

Tarsus

The Future of Eye Care Begins Here

January 2025

Forward-Looking Statements

Statements in this presentation about future expectations, plans and prospects, as well as any other statements regarding matters that are not historical facts, may constitute "forward-looking statements." These include statements regarding the potential commercial success and growth of XDEMVY in Demodex blepharitis, including market size, acceptance, demand, prescription fill rate and adoption rate for XDEMVY; our ability to successfully implement our sales force expansion and new direct-to-consumer campaign including network TV; our ability to achieve distribution and patient access for XDEMVY; our ability to continue to educate the market about Demodex blepharitis; our ability to continue to drive a successful launch of XDEMVY and become an eye care leader; anticipated regulatory and development milestones including potential regulatory pathways for approval of XDEMVY in Europe, China, and Japan; the market size and opportunity for our pipeline products including TP-04 for the potential treatment of Ocular Rosacea and TP-05 for the potential prevention of Lyme disease; the timing of initiation and results of our clinical studies including additional studies on the impact of XDEMVY, a Japan DB prevalence study, TP-04, and TP-05; the potential regulatory pathways and timing of discussions with regulators including the FDA; the impact of our new sales force representatives on XDEMVY sales; our ability to continue investing in our business, and the guotations of Tarsus' management. The words, without limitation, "believe," "contemplate," "continue," "could," "estimate," "expect," "intend," "may," "might," "plan," "potential," "predict," "project," "should," "target," "will," or "would," or the negative of these terms or other similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these or similar identifying words. Further, there are other risks and uncertainties that could cause actual results to differ from those set forth in the forward-looking statements and they are detailed from time to time in the reports Tarsus files with the Securities and Exchange Commission, including Tarsus' Form 10-K for the year ended December 31, 2023 filed on February 27, 2024 and the most recent Form 10-Q guarterly filing filed with the SEC filed on November 13, 2024, which Tarsus incorporates by reference into this press release, copies of which are posted on its website and are available from Tarsus without charge. However, new risk factors and uncertainties may emerge from time to time, and it is not possible to predict all risk factors and uncertainties. Accordingly, readers are cautioned not to place undue reliance on these forward-looking statements. Any forward-looking statements contained in this presentation are based on the current expectations of Tarsus' management team and speak only as of the date hereof, and Tarsus specifically disclaims any obligation to update any forward-looking statement, whether as a result of new information, future events or otherwise.

Expert Leadership Team with Broad Range of Eye Care and Biotech Experience

Bobby Azamian, MD, PhD CEO & Chairman



Jeff Farrow Chief Financial & Strategy Officer

Aziz Mottiwala Chief Commercial Officer



Sesha Neervannan, PhD Chief Operating Officer



Bryan Wahl, MD, JD General Counsel



Dianne Whitfield Chief Human Resources Officer



Elizabeth Yeu, MD Chief Medical Officer



1. Year to date through Q3 2024. 2. Source on file

Tarsus The Next Leader in Eye Care

Proven Blueprint For Creating New Categories

- **Expert leadership team** with an expansive range of eye care and biotech experience
- Unwavering focus on 4 key principles:
 - Evidence generation
 Education
 - Ease of access
 Execution

XDEMVY® Accelerating Toward Potential Blockbuster Status

- The first and only FDA-approved therapy for *Demodex* blepharitis (DB)
- >104K bottles delivered and >\$113M in net product sales¹

Ocular Rosacea A New Potentially Transformational Opportunity

- Another impactful eye disease impacting ~15-18M Americans²
- Pioneering the potential standard of care in a new category with TP-04





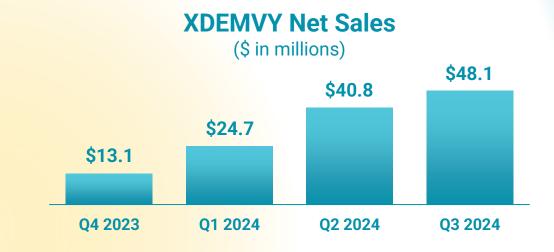
XDEMVY®

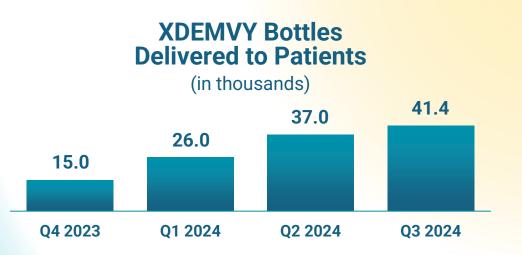
Accelerating Toward Potential Blockbuster Status

