ORIGINAL RESEARCH A Comparative Study Between the Goldmann Applanation Tonometer and the Non-Contact Air-Puff Tonometer (Huvitz HNT 7000) in Normal Eyes

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Objective: To assess whether the non-contact air-puff tonometer (NCT) is an appropriate alternative to the Goldmann applanation tonometer (GAT) for measuring intraocular pressure (IOP).

Patients and Methods: An observational, cross-sectional, and comparative study with a quantitative approach was carried out. Two techniques for IOP measurements using the standard GAT and the NCT were compared. A total of 180 eyes from 90 patients were included in the study.

Results: The total mean IOP according to NCT measurements was 14.12 mmHg, and the total mean IOP according to the GAT was 12.98 mmHg; these values were significantly different (p=0.0018). When dividing the participants into three groups according to the measurement range obtained and comparing the mean NCT and GAT measurements in each group, in Group 1 (10-15 mmHg), no statistically significant difference was found between the means of the two tonometers (p=0.3100), a difference was observed between Group 2 (16-19 mmHg) and Group 3 (20 mmHg or more) (p<0.001). When dividing the participants by age group, the means obtained by the two tonometers also differed significantly between Group 4 (40-59 years) and Group 5 (60 years or more) (p<0.0001). In all groups, the mean measurements by the NCT were higher than those by the GAT.

Conclusion: The NCT presented an approximate mean of the measures with the GAT in group 1 but was overestimated in the measurements of the groups 2 and 3.

Keywords: ocular tonometry, glaucoma, ocular hypertension, intraocular pressure, efficiency

Plain Language Summary

Glaucoma is one of the major causes of blindness in the world even though, in most cases, it is a controllable desease. One method of detecting the disease is measuring high intraocular pressure using tonometers, one of the manifestations of glaucoma, although there are others, also important, such as changes in the visual field and optic nerve. Some tonometers directly contact the eyes, requiring instillation of anaesthetic eye drops and fluorescein for their use, while others do not contact the eyes and are easier to use but are less accurate. The ease of obtaining intraocular pressure measurements using non-contact air-puff tonometers for the early diagnosis of glaucoma motivates studies on how these devices perform relative to the standard measuring instrument-the Goldmann tonometer. The non-contact tonometer overestimates the measurement at higher pressures, does not replace the Goldmann tonometer,

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but can be used for screening due to its easier use when compared to the Goldmann tonometer.

Introduction

The decision to treat glaucoma is directly influenced by intraocular pressure (IOP) measurements, emphasizing the need to obtain accurate measurements. Decisions based on false-positive or false-negative diagnoses of glaucoma can have severe consequences because the damage caused to the optic nerve is irreversible.^{1(p. 12)} On the other hand, unnecessary exposure to potentially harmful medications can generate inappropriate anxiety and waste resources, rendering the discussion about the modalities of medical equipment that best support the diagnosis of glaucoma very timely.

IOP is the hydrostatic pressure exerted by the aqueous humour in the intraocular tissues as a result of the balance between production and drainage. IOP varies between individuals and between the eyes of the same individual. The mean IOP in adult populations is estimated to be between 15 mmHg and 16 mmHg, with a standard deviation (SD) of 2.5 mmHg to 2.8 mmHg²; however, values below 21 mmHg are considered normal.^{1(p. 33)} IOP measurement is part of the basic ophthalmological evaluation since ocular hypertension is the main risk factor for the onset and development of glaucoma, which is the only modifiable risk factor.^{3(p. 128),4(p. 20)} Lowering IOP is a proven method to delay or stop the progression of glaucoma and thereby reduce visual loss, thus preserving patients' quality of life.⁵

The slit lamp-mounted Goldmann applanation tonometer (GAT) is the most commonly used instrument to measure IOP and is currently considered the reference instrument, ie, the most accurate and recommended standard to which all other tonometers are compared.^{1(p. 33)} However, it should be noted that even the measurement made with the standard tonometer (GAT) is not precise since its values are significantly lower than the intracameral pressure.⁶

On the other hand, the non-contact air-puff tonometer (NCT) is simpler to operate, is non-invasive, does not reduce IOP by the massage effect, and does not require prior anaesthetic instillation, all of which contribute to increased comfort, reduced damage to the corneal surface, and a lower risk of contamination.^{2,6} The measurement requires only a few fractions of a second and is completed when the sensor receiver detects the light being reflected

in the cornea; the measured value is shown on an LCD screen or a touch screen depending on the model.

