High Frequency of Latent Conjunctival C. trachomatis, M. hominis, and U. urealyticum Infections in Young Adults with Dry Eye Disease

Ernest V. Boiko, Alexei L. Pozniak, Dmitrii S. Maltsev, Alexei A. Suetov, and Irina V. Nuralova

Department of Ophthalmology, Military Medical Academy, 5 Klinicheskaya Street, St. Petersburg 194044, Russia

Correspondence should be addressed to Ernest V. Boiko; boiko111@list.ru

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1. Introduction

The Dry Eye WorkShop (DEWS) definition of dry eye disease (DED) emphasizes the role of inflammation in the pathogenesis of this disease [1], which is reflected in the therapeutic strategies that have been used recently to treat DED [2, 3]. Some long-term clinical manifestations of inflammation, as conjunctival hyperemia, edema, and insignificant infiltration, are shared by both DED and chronic conjunctivitis. Between DED and chronic conjunctivitis, any significant diagnostic distinctions that can be revealed without special examination techniques are absent. Furthermore, the connections that have been revealed between some forms of conjunctivitis (in particular, the allergic one) [4] and DED indicate that chronic conjunctivitis may possibly result in DED. However, besides allergy, chronic inflammation of the conjunctiva also may be caused by persistent infection that, thus, leads to the development of DED. C. trachomatis, M. hominis, and U. urealyticum are the most common pathogenic microorganisms capable of persisting in tissues of human body for long time periods and causing not acute but mostly chronic low-grade nonspecific inflammation [5–7].

Because these infectious agents are those of sexually transmitted diseases, they are predominantly found in young adults [8–10]. Moreover, evidences of conjunctival localization with possible development of conjunctivitis have been reported for these pathogens and closely related species [11, 12].

Therefore, the aim of this study was to determine the frequency of detection of conjunctival C. trachomatis (CT), M. hominis (MH), and U. urealyticum (UU) infections in young adults with dry eye disease (DED), since these infections may potentially produce the chronic subclinical inflammation characteristic of DED. Materials and Methods. The study included subjects of 25–45 years of age, divided into the DED (𝑛=114) and nondry eye control (𝑛=98) groups, with the diagnosis based on self-reported complaints, biomicroscopy, the Schirmer I test, and break-up time. All patients had conjunctival scrapings taken to detect CT, MH, and UU with direct fluorescent-antibody assay kits. Results. At least one of the three microorganisms was found in 87.7% of the DED patients versus 8.2% of the controls. Of all the DED patients, 63.2%, 50.8%, and 42.1% were found to be infected with CT, MH, and UU, respectively. Multiple pathogens were identified in 65% of the DED patients found to be infected. CT infection was detected in 6.1% of the controls.

Conclusion. C. trachomatis, M. hominis, and U. urealyticum were detected with high frequency in the conjunctiva of young adults with DED and may be an important risk factor for DED in them.
2.2. Patients. The study included 212 subjects divided into two groups, the DED group (n = 114) and non-dry eye control group (n = 98). The inclusion criteria for DED group were age from 25 to 45 years, complaints of dryness, sensation of sand and/or foreign body sensation in the eye, insignificant conjunctival discharge and tearing (alone or in combinations), a Schirmer I test of 11 mm or less, and tear film breakup time (BUT) of 5 seconds or less. The nondry eye control group included nondry eye subjects of the same age range. Exclusion criteria included acute conjunctivitis, pathological lacrimal passages, contact lens wear, history of refractive surgery, and DED secondary to systemic diseases (Sjogren’s syndrome, Reiter’s syndrome, Stevens-Johnson syndrome, etc.), endocrine diseases, systemic diseases of connective tissue, current administration of antibiotic, anti-inflammatory, cytostatic, or hormonal agents, either locally or systemically, administration of oral contraceptives, and smoking.

2.3. Ophthalmic Examination. All patients underwent complete ophthalmic examinations and had conjunctival scrapings taken for direct fluorescent assay (DFA). Duration of the disease was self-reported by patients; Schirmer’s I and BUT tests were performed to assess the severity of the disease. These are widely used and the most available dry eye diagnostic tests, with the sensitivity and specificity of the Schirmer I test reaching 85% and 100%, respectively, and those of the BUT test reaching 83% and 85%, respectively [1]. Because the DEWS recommends these two tests, along with clinical history, symptom questionnaires, and ocular surface staining grading, as those of the first five in “a practical sequence of tests” [1] for dry eye, they were used to detect DED in this study.

Schirmer’s I test was performed by placing a Schirmer strip in the lateral lower conjunctival sac after instillation of one drop of topical 0.5% proxymetacaine (Alcaine, Alcon-Couvreur, Puurs, Belgium) for 5 minutes. The amount of wetting was measured. To measure tear BUT, after instillation of a drop of sodium fluorescein dye (BioGlo Sterile Fluorescein Strips, HUB pharmaceutical, Rancho Cucamonga, CA), the tear film was observed under cobalt-blue filtered light of the slit lamp biomicroscope, and the interval between the last blink and appearance of the first break in the tear film was noted. Individual average BUT values were calculated from three repeated measurements.

2.4. Sampling. After instillation of one drop of topical 0.5% proxymetacaine (Alcon-Couvreur), each patient had conjunctival epithelial scraping taken from both eyes in a standardized manner, with the samples collected from tarsal conjunctiva and passed firmly four times across the conjunctiva. Then, the material obtained from a conjunctival scraping was spread on a slide and fixed in 70% cold methanol.

2.5. Direct Fluorescent Assay (DFA). The method is based on binding of antibodies to an epitope (a specific trisaccharide component (aKdo-(2-8)-aKdo-(2-4)-aKdo) of cell wall lipopolysaccharide (LPS) for C. trachomatis, a surface protein antigen for M. hominis, and a surface protein antigen for U. urealyticum); currently, DFA tests are the only tests cleared by the Food and Drug Administration for the detection of ocular C. trachomatis infections [15]. Moreover, the DFA is of relatively low cost, easy, rapid, and suitable for routine use. The reported sensitivity and specificity of the DFA varies between 86% and 92% and 96% and 99% [16–18], respectively, in urogenital specimens, and approaches 100% and varies between 96% and 99%, respectively, in conjunctival scrapings [19]. The high rates of sensitivity and specificity of the DFA in the detection of ocular infection are attributed to the relative “purity” of conjunctival scrapings compared to urogenital specimens, and this is why the DFA actually conforms better to the detection of infection in the former than in the latter. For this reason, DFA method was chosen in this study.

The polyclonal antibody based C. trachomatis, M. hominis, and U. urealyticum direct specimen kits, ChlamyScan, MicroScan, and UreaScan (LABDiagnostika, Moscow, Russia), respectively, were used for the detection of proper antigens according to the manufacturer’s instructions. Briefly, conjunctival scrape smears were covered with 30 microliters of Evans blue counterstain containing solution of fluorescein-isothiocyanate- (FITC-) conjugated antibodies for 20 min at 20°C in a dark, humidified chamber. After being washed in phosphate buffer saline (PBS) and twice in distilled water, dried, and coverslipped with 10% glycerin solution in PBS, specimens were examined on Leica DM2500 microscope (Leica Microsystems GmbH, Wetzlar, Germany) (excitation wavelength, 490 nm; mean emission wavelength, 520 nm) equipped for FITC fluorescence. In C. trachomatis diagnostic tests, the positive-control was heteroploid line of L929 mouse fibroblasts (provided with ChlamyScan kit) infected with C. trachomatis strain L2 (Figure 1(d)). In M. hominis and U. urealyticum diagnostic tests, the positive-control (provided with MicroScan and UreaScan kits, resp.) contained suspension of HeLa cell culture separately infected with different strains of M. hominis and U. urealyticum, respectively (Figures 1(e), 1(f)). In C. trachomatis, M. hominis, and U. urealyticum diagnostic tests, the negative control contained conjunctival scrape smears of nondry eye patients and pathogens-free suspension culture of heteroploid L929 mouse fibroblasts (Figures 1(g)–1(l)). Evaluation was performed if the amount of epithelial cells in a scrape sample was at least 50. Loci of specific fluorescence were visualized at a magnification of ×400, with identification confirmed at a magnification of ×1000. The following was considered as a specific pattern: (1) small, well-defined, round, apple-green loci of fluorescence, located intracellularly or extracellularly or (2) large, moderate bright green loci of fluorescence, located intracellularly, corresponding to solitary cells and to intracellular inclusions of the pathogens, respectively (Figures 1(a)–1(c)). This pattern has been described as specific by the manufacturer and presented in some works [20, 21]. A sample was considered positive if at least 10 loci of specific fluorescence were identified, because this criterion has been found to provide an optimal ratio of sensitivity to specificity and used in a number of works [16–18]. If a unicellular infection was found, a patient was considered positive for infection.

2.6. Statistical Analysis. Nonparametric data analysis was performed with Statistica for Windows 6.0 software (Statsoft,
Figure 1: Direct fluorescence assay (DFA) staining for the detection of *C. trachomatis*, *M. hominis*, and *U. urealyticum* infection in conjunctival scrapesmears of a dry eye patient ((a)–(c)), positive control slides ((d)–(f)), conjunctival scrape smears of a nondry eye patient ((g)–(i)), and negative control slides ((j)–(l)). Note the specific DFA staining patterns (small, well defined, round, apple-green or large, moderate bright green loci of fluorescence) in panels (a)–(f) (arrowheads) and absence of specific fluorescence in panels (g)–(l). DFA with Evans blue counterstain, original magnification ×400.
3. Results

3.1. Characteristics of Patients and Results of Ophthalmic Examination. No statistically significant differences were noted between the DED and control groups in demographic characteristics (Table 1). In all patients of the DED group, consistent with DED symptoms (conjunctival hyperemia, complaints of dryness, smarting eyes, burning sensation, and foreign body sensation in the eye), Schirmer’s I and BUT tests showed reduced tear production and destabilization of the tear film, respectively. In all patients of the control group, these characteristics were within normal ranges. These patients had neither complaints nor symptoms related to DED. In the DED group, the mean duration of the disease reported by 90.2% of the patients was 41.16 ± 9.12 months (range 37 to 58 months), with slow increase in the level of symptoms reported over time, whereas that reported by 9.8% of the patients was 22.92 ± 6.60 months (range 12 to 26.4 months).

3.2. DFA Results. At least one of the three microorganisms investigated in this study was found in 100 (87.7%) patients of the DED group versus 8 (8.2%) patients of the control group (Figure 2). Of all infected DED cases, only 35% were found infected with a single agent. Interestingly, of the DED patients infected with at least two pathogens, 86.2% were coinfected with C. trachomatis, which was found to be the most common infectious agent (72% in all infected study patients and 63.2% in the DED group). Besides, C. trachomatis, either alone or in association with other species, was identified in 8 (6.1%) patients of the nondry eye control group. During ophthalmic examination, no signs of chronic conjunctivitis or dry eye were found in the infected controls.

4. Discussion

This study showed that a large share of persons aged 25–45 years, with reduced tear production, destabilization of the tear film, conjunctival hyperemia, and complaints characteristic for DED, have chronic infectious conjunctivitis which might be caused by C. trachomatis, M. hominis, and U. urealyticum infections, either alone or mixed. This is in agreement with the statement that mild conjunctivitis is often
associated with dry-eye patients [22] and suggests that, in
persons of this age group, latent conjunctival infection is
another important risk factor for DED.

The development of DED in young adults without any
apparent risk factors for DED (age, history of refractive
surgery, contact lens wear, systemic diseases or specific
drug therapy, and obvious occupational risks) has no other
possible explanation except for the action of a risk factor
that has not yet been established (e.g., infectious agents). The
complaints and clinical picture do not completely correspond
to the conjunctival inflammation being characteristic for
infectious damage, and this is the very reason why this
chronic conjunctival infection is diagnosed as DED and not
as conjunctivitis. Such cases of latent conjunctival infection
might account for a part of the incidence of DED and require
specific diagnostic and management approaches.

In this study, clinical manifestations of Chlamydia trachomatis-, M. hominis-, and U. urealyticum-induced chronic conjunctival
inflammation were completely masked by DED symptoms and
differed from manifestations of acute conjunctivitis
(acute conjunctivitis was an exclusion criterion for enrollment).
However, what needs to be explained is the fact that
not all the patients found to be infected suffered from DED (in
particular, infected controls had no manifestations of DED).
Two of the most possible causes are (1) early stage of the
disease and (2) genetically determined features accounting
for intensity of the host conjunctival inflammatory response
[23, 24]. Another possible cause is genetic variability in a
pathogen, which is mostly a characteristic of C. trachomatis
[23]. A limitation of the study is that serotyping was not
performed. Thus, we do not know for sure whether association
with DED cases is characteristic of the C. trachomatis species
or of individual serovars within it [23]. In addition, other con-
junctival bacterial microflora that might be of some potential
value as a risk factor for DED was not investigated. Normal
(saprophytic) conjunctival bacterial microflora may include
a number of microorganism species that cause no inflamma-
tion and, therefore, are unlikely to have any value as a risk
factor for DED. Most of the infectious agents being pathogenic
for the conjunctiva, on the other hand, cause either acute or
subacute conjunctivitis with a characteristic clinical picture
that was not observed in DED patients of the study. Because
these microorganism species were unlikely to play a role as
a risk factor for DED in these patients, the control of the
conjunctival bacterial microflora in them was not performed,
and this could be considered a limitation of the study.

The inflammation associated with DED has the potential
to promote conjunctival colonization, although predomin-
antly by nonpathogenic and opportunistic microorganisms.
The occurrence of rather contagious obligate pathogens such as
C. trachomatis suggests that secondary colonization of
already inflamed conjunctiva is not the case but indicates
rather that these pathogens may play a primary role in
the development and maintenance of inflammation; these
issues, however, require further investigation. Association has
already been established between DED and a number of
infectious agents relating to such viral infections as human
T-cell lymphotropic virus, human immunodeficiency virus,
the Epstein-Barr virus, and hepatitis C virus [25]. These
chronic viral infections trigger autoimmune reactions either
initiating or contributing to lacrimal gland dysfunction in
Sjogren's syndrome [25]. In those studies (reviewed by Alves et al. [25]), the subject of discussion has been autoimmune
mechanisms and not the direct conjunctival or lacrimal gland
damage induced by infectious agents. Yet, there is still a
lot to be understood about the association between chronic
conjunctival infections and non-Sjogren's dry eye, with the
latter accounting for the major part of the incidence of DED
[26]. Recently, the connection between DED and Chlamy-
dophila pneumoniae infection in simultaneous clinical signs
of follicular conjunctivitis has been reported, and, in that
case, conjunctival localization of the agent as well as partial
efficacy of etiotropic therapy has been proved [27]. Similar
connection can be observed in infection with C. trachomatis,
with the latter being a known cause of chronic conjunctival
inflammation [28]. The role of C. trachomatis in the patho-
genesis of DED may result from its high prevalence [9] and
potential for persistence and support of chronic inflamma-
tion [5, 29]. These biological features of the infectious agent
play a key role in the pathogenesis of endemic trachoma,
which is caused by serovars A, B, Ba, and C only, whereas it is
conjunctivitis that is caused by widespread serovars (D to K)
of C. trachomatis [28]. In trachoma, C. trachomatis-induced
conjunctival damage is characterized by marked alteration in
the conjunctival tissue, lymphocytic infiltration, and scarring
[24]. The same processes underlie the DED associated with
infection, but in this case they are less active and result in
either a gradual decrease in basal tear production or change
in tear composition (due to the accompanying damage to
accessory lacrimal glands and goblet cells). This explains why,
in most (90.2%) of the patients, the mean duration of the
disease was at least 3 years, with slow increase in the level
of symptoms reported over time. And these are the long-
duration cases of clinically asymptomatic disease showing
no tendency to resolve spontaneously that are attributable
to latent infection. Although persistence of the pathogen has
been shown to be accompanied by changes in its morphology
and epitope expression [30], this evidence is not used to
confirm latent infection in clinical practice. The cellular
morphology of the conjunctiva might also undergo changes
during latent infection; assessing these changes was not the
aim of the study.

Because localization of the infectious agent in only one
of the two eyes of a patient is deemed unlikely, we did
not study the association of unilateral detected infection
with manifestations of DED. Part of the reason for this
unlikelihood is that interpretation of DFA is specific, with
DFA positivity requiring detection of at least a cutoff number
of loci of specific fluorescence and with ensuing false-positive
results (e.g., those for a contralateral eye) [16].

The share of an infectious agent in general prevalence
of DED may vary depending on the prevalence of this
agent in the population. A rather high prevalence of ocular
(conjunctival) C. trachomatis infection in persons aged 25–
45 years may be related to the increased risk for urogenital
infections for this age group [8–10]. Here, the infection can be
transmitted to the conjunctiva by contact or hematogenously
[20]. Our study provides evidence that M. hominis and
U. urealyticum are two other infectious agents associated with chronic conjunctivitis and DED in persons aged 25–45 years. Although Mycoplasmataceae family members are also capable of damaging the conjunctiva, the clinical value of this fact has been unknown [12]. Because M. hominis and U. urealyticum also cause urogenital diseases and are of high prevalence in persons aged 25–45 years [7, 10], they might be one of the causes of low-grade conjunctival inflammation and DED in this age group. In this study, microbial coinfestions were found more frequently (65%) than mono infections, which agrees with frequent detection of these coinfestions in urogenital infections and supports the association of urogenital diseases with chronic conjunctivitis in patients of this age group. Moreover, because the association of chlamydial urogenital infection with chlamydial ocular diseases has been repeatedly reported [31], the association of urogenital diseases caused by these infectious agents with chronic conjunctivitis is also possible, but this issue has not been studied in this work and needs further investigation.

According to the International Dry Eye WorkShop, the disease comprises two subgroups: (1) evaporative dry eye and (2) aqueous-deficient dry eye; nevertheless, the etiopathogenetic subcategory of DED described in this study can be attributed to both of them [1]. Thus, this subcategory can belong to two DEWS classification categories, ocular surface disease (with the latter involving, e.g., allergy) and lacrimal deficiency (due to inflammatory infiltration and the ensuing reduction in basal tear production).

5. Conclusion

Latent C. trachomatis, M. hominis, and U. urealyticum infections are detected with high frequency in the conjunctiva of young adults with DED and may be an important risk factor for this disease in persons aged 25–45 years. This is associated with their potential for long-term damage to the conjunctiva and with high prevalence of these infectious agents among this age group. Therefore, it is deemed appropriate to conduct an examination for latent infections and, possibly, further antimicrobial treatment in some patients with DED.

Disclosure

The authors have no proprietary or financial interest in any aspect of this report.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

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