Meibomian gland dysfunction: an overlooked eyelid disease

Abstract

Meibomian gland dysfunction is a multifactorial and chronic disease of the eyelids, leading to eye irritation, inflammation, evaporative and aqueous-deficient dry eye and negatively affecting the quality of life. MGD is often overlooked clinically. This review presents a general and practical guide for MGD diagnosis and management.

Keywords: Meibomian gland, dysfunction, dry eye, hypersecretory, hyposecretory, duct obstruction, lipid layer, tear film, ocular surface disease

Introduction

Meibomian glands (MGs) are large sebaceous glands, vertically arranged in the tarsal plates of the upper and lower eyelids and produce the lipids of the outermost layer of the preocular tear film. The tarsal glands are firstly described by Heinrich MEIBOM (1638-1700), a professor of medicine at the university town of Helmsted, and afterward, these glands were called as MGs.1

Definition

The term “Meibomian gland disease” is used to describe various disorders of MGs such as congenital lack, neoplastic and inflammatory disorders, replacement distichiasis and Meibomian gland dysfunction (MGD). The term has been first coined in the ophthalmic literature by Korb and Henriques in 1980s and it has been defined as a chronic MGD resulting in decreased secretion or poor quality of meibum. Lastly, MGD has been defined by subcommittee of The International Workshop on MGD (IWMGD) in 2011 as “a chronic, diffuse abnormality of the MGs, commonly characterized by terminal duct obstruction and/or qualitative/quantitative changes in the glandular secretion which may result in alteration of the tear film, clinical apparent inflammation, OSD and symptoms of eye irritation”.2,4

The importance of the functions of MGs

The functions of Meibomian lipids include providing a smooth optical surface for the cornea, reduction evaporation of the tear film during waking hours, lubrication during blinking, decreasing the surface tension, enhancing the stability and the spreading of the tear film, formation of a barrier to prevent bacteria from entering the tear film, prevention of spillover of tears from the lid margin and the contamination of the tear film by sebum and sealing the opposing lid margins during sleep.4 MGD is one of the most common diseases of MGs. However, it is often overlooked clinically. MGD is a multifactorial and chronic disease of the eyelids, leading to eye irritation, inflammation and ocular surface disease (OSD). Additionally, it is well-documented that MGD is the leading cause of evaporative dry eye and also the most common underlying pathology in the cases with the aqueous-deficient dry eye. Additionally, MGD can negatively affect the quality of life.5–9

Risk factors

The risk factors for MGD identified by the epidemiology and risk factors identification committee of International Meibomian Gland Dysfunction Study group are divided into three group as ophthalmic, systemic and therapeutic (Table 1). Ophthalmic risk factors include aniridia, chronic blepharitis (anterior or posterior), contact lens wearing, Demodex infestation of the eyelid, floppy lid syndrome, giant papillary conjunctivitis, ichthyosis, Salzmann nodular degeneration, and trachoma. In the other hand, systemic risk factors include especially in older age (increasing the amount of meibomian cells expressing estrogen receptors in the lower lid), ocular rosacea, hormonal disorders such as androgen insufficiency, Sjogren’s syndrome, Stevens-Johnson syndrome, atopy, cicatricial pemphigoid and ectodermal dysplasia. The main therapeutic risk factors for MGD are reported as the usage of antiandrogen, antidepressant and antihistaminic drugs, retinoic acid for acne, drugs for benign prostatic hyperplasia and postmenopausal hormone therapeutics.10–13

Symptoms

The cases with MGD may be symptomatic or asymptomatic due to severe involvement of MGs. MGD is associated with various ocular symptoms may not only occur depending on the MGD itself but also on the secondary dry eye and ocular surface damage. The most common symptoms in the admission are fatigue, grittiness, dryness, burning or heavy sensation, stinging and itching in the eyes, redness and swelling on the eyelids and sometimes transient visual blurring.14–17 In case of suspected cases, detailed inquiries of the risk factors for MGD would be useful.

