

XDEMVIY® gives you Might Over Mites™

XDEMVIY is here



XDEMVIY is the first and only FDA-approved treatment for patients troubled with *Demodex* blepharitis, an eyelid disease estimated to impact approximately 25 million eye care patients in the United States.¹⁻³

We invite you to join us and a select group of your peers for one of our *DemodeXperience* discussions! Your host will be Andrew Morgenstern, OD, who will discuss the data and clinical results behind XDEMVIY.

INDICATIONS AND USAGE

XDEMVIY (lotilaner ophthalmic solution) 0.25% is indicated for the treatment of *Demodex* blepharitis.

IMPORTANT SAFETY INFORMATION:

WARNINGS AND PRECAUTIONS

Risk of Contamination: Do not allow the tip of the dispensing container to contact the eye, surrounding structures, fingers, or any other surface in order to minimize contamination of the solution. Serious damage to the eye and subsequent loss of vision may result from using contaminated solutions.

Please see additional Important Safety Information on next page.

DEMODEXPERIENCE
Demodex blepharitis Dialogues

XDEMVIY and the *DemodeXperience*: *Demodex* Blepharitis Dialogues

Live Virtual Webconference
Hosted by Tarsus Pharma
HQ: Irvine, CA 92618
Program ID: 890
Date: Tuesday, July 22, 2025
Time: 8:00 PM ET

Presented by Key Thought Leader:



Andrew Morgenstern, OD
Optometrist
Walter Reed National Military
Medical Center
Bethesda, MD 92618

**Join us at the *DemodeXperience* to
elevate your expertise!**

**Click [HERE](#) to register for this
virtual event.**

IMPORTANT SAFETY INFORMATION (cont'd)

Use with Contact Lenses: XDEMVY contains potassium sorbate, which may discolor soft contact lenses. Contact lenses should be removed prior to instillation of XDEMVY and may be reinserted 15 minutes following its administration.

ADVERSE REACTIONS: The most common adverse reaction with XDEMVY was instillation site stinging and burning which was reported in 10% of patients. Other ocular adverse reactions reported in less than 2% of patients were chalazion/hordeolum and punctate keratitis.

To report SUSPECTED ADVERSE REACTIONS, contact Tarsus Pharmaceuticals, Inc. at 1-888-421-4002 or the FDA at 1-800-FDA-1088 (www.fda.gov/medwatch).

Please see full Prescribing Information.

Available by prescription only.

For additional information, please see www.xdemvyhcp.com.

References: **1.** XDEMVY [prescribing information]. Tarsus Pharmaceuticals, Inc; 2023. **2.** O'Dell L, Dierker DS, Devries DK, et al. Psychosocial impact of *Demodex* blepharitis. *Clin Ophthalmol*. 2022;16:2979-2987. **3.** Trattler W, Karpecki P, Rapoport Y, et al. The prevalence of *Demodex* blepharitis in US eye care clinic patients as determined by collarettes: a pathognomonic sign. *Clin Ophthalmol*. 2022;16:1153-1164.



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Our hope is that this *DemodeXperience* will provide the information you need to begin important conversations with your patients about XDEMVY and its compelling approach to treating *Demodex* blepharitis.

If you have any questions or would like additional details about this meeting, please contact me at the information provided below.

We look forward to seeing you there!

Sincerely,

Tarsus Representative:

Cathy Paulson

Brand

Cell: (414) 704-8505

Email: cpaulson@tarsusrx.com