling was done about the procedure and results.

Twenty-five patients (50 eyes) were included in this study out of which 15 were males and 10 were females. Mean age of the participants was 30.36 ± 5.376 years. The sample size was calculated using WHO sample size calculator on the basis of recent study.¹¹ All patients were enrolled from ophthalmology out-patient department (OPD), Mohi-ud-Din teaching hospital, the affiliated hospital of Mohi-ud-Din Islamic Medical College, Mirpur Azad Kashmir. Any patient with central corneal opacity, corneal astigmatism, nystagmus, ocular surface disease i.e. dry eyes and conjunctivitis, ocular trauma was excluded from the study. History of contact lens use and refractive or intraocular surgery was also taken into consideration while setting exclusion criteria.

All enrolled patients were examined using Slit lamp. IOP measurements were taken using Airpuff – iCare – GAT sequence while in the sitting position. All tonometers were properly calibrated before taking readings and there was a gap of 10 minutes maintained between each IOP measurement to reduce any after measurement fluctuation.

Tonometry was first performed in each patient with an Airpuff tonometer (Nidek RKT 7700, Nidek Corporation, Japan). Three consecutive readings were taken in each eye. If there was difference of 2 mm Hg or more among the readings, measurement was repeated. Once we got three readings, their average was taken and analyzed. After ten minutes, IOP measurement was done with iCare pro (iCare Finland Oy, Helsinki, Finland). It has a single use/disposable probe, loaded into the instrument and was aligned 4–8 mm vertical to the central corneal surface. After six measurements were taken, the highest and lowest values were automatically discarded and the average IOP was calculated from the remaining four values by the built-in software. At the end Goldmann Applanation Tonometry was performed using GAT AT900, Haag Streit, Koniz, Switzerland installed on a slit-lamp biomicroscope. Three IOP readings were taken after instillation of a drop of 0.5% proparacaine hydrochloride (Alcaine) and 0.25% fluorescein in each eye. The final IOP was taken from the average value of these three measurements provided that there was no difference of 2 mm Hg or more among these three values. All the measurements were taken by same

ophthalmologist. Measurements by Airpuff Tonometry, iCare Tonometer, Goldmann Applanation Tonometer (GAT) were analyzed and compared with each other.

Data was entered and analyzed using Statistical Package for Social Sciences (SPSS) version 21. Mean and standard deviation were calculated for quantitative variable i.e. age. Qualitative variable like gender and eye involved were calculated by frequency and percentage. One way Anova and Tukey test were used to see the difference in intraocular pressure readings in all measurement methods. p-value of < 0.05 was considered significant.

RESULTS

A total of 25 patients (50 eyes) were included in this study. Mean age of the patients was 30.36 ± 5.376 years (21 – 39). There were 15 male patients and 10 female patients in the study (Figure 1). Out of 50 eyes, 25 were right eyes and 25 were left eyes. Overall the IOP measured by all instruments was 15.02 ± 2.607 mm of Hg.

The IOP averages measured with all instruments are shown in Table 1. Mean IOP reading with Airpuff tonometer 15.84 ± 2.736 mm Hg, with iCare Tonopen it 14.48 ± 2.435 mm Hg and with Goldman applanation tonometer was 14.74 ± 2.489 mm Hg.

Table 1: Average IOP in each method with One Way Anova Test (p value = 0.021).

Method	n	IOP (mm of Hg)	p Value
Airpuff Tonometer	50	15.84 ± 2.736 (10 – 21)	0.021
Goldmann Applanation Tonometer	50	14.74 ± 2.489 (10 – 20)	
iCare Tonopen	50	14.48 ± 2.435 (9 – 19)	
Total	150	15.02 ± 2.607 (9 – 21)	

Table 2: Differences in intraocular pressure (IOP) measurement methods using Tukey Test.

Variable	Mean Difference (mm of Hg)	p Value
Airpuff Tonometer – Goldman Applanation Tonometer	1.10	0.08
Goldman Applanation Tonometer - iCare Tonopen	0.26	0.87
iCare Tonopen – Airpuff Tonometer	-1.36	0.02

Overall, the difference was statistically significant with p value = 0.021. Table 2 shows the differences between IOP measurement methods.