Clinical findings and diagnosis

MGD can present with thickening of the MGs secretions and eyelid margin, turbid secretions, plugged, deformed or cystoid and scarred
MGs orifices, irregularity or deformation or a notch in the lower eyelid margin, anterior or posterior displacement of the mucocutaneous junction, telangiectatic/fine vessels crossing the MGs margins or orifices, vascular dilatation with or without inflammation, increased vascularity at the posterior border of the lid margin, madarosis or trichiasis, frothy or foamy tears, and bubbles in the tear film.2-8 A normal secretion from healthy MGs typically should seem clear and somewhat viscous similar to the olive or baby oil. A turbid or cloudy secretion may be an early sign of MGD while as plugged/obstructed/scarred orifices and notch in the lower lid margin are the signs of advanced MGD. It is considered that the obstruction in orifices and terminal ducts of the MGs is identified as the most prominent aspect of MGD. On the ocular surface, evidence of dry eye and damage can be seen. Applying pressure with the finger on the region of MGs in the middle of the lower or upper eyelid may provide to reveal the absence of excretion of the meibum, to check the emptying of the MG content and to check the quality of the excretion. However, it is recommended the distinguishing evaporative dry eye from aqueous insufficiency due to MGD. For this purpose, initially, OSD indexing (OSDI) scoring should be performed. Then, the evaluation of blinking frequency and the time between two blink (blinking interval), measurements of tear film meniscus height on the lower eyelid conjunctiva, the tears osmolarity, fluorescein break up time or the ratio of break up time to blinking interval (Ocular protection index, N>1), the evaluation of corneal staining with fluorescein and conjunctival staining with lissamine green and aqueous layer with Schirmer test. Following these evaluations revealed the dry eye, MGs functions, meibum excretion and quality with finger pressing and grading of glandular morphology and loss with meibography should be evaluated.2-9,14-17

In the dry eye-related tests, if the tears flow and volume are in the normal limits, an evaporative dry eye is considered, and the presence and severity of MGD can be determined by evaluating the MG morphology and functions. Meibography can reveal acinar atrophy and atrophic degeneration of the MGs. Additionally, non-contact meibography and OCT-meibography have provided important benefits in the diagnosis and follow-up of MGD. Meibometry, interferometry, evaporimetry, fluorophotometry, and meniscometry are other diagnostic tests which were used for etiological and detailed research in the diagnosis of MGD. Meibometry is a test to measure the amount of meibomian lipid at the edge of the steady state lid. Interferometry is a test allowing the visualization of lipid layer using optical principles. Based on the principle of elimination of the speed of disappearance of a dye applied to the ocular surface, fluorophotometry measures the production and volume of tears and the speed of regeneration. Evaporimetry shows the evaporation rate from the ocular surface. Meniscometry provides the objective measurement of height, radius, and volume of the tear film meniscus. The diagnosis of MGD is performed with evaluation of ocular symptoms, amount and quality of MG secretion, and lid margin abnormalities.2-9,14-17

**Stages**

**International Working Group of Meibomian Gland Dysfunction** has graded four stages MGD (Stage 1-Stage 4) to meibomian gland findings, ocular surface findings and the severity of the symptoms (Table 2).1

**Classification and pathophysiology of MGD:** The disturbed functions in MGD may be due to anatomic or secretory abnormalities of MGs and these leads to decreased tear film stability (evidenced by increased evaporation, increased surface tension, contamination with sebum, unsealed lid during sleep) and/or symptoms. MGD may be classified based on the severity, pathophysiologic or anatomic changes of the disease. In the last classification described by the International Meibomian Gland Dysfunction Task Group Identification and Classification Committee, MGD is divided into two groups according to the amount of secretion of the glands.2-9,15