Early identification of patients with glaucoma can delay their blindness process from 23 to 35 years, supporting the adoption of methods that facilitate IOP measurement on a large scale as is possible with the NCT.² In this context, the present study aimed to analyse whether the NCT is an appropriate alternative to the GAT to measure IOP.

Patients and Methods

A retrospective, cross-sectional, and comparative study with a quantitative approach was performed. Two IOP measurement techniques were compared: the standard GAT and the NCT.

The study was conducted at the *Centro Oftalmológico Belloto Stock* (Belloto Stock Ophthalmological Centre) – COBS during routine ophthalmologic visits and included 180 eyes of 90 patients. Before data collection, the participants received verbal information about the procedures involved. All patients signed an informed consent.

The medical devices used in the study were the NCT Huvitz HNT 7000 (Republic of Korea) and the GAT HS Haag-Streit Diagnostics (Switzerland). For the examination, Anestalcon anaesthetic eye drops (5 mg/mL proxymetacaine hydrochloride) were used. The medical equipment was calibrated prior to the examination.

The target population consisted of patients aged 40 years or older with healthy eyes. The exclusion criteria were patients with previous eye diseases, glaucoma patients or eye with 20mmhg or more measured by TAG, eye drop users, and patients diagnosed with some systemic disease, such as diabetes or rheumatic diseases, during the first visit.

The IOPs of both eyes of each patient were evaluated, first on the right and then on the left and each eye was used as one isolated data. The NCT measurement was always performed before the GAT measurement to eliminate the effect of ocular massage, which has been described when using the GAT and is absent with the NCT.⁶ The patients were selected randomly and sequentially from February 2019 to February 2020.

After performing tonometry with both devices, the values were tabulated and statistically analysed using the statistical programme Bioestat 5.3. For data analysis, the Mann–Whitney test was used for the difference between the IOP measures at a significance level of 5%. The groups were considered different when the comparison of the mean IOP measurements resulted in p≤0.05.

Results

A total of 180 eyes from 90 patients with healthy eyes were included in the study. Of the 90 patients, 51 were female and 39 were male. The total mean IOP with the NCT was 14.12 mmHg, with an SD of \pm 3.41, a minimum value of 8 mmHg, and a maximum value of 23 mmHg. The total mean IOP with the GAT was 12.98 mmHg, with an SD of \pm 2.93, a minimum IOP value of 9 mmHg, and a maximum value of 19 mmHg.

The IOPs measured by the NCT and GAT were divided into three groups. For Group 1 (Table 1), IOPs between 10 and 15 mmHg were selected, totalling 123 eyes. In this group, the total mean IOP measured by the NCT was 12.22 mmHg (SD \pm 1.85), and that measured by the GAT was 12.12 mmHg (SD \pm 2.50). In Group 2 (Table 2), IOPs between 16 and 19 mmHg were selected, with total mean IOPs of 17.36 mmHg (SD \pm 1.19) by the NCT and 14.39 mmHg (SD \pm 2.83) by the GAT, totalling 47 eyes. In Group 3 (Table 3), patients with IOPs of 20 mmHg or more were included, with total mean IOPs of 21.3 mmHg (SD \pm 1.05) by the NCT and 17.5 mmHg (SD \pm 1.08) by the GAT, totalling 10 eyes.

According to age, patients were divided into two groups: Group 4, patients aged between 40 and 59 years, totalling 96 eyes, and Group 5, patients aged 60 years or older, totalling 84 eyes. In Group 4, the mean IOPs were 14.29 mmHg (SD \pm 2.82) by the NCT and 12.85 mmHg (SD \pm 2.50) by the GAT. In Group 5, the mean IOPs were 14.03 mmHg (SD \pm 3.87) by the NCT and 13.27 mmHg (SD \pm 3.10) by the GAT.

