The effect of conjunctivochalasis detected by anterior segment optical coherence tomography on tear function in an elderly population

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Abstract
Introduction: The aim of this study was to evaluate conjunctivochalasis (CCH) and its relationship with tear meniscus and tear function in an elderly population.
Materials and methods: This prospective, observational study included 144 eyes of 144 patients aged >65 years who were referred to our clinic for various reasons. The patients were separated into group 1 including 64 eyes of 64 patients with CCH and group 2 including 80 eyes of 80 patients without CCH. All patients in both groups underwent a full ophthalmological examination, and the presence of CCH, fluorescein tear break-up time (FTBUT) test, Schirmer test, ocular surface staining [Oxford grading score] and OSDI [Ocular Surface Disease Index] test results were recorded. Measurements of the conjunctivochalasis area (CCHA), tear meniscus height (TMH) and tear meniscus area (TMA) were taken using anterior segment optic coherence tomodraphy (AS-OCT).
Results: Group 1 comprised 34 females and 30 males with a mean age of 71.15 ± 12.34 years. Group 2 comprised 43 females and 37 males with a mean age of 68.16 ± 6.05 years (p = 0.122). The CCH rate was 44.4% in all of the examined patients. The OSDI score and the ocular surface staining test were significantly higher (p < 0.05), and the FTBUT, TMH and TMA were significantly lower (p < 0.05) in group 1 than in group 2. The Schirmer I test results were not significantly different between the two groups.
Conclusion: The prevalence of CCH is quite high in elderly individuals and may disrupt tear function in these patients.

Keywords: conjunctivochalasis, elderly population, tear meniscus

Introduction
Conjunctivochalasis (CCH) is a common clinical condition characterized by redundant, loose, non-oedematous inferior bulbar conjunctiva interposed between the globe and the lower eyelid causing obliteration of the inferior tear reservoir. It tends to be bilateral and is more prevalent in older populations. It can be seen in different areas such as nasal, temporal and central regions of the upper and lower eyelids. CCH can cause a number of eye problems ranging from dryness in the early mild stage to disturbance of tear outflow in the moderate stage and exposure problems as it becomes severe.

 Conjunctiva with CCH may cause dry eye, interfering with tear distribution due to the mechanical effect of sliding on the cornea and the disruption of integrity of breakdown of continuity tear meniscus in addition to lacrimal punctum occlusion. Intermittent epiphora, prolongation of tear clearance and an accumulation of inflammatory components on the ocular surface as a consequence of ocular surface disease and conjunctival...
epithelium changes can be caused by punctum occlusion. Any inflammation can be improved with the long-term use of protective drops providing symptom relief. Long-term changes in the corneal epithelium may cause displacement in the grey line and occlusion of meibomian gland orifices. This process leads to further instability of the tear film, aggravating the symptoms.

Anterior segment optic coherence tomography (AS-OCT), which provides high-resolution cross-sectional images of anterior segment structures, including the anterior chamber angle, cornea, conjunctiva and tear meniscus, has recently become more widely used. The uniform distribution of the tear film on the ocular surface is important because the distribution of the tear film in the inferior tear meniscus when blinking ensures good optic quality on the ocular surface. A reduction in inferior tear meniscus is increasingly prevalent in the diagnosis of dry eye. Therefore, the measurement of both tear meniscus height (TMH) and area using AS-OCT is becoming more widespread. The measurement of tear meniscus area (TMA) is important, as tear distribution may change in CCH. AS-OCT has been reported as an objective means of the measurement of TMA for the diagnosis of CCH.

The aim of this study was to determine TMH, TMA and conjunctivochalasis area (CCHA) values using AS-OCT in an elderly population with CCH and to evaluate the effects on tear function.

**Method**

The study was conducted in accordance with the principles of the Helsinki Declaration. Approval for the study was granted by the Ethics Committee of Ankara Numune Research and Training Hospital, where the study was carried out (approval number E-18-2097).

This prospective, observational study included 144 eyes of 144 individuals. The patients were divided into two groups according to the presence of CCH. Group 1 included 64 eyes of 64 patients (34 females and 30 males), and group 2 included 80 eyes of 80 patients (43 females and 37 males).

All participants were questioned regarding symptoms of epiphora, pain, burning and itching sensations in the eyes.