DISCUSSION

Goldmann applanation tonometry has been considered as a gold standard for IOP measurement for a long time.^{12,13} However, there are some instances in which IOP measurement by GAT is not possible or very difficult, for example in children and in bed ridden cases.¹⁴ Recent advancements have introduced many instruments and methods for measuring IOP.¹⁵ The reliability and accuracy of these instruments is comparable to GAT.¹⁶ Airpuff tonometer and a recently introduced iCare tonometer are some alternatives which can be used to measure IOP in challenging cases.¹⁷

In a recent study by Demirci et al¹⁶, measurements by rebound tonometer, non-contact airpuff tonometer and Goldmann applanation tonometer were compared in three groups of healthy subjects based upon the age i.e. group 1 (7 – 17 years), group 2 (18 – 40 years) and group 3 (41 – 75 years). Central corneal thickness was also measured by ultrasonic pachymeter. According to their study, Airpuff tonometer readings were significantly higher than both Goldmann applanation tonometer and rebound tonometer measurements in all groups. There was no statistically significant difference between Goldmann applanation tonometer and rebound tonometer measurements in group 1 ($p = 0.248$), group 2 ($p = 0.63$), and group 3 ($p = 0.126$). In our study, we did not group the subjects on the basis of age. Rather we only included subjects aged between 21 and 39 years with healthy eyes. Moreover, we did not perform pachymetry in our subjects.

Erdogan et al¹⁸ conducted a study to compare intraocular pressure (IOP) measurements by noncontact tonometer (NCT), Goldmann applanation tonometry with fluorescein (fGAT), and GAT without fluorescein (nGAT). They also assessed the effect of central corneal thickness (CCT) and keratometric values on IOP. One hundred and eighty eight eyes of 94 healthy subjects were included in the study. Their study showed that there were statistically insignificant differences in IOP values by nGAT and fGAT ($P > 0.05$), and were correlated positively with NCT readings. IOP readings were independent of CCT and keratometry readings. Whereas in our study we did not

compare the IOP values obtained by GAT done with and without fluorescein. Our sample size was 25 as compared to 94 in their study.

In another study, Grewal et al¹⁹ investigated the correlation among intraocular pressure (IOP) values, in 50 eyes of 50 patients who already had undergone vitreoretinal surgery, obtained by iCare rebound tonometer, Tonopen, and GAT. However, in our study, we only included subjects with healthy eyes with no previous history of trauma or surgery. Mean IOP values obtained by iCare, Tonopen, and Goldmann were 15.9 ± 8.9 , 16.9 ± 6.2 , and 16.0 ± 7.3 mm Hg, respectively. They concluded that, post vitreoretinal surgery, there is excellent agreement among IOP values obtained by iCare rebound tonometer, Tonopen, and GAT. They also concluded that iCare overestimated IOP when IOP was ≥ 23 mm Hg and underestimated the IOP when IOP was < 10 mm Hg. We, in our study, did not assess the effect of low or high IOP on iCare tonopen values.

In another study by Takenaka et al²⁰, comparison was done among the IOP values measured by NCT, GAT, iCare tonometer and the Tonopen XL while wearing soft contact lenses (SCLs). IOP was measured in twenty-six healthy subjects, wearing SCLs of -5.00 D, -0.50 D and +5.00 D, using NCT, GAT, iCare, and the Tono-Pen XL. Using GAT, while wearing +5.00 D lenses, IOP readings were higher than those of the naked eyes. Whereas, when measured by iCare, IOP readings were almost same as measured over SCLs ranging from -5.00 D to +5.00 D and were also comparable with values obtained by GAT in the naked eyes. Thus, it was concluded that to measure IOP through SCLs, NCT and iCare were best alternatives. Tonopen XL was not included in our study.

Raina et al²¹ in their study compared the IOP values obtained by GAT, Tono-Pen and noncontact tonometer in children. They had a sample size of 200 eyes of Indian children (aged 8 – 18 years). IOP was measured by using above-mentioned three tonometers. Effect of CCT on IOP was also analyzed. The mean IOP was 14.38 with NCT, 15.63 with Tonopen, and 12.44 mm Hg with GAT i.e. lowest with GAT and highest with tonopen. These results contradict the results of our study as mean IOP in our study was 15.84 ± 2.736 mm Hg with Airpuff Tonometer, 14.48 ± 2.435 mm Hg with iCare Tonopen and 14.74 ± 2.489 mm Hg with GAT i.e. lowest with iCare tonopen and highest with Airpuff NCT. Therefore, there was some ambiguity regarding the IOP values obtained by

different tonometers.

Limitation of our study was that we did not assess the effect of central corneal thickness (CCT) on IOP values obtained by different tonometers. The effect of very low and high IOPs on the measurements was also not investigated with three tonometers.

CONCLUSION

IOP readings taken by iCare Tonopen and Airpuff Tonometer are comparable to those taken by Goldmann Applanation Tonometer and iCare Tonopen underestimates the IOP when compared with Airpuff Tonometer. Thus iCare rebound tonometry and Airpuff tonometry are reliable alternatives to Goldmann applanation tonometry and both of these can be used to measure IOP in challenging cases.

Ethical Approval

The study was approved by the Institutional review board/ Ethical review board.

(1-2/20-MIMC/ERB/0018)

Conflict of Interest

Authors declared no conflict of interest.

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