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Ocular surface inflammation impairs structure and function of meibomian gland[★]

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Abstract

Dysfunction of the meibomian glands alters secreted meibum quantitatively and qualitatively that can lead to damage to the ocular surface epithelium. In response to an unstable tear film cause by meibomian gland dysfunction, ocular surface epithelium is damaged and expresses inflammatory cytokines leading to secondary ocular inflammation. In turn, inflammatory disorders of the palpebral conjunctiva and lid margin may affect the structure and function of meibomian gland. The disorders include allergic conjunctivitis, long-term usage of contact lenses, dermatological diseases that affect conjunctival homeostasis, Stevens-Johnson's syndrome or chemical burning of the ocular surface and lid margin.

Keywords

Meibomian gland; Ocular surface inflammation; Contact lens; Conjunctivitis; Rosacea; Chemical burn

1. Introduction

Meibomian glands locate to the tarsal plates of the upper and lower eyelids and the orifice of the gland is opened at the lid margin. The glands play a critical role in ocular surface homeostasis. Lipid components secreted by the gland function as a barrier to prevent excess evaporation of the water component of the tear fluid. Therefore, dysfunction of the meibomian glands (MGD) leads to an impairment of ocular surface lubrication and wetness (dry eye syndrome) as well as induction of local inflammatory responses in the ocular surface. In turn, Suzuki et al. (2015) proposed a concept of one unit, i.e., the “meibomian

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gland and ocular surface” by encountering patients with meibomitis (an inflammatory form of MGD) of which successful management is essential to the treatment of inflammation in the ocular surface.

The etiology of MGD includes terminal duct obstruction and/or qualitative or quantitative changes in the glandular lipid secretion, impairing ocular surface homeostasis. In turn, we have noticed that primary ocular surface inflammation may also effect the function of the meibomian glands. In the present article we review the abnormalities of the meibomian glands in association with primary ocular surface inflammatory diseases.

2. Conjunctivitis and meibomian gland

It was reported that conjunctivitis affects the structure of meibomian glands as revealed by non-invasive meibography. Distortion of meibomian gland ducts were found to be significantly greater in patients with allergic conjunctivitis (Fig. 1, case 1) or atopic-keratoconjunctivitis (Ibrahim et al., 2012). Conjunctival inflammation could be associated with the meibomian gland abnormalities in eyelids of the patients with vernal keratoconjunctivitis (Wei et al., 2015). Continuous mechanical stress to the tarsal tissue by eye rubbing might be one of the causes (Arita et al., 2010; Ibrahim et al., 2012). Rynerson and Perry proposed a clinical entity named as DEBS (dry eye and blepharitis). Blepharitis caused by bacterial colonization in the lid margin could cause follicular inflammation, then MGD (Rynerson and Perry, 2016). Blepharokeratoconjunctivitis also causes loss of the gland as evaluated by non-invasive meibography (Yin and Gong, 2016). It was reported that treatment of posterior blepharitis and conjunctival inflammation due to MGD were treated with topical 1% azithromycin suppressed expression levels of IL-1 β , IL-8, and MMP-9 mRNA. On the other hand, expression of TGF β 1 increased during treatment and does not decline after drug withdrawal, suggesting this growth factor contribution to the anti-inflammatory activity of azithromycin in MGD (Zhang et al., 2015). Because in other tissue TGF β 1 could be pro-inflammatory, it is to be further investigated if cytokines produced by inflammatory cells in the conjunctiva/sub-conjunctiva directly damage the acini via palpebral conjunctiva.

3. Contact lens wearing

Contact lens wearing impairs the function of meibomian gland. Mechanical stress and/or chronic, sub-clinical inflammation could be proposed as underlying mechanism, although it is to be further investigated detailed pathobiology of conjunctival inflammation associated with long-term contact lens wearing. One of the commonly observed characteristics in contact lens-related meibomian gland pathology was orifice pouting/plugging (Cox et al., 2016). Inflammation in the meibomian glands can be detected as punctate hyperreflective components by using in vivo laser scanning confocal microscopy (Fasanella et al., 2016). This technique of observation showed the presence of gland dropout, duct obstruction, and glandular inflammation in meibomian glands of contact lens wearers (Villani et al., 2011).