The diagnosis of CCH was based on slit lamp examination findings according to the size of the conjunctival folds compared with the height of the normal TMH and number of folds. Grade 0 (no CCH) without persistent fold, grade 1 with a single small fold, grade 2 with two or more folds but not higher than the tear meniscus and grade 3 with multiple folds that are higher than the tear meniscus were evaluated.

After full ophthalmological examination and OSDI (Ocular Surface Disease Index) test, tear function tests were performed in the following order: the measurement of TMH and TMA, Schirmer I test without anaesthesia, fluorescein tear break-up time (FTBUT) and ocular surface staining. Each evaluation was performed by an experienced and masked physician in the early phase.

The OSDI is a questionnaire of 12 items to evaluate the symptoms of eye-related irritation and the effect on visual acuity. The questionnaire was administered into Turkish by a single doctor (D.O.). The total OSDI score of each patient was calculated as follows: OSDI score = Total score of all answered questions × 100/Total number of questions answered × 4. A total score of ≥33 was evaluated as severe.

A standard Schirmer’s filter paper strip (Bio Schirmer Strips; Bio-Tech Vision Care Pvt Ltd, Gujarat, India) was applied to the lower third of the lateral bulbar conjunctiva without topical anaesthesia, left for 5 minutes, then the patient was instructed to blink normally. The results were recorded in millimetres (mm).

The FTBUT test was applied with biomicroscope under cobalt blue. Fluorescein was dropped onto the tear film of the patient, who was then told to blink only once. The FTBUT is recorded as the time in seconds from the blink until the appearance of the first break spot in the fluorescein under cobalt blue illumination. The test was repeated for a total of three times, and the average was taken as the result.

The ocular surface staining with fluorescein test was applied to the lower eyelid of the patients. Fluorescein paper diluted with unpreserved artificial tears was placed in the lateral third of the bulbar conjunctiva. Biomicroscopy was evaluated by a single doctor (D.O.) for the fluorescein staining score, and the ocular surface was graded using the Oxford scoring system (grade 0 = absent and grade 5 = the most severe).
Exclusion criteria were any previous ocular surgery, meibomian gland disease which may cause dry eye; inflammatory, autoimmune, haematological or neoplastic disease; diabetes mellitus; or thyroid disease.

**Determination of the tear meniscus area and conjunctivochalasis area using anterior segment optic coherence tomography**

TMA and CCHA measurements were taken using the Fourier-domain OCT (RTVue, software version 2.7; Optovue, Inc., Fremont, CA, USA). This uses a light source with a wavelength of 1310 nm, a high scanning speed of 30 000 A-scans/second and measures 256 B-scans over the cornea, which enables real-time three-dimensional (3D) image acquisition. The Cornea–Anterior Module is additional software on the device, for anterior segment imaging and the measurement of tear meniscus parameters, including TMH, TMA and CCHA. In the current study, the images were taken after connecting the Cornea–Anterior Module lens to the OCT. The lower tear meniscus and conjunctiva in each eye were imaged with vertical scans centred on the inferior cornea and the lower eyelid, and were evaluated each eye with a digital caliper. TMH, TMA and CCHA were defined as the triangular-shaped wedge of tear film between the lower lid margin and ocular surface (cornea/conjunctiva). TMA and CCHA are shown in Figures 1 and 2.

All measurements of the patients were taken by one individual (D.O.) with three separate evaluations, and the mean of all the measurements was calculated.

**Statistical analyses**

All data were analysed using Statistical Package for Social Sciences (SPSS, Inc., Chicago, IL, USA) 15.0 program for Windows. Continuous variables were expressed with mean ± standard deviation (SD) and categorical variables with number (n). The independent-samples t test was used in the comparison of the means of the patient and control groups. Pearson correlation analysis was used to determine the relationship between TMA and OSDI, fluorescein tear break-up time (FTBUT), ocular surface staining, Schirmer tests and CCHA parameters. A p value of <0.05 was considered as statistically significant.

**Results**

In this study, all patients aged >65 years, CCH was determined in 44.4%. Group 1 comprised 34 females and 30 males, and group 2 comprised 43 females and 37 males. Mean age of patients was 71.15 ± 12.34 years in group 1 and 68.16 ± 6.05
years in group 2. There were no significance between two groups in terms of the age distribution ($p=0.122$). The mean CCH score was $2.12 \pm 0.23$ in group 1. Distribution of all the symptoms in both groups is shown in Table 1. Group 1 patients were significantly more symptomatic than group 2, and burning and itching sensations were the most frequent symptoms. The results of the tear function tests, nasal temporal, central TMH, TMA and CCHA are

Figure 2. Conjunctivochalasis area measurement.