Table I Comparison of the Mean and Standard Deviation of IOPBetween 10 and 15 mmHg Measured by the Non-Contact Air-Puff Tonometer (NCT) Relative to the Goldmann ApplanationTonometer (GAT)

	NCT (mmHg)	GAT (mmHg)	p value	
Mean Stead and deviation	12.22	12.12	0.3100	
Standard deviation	1.85	2.50		

Table 2 Comparison of the Mean and Standard Deviation of IOPBetween 16 and 19 mmHg Measured by the Non-Contact Air-Puff Tonometer (NCT) Relative to the Goldmann ApplanationTonometer (GAT)

	NCT (mmHg)	GAT (mmHg)	p value
Mean	17.36	14.39	<0.0001
Standard deviation	1.19	2.83	

Table 3 Comparison of the Mean and Standard Deviation ofIntraocular Pressure (IOP) of 20 mmHg or More Measured bythe Non-Contact Air-Puff Tonometer (NCT)Relative to theGoldmann Applanation Tonometer (GAT)

	NCT (mmHg)	GAT (mmHg)	p value
Mean	21.3	17.5	<0.001
Standard deviation	1.05	1.08	

Discussion

When comparing the total mean IOP, the value obtained by the NCT (12.12 mmHg) was greater than the mean obtained by the GAT at 1.98 mmHg, and this difference was statistically significant (p=0.0018). However, when dividing the patients into three groups according to the pressure obtained, in Group 1 (10–15 mmHg), no statistically significant difference was found between the total mean IOP measurements obtained with the NCT and the GAT (p=0.3100); that is, the IOP measurements with both devices were similar in the range of 10 to 15 mmHg (Table 1).

A similar result was found in three other studies, which did not divide the participants by pressure range and obtained NCT results comparable to those of the GAT except at extreme pressures, although the mean NCT values were slightly higher.^{7–9} However, a study conducted with the same device (Huvitz HNT-7000) found lower values relative to GAT measurements in healthy participants aged between 20 and 25 years, but the authors warned that other studies found higher values, as observed in the present study and that the measurements may be influenced by both human factors and the models (as well as the units) of the devices used.¹⁰

However, the correlation between the mean IOPs obtained by the NCT in Group 2 (16–19 mmHg) and in Group 3 (20 mmHg or more) was higher than that found with the GAT, and the differences were statistically significant (p<0.0001), showing that in these groups, the NCT overestimated the IOP (Tables 2 and 3). This result is consistent with two other studies that found results similar to those of the GAT at measurements between 10 and 20 mmHg, although the accuracy decreased at the higher values.^{11,12}

NCT measurements are influenced by corneal thickness but are highly related to GAT measurements as shown by another study in both patients with ocular hypertension (p=0.086) and patients with primary open-angle glaucoma, with no significant difference between groups (p=0.1112), although the NCT measurements were slightly higher than the GAT measurements, and a difference in the measurements of the control group was found (p=0.033).¹³ Regarding thickness, a study showed that NCT values were higher than GAT values in thin and normal corneas, but that IOP was overestimated in thicker corneas.¹⁴

Glaucoma is prevalent among older people. Therefore, an analysis by age group was performed. In Group 4 (40–59 years) and Group 5 (60 years or more), as in Group 2 (16–19 mmHg) and Group 3 (over 19 mmHg), a statistically significant difference (p<0.0001) was found between the total mean IOP measurements obtained by the NCT, which were higher, and the total mean IOP measurements obtained by the GAT, which were lower. This result is consistent with a study that found a significant difference and an increase in IOP measurements for both devices with increasing age, which were not found in the present study.¹⁵

However, the equipment does not perform equally, and NCT measurements may vary according to the brand and model of the device used. Therefore, the results were compared by the brands of devices studied in articles containing information about brands, results, statistical analyses, and participant characteristics to build the theoretical framework (Table 4).