Soft contact lens could induce allergic reaction in the ocular surface epithelium or palpebral conjunctiva. This might also be the case in contact lens-related allergic conjunctivitis; the

condition is associated with an increase in meibomian gland distortion (Fig. 1, case 2). The authors conclude that the allergic reaction, rather than contact lens wear, appears to be responsible for abnormal configuration of meibomian gland in patients with contact lens-related allergic conjunctivitis (Arita et al., 2012).

4. Medication-related meibomian gland diseases

Topical instillation of drugs might also cause ocular surface inflammation, leading secondarily to the damage of meibomian gland. For example, obstructive type of meibomian gland disorders were observed in patients treated with topical anti-glaucoma prostaglandin derivatives (Fig. 1, case 3) (Mocan et al., 2016). Although the exact mechanism underlying the condition is to be investigated, it is known that prostaglandin derivatives could damage the ocular surface epithelium resulting in the induction of local inflammation.

Currently, there is no evidence showing systemic medication induced MGD is due to inflammation. However, it was suggested that systemic medication also could damage the gland or impair its secretory function. We reported patients with loss of meibomian glands with superficial punctate epitheliopathy during treatment of cancer with an anticancer TS-1[®] combination capsules of tegafur, gimeracil, and oteracil potassium (Taiho Pharmaceutical Co. Ltd, Japan) (Mizoguchi et al., 2015) (Fig. 2). The possible mechanism underlie TS-1[®]-related MGD might be similar to the obstruction of drainage system or corneal epithelial disorders associated with systemic TS-1, but not via direct toxicity of the drug to the acinar cells. 5-FU leaked to tear fluid might cause local inflammation and obstruction of the orifice of the gland. On the other hand, the influence of inflammatory fibrotic diseases to the structure or function of meibomian glands is to be further examined. Another case is long-standing consequences of isotretinoin use. Without a significant finding in either gland or eyelid margin the secretory function of these glands is impaired, causing dry eye symptoms (Moy et al., 2015).

5. Dermatological disorders and meibomian gland pathology

Rosacea is a skin disorder of which etiology has not yet been uncovered. Abnormality in the innate immune system, dysregulated growth of commensal skin organisms might have roles in the pathogenesis of rosacea (Two et al., 2015). Microbial agents including commensal *Demodex* mites, *Demodex*-associated bacteria, *Staphylococcus epidermidis*, etc, are suggested to be involved in the pathological mechanism of rosacea. Acne rosacea could cause inflammation in the eyelid and ocular surface. It was reported that rosacea is accompanied with significant loss of meibomian glands highly associated with eyelid margin inflammation. (Palamar et al., 2015; Machali ska et al., 2016; Liang et al., 2016). Two *Demodex* species (*folliculorum* and *brevis*) have been identified as causative mites of rosacea. The former one is primarily found in clusters around the root of lashes and lash follicles, whereas the latter is detected in the meibomian (and sebaceous) glands and is much closely related to recurrent chalazia, refractory keratoconjunctivitis or MGD (Cheng et al., 2015; Zhao et al., 2012; Liang et al., 2014). Possible mechanisms of induction of blepharitis and obstruction of meibomian glands include irritation of local tissue inflammation by mite wastes and debris as well as their eggs. However, detailed mechanism of the onset of

blepharitis upon *Demodex* infection is still to be investigated (Czepita et al., 2007). The notion of involvement of rosacea in the health of Meibomian gland is further supported by the fact that rosacea is often associated with chalazion (Chamaillard et al., 2008). We also reported a case of multiple chalazion in a rosacea patient, suggesting the presence of involvement of meibomian gland (Shirai et al., 2015) (Fig. 3).

Psoriasis vulgaris is a skin disease characterized by chronic inflammation and keratinocyte hyperproliferation (hyperkeratinization) presumed to be associated with dysregulation of immune system. Jester et al. suggested that hyperkeratinization of the duct epithelium causes plugging, resultant dilation of the ducts and disturbance of the secretory function of the glands (Jester et al., 1989). Thus, hyperkeratinization in eyelid margin could affect the structure and function of meibomian glands by plugging the duct and impairing meibum secretion in patients with Psoriasis vulgaris (Zengin et al., 1996).