The **low delivery group** includes hyposecretory (Meibomian sicca) and obstructive-type MGD. Hyposecretory type is characterized by decreased lipid secretion without gland obstruction. Primer hyposecretion is usually associated with glandular atrophy. Obstructive MGD may occur in the case of a low secretion level. Histopathologic reaction on obstructive MGD have shown squamous metaplasia and keratinized changes at MG orifices, thickening of meibum and cellular debris within dilated ducts, MG hypertrophy, or atrophy. The reduction in the number of functional MGs is often associated with the wearing duration and use of contact lenses. Obstruction in MGs may be seen in older patients and patients who have previously used retinoid therapy for acne, and also the lack of androgen deficiency or androgen receptor. Obstructive MGD may occur as cicatricial or non-cicatricial types. In cicatricial type, MG ducts and orifices displace mucosa backward while as, in non-cicatricial type, they locate at their normal anatomic positions. The main causes of cicatricial MGD are trachoma, ocular cicatricial pemphigoid, erythema multiforme, and atopic eye diseases. Noncicatricial MGD may occur in seborrheic dermatitis, acne rosacea, Sjogren’s syndrome, atopy and psoriasis.2-9,15 MG obstruction is the most common form of MGD. In initial stages of obstructive MGD, MGs continue to produce meibum. However the meibum cannot excrete due to the obstruction in MG duct or orifices. This cause to increase intraglandular pressure within the MGs and consequently, the dilation of the duct and the acini, acinar degeneration and atrophy which may cause to the loss of meibocytes. Additionally, prolonged obstruction causes to bacterial colonization in MGs and the release of inflammatory mediators and lipolytic enzymes from bacteria. These lipolytic enzymes released by bacteria cause very irritating free fatty acids to breakdown the lipids in the tear film. Eventually, the loss of tear film integrity and stability results in an increased aqueous tear evaporation and evaporative dry eye.2-9,15

High delivery/secretory or seborrhoeic-type MGD (Meibomian seborrhoea) is characterized by a high amount release of meibomian lipid at the lid margin when finger pressure applied on tarsae. It has main histopathologic changes including hypertrophy of the ductal epithelium and keratinization of the orifice epithelium. Hypersecretion of meibum into the tear film triggers an inflammatory reaction at the ocular surface. In primary hyposecretory type, there is no any associated disease. Secondary hyposecretory MGD can occur usually with seborrhoeic dermatitis, and sometimes in atopy and acne rosacea.2-9,15

**Differential diagnosis**

The term “diffuse abnormality of the MGs used by IMGSG in the definition of MGD is critical. Because, in chalazia, a confused disease, the MGs are involved locally and, the involvement does not lead the tear film abnormalities or ocular surface epithelia. In the other hand, some inflammatory entities may be confused with MGD. However, the most common overlap in terminology is between MGD and posterior blepharitis. Meibomitis/meibomianitis is defined as a subset of disorders with inflammation of the MG orifices. Anterior blepharitis is an inflammation of the lid margin anterior to the gray line and concentrated around the lashes. It may be accompanied by squamous
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Aniridia

Antiandrogens

20–22

Older age

20

describes an inflammation of the posterior lid margin, which may have different causes, including MGD, conjunctival inflammation (allergic or infective), and/or other conditions, such as acne rosacea. Marginal blepharitis is inflammation in eyelid margin including both anterior and posterior blepharitis.2–915 Meibomian keratoconjunctivitis has been defined as often associated with anterior blepharitis, with the most prominent changes centered on the meibomian glands. It is usually associated with some form of skin disease and is characterized by tear film instability, ocular surface inflammation, and ocular surface damage. It is an important cause of symptoms in severe chronic blepharitis.16,19 In MGD, inflammation is not essential; the term does not include neoplasia or congenital disease. The role of inflammation in the etiology of MGD is not clearly understood. However, associations between meibomian gland dropout and ocular surface inflammatory diseases, such as chronic blepharitis, giant papillary conjunctivitis, and Sjogren syndrome, histopathologic lipogranulomatous inflammation around the gland lobules and increased vascularization of the posterior lid margin support to be have a role of inflammation in pathophysiology.2–9