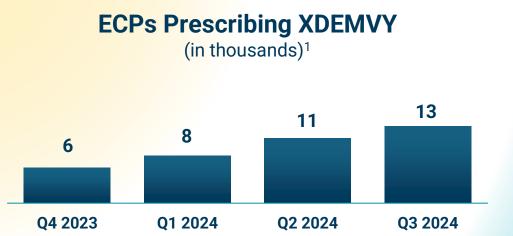
The Tarsus Opportunity is Real

XDEMVY

On-track to potentially be one of the best launches in eye care







1.ECPs prescribing in each listed quarter are cumulative launch-to-date numbers announced at the respective quarterly earnings dates, and as of February 23, 2024 (Q4 2023); as of May 3, 2024 (Q1 2024); as of August 7, 2024 (Q2 2024); and as of November 13, 2024 (Q3 2024)

XDEMVY Launch YTD Q3 2024



Driving One of the Fastest Growing Categories in Eye Care



In Just One Year, XDEMVY has Become One of the Best Launches in Eye Care

Abby, an XDEMVY patient. ECPs = Eye Care Providers

Demodex Blepharitis (DB): A Pervasive and Damaging Eyelid Disease





The result of an infestation of *Demodex* mites, DB can cause eyelid inflammation, redness and irritation

Easily diagnosed during a routine eye exam through the identification of collarettes Potential for serious clinical implications if left untreated

(pictured: corneal opacity^{3,4})

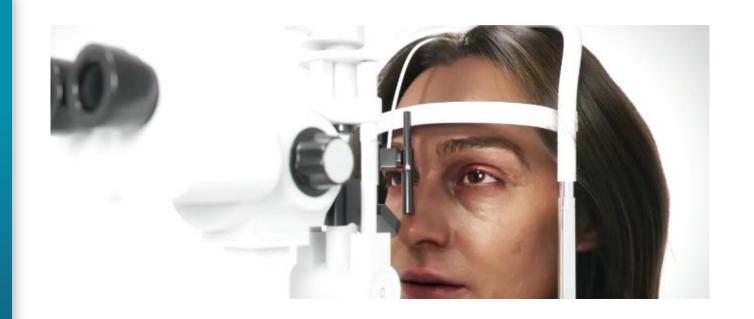
~25M Americans Impacted^{1,2}

1. Wilson J Ophthalmology 2015, 435606, 2014; 2. Titan collarette prevalence study. 3. Liu J et al. Curr Opin Allergy Clin Immunol. 2010;10(5):505-510. 4. Cheng AM et al. Curr Opin Ophthalmol. 2015;26(4):295-300.