Two studies comparing the same brand of device, but different models (Topcon CT-60¹³ and CT-80)⁹ showed more accurate measurements with the latest model (CT-80) without a significant difference relative to GAT measurements; however, the participants in this group (CT-80) were young, between 20 and 27 years old, with lower mean pressures, which may have influenced the accuracy. In another study with a Topcon CT 80 tonometer including patients with all pressure levels, the mean IOP measured by the NCT was higher than that measured by the GAT (15.91 mmHg x 13.01 mmHg), and the difference was significant, suggesting that an increase in IOP increases the inaccuracy of the NCT (Table 4).¹⁶

A study using the Nidek NT-530 model in participants with a mean IOP below 21 mmHg showed no difference relative to the GAT (p=0.998) as observed in Group 1 in the present study (Tables 1 and 4).⁸ However, two other studies using the Nidek SL-300 and Tonoref Nidek II models in patients with all IOP levels found a significant difference between the NCT and GAT (p \leq 0.001 and p<0.01, respectively) as observed in Groups 2 and 3 in the present study (Tables 2 and 3), showing that participant

characteristics and the model of devices of the same brand can influence the results (Table 4).^{15,17}

Reliable results relative to GAT results for Goldmanncorrelated intraocular pressure (IOPg) were obtained with the Reichert PT-100 (p=0.64)⁷ and Reichert CR7 (p=0.52) even in patients with all pressure levels, although lower accuracy was achieved when considering the cornealcompensated IOP (IOPcc) modality in Reichert CR7 (p<0.01) (Table 4).¹⁸

A study with the Corvis ST tonometer including patients with all pressure levels obtained means similar to those of the GAT, with no significant difference between the tonometers (p=0.1162).¹⁹ However, these results do not agree with the findings of another study with glaucoma patients with all pressure levels, which obtained lower values with a device of the same brand and model (Corvis ST) for both the corrected IOP based on corneal thickness (IOPpachy) and IOPcc measurements (p<0.01), perhaps because these units do not operate exactly the same, and technical or human factors affect the measurements (Table 4).¹⁸

A recent study confirmed that no significant difference existed between the Tomey FT-1000 and the GAT for measurements below 21 mmHg (p=0.151), which is similar to the observations in Group 1 in the present study, but when measurements were equal to or greater than 21 mmHg, the NCT obtained significantly lower values than the GAT (p=0.016) in contrast to the observations in the present study.¹⁸ In another study conducted with a different NCT model (Thomey NCT) in which participants were not divided by pressure group, a difference relative to GAT was found (p<0.001), with the NCT yielding lower values.²⁰

A study analysing the Canon Full Auto Tonometer TX-F in patients aged 16 to 78 years without stratification by pressure group also found a higher mean value obtained by the NCT than by the GAT, with values of 18.17 mmHg and 15.59 mmHg, respectively, which may have contributed to the significant difference between groups (p=0.003) (Table 4).^{20,21} Accordingly, a study evaluating the repeatability and reproducibility of the Reichert PT-100 tonometer in young patients showed exactly the same mean results between the NCT and GAT in the first measurement (a mean of 15 mmHg for both instruments), corroborating the finding that NCT results are more reliable at lower IOP levels.⁶

In general, as shown in Table 4, seven of the thirteen devices showed no significant differences relative to the