Table 1. Symptoms of the subjects.

| Symptoms                        | Group 1 ($n=64$) | Group 2 ($n=80$) | $p$   |
|---------------------------------|------------------|------------------|-------|
| Epiphora                        | 16               | 10               | 0.054 |
| Pain                            | 23               | 17               | 0.103 |
| Burning and itching sensations  | 38               | 29               | 0.046*|
| Asymptomatic                    | 20               | 42               | 0.045*|

*Statistically significant, $p < 0.05$ is recorded as significant.

Table 2. Results of the tear function tests.

|                              | Group 1          | Group 2          | $p$   |
|                              | $44.13 \pm 16.58$| $18.09 \pm 9.61$| 0.021*|
| OSDI                           | $5.32 \pm 4.12$  | $12.35 \pm 5.16$| 0.041*|
| FTBUT                          | $1.18 \pm 0.43$  | $0.65 \pm 0.32$  | 0.185 |
| Ocular surface staining (0–5)  | $11.25 \pm 5.98$ | $13.57 \pm 2.50$| 0.980 |

*OSDI, Ocular Surface Disease Index; FTBUT, fluorescein tear break-up time. *$p < 0.05$ is recorded as significant.
shown in Table 2. Significant differences were found between the groups with respect to OSDI, FTBUT, nasal TMH, and central TMA and nasal TMA (p = 0.021, p = 0.041, p = 0.021, p = 0.035 and p = 0.023, respectively; Tables 2 and 3). We selected lowest TMA and highest CCHA for each patient and applied correlation tests. TMA was found to be statistically correlated with age, OSDI score, FTBUT and CCHA (p = 0.046, p = 0.025, p = 0.022 and p = 0.021, respectively). CCHA was found to be statistically correlated with age, OSDI score, FTBUT

|                | Group 1          | Group 2          | p    |
|----------------|------------------|------------------|------|
| TMH (µm) nasal | 221.19 ± 12.41   | 301.26 ± 56.47   | 0.021* |
| TMH (µm) central | 289.19 ± 9.22   | 296.25 ± 15.51   | 0.143 |
| TMH (µm) temporal | 305.28 ± 8.14   | 299.15 ± 18.35   | 0.124 |
| TMA (mm²) nasal | 0.043 ± 0.014    | 0.081 ± 0.026    | 0.023* |
| TMA (mm²) central | 0.051 ± 0.009   | 0.089 ± 0.011    | 0.035* |
| TMA (mm²) temporal | 0.063 ± 0.021   | 0.085 ± 0.038    | 0.121 |
| CCHA (mm²) nasal | 0.28 ± 0.09     | –                |      |
| CCHA (mm²) central | 0.19 ± 0.10     | –                |      |
| CCHA (mm²) temporal | 0.17 ± 0.06     | –                |      |
| LIPCOF grade nasal | 2.12 ± 0.23    | –                |      |
| LIPCOF grade central | 1.22 ± 0.01    | –                |      |
| LIPCOF grade temporal | 0.91 ± 0.13    | –                |      |

CCHA, conjunctivochalasis area; TMA, tear meniscus area; TMH, tear meniscus height.
* p < 0.05 is recorded as significant.

|                | TMA [r]          | CCHA [r]         |
|----------------|------------------|------------------|
| r              | p                | r                | p                |
| Age            | -0.610           | 0.046*           | 0.678            | 0.031*           |
| OSDI score     | -0.751           | 0.025*           | 0.571            | 0.043*           |
| FTBUT (sn)     | -0.586           | 0.022*           | 0.533            | 0.044*           |
| Schirmer I test (mm/5sn) | 0.171     | 0.113            | 0.679            | 0.108            |
| Ocular surface staining | 0.183     | 0.121            | 0.121            | 0.132            |
| CCHA           | -0.779           | 0.021*           | –                | –                |
| TMA            | –                | –                | 0.779            | 0.021*           |

CCHA, conjunctivochalasis area; FTBUT, fluorescein tear break-up time; OSDI, Ocular Surface Disease Index; TMA, tear meniscus area.
* p < 0.05 is recorded as significant.
and TMA ($p=0.031$, $p=0.043$, $p=0.044$ and $p=0.021$, respectively; Table 4).

