Effect of mydriatic and cycloplegic drugs in glaucomatous and nonglaucomatous eyes using ultrasound biomicroscopy

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SUMMARY

**Purpose:** To study the anatomical relationships of the anterior segment of the eye, using ultrasound biomicroscopy in patients with chronic simple glaucoma and nonglaucomatous eyes, after mydriatic and cycloplegic instillation.

**Patients and Methods:** Thirty eyes with chronic simple glaucoma and 30 nonglaucomatous eyes were studied. Anterior chamber depth, iris-lens contact and iris-zonule distance were measured, in both groups, using ultrasound biomicroscopy, in the three following conditions: without mydriatic and cycloplegic instillation, with 1% tropicamide alone and with 1% tropicamide plus 10% phenylephrine chlorhydrate.

**Results:** Ultrasound biomicroscopy examinations of glaucomatous and nonglaucomatous eyes, without mydriatic and cycloplegic instillation, showed no statistically significant difference in the central anterior chamber depth, iris-lens contact and iris-zonule distance. The difference between examinations without mydriatic and cycloplegic instillation, with 1% tropicamide and with 1% tropicamide plus 10% phenylephrine chlorhydrate was statistically significant regarding the variables measured in both groups.

**Conclusions:** No anatomical difference was observed between the anterior segment of glaucomatous and nonglaucomatous eyes after mydriatic and cycloplegic instillation. The results of this clinical investigation, using ultrasound biomicroscopy were innovative, allowing a dynamic and quantitative evaluation of the anatomical relationships between intraocular structures when submitted to mydriasis and cycloplegy, which up to now were only qualitative.

**Key words:** Anterior segment of the eye; Ultrasonography; Glaucoma; Open-angle; Mydriatics; Cycloplegics.

INTRODUCTION

The advent of ultrasound biomicroscopy (UBM) expanded echographic studies *in vivo* of the anterior segment structures of the eye and their anatomical relationships with maximum depth at the ciliary body, peripheral retina and vitreous 1,2. The UBM examination using a 50 MHz transducer, allows a 5 mm penetration but with a 37 µm microscopic resolution 3.

The UBM technique turned to be useful in research and clinical practice of glaucoma, because serial observations of different stages of the disease can be obtained 4. Extending this method to glaucomatous eyes allows to
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get a view and to measure the relationship between the anatomical structures of the anterior segment among these eyes, since the cause of several types of glaucoma is a structural component and possibly may be explained using this technique.

The mydriatic and cycloplegic solutions have been used for pupil dilation as a routine in ophthalmic practice. It is therefore significant to note that these drugs also have the ability to raise the intraocular pressure (IOP) in eyes with open angles specially when they are glaucomatous. Some eyes with open angle glaucoma may develop increased intraocular pressure while under mydriasis even though no angle narrowing occurs. No matter what the true mechanism of this response, it is obvious that significant increases in intraocular pressure under cycloplegia are not the province of angle closure glaucoma alone.

The objective of this study is to evaluate, by means of ultrasound biomicroscopy, the anterior segment of the eye of the patient with chronic simple glaucoma and nonglaucomatous eyes after mydriatic and cycloplegic instillation.

PATIENTS AND METHODS

Sixty eyes were selected and divided into two groups: group G consisting of 30 eyes with chronic simple glaucoma (CSG) and group NG, consisting of 30 nonglaucomatous eyes. Group G presented IOP ≥21 mmHg; open and wide anterior chamber angle; alterations of the optic disk and visual field; severe hypertension, arrhythmia, hyperthyroidism, myocardopathies and arteriosclerosis; hypersensitivity to the used drugs; corneal abnormalities; previous eye surgeries; ametropia and was under recent medical control with 0.5% timolol maleate eyedrops associated or not with 2% pilocarpine chlorhydrate. Patients of both groups were white and their irises were brown. The protocol of this investigation was approved by the Medical Ethics Committee of the Federal University of São Paulo, Brazil.

Patients were excluded when they presented: evidence of diseases which would alter pupillary response; high risk of CSG; IOP without clinical control; and/or excavation/disk ratio greater than -1.25 diopters; open and wide anterior zonule, at its junction with the ciliary process.

UBM examination (C2) was performed after using 1% tropicamide. On the same day, after using 1% tropicamide, a second examination was carried out (C1). A week later, a third UBM examination (C2) was performed after using 1% tropicamide plus 10% phenylephrine.

