

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): **April 14, 2026**

PELTHOS THERAPEUTICS INC.
(Exact name of registrant as specified in its charter)

Nevada (State or other jurisdiction of incorporation)	001-41964 (Commission File Number)	86-3335449 (IRS Employer Identification No.)
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4020 Stirrup Creek Drive, Suite 110 Durham, NC (Address of registrant's principal executive office)	27703 (Zip code)
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Registrant's telephone number, including area code: **(919) 908-2400**

(Former name or former address, if changed since last report.)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, par value \$0.0001 per share	PTHS	The NYSE American LLC

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company x

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. o

Item 7.01. Regulation FD Disclosure.

On April 14, 2026, Pelthos Therapeutics Inc. (the “Company”) issued a press release announcing that its Chief Executive Officer, Scott Plesha, will participate in one-on-one investor meetings at the Piper Sandler Spring Biopharma Symposium on April 16, 2026 in Boston, Massachusetts. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

On April 14, 2026, the Company also made available a presentation on its website. A copy of the presentation is attached hereto as Exhibit 99.2. Information contained on the Company’s website is not incorporated by reference into and should not be considered to be part of this Current Report on Form 8-K.

The information contained in this Item 7.01, including Exhibit 99.1 and Exhibit 99.2 attached hereto, which is incorporated into this Item 7.01 by reference, is being furnished and shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended (the “Securities Act”), or the Exchange Act, except as expressly set forth by specific reference in such filing. The information set forth in this Item 7.01 and Exhibit 99.1 and Exhibit 99.2 attached hereto shall not be deemed an admission as to the materiality of any information in this Current Report on Form 8-K that is required to be disclosed solely to satisfy the requirements of Regulation FD.

Forward-Looking Statements

Exhibit 99.2 attached hereto contains, and may indicate, forward-looking statements within the meaning of Section 27A of the Securities Act, Section 21E of the Exchange Act and as defined in the U.S. Private Securities Litigation Reform Act of 1995. Forward-looking statements include, but are not limited to, statements that express the Company’s intentions, beliefs, expectations, strategies, predictions or any other statements related to the Company’s future activities, or future events or conditions, including without limitation, those statements relating to the success of its products and product candidates, timing, progress and results of any preclinical and clinical trials, its estimates regarding the potential market opportunity for its products and product candidates, its ability to develop its pipeline, its ability to protect its intellectual property and enforce its intellectual property rights, its ability to procure new customers and partners, and its ability to execute its development strategy and sustain its competitive position. Actual future results and trends may differ materially depending on a variety of factors, including, but not limited to, the Company’s limited operating history, its ability to establish its market development capabilities to commercialize its products and generate any revenue, its ability to secure and execute financing transactions, and its ability to maintain regulatory approval of certain of its products, which can be identified by terminology such as “anticipate,” “believe,” “continue,” “could,” “estimate,” “expect,” “intend,” “may,” “plan,” “potential,” “predict,” “project,” “seek,” “should,” “target,” “will,” “would” and other similar expressions intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words. These statements are not historical facts and are based on current expectations, estimates and projections about the Company’s business based, in part, on assumptions made by its management. These statements are not guarantees of future performance and involve risks, uncertainties and assumptions that are difficult to predict, many of which are beyond the Company’s control. Any forward-looking statements speak only as of the date on which they are made, and the Company undertakes no obligation to update any forward-looking statement to reflect events or circumstances after the date of this Form 8-K, except as required by applicable law.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits:

<u>Exhibit No.</u>	<u>Description</u>
99.1	Press Release dated April 14, 2026
99.2	Company Presentation
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

Date: April 14, 2026

Pelthos Therapeutics Inc.

By: /s/ John M. Gay

Name: John M. Gay

Title: Chief Financial Officer



Pelthos Therapeutics to Participate in Piper Sandler Spring Biopharma Symposium

DURHAM, N.C., April 14, 2026 — Pelthos Therapeutics Inc. (NYSE American: PTHS), a biopharmaceutical company committed to commercializing innovative therapeutic products for unmet patient needs (“Pelthos”), today announced that CEO Scott Plesha will participate in one-on-one investor meetings at the Piper Sandler Spring Biopharma Symposium on April 16, 2026 in Boston. Investors interested in arranging one-on-one meetings should contact their Piper Sandler representative.

About Pelthos Therapeutics

Pelthos Therapeutics is a commercial-stage biopharmaceutical company focused on building and advancing a portfolio of differentiated cutaneous infectious disease products that address unmet patient needs. ZELSUVMI™ (berdazimer) topical gel, 10.3%, the company’s lead product, is the first and only prescription therapy approved for use at home by patients, parents, and caregivers to treat *Molluscum contagiosum*. The company’s portfolio of assets includes Xepi® (ozenoxacin) Cream, 1%, a topical treatment for impetigo, and Xeglyze® (abametapir), a topical treatment for head lice. More information is available at www.pelthos.com. Follow Pelthos on [LinkedIn](#) and [X](#).