Management

The management of MGD is performed based on MGD staging according to the severity of the disease and possible accompanying diseases (Table 2).20–22 The main purpose of the management should improve the symptoms and quality of life of the patient. The cases in Stage 1 should be informed about the chronic nature of the disease and the lifestyle changes, the potential impact of diet and the effect of work/home environments on tear evaporation, and the possible drying effect of certain systemic medications. In this stage, eyelid hygiene including warming/expression/baby shampoo or eyelid wipes and removing the debris or collarettes around the lashes, and inflammation may spill onto the posterior lid margin. Anterior blepharitis may simultaneously associate to MGD and in this case, concurrent treatment should be performed. Posterior blepharitis describes an inflammation of the posterior lid margin, which may have different causes, including MGD, conjunctival inflammation (allergic or infective), and/or other conditions, such as acne rosacea. Marginal blepharitis is inflammation in eyelid margin including both anterior and posterior blepharitis.2–915 Meibomian keratoconjunctivitis has been defined as often associated with anterior blepharitis, with the most prominent changes centered on the meibomian glands. It is usually associated with some form of skin disease and is characterized by tear film instability, ocular surface inflammation, and ocular surface damage. It is an important cause of symptoms in severe chronic blepharitis.16,19 In MGD, inflammation is not essential; the term does not include neoplasia or congenital disease. The role of inflammation in the etiology of MGD is not clearly understood. However, associations between meibomian gland dropout and ocular surface inflammatory diseases, such as chronic blepharitis, giant papillary conjunctivitis, and Sjogren syndrome, histopathologic lipogranulomatous inflammation around the gland lobules and increased vascularization of the posterior lid margin support to be have a role of inflammation in pathophysiology.2–9

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Plus disease is the stage which diseases to the eyelid and ocular surfaces (causative or secondary to MGD or incidental) such as other exacerbated inflammatory OSD, mucosal keratinization, phlyctenular keratitis, trichiasis, cicatricial conjunctivitis, ocular cicatricial pemphigoid, chalazion, anterior blepharitis and Demodex-related anterior blepharitis with cylindrical dandruff associated to MGD. Simultaneously treatment modality to stage and specific treatment of these diseases should be considered with the use of pulsed soft steroid, bandage contact lens/scleral contact lens, topical N-acetylcysteine and cyclosporine A, steroid therapy, epilation, cryotherapy, intraleralional steroid or excision, topical antibiotic or antibiotic/steroid and Tea tree oil scrubs (specifically the terpenin-4-ol component) (Table 2).20–22 Except for above-mentioned treatment modalities, in selected cases, other manual and mechanical but invasive techniques such as lid margin debridement, intraductal meibomian gland probing, MG probing, Intense Pulsed Light and LipiFlow® thermal pulsation system may be tried.20–24

Table 1 Risk factors for MGD

| Risk factors for MGD | Ocular Risk factors | Systemic Risk factors | Therapeutic Risk factors |
|----------------------|--------------------|-----------------------|-------------------------|
|                      | Aniridia           | Older age             | Antiandrogens           |
|                      | Chronic blepharitis (anterior or posterior) | Rosacea               | Antidepressants         |
|                      | Contact lens wearing | Androgen insufficiency | Antihistaminics         |
|                      | Demodex infestation of eyelid | Sjogren's syndrome | Retinoic acid           |
|                      | Floppy lid syndrome | Stevens-Johnson syndrome | Drugs for benign prostate hyperplasia |
|                      | Giant papillary conjunctivitis | Atopy | Postmenopausal hormone therapeutics |
|                      | Ichthyosis          | Cicatricial pemphigoid |                         |
|                      | Salzmann nodular degeneration | Ectodermal dysplasia |                         |
|                      | Trachoma            |                        |                         |