Year	Authors	NCT Brand/ Model	NCT Measurement in mmHg	GAT Measurement in mmHg	Þ	Participants
2021	Stock et al (This research)	Huvitz HNT 7000 ^a	12.22	12.12	0.3100	Group 1: 10–15 mmHg
			17.36	14.39	<0.0001	Group 2: 16–19 mmHg
			21.30	17.50	<0.001	Group 3: ≥20 mmHg
2020	Trakanwitthayarak	Tomey FT-1000ª	14.14	13.60	0.151	Group 1:<21 mmHg
	S		30.21	35.12	0.016	Group 2: ≥21 mmHg
2019	Luebke et al	Corvis ST ^a	18	17.6	0.1162	Outpatients with a mean age of 66.9
		Thomey NCT	16.3	17.6	<0.001	years
2017	Nakao et al	Corvis ST- IOPpachy	10.3	13.6	<0.01	Patients with primary open-angle glaucoma with a mean age of 61.94 years
		Corvis ST-IOPcc	9.7	13.6	<0.01	
		Reichert 7CR- IOPg ^a	13.7	13.6	0.52	
		Reichert 7CR- IOPcc	15.5	13.6	<0.01	
2017	Jacob et al	Tonoref Nidek II ^b	<13 mmHg	> GAT	<0.001	Patients aged 20 to 60 years
			≥I3 mmHg	<gat< td=""><td></td><td></td></gat<>		
2016	Mahsud et al	Canon Full Auto Tonometer TX-F	18.17	15.59	0.003	Patients aged 16 to 78 years
2016	Tolomei et al	Nidek SL-300	22.77	17.79	<0.001	Patients with a mean age of 51.6 years
2014	llmaz et al	Nidek NT-530ª	16.1	15.5	0.998	Patients without glaucoma, with IOP <21 mmHg
2013	Farhood QK	Topcon CT 80	15.91	13.06	0.001	Outpatients aged 15 to 84 years
2009	Salim et al	Reichert PT-100 ^a	15.0	14.3	0.64	Patients without glaucoma with a mean age of 62.6 years
2006	Ogbuehi CK	Topcon CT 80 ^a	13.6	13.4	>0.05	University students aged 20 to 27 years
2005	Ko et al	Topcon CT-60	16.6	15.5	<0.001	Patients older than 18 years

 Table 4 Comparative Results of the Non-Contact Air-Puff Tonometer (NCT) Relative to the Goldmann Applanation Tonometer (GAT) by the Brand/Model of Device from 2005 to 2020

Notes: ^aDevices that showed no difference relative to the Goldman tonometer in at least one of their measurement methods or one of the groups of participants. ^bFor values below 13 mmHg, the NCT obtained higher values than the GAT, and for values equal to or above 13 mmHg, the NCT values were lower than the GAT values.

GAT (p<0.05) in at least one of its measurement methods (when the device had more than one) as observed in Group 1 in the present study, which had lower pressure values, contributing to the similarity of the results.

However, six NCT devices showed differences relative to the GAT (p>0.05) as observed in the other groups in the present study, suggesting that in addition to the technical, human, pressure, and corneal thickness factors involved in the measurements, the NCT brand, unit, and model can also influence the results. On the other hand, in addition to the risk of infection, inaccuracies can also occur during measurements with the GAT due to calibration defects, incorrect techniques, a tight collar or tie, the Valsalva manoeuvre, breath holding, constricting or touching the eyelids, measurement repetitions, and alteration of the tear film.^{1(p. 33),6(p. 33),22}

Conclusion

The results of this study indicate that the NCT used was a valid alternative to the GAT, which is considered the gold standard, only in Group 1 (eyes with an IOP between 10 and 15 mmHg). In Groups 2 and 3 (eyes with an IOP equal to or greater than 16 mmHg), as well as in Groups 4 (patients aged 40 to 59 years) and 5 (patients aged 60 years or older), the NCT overestimated the IOP.

The non-contact tonometer overestimates the measurement at higher pressures, does not replace the Goldmann tonometer, but may be a good screening tool that helps the measurement of IOP, especially in eyes with less than 15mmhg, due to its ease of use. The results of the NCT depend on the brand, unit, and model used, and most NCT devices are more reliable for measuring lower eye pressures as observed in the present study, emphasizing the need to develop devices with greater accuracy at higher pressures to increase the level of reliability.

The limitation of the study was the performance of NCT with only one device brand, and a strength of the study was the stratification of participants by pressure range. Future studies comparing different non-contact devices (brand and model) with participants stratified by eye pressure range will be necessary to better understand their performance compared to the Goldmann tonometer in the measurement of intraocular pressure.

Abbreviations

IOP, intraocular pressure; GAT, Goldmann applanation tonometer; NCT, Non-contact air-puff tonometer.

Ethics Approval

Ethics approval was obtained from the research ethics committee of the Universidade do Oeste de Santa Catarina by number 3,412,730.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

The authors declare no conflicts of interest for this work.

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