Contacts

Investors:

LifeSci Advisors, LLC

Mike Moyer, Managing Director

mmoyer@lifesciadvisors.com

Media:

KWM Communications

Kellie Walsh

pelthos@kwmcommunications.com

(914) 315-6072



Corporate Presentation

APRIL 2026

Legal Disclaimer

This presentation of Pelthos Therapeutics Inc. (“we”, “us”, “our” or the “Company”) contains “forward-looking statements” within the meaning of the Private Securities Litigation Reform Act and other securities laws. Words such as “anticipate,” “believe,” “continue,” “could,” “estimate,” “expect,” “intend,” “may,” “plan,” “potential,” “predict,” “project,” “seek,” “should,” “target,” “will,” “would” or similar expressions and the negatives of those term are intended to identify forward-looking statements. Forward-looking statements reflect management’s current expectations, are based on judgments and assumptions, are inherently uncertain and are subject to risks, uncertainties and other factors, which could cause the Company’s actual results, performance or achievements to differ materially from expected future results, performance or achievements expressed or implied in those forward-looking statements. Examples of these forward-looking statements and the related risks, uncertainties and other factors include, but are not limited to, the following: the success of the launch of the products in our portfolio, timing, progress and results of any preclinical and clinical trials, its estimates regarding the potential market opportunity for our products, its ability to develop its pipeline, its ability to protect its intellectual property and enforce its intellectual property rights, and its ability to execute its development strategy and sustain its competitive position. Actual future results and trends may differ materially depending on a variety of factors, including, but not limited to, the Company’s limited operating history, the Company’s ability to establish its market development capabilities to commercialize its products and generate any revenue, and the Company’s ability to maintain regulatory approvals of products in our portfolio.

Forward-looking statements are provided to allow potential investors the opportunity to understand management’s beliefs and opinions in respect of the future so that they may use such beliefs and opinions as one factor in evaluating an investment. These statements are not guarantees of future performance and undue reliance should not be placed on them. Any forward-looking statement in this presentation, in any related presentation supplement and in any related free writing presentation reflects our current view with respect to future events and is subject to these and other risks, uncertainties and assumptions relating to our business, results of operations, industry and future growth. You should read this presentation with the understanding that our actual future results may be materially different from any future results expressed or implied by these forward-looking statements. Except as required by law, we assume no obligation to update or revise these forward-looking statements for any reason, even if new information becomes available in the future.

Investment Highlight

- ✓ Commercial biopharmaceutical company focused on growing, differentiated cutaneous infections product portfolio
- ✓ Highly synergistic Xepi and Xeglyze product acquisitions leverage Zelsuvmi's current commercial and market access team and infrastructure
- ✓ Strong potential revenue streams with very attractive gross to nets
- ✓ Disciplined, accretive, cost-efficient product acquisition model and experienced management team to manage execution

Product Portfolio



- Large addressable market with \$2,008.50 wholesale acquisition cost ("WAC")
- **Launched July 2025**



- Modest acquisition cost, unencumbered future revenue stream
- **Expected launch in late 2026**



- 6-12 million U.S. cases annually
- **Expected launch during the first half of 2027**

Corporate Profile

Pelthos is a competitive drug portfolio company — committed to commercializing innovative, safe, and efficacious therapeutic products to help patients with unmet dermatological treatment burdens

Zelsuvmi: Launched in July 2025.

- First and only at home treatment addressing *Molluscum contagiosum* (“MC”), a large, underserved market treating contagious viral disease

Recent portfolio acquisitions: Two FDA-approved complimentary dermatological acquisitions, will leverage Zelsuvmi commercial infrastructure buildout

- **Xepi** (ozenoxacin) Cream 1% - novel topical treatment for impetigo
 - First line impetigo treatment addresses antimicrobial resistance in pediatric dermatology, drug relaunch expected in late 2026
- **Xeglyze** (abametapir) Lotion 0.74% - novel topical treatment for head lice
 - Commercial launch expected in 2027

Experienced management team: Over 20 successful prior drug launches, including Cosentyx, Otezla, Ohtuvayre, Xifaxan

Current peak Net Revenue forecast of \$175M per annum based on Zelsuvmi alone by 2028

Key Data Points (as of 03/16/26, except where noted)	
Ticker	PTHS
Stock Price	\$24.00
O/S Shares of Common Stock (on an as converted basis)	8.9M
Market CAP	~\$215M
Avg. Daily Trading Volume	11,900 shares (Yahoo)
Cash at end of 2025	\$18.0M (not including \$30.0M in term notes issued in Jan. 2026)
Investment to date	>\$400M

Management Team



Scott Plesha | Chief Executive Officer

- >30 years of experience in the pharmaceutical industry, including two decades building and leading specialty pharma commercial efforts
- President and Chief Commercial Officer at BDSI until it was acquired by Collegium Pharmaceutical in 2022
- Grew BDSI sales from \$5 million to \$160 million
- Previously served as Senior Vice President of Gastrointestinal Sales at Salix Pharmaceuticals. During fifteen-year tenure at Salix, led nationwide salesforce that grew product sales to more than \$1.5 billion annually
- Earned a BA in Pre-Medicine and Pre-Medical Studies at DePauw University and pursued graduate studies in Dentistry at Indiana University Dental School



John M. Gay | Chief Financial Officer

- Over 25 years of experience in public company finance and accounting for life science and technology companies, including R&D and commercial organizations
- Raised nearly \$300 million in capital and managed various acquisitions, divestitures, integrations and initial public offerings
- Corporate Controller for Furiex Pharmaceuticals, Inc., from its initial spin-out transaction, prior to the sale to Forest Labs for \$1.1 billion
- Earned BAs in Economics and History, and a Master's in Accounting from the University of North Carolina at Chapel Hill



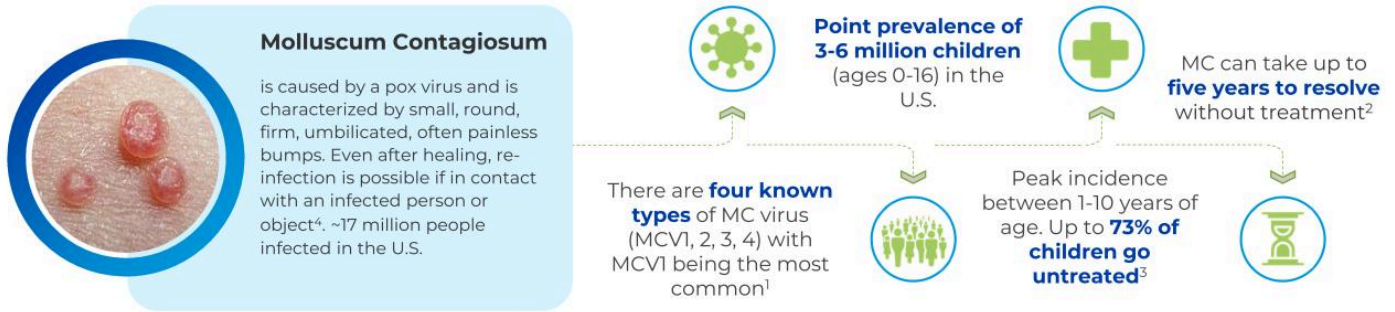
Sai Rangarao | Chief Commercial Officer

- >18 years of experience leading, launching, and marketing pharmaceutical products
- VP of Marketing & Head of Neurology Sales at Collegium Pharmaceutical
- VP of Marketing & Commercial Operations at BDSI, until it was acquired by Collegium in 2022
- Head of US Dermatology Marketing for Otezla at Celgene leading to acquisition by Amgen for \$13 Billion
- Member of the commercial and marketing organization at Novartis Pharmaceuticals that launched COSENTYX® in the U.S
- Earned an MS in Bioscience Regulatory Affairs from The Johns Hopkins University, an MBA and MS from the New Jersey Institute of Technology, and a BS in Computer Science from Indiana University of Pennsylvania

Molluscum & Zelsuvmi Overview

Molluscum Contagiosum

A highly infectious viral condition primarily affecting children 1 year of age or older



Untreated Molluscum Contagiosum Has Severe Effects

Infection, Persistence, and Spread

Auto-inoculation²

Highly contagious to others

+ risk of secondary bacterial infections²

Pain & Skin Irritation

Potential worsening of atopic dermatitis

Itching, redness

Visible and Psychological Impacts

Inflammation

Anxiety

Social withdrawal

1) Hebert AA, Bhatia N, Del Rosso JQ. Molluscum Contagiosum: Epidemiology, Considerations, Treatment Options, and Therapeutic Gaps. J Clin Aesthet Dermatol. 2023 Aug;16(8 Suppl 2):S4-S11. PMID: 37636018; PMCID: PMC10453394. 2) Ludmann P. American Academy of Dermatology. Molluscum contagiosum. 4 October 2023. 3) Basdag H, Rainer BM, Cohen BA. Molluscum contagiosum: to treat or not to treat? Experience with 170 children in an outpatient clinic setting in the northeastern United States. Pediatr Dermatol. 2015;32(3):353-357. doi:10.1111/pde.12504. 4) Schaffer JV, Berger EM. Molluscum Contagiosum. JAMA Dermatol. 2016;152(9):1072. doi:10.1001/jamadermatol.2016.2367. 5) CDC. Clinical Overview of Molluscum Contagiosum. Jan 2025.

Zelsuvmi™ Has the Potential to Shift MC Treatment Paradigm

The 1st & Only At Home Prescription Treatment

Previous Treatment Options



- Other available topical treatment **requires in-office visits every 3 weeks**²



- **Painful, destructive** treatments³



- Necessitates travel to HCP offices, adding to the **time burden for MC patients and caregivers**²



- Remaining treatment options such as off-label drugs / natural remedies have **unproven efficacy**⁴



- **Daily** application that can be **started immediately**

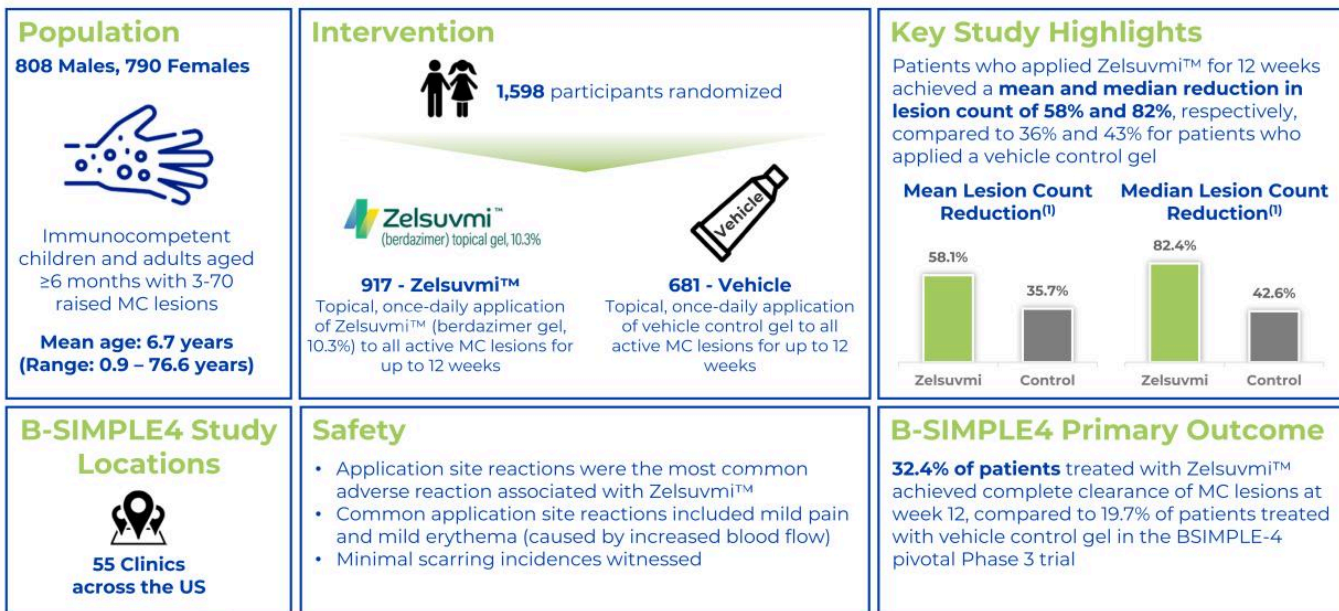
- **Attractive safety profile** demonstrated in clinical trials with no / minimal scarring^{5,6}

- **First FDA approved medication** for molluscum that can be applied at home by patients or caregivers⁵

- **Demonstrated, proven efficacy** across key primary and secondary endpoints in clinical trials⁶

¹Least-squares mean count reduction. See Figure 9; Browning JC, Hebert A, Erloe C, Cartwright M, Maeda-Chubachi T. Berdazimer Gel 10.3% is a Clinically Meaningful Therapeutic Intervention for Molluscum Contagiosum. Abstract and poster presented at Fall Clinical 2024, Las Vegas NV, October 26-27, 2024. ² Eichenfield LF, Kwong P, Gonzalez ME, et al. Safety and Efficacy of VP-102 (Cantharidin, 0.7% w/v) in Molluscum Contagiosum by Body Regions: Post Hoc Pooled Analyses from Two Phase III Randomized Trials. J Clin Aesthet Dermatol. 2021;14(10):42-47. ³ Hebert AA, Bhatia N, Del Rosso JQ. Molluscum Contagiosum: Epidemiology, Considerations, Treatment Options, and Therapeutic Caps. J Clin Aesthet Dermatol. 2023;16(8 Suppl 1):S4-S11. ⁴ Ong SK, Hoft I, Siegfried E. Analysis of over-the-counter products marketed to treat molluscum contagiosum. Pediatr Dermatol. 2023;38(5):1400-1403. doi:10.1111/pde.14776. ⁵ Zelsuvmi Package Insert. ⁶ Sugarman JL, Hebert A, Browning JC, et al. Berdazimer gel for molluscum contagiosum: An integrated analysis of 3 randomized controlled trials. J Am Acad Dermatol. 2024;90(2):299-308. doi:10.1016/j.jaad.2023.09.066Ong

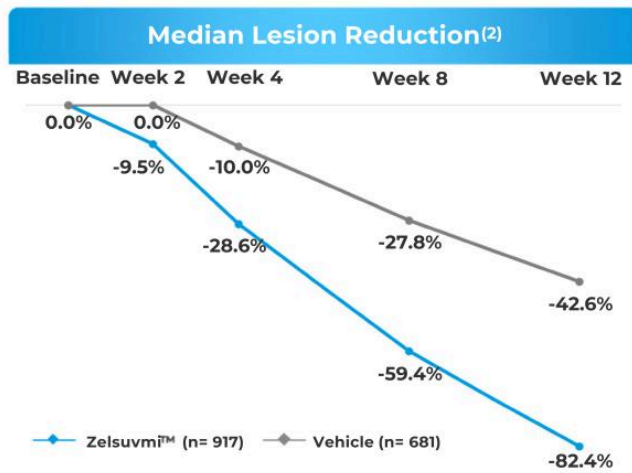
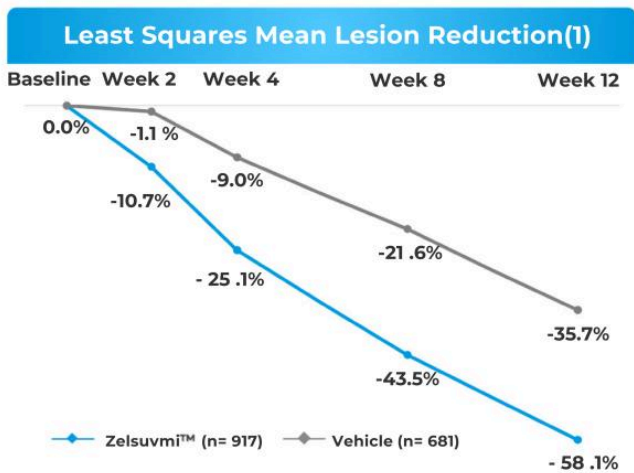
Zelsuvmi™ Efficacy Shown in Phase 3 Clinical Trials Drives Commercial Launch



¹⁾ p-value <0.0001, favoring Zelsuvmi™.
Source: Sugarman JL, Hebert A, Browning JC, Paller AS, Stripling S, Green LJ, Cartwright M, Enloe C, Wells N, Maeda-Chubachi T. Berdazimer gel for molluscum contagiosum: An integrated analysis of 3 randomized controlled trials. J Am Acad Dermatol. 2023 Oct 5:50190-9622(23)02890-6. doi:10.1016/j.jaad.2023.09.066.Epub ahead of print. PMID: 37804936.

Phase 3 Trial Results

Zelsuvmi™ showed statistically significant benefit vs. vehicle after 2 weeks of therapy and through out the entire 12-week length of the Phase 3 studies

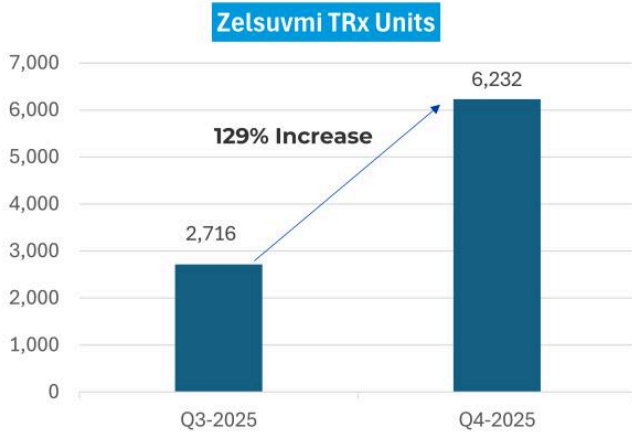


P<0.0001 at all time points, favoring Zelsuvmi™

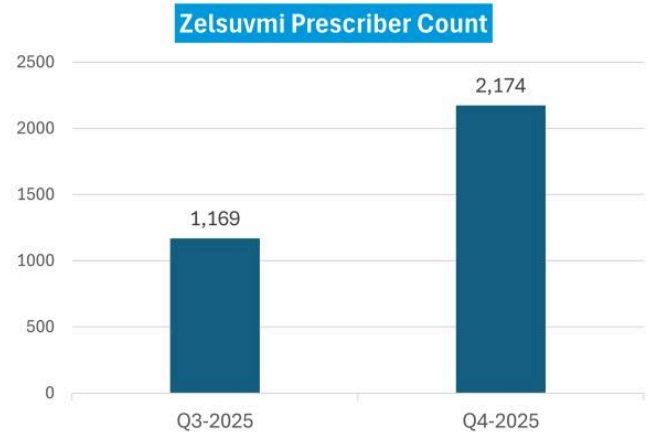
1) Figure 9: Browning JC, Hebert A, Enloe C, Cartwright M, Maeda-Chubachi T. Berdazimer Gel 10.3% is a Clinically Meaningful Therapeutic Intervention for Molluscum Contagiosum. Abstract and poster presented at Fall Clinical 2024. Las Vegas, NV. October 24-27, 2024. 2) Figure 10: Browning JC, Hebert A, Enloe C, Cartwright M, Maeda-Chubachi T. Berdazimer Gel 10.3% is a Clinically Meaningful Therapeutic Intervention for Molluscum Contagiosum. Abstract and poster presented at Fall Clinical 2024. Las Vegas, NV. October 24-27, 2024.

Zelsuvmi Commercial Overview

Strong Quarter over Quarter Growth during Zelsuvmi Launch



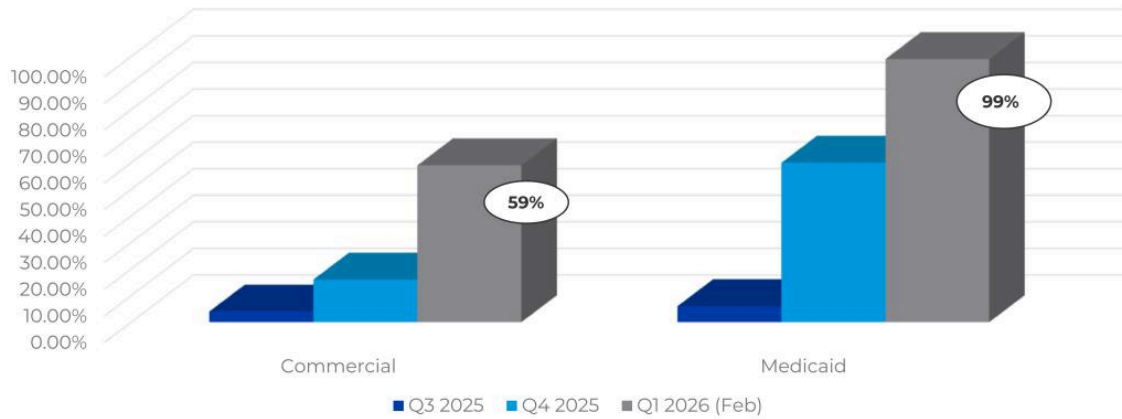
Total of 8,948 Prescribed Units in Q3 & Q4 2025



Total of 2,712 Unique Prescribers in Q3 & Q4 2025

Data Source: Symphony Health- Metys Data

Zelsuvmi Covered Lives by Quarter*

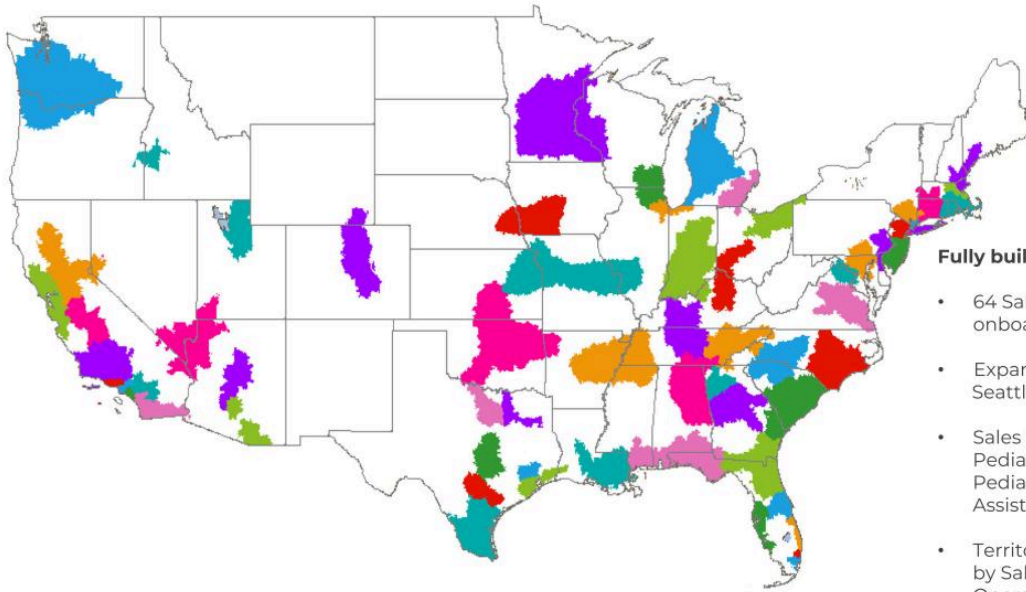


- Selective contracting strategy
- 70% combined Medicaid/commercial coverage with 1 contract
- Favorable Gross to Nets
- Favorable approval rates with all payers

- Commercial, Cash and Assistance Programs - 74% of TRxs
- Managed and FSS Medicaid - 26%
- Medicare <1% of TRxs

*Clarivate (DRG) Fingertip Formulary

Sales Team Expansion Complete



Fully built out commercial team:

- 64 Sales Territories Expansion completed & onboarded ~53% of MC Claims
- Expanded territories include sizable markets: Seattle, Las Vegas, San Francisco, Salt Lake, etc.
- Sales Team targeting: Pediatric/Adult Dermatologists, General Pediatrics, Nurse Practitioners & Physicians Assistants
- Territory managers supported by Sales Training, Marketing, Commercial Operations & Market Access teams

Robust Zelsuvmi Tactical Execution



National & Regional Conference Presence



YouTube Promotional Commercial



Live & Virtual Educational Speaker Development



Digital Marketing



New Patient Testimonials & Information



ZELSUVMI GO Patient Support Program

Xepi: New Product Acquisition



Xepi (ozenoxacin) Cream for the treatment of Impetigo



(ozenoxacin) Cream, 1%

Acquired from BioFrontera
in October 2025

FDA Approved in 2017

Exclusivity until **2032**

Xepi Clinical Story

- Ozenoxacin cream 1% developed as first line treatment in patients aged 2 months and older
- 15 clinical studies in Phase 1 & 2 conducted
- Two Pivotal Phase 3 studies conducted in both adult & pediatric patients with impetigo 2 months old and up
- Ozenoxacin demonstrated superior clinical and bacteriological outcomes vs. vehicle control

Impetigo Facts¹

- #1 bacterial infection seen in pediatrician offices, represents 1-2% of all visits to Pediatricians in the US, with **135M** children suffering worldwide
- Impetigo is a highly contagious bacterial skin infection, most often caused by Staphylococcus aureus and/or Group A Streptococcus (Streptococcus pyogenes)
- Mupirocin resistance is growing significantly in the US

Pelthos Opportunity

- Strong synergy with existing commercial infrastructure for Zelsuvmi
- Significant overlap between Xepi & Zelsuvmi HCP call points
- Promotional alignment across Sales, Marketing & Commercial Operations
- Anticipated Commercial Launch: Late 2026

Sources: CDC Website, Xepi Pack Insert, FDA.gov

¹<https://www.cdc.gov/group-a-strep/about/impetigo.html>

Xeglyze: New Product Acquisition



Xeglyze (abametapir) Lotion for the Treatment of Head Lice



Acquired from Hatchtech in December 2025

FDA Approved in 2020

Exclusivity until **2034**

Xeglyze Clinical Story

- Abametapir lotion 0.74% developed as first line treatment in patients aged 6 months of age and older
- Phase 2b study completed in 2014 demonstrated 100% ovicidal efficacy
- Two Pivotal Phase 3 studies demonstrated that a single, ten-minute application of Xeglyze® results in a statistically significant increase in the proportion of subjects who are cleared of lice versus vehicle.

Head Lice Facts¹

- **100m+** infestations globally, with **6-12m cases** in the US, each year with substantial social cost
- Increasing resistance to current products containing pyrethrin, permethrin & malathion
- Existing products are only effective against lice and not eggs, and most require repeat treatments to break life cycle of infestation, leading to poor compliance and reduced efficacy

Pelthos Opportunity

- Strong synergy with existing commercial infrastructure for Zelsuvmi and Xepi
- Significant overlap between Xeglyze & Zelsuvmi HCP call points
- Promotional alignment across Sales, Marketing & Commercial Operations
- Anticipated Commercial Launch: First half 2027

Sources: CDC Website, Xepi Pack Insert, FDA.gov

¹https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/2089451bl.pdf

Summary Financial Statements

Summary Balance Sheet

	December 31, 2025	December 31, 2024
Cash and cash equivalents	\$ 17,973	\$ 513
Accounts receivable, net	8,858	—
Inventory, net	23,574	—
Total current assets	53,410	1,369
Total assets	130,397	1,369
Accounts payable	\$ 2,986	\$ 1,897
Accrued expenses	15,364	—
Total current liabilities	25,993	4,083
Total liabilities	91,516	4,083
Total stockholders' equity (deficit)	38,881	(2,714)
Total liabilities and stockholders' equity (deficit)	130,397	1,369

Notes

- Cash balance does not include \$30 million raised in January 2026 with issuance of 5-year term notes; additional \$20 million available on hitting certain milestones
- Strong stockholder equity position and cash balance provide resources to execute on our business plan
- Active \$200 million shelf registration statement

Summary Income Statement

	Quarters Ended	
	December 31, 2025	September 30, 2025
Revenue		
Net product revenues	\$ 9,094	\$ 7,112
License and collaboration revenues	295	294
Total revenue	9,389	7,406
Operating expenses		
Cost of goods sold	1,672	2,316
Selling, general and administrative	18,469	19,628
Research and development	374	145
Amortization of intangible assets	877	679
Total operating expenses	21,392	22,768
Operating loss	(12,003)	(15,362)
Other (expense) income		
Interest expense	(1,314)	(1,346)
Impairment of intangible assets	(285)	—
Change in fair value of convertible debt	(14,984)	—
Interest income and other income	—	5
Total other (expense) income	(16,583)	(1,341)
Net loss before provision for income taxes	(28,586)	(16,703)
Provision for income taxes	(6,922)	(465)
Net loss and comprehensive loss	\$ (21,664)	\$ (16,238)

Notes

- Revenue increased 28% quarter over quarter
- SG&A expenses declined 5% quarter over quarter and operating loss improved
- Adjusted EBITDA loss – removing non-cash, royalties and one-time items – improved 22% quarter over quarter
- Expect personnel expenses to rise approximately \$1 million per quarter in 2026 with expanded sales team and minor headcount additions for Xepi and Xeglyze launches

Appendix

Nitricil Platform Pipeline*

Asset Description	Asset Description	Approximate Time to NDA Filing	Market Potential
SB414 (AD/Psoriasis)	Berdazimer topical cream, dose TBD, for treatment of mild to moderate atopic dermatitis. Phase 1/2 Clinical stage.	7.5 years	\$\$\$ (AD) \$\$ (Psoriasis)
SB208 (Tinea Pedis -> Onychomycosis)	Low alcohol berdazimer topical gel for treatment of athlete's foot with label expansion for onychomycosis following initial approval. Phase 2/3 Clinical stage.	5 years (T. Pedis) 6.5 years (Onychomycosis)	\$\$\$\$\$
SB208 (Tinea Pedis + Onychomycosis)	Low alcohol berdazimer topical gel for treatment of both athlete's foot and onychomycosis. Phase 2/3 Clinical stage.	6.5 years	\$\$\$\$\$
SB207 (EGW/PAW)	Berdazimer topical gel, 10.3% for treatment of external genital and perianal warts. Same active gel (Tube A) as Zelsuvmi but different hydrogel (Tube B) formulation. Phase 3 clinical stage.	6.5 years	\$

*Pelthos has contractual rights to SB207 and would need to enter into a separate license for other indications set forth herein

NaV1.7 Pipeline

Product/ Indication	Asset Description	Approximate Time to NDA Filing	Market Potential
CT2000 Eye Drops Chronic Ocular Pain	CC8464 1%, 1.25% and 1.5% ophthalmic solution Phase 1-2a ready	3-4 years	\$8 billion globally
CT2000 Eye Drops Acute Ocular Pain	CC8464 1%, 1.25% and 1.5% ophthalmic solution Phase 1-2a ready	2-3 years	\$400 million globally
CT3000 depot Nerve Blocks	CC8464 5% and 10% depot injectable Preclinical Stage	5+ years	\$300-570 million globally
CC8464 Oral Erythromelalgia	CC8464 melt-granulation capsules 50mg, 100mg, 400mg Phase 2 Ready	5+ years	\$2.4 billion globally
CC8464 Oral Small Fibre Neuropathy	CC8464 melt-granulation capsules 50mg, 100mg, 400mg Phase 2 Ready	5+ years	\$50 million – 100 million
CC8464 Oral Acute Pain	CC8464 melt-granulation capsules 50mg, 100mg, 400mg Phase 2 Ready	5+ years	\$20 billion globally

Board of Directors



Peter Greenleaf, Chairman



Richard Baxter



Todd Davis



Andrew Einhorn



Ezra Friedberg



Richard Malamut, MD



Matt Pauls



Scott Plesha



Key Highlights

- 
Portfolio of FDA Approved Products

Differentiated portfolio of novel, cutaneous infectious disease products, including Zelsuvmi, Xepi and Xeglyze for the treatment of MC, impetigo and head lice, respectively
- 
Significant Unmet Need and Large Market Opportunities

Each Pelthos product is differentiated from existing treatment options with considerable market opportunities
- 
Barriers to Entry

Strong patent portfolio, along with complex, proprietary manufacturing process for Zelsuvmi and complex, multi-step manufacturing process for Xepi provides hefty market protection
- 
Operating Leverage

All three products utilize the same sales team, with largely overlapping call points, provides greater operating and financial leverage with very little dedicated overhead
- 
Strong Financial Position

Current balance sheet, revenue growth and strong existing investor support with substantial investable cash provides robust foundation for growth
- 
Biopharmaceutical Platform Poised for Growth

Strategically positioned to explore and integrate synergistic acquisitions, serving as a platform for investors seeking a strong foothold in the specialty biopharmaceutical market
- 
Pipeline

Opportunity to exploit legacy Channel clinical programs and work with Ligand to execute on clinical stage programs based on the same Nitricil platform as Zelsuvmi



Thank You



Pelthos
Therapeutics