Easily Diagnosed Through the Presence of Collarettes

100%

Of patients with collarettes have *Demodex* blepharitis¹

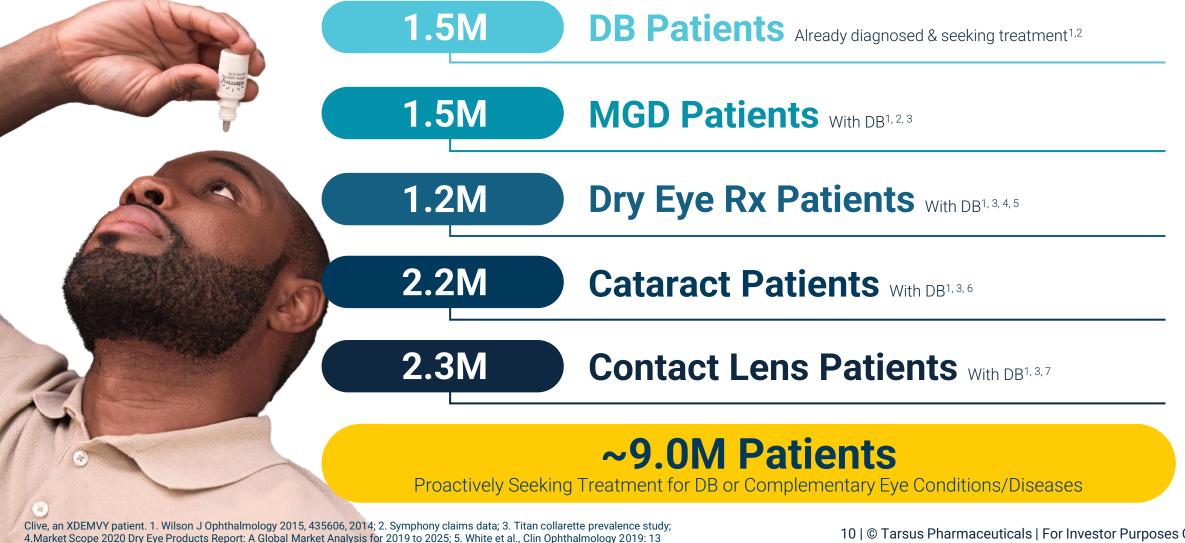


Watch Video

Demodex Blepharitis

A Multi-Billion-Dollar Potential Opportunity

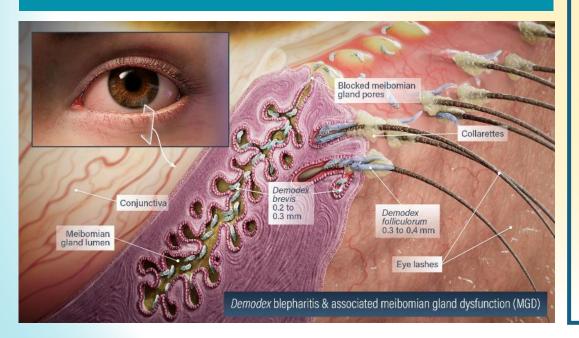




2285-2292 6. AAO/ASCRS Statement on Cataract Surgery, July 2021; 7. Refractive Surgery Council August 2021

XDEMVY: First Pharmacologic Treatment To Demonstrate Groundbreaking Improvements in MGD and Patient Symptoms in DB Patients

Combined data¹ in patients with DB and MGD demonstrated **statistically significant** and clinically meaningful **improvements in objective measures** and patient symptoms



Improvement in Objective Measures of Meibomian Gland Disease (MGD)

- The presence and quality of liquid secretion as measured by the Meibomian Gland Secretion Score
- The number of glands secreting normal (clear) liquid
- The number of glands yielding any liquid

Improvements in Patient Symptoms – Look, Feel, and See

- Fluctuating Vision
- Itching
- Redness
- Burning

1.Combined analysis of two separate pilot studies, ERSA and RHEA, after establishing between-group baseline equivalencies; Tarsus data on file; individual patient outcomes may vary.



XDEMVY: Delivering for Patients



Patient outcomes and experiences may very

12 | © Tarsus Pharmaceuticals | For Investor Purposes Only

Fully Deployed Sales Force Propelling Continued Growth



Driving Depth of Adoption

Continued Evidence Generation Seminal MGD data and

additional DB evidence on the horizon

Ease of Access

>80% of Commercial and Medicare covered lives

Optimized Sales Force

Deepening utilization across all DB patient segments

Making XDEMVY a Household Name Direct-To-Consumer Campaign Driving Patients to ECPs







The Next Impactful Area of Unmet Need Ocular Rosacea



Demodex Blepharitis (DB)

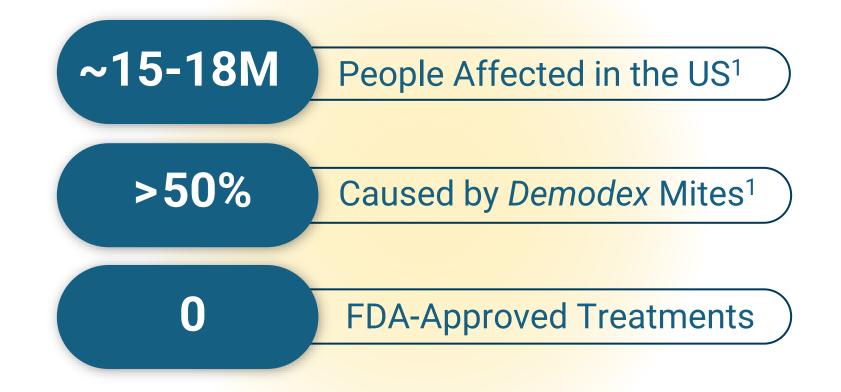
- Caused by *Demodex* mites
- Eyelid inflammation, redness, irritation, collarettes
- Quick and simple slit lamp diagnosis
- Pervasive and damaging eyelid disease

Ocular Rosacea (OR)

- Caused by *Demodex* mites
- Eye and eyelid inflammation and redness, prominent and visible blood vessels
- Quick and simple slit lamp diagnosis
- Pervasive and damaging periocular disease