**Discussion**

Although CCH is a common condition in clinical practice, particularly in elderly patients, it is usually an overlooked finding. These patients are often treated with artificial tears because they usually present with dry eye symptoms. However, treatment of dry eye is not satisfactory. CCH tends to get ignored, but it should be considered in the differential diagnoses of ocular surface disease and dry eye. In previous studies, CCH frequency has been reported to range between 8.6% and 98.3% in elderly populations. In the current study of an elderly population, CCH was present in 44.4% of the sample. Diseases such as meibomian gland disease, which are common in the society, and other causes of dry eye were not enrolled in the current study. Thus, our frequency of CCH was lower than previous studies.

It is known that conjunctival flexibility decreases with ageing and this can lead to increased CCH. Ageing is one of the most significant risk factors for the development of CCH. In the current study of an elderly population, an increase was determined in the mean CCHA when the TMA was decreased because the protrusion into tear meniscus created by CCH caused blockage and reduced the TMA.

Dry eye symptoms increase with the severity of CCH. Particularly in the middle and nasal regions, the symptoms associated with the punctum occlusion and disruption of tear meniscus are greater in CCH. FTBUT is more important than the Schirmer test in the diagnosis of dry eye in CCH, because the problem is related to the distribution of tears on the ocular surface, and false-positive rates may be high in the Schirmer test because of the prolongation of tear clearance. Dalianis and colleagues reported that the mean FTBUT values were 5.5 minutes in all patients and 2.8 minutes in symptomatic ones, and CCH in the temporal region caused symptoms, particularly epiphora. In the current study, FTBUT was significantly decreased in patients with CCH ($5.32 \pm 4.12$ in group 1 and $12.35 \pm 5.16$ in group 2; $p=0.041$). FTBUT decreased when the TMA decreased and the CCHA increased, suggesting that CCH caused a decrease in FTBUT through the disruption of tear distribution and a decrease in the TMA by blocking the tear meniscus.

The TMA and CCHA are distributed variably along the inferior lid margin; thus, TMA and CCHA should be measured in the central nasal and temporal parts of the eyelid. CCH severity can be evaluated using a slit lamp or by AS-OCT. The assessment of the severity of CCH with AS-OCT provides more objective information than biomicroscopic examination. The measurement of CCHA and TMA with AS-OCT provides more accurate interpretation about the tear function. Measurement of CCHA and TMA with caliper can be reproduced. Thus, these measurements can be taken as an objective criteria for the follow-up and regulation of treatment of patients with CCH.

In the current study, an evaluation was made with the addition of the Cornea–Anterior Module software on the diagnostic device to measure the area of conjunctival protrusion into the TMA in patients. CCH was found at a rate of 44.4% in a sample aged >65 years, and the CCHA was measured as $0.28 \pm 0.09 \text{mm}^2$ in the nasal region, $0.15 \pm 0.10 \text{mm}^2$ in the central region and $0.17 \pm 0.06 \text{mm}^2$ in the temporal region of group 1. The nasal TMA was $0.043 \pm 0.014 \text{mm}^2$ in group 1 and $0.081 \pm 0.026 \text{mm}^2$ in group 2, and the central TMA was $0.051 \pm 0.009 \text{mm}^2$ in group 1 and $0.089 \pm 0.011 \text{mm}^2$ in group 2, and the difference was statistically significant ($p=0.023$ and $p=0.035$, respectively). Previous studies have shown that the TMA is associated with dry eye. The results of the current study also showed a statistically significant correlation between the TMA and age, OSDI score, and FTBUT ($p=0.046$, 0.025 and 0.022, respectively). However, no correlation was found between the TMA and the Schirmer I test and ocular surface staining ($p=0.113$ and 0.121, respectively) in group 1. Thus, it was established that CCH disrupted the tear distribution but did not affect the amount of tears.

**Limitations**

The major limitation of this study was its small sample size. CCH may be treated with surgical procedures when it is refractory to medical treatment. However, none of the patients were examined after surgery. The tear function of the patients after surgery or medical therapy should be evaluated in further studies.

**Conclusion**

In conclusion, CCH is found at a high rate in the elderly population and it is mostly neglected during
ophthalmologic examination. When elderly patients present with complaints such as burning, itching sensations and epiphora, they should be investigated for CCH, and if diagnosed, tear function tests and AS-OCT evaluation of CCH should be performed. CCH evaluation using AS-OCT is a useful method for determining its effect on the TMH and TMA.

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