No significant difference was found (Mann Whitney test, p>0.05) between the anterior chamber depth of groups G and NG at the previous examinations (Prev). A significant increase in the anterior chamber depth of both groups was observed (Friedman test, p<0.05) when comparing the previous examination (Prev), with the examination after instillation of 1% tropicamide (C1) and after instillation of 1% tropicamide plus 10% phenylephrine chloride (C2) (Table 1).

No significant difference of the iris-lens contact and iris-zonule distance was observed (Mann Whitney test, p>0.05) between the groups G and NG in the 12, 3, 6 and 9 regions at the previous examination (Prev). A statistically significant decrease in iris-lens contact and iris-zonule distance in the 12, 3, 6 and 9 regions was observed in the studied groups, comparing the previous examinations (Prev), with those using 1% tropicamide (C1) and with those using 1% tropicamide plus 10% phenylephrine chloride (C2) and also on comparison of the two latter (Friedman test, p<0.05) (Table 2 and 3).
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There was no significant difference between the measurements of ocular structures, such as, iris-lens contact and iris-zonule distance at the previous examinations (Prev), with 1% tropicamide (C1) and with 1% tropicamide plus 10% phenylephrine chlorydrate (C2) in the 12, 3, 6 and 9 regions of each studied eye (Friedman test, p>0.05).

Table 1. Results of the mean of the anterior chamber depth measurements (µm) of glaucomatous (G) and nonglaucomatous (NG) eyes at the previous examinations (Prev), with 1% tropicamide eyedrops (C1) and with 1% tropicamide plus 10% phenylephrine chlorydrate (C2).

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<th>G (µm)</th>
<th>NG (µm)</th>
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<tbody>
<tr>
<td>Prev</td>
<td>3046</td>
<td>3019</td>
</tr>
<tr>
<td>C1</td>
<td>3111</td>
<td>3114</td>
</tr>
<tr>
<td>C2</td>
<td>3097</td>
<td>3085</td>
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Table 2. Results of the mean of the iris-lens contact measurements (µm) in the 12, 3, 6 and 9 regions of glaucomatous (G) and nonglaucomatous (NG) eyes in the previous examinations (Prev), with 1% tropicamide eyedrops (C1) and with 1% tropicamide plus 10% phenylephrine chlorydrate (C2).

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<th></th>
<th>G (µm)</th>
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<tr>
<td>Prev</td>
<td>1287</td>
<td>1203</td>
</tr>
<tr>
<td>C1</td>
<td>1269</td>
<td>1249</td>
</tr>
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<td>C2</td>
<td>1292</td>
<td>1220</td>
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Table 3. Results of the mean of the iris-zonule distance measurements (µm) in the 12, 3, 6 and 9 regions of glaucomatous (G) and nonglaucomatous (NG) eyes, at the previous examinations (Prev), with 1% tropicamide eyedrops (C1) and with 1% tropicamide plus 10% phenylephrine chlorydrate (C2).

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<th></th>
<th>G (µm)</th>
<th>NG (µm)</th>
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<tr>
<td>Prev</td>
<td>622</td>
<td>617</td>
</tr>
<tr>
<td>C1</td>
<td>581</td>
<td>519</td>
</tr>
<tr>
<td>C2</td>
<td>371</td>
<td>344</td>
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Figure 1 - A) Central measurement of anterior chamber depth of a glaucomatous eye at the previous examination. The vector (arrow) is positioned at (a) the central area of the anterior chamber. Mode A was used to measure the distance from the peak (b) referring to the Descemet membrane /endothelium to (c) the peak referring to the anterior surface of the lens and the result being X = 2.726 mm or 2726 µm. B) Measure of the iris-lens contact of a glaucomatous eye at the previous examination obtained along the extension of the surface of the iris-lens contact (arrow), the result being length (L) = 0.848 mm or 848 µm. C) Measure of the iris-zonule distance of a glaucomatous eye at the previous examination obtained by drawing a perpendicular line from the anterior zonule to the posterior face of the iris (arrow), the result being length (L) = 0.622 mm or 622 µm.
Elevation of IOP by many mydriatic and cycloplegic drugs has been reported to occur in eyes in which the filtration angle remained open during mydriasis. The incidence of pressure elevation of 6 mmHg or more is 23% in a population with open angle glaucoma and 2% in apparently normal population.

Conventional 10 MHz ultrasonography is capable of measuring relatively large distances such as anterior chamber depth; however, ultrasound biomicroscopy increases accuracy because of finer positioning endpoints and an improved exact measurement position.