Ocular Rosacea

Another Clear, Large and Underserved Patient Population



Opportunity to Leverage Our Proven Blueprint

TP-04 Potential to Pioneer the Standard of Care for Ocular Rosacea



Targets Root Cause of Disease

Demodex mites

Uniquely Tailored Sterile Ophthalmic Gel

Specifically designed to be applied to the eyelid and surrounding tissue

Best-in-Class Molecule

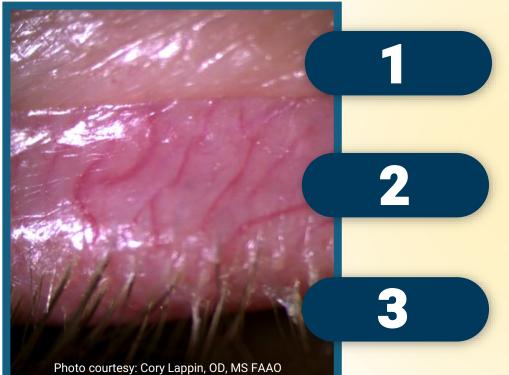
Lotilaner - poised to deliver another transformative eye care therapeutic

Strong IP

Patent exclusivity expected through at least 2038

TP-04

Clear FDA Guidance and a Defined Path Forward



Plans to initiate Phase 2 in 2H 2025 with focus on objective measures of disease – reduction in visible blood vessels and redness

Proof-of-concept results expected next year

Regulatory path that leverages blueprint for success and proven commercial strategy

Upcoming Potential Value Driving Catalysts

TP-04 for Ocular Rosacea: Plans to initiate Ph 2 study in 2H 2025

U.S.

TP-05 for Lyme Disease Prevention: Update planned on 2024 Q4/YE call **XDEMVY for DB:** On-track for potential regulatory approval in 2027

Europe

TP-03 for DB: Initial results from a prevalence study anticipated in 2H 2025

Japan

Greater China*

TP-03 for DB: New Drug Application accepted in China; potential approval anticipated in 2027

* Submitted by Tarsus's partner Grand Pharma

Lyme Disease – A Growing Public Health Crisis No FDA-Approved Prophylaxis

Lyme Disease

A tick-borne infection caused by the transmission of *Borrelia burgdorferi*



Americans at high-tomoderate infection risk¹ \$1.3B

Impact to U.S. healthcare system²



1. CDC Estimate and Corsica Market Research. 2. Adrion E, et al, PLoS One, Feb 2015, Vol. 10(2):e0116767.

TP-05: Potential to Be the First and Only Durable, On-Demand Oral Prophylaxis for Lyme Disease

TP-05

ONGOING Carpo Phase 2a Study

To inform: Safety • Pharmacokinetics • Tick-kill efficacy

Prevention is Key:

Strong physician/patient interest in a non-vaccine option that targets the tick – preventing exposure to the bacteria that causes Lyme Disease

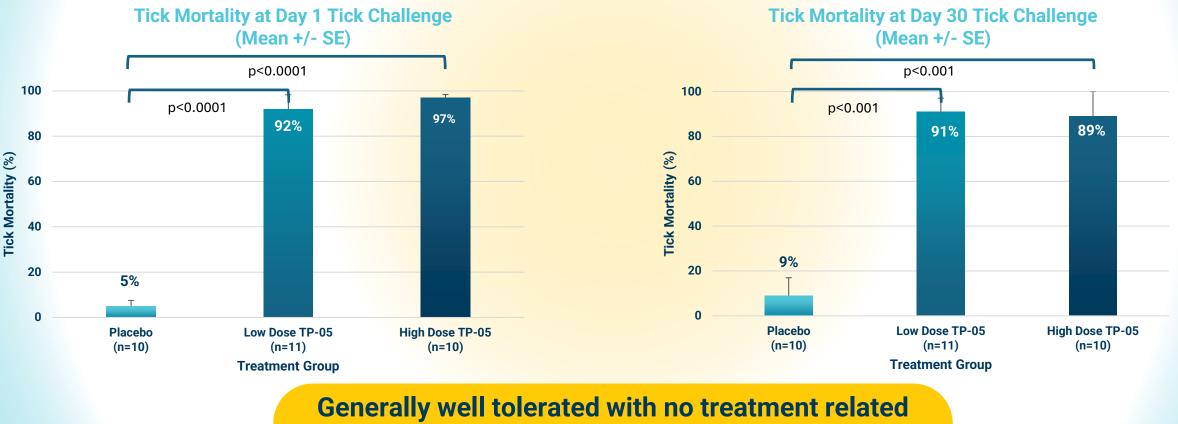
Patient Impact:

Difficult to manage; long-term sequelae can progress to severe joint, CNS and cardiac complications

TP-05:

Fast- and long-acting, with the potential to provide protection throughout the entire tick season

Carpo Phase 2a Trial for Lyme Disease Prevention TP-05: Statistically Significant Tick Mortality Observed at Day 1 and Day 30 Compared to Placebo



discontinuations or serious adverse events



Thank You

Sulma, an XDEMVY® Patient