6. Stevens-Johnson syndrome and related diseases

Stevens-Johnson syndrome and its more severe form, toxic epidermal necrolysis are a set of diseases characterized by severe immune-based inflammation in mucosal tissue including conjunctiva. The onset of the disease is triggered by systemic medications or kinds of infection. Ocular surface is severely damaged first by acute inflammation, being followed by a chronic, longstanding complication of conjunctival scarring or conjunctivalization of the corneal surface with severe neovascularization. Detailed examination of the meibomian glands in patients of acute phase of Stevens-Johnson syndrome is not easy due to severe discharge and ocular surface pain. A similar damage to the ocular surface was recognized in patients with chronic graft-versus-host disease (Engel et al., 2015). In the later phase of the chronic stage of the disease, the ocular surface homeostasis is impaired with deficiency of limbal stem cells and dry eye caused by the loss of conjunctival goblet cells (Sotozono et al., 2007). It was reported that lid margin keratinization could lead to lipid tear deficiency (Di Pascuale et al., 2005). Meibography showed the total loss of meibomian glands in chronic, long-standing, cases of Stevens-Johnson syndrome. (Alsuhaibani et al., 2011). However, acute influence of severe conjunctival inflammation on the structure of meibomian glands has not been reported.

As mentioned above, examination of pathology of meibomian glands in patients of acute phase of Stevens- Johnson syndrome seems rare in daily medical service. On the other hand, an animal model of Stevens- Johnson syndrome has not been established, although animal experiments might be helpful in identifying alterations of the glands in the earlier phase of the disease after its onset. Chemical burn in the total ocular surface could mimic the situation of severe inflammation in the both cornea and conjunctiva of Stevens-Johnson syndrome, although the underlying mechanism of the inflammation differs from each other. We therefore employ this model of alkali burn of the ocular surface to investigate the acute influence of the conjunctival inflammation to the meibomian gland structure, although alkali might also directly damage the tissue.

7. Chemical burn in the ocular surface and meibomian gland

Chemical burn in the ocular surface often results in the formation of conjunctivalization of the corneal surface and neovascularization with severe dry eye symptoms (Saika et al., 2008; Bunker et al., 2014; Baradaran-Rafii et al., 2017; Saika et al., 2016). The loss of goblet cells and limbal stem cells are believed to largely contribute to these ocular surface changes. However, chemical burn in the area including eyelid margin could damage the orifice of the meibomian gland leading to obstruction of the orifice and impaired secretion of the meibum from the gland. Moreover, because the gland acini are located just beneath the conjunctiva, alkali exposure may also directly damage the gland or indirectly affect the tissue via induction of local inflammation. Fig. 4 shows a case of alkali burn in the eyelid margin. In this case the glands within the burned area were depleted. However, detailed evaluation of the changes in the meibomian gland following chemical injury needs further investigation.

To address this issue, we developed a model of ocular surface alkali burn in C57BL/6 mice (Mizoguchi, 2015). Severe inflammation in ocular surface including lid margin epithelium is induced by an exposure to alkali. The direct effects of alkali to the tissue and infiltration of inflammatory cells are considered to damage the tissue surrounding the orifice, resulting in the retention of the meibum in the duct. Scanning electron microscopy shows that the orifice appeared plugged with debris-like material (Fig. 5). The dilation of the ducts and partial loss of acini were observed in the upper eyelid at day 20 (Fig. 6A). On the other hand whole gland structure including ducts and acini were lost in the lower eyelid (Fig. 6B). Histology in the samples shown in Fig. 6A and B shows that the ducts were extensively dilated and the acinar cells seemed to be scattered among the dilated ducts (Fig. 6C and D). However, melanin pigments in the tissue in C57BL/6 mice disturbed the histological or immunohistochemical analysis of detailed mechanism of the tissue damage in the eyelid and meibomian glands, and an investigation in Balb c mice was performed with similar results (Mizoguchi et al., 2017). In this study we showed clearly the dilation of the ducts was occupied meibum accumulation. In the lower eyelid apoptosis-related acini destruction predominated with neutrophils infiltration (Mizoguchi et al., 2017). Three-dimensional reconstruction image shows that the unaffected meibomian gland consists of a duct and many acinar portion, and shows the loss of acini and dilation of ductal portion in day 20 specimen following an alkali exposure (Fig. 7) as compared with an uninjured gland (Parfitt et al., 2012).

The reason underlying the difference in damage to the meibomian gland between upper and lower eyelids is not known. One explanation may include differences in the degree of inflammation within the conjunctival/subconjunctival tissue due to the level of exposure to the alkali. Tear meniscus between ocular surface and lower eyelid might retain more alkali solution as compared with the space between upper eyelid and ocular surface. Moreover, tear fluid could wash out the alkali in the upper eyelid. Further study is to be conducted to uncover the reason of the difference of the damage patten between upper and lower eyelids.

8. Concluding remarks

The effect of ocular surface inflammation on meibomian gland function is not well understood. In this review we show that ocular surface and palpebral conjunctival including eyelid margin inflammation is associated with alteration in meibomian gland structure and potentially function. In the clinical setting it is recommended to examine the meibomian glands in patients with ocular surface inflammation and infection during the disease as well as after resolving the condition. Systemic and topical anti-inflammation treatment is to be included in strategies to maintain the function of meibomian glands besides targeting ocular surface inflammation.

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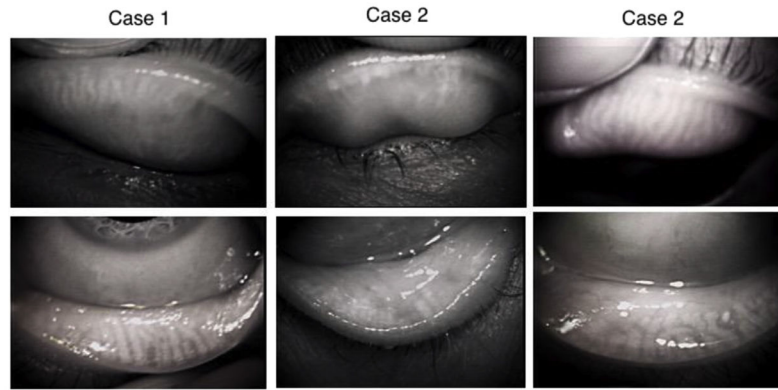


Fig. 1. Morphologically abnormal configuration of meibomian glands in a 35-year-old male patient with chronic allergic conjunctivitis (case 1), a long-term (approximately 20 years) female user (39-year-old) of a soft contact lens (case2) and a 74-year-old male patient with a long-term application of antiglaucoma medications (details unknown) (case 3).

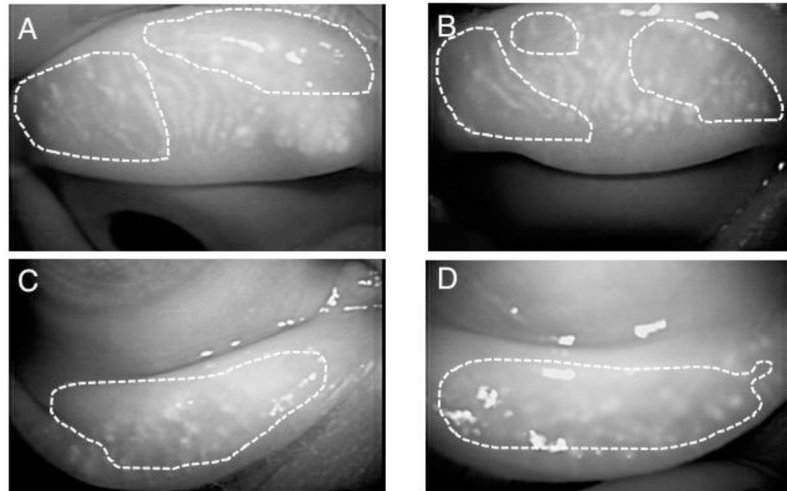


Fig. 2. Infra-red noninvasive meibography finding of a 59-year-old patient with systemic TS-1 anti-cancer treatment for gallbladder cancer. The meibo-score (Arita et al., 2008) are 3 in all of upper and lower eyelids of both eyes. A, B: right; C, D: left; A, C: upper eyelids; B, D: lower eyelids (reproduced from Mizoguchi et al., 2015).