Citation: Turgut B, Çatak O, Demir T. Meibomian gland dysfunction: an overlooked eyelid disease. Adv Ophthalmol Vis Syst. 2018;8(3):168–172. DOI: 10.15406/aovs.2018.08.00295
Table 2: A brief guide for MGD diagnosis and treatment

NOTE: Adapted from the relevant table in "Report of the Subcommittee on Management and Treatment of Meibomian Gland Dysfunction" of The International Workshop on Meibomian Gland Dysfunction.†

Source: https://doi.org/10.1167/iovs.10-6997g

| Stage | Severity of MGD | Ocular discomfort itching or photophobia | Staining on the ocular surface | Management |
|-------|----------------|------------------------------------------|-------------------------------|------------|
| 1     | The minimally altered amount and quality of secretions | None | None | Inform patient about MGD the potential impact of diet and the effect of work/home environments on tear evaporation and the possible drying effect of certain systemic medications. Consider eyelid hygiene including warming/expressing. |
|       |               |              |                              |            |
| 2     | The mildly altered amount and quality of secretions | Minimal | None or limited | Advise patient on improving ambient humidity; optimizing workstations and increasing dietary omega-3 fatty acid intake. Institute eyelid hygiene with eyelid warming (a minimum of four minutes once or twice daily) followed by moderate to firm massage and expression of MG secretions. All the above plus: Artificial lubricants (for frequent use nonpreserved Preferred). Topical emollient lubricant or liposomal spray. Topical azithromycin. Consider oral tetracycline derivatives. |
|       | Minimal to mild MGD clinical signs |                              |                              |            |
|       | Scattered lid margin features |                              |                              |            |
| 3     | The moderately altered amount and quality of secretions | Moderate | Mild to moderate conjunctival and peripheral corneal staining often inferior | All the above plus: Oral tetracycline derivatives. Lubricant ointment at bedtime. Anti-inflammatory therapy for dry eye as indicated. |
|       | Moderate MGD clinical signs | Moderate |                        |            |
|       | Increased lid margin features: plugging vascularity |                         |                              |            |
| 4     | Marked/severely altered amount and quality of secretions symptoms of ocular discomfort itching or photophobia with definite limitations of activities | Marked with definite limitations of activities | Moderate inflammatory signs such as conjunctival hyperemia and phlyctenules | All the above plus: Anti-inflammatory therapy for dry eye. |
|       | Severe MGD clinical signs |                              |                              |            |
|       | Increased lid margin features: dropout displacement |                              |                              |            |
|       | Plus |                              |                              |            |
|       | Associated Diseases to the eyelid and ocular surface (causative or secondary to MGD or incidental) |                              |                              |            |
|       | 1. Exacerbated inflammatory ocular surface disease |                              |                              |            |
|       | 2. Mucosal keratinization |                              |                              |            |
|       | 3. Phlyctenular keratitis |                              |                              |            |
|       | 4. Trichiasis (cicatricial conjunctivitis ocular cicatricial pemphigoid) |                              |                              |            |
|       | 5. Chalazion |                              |                              |            |
|       | 6. Anterior blepharitis |                              |                              |            |
|       | 7. Demodex-related anterior blepharitis with cylindrical dandruff |                              |                              |            |

Acknowledgments
None.

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

Funding
The authors received no financial support for the research, authorship, and/or publication of this article.

Authorship contributions
Concept, design and data collection: Burak Turgut; literature Search: Onur Çatak, Tamer Demir; writing, analysis, interpretation:
Burak Turgut, Onur Çatak, Tamer Demir.

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Citation: Turgut B, Çatak O, Demir T. Meibomian gland dysfunction: an overlooked eyelid disease. Adv Ophthalmol Vis Syst. 2018;8(3):168–172. DOI: 10.15406/aovs.2018.08.00295