In a biometry study of cataractous eyes the mean anterior depth in the phakic eyes was 3.24 ± 0.44 mm.

The UBM technique produces sections of the living human eye, at microscopic resolution without violating the integrity of the globe. This technique allowed to define the profile of the anterior chamber extension. The measurement of the anterior chamber depth is not restricted to the axial position and corresponds to any point from the endothelial face to the surface of the iris or lens.

In this study, the axial anterior chamber depth was similar in the previous examinations (Prev) of groups G (mean, 3046 μm) and NG (mean, 3019 μm). This result agrees with that found in nine normal individuals in which the axial measurement of the anterior chamber depth using UBM was 3128 ± 372 μm. An increase in the anterior chamber depth was observed by the UBM technique after the use of mydriatics and cycloplegics in groups G and NG, corroborating the findings of the literature. But the difference of the anterior chamber depth between the examinations with 1% tropicamide (C1) and with 1% tropicamide plus 10% phenylephrine chloride (C2) was not significant. In the investigated literature, no studies using a similar method which would allow a comparison with our results were found.

The iris is positioned on the lens at the pupillary border and this contact area may be measured using UBM examination. However, any space which may exist between the iris and the lens is too small to be detected. On the other hand, the aqueous humor could not escape from the posterior chamber of the eye if there would be a constant iris-lens contact in all meridians. It is assumed that the iris-lens separation is well below the power of resolution of the equipment used in the UBM and, therefore, is not observed.

In this study, no significant difference of the iris-lens contact was observed in the 12, 3, 6 and 9 regions of each studied eye. The small differences may, on the other hand, show the dynamics of the mydriatic and cycloplegic status between the several regions of the eye at the same examination. The variable of the anterior chamber depth was measured only in the axial position but all the previous conditions established for each examination were observed.

The results obtained in this investigation agreed with gonioscopic and histological findings of the literature, corroborated the pharmacological and biomechanical action of the studied mydriatic and cycloplegic drugs as well as allowed a dynamic and quantitative evaluation of the anatomical relationships of intraocular structures in normal and glaucomatous eyes.

The UBM equipment is valuable in the conduction of clinical investigation of glaucoma. Eyes under cycloplegia whose angles remained open but whose intraocular pressure rose also displayed a simultaneous decrease in outflow facility. Some authors speculated that such a decrease in aqueous outflow values produced by cycloplegia is due to decrease in the tonus of the ciliary muscle. However, in this study, no anatomical difference was observed between the anterior segment of glaucomatous and nonglaucomatous eyes after mydriatic and cycloplegic instillation.

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**RESUMO**

Objetivo: Estudar por meio da biomicroscopia ultra-sônica, as relações anatômicas das estruturas do segmento anterior dos olhos de pacientes portadores de glaucoma crônico simples e não-glaucomatosos, após a instalação de miôdriáticos e cicloplégicos.

Pacientes e métodos: Foram estudados 30 olhos com glaucoma crônico simples e 30 olhos não-glaucomatosos. A profundidade da câmara anterior, o toque iris-cristalino...
e a distância iris-zônula foram analisados pela biomicroscopia ultra-sônica sem a instilação demidriáticos e cicloplégicos, com tropicamida a 1% e com associação de tropicamida a 1% mais cloridrato de fenilefrina a 10%.

Resultados: Os exames de biomicroscopia ultra-sônica dos olhos glaucomatosos e não-glaucomatosos, sem a instilação de cicloplégicos e midriáticos, não mostraram diferença significante da profundidade da câmara anterior, do toque iris-cristalino e da distância iris-zônula. A diferença entre os exames sem a instilação de midriáticos e cicloplégicos, com tropicamida a 1% e com associação de tropicamida a 1% mais cloridrato de fenilefrina a 10% foi estatisticamente significante nas variáveis estudadas em ambos os grupos.

Conclusões: Não foram observadas diferenças anatômicas entre o segmento anterior dos olhos glaucomatosos e não-glaucomatosos após o uso de drogas midriáticas e cicloplégicas. Os resultados desta investigação clínica usando a biomicroscopia ultra-sônica foram inovadores, permitindo uma avaliação dinâmica e quantitativa das relações anatômicas entre as estruturas intra-oculares quando subme-tidas a midriase e cicloplégia, que até então eram somente qualitativas.

**Palavras-chave:** Segmento anterior do olho; Ultrasonografia; Glaucoma; Ângulo aberto; Midriáticos; Cicloplégicos.

**REFERENCES**


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