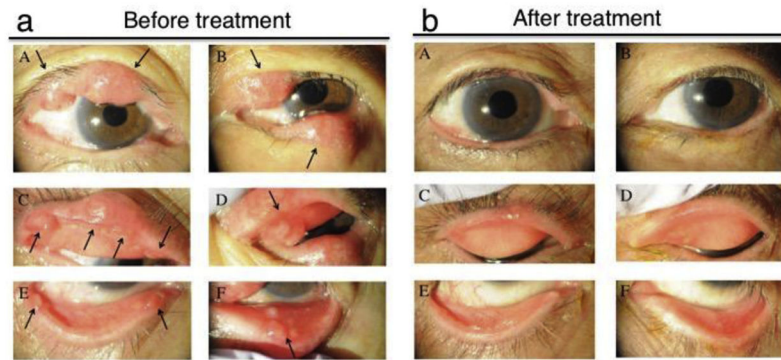


Fig. 3. Multiple chalazion in a patient with acne rosacea. Chalazion are observed in upper and lower eyelids of both eyes of a 79-year-old woman suffering from acne rosacea (a). Black arrows indicate the chalazion. The patient was treated with ophthalmic ointment of erythromycin lactobionate, colistin sodium methanesulfonate and systemic administration of minocycline hydrochloride (b). A, C and E: right eye; B, D and F: left eye (reproduced from Shirai et al., 2015).

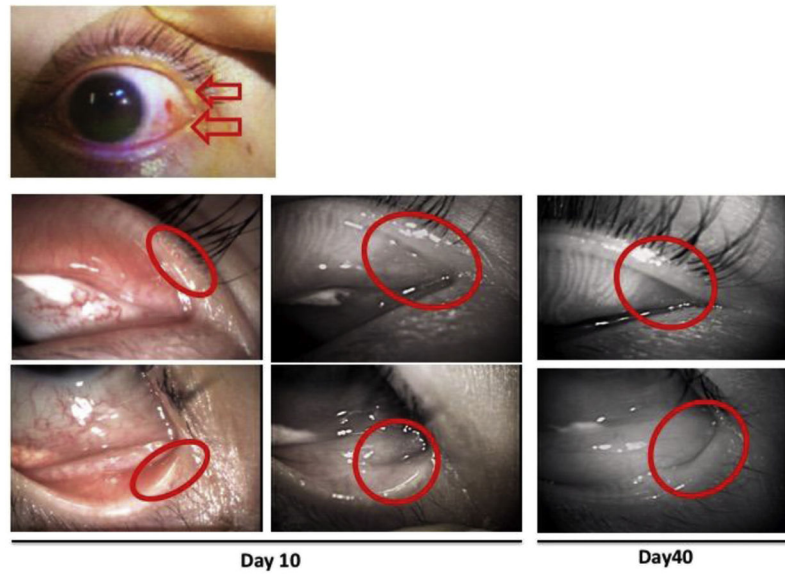


Fig. 4. Alkali burn with an exposure to slaked lime (Calcium hydroxide) of the eyelid of a 33-year-old man. Meibomian glands of which orifices locate to the burned area (red circles) gradually disappeared at day 10 and 40 (reproduced from Mizoguchi, 2015).

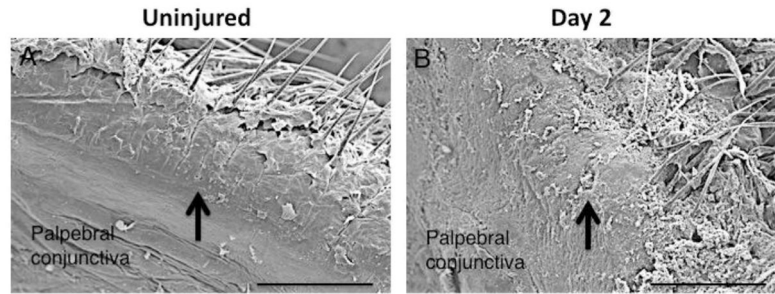


Fig. 5. Scanning electron microscopy shows the orifice of the meibomian gland in the mouse upper eyelid (a). Two days after the alkali burn in ocular surface the orifice is found to be plugged with debris-like material (b).

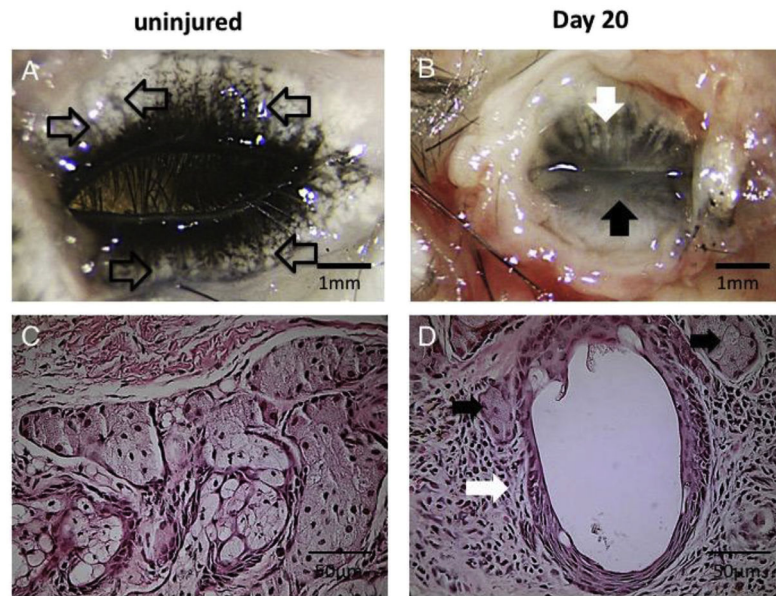


Fig. 6. A. Eyelids of a C57BL/6 mouse. Eyelids were excised and observed from the behind. Acini were observed (open arrows). B. Meibomian glands of a C57BL/6 mouse 20 days after ocular surface alkali burn. Loss of the acini and dilated ducts were observed in the upper eyelid (white arrow). In the lower eyelid meibomian gland was not observed (black arrow). C. Histology of meibomian glands of a C57BL/6 mouse. D. Histopathology of meibomian glands in the upper eyelid of a C57BL/6 mouse 20 days after ocular surface alkali burn. Largely dilated duct (open arrow) of the meibomian glands is observed, acinar cells (black arrows) are also seen among hypercellular mesenchyme (reproduced from Mizoguchi, 2015).

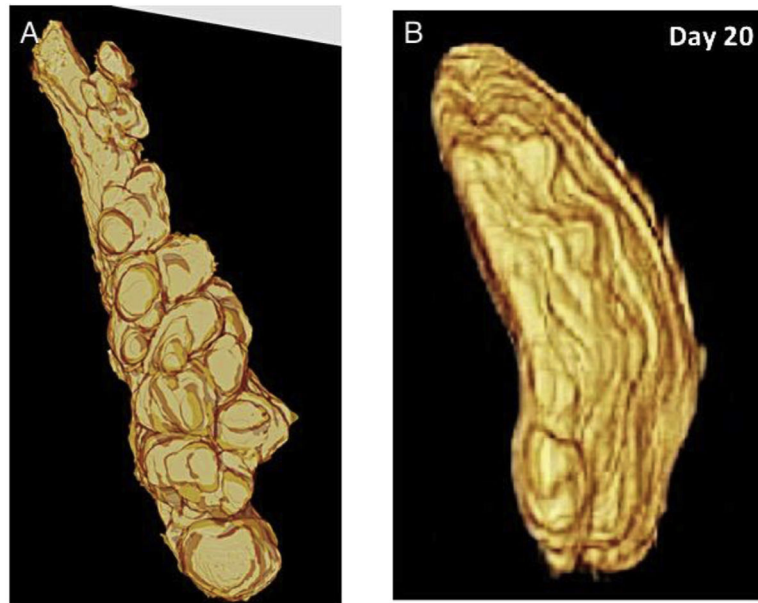


Fig. 7. Three-dimensional reconstruction images of an uninjured meibomian gland (a) and the dilated duct of an eyelid specimen at 20 days after ocular surface alkali burn (b). Marked dilation with decreased number of acini are observed in a day 20 sample (a, reproduced from Parfitt et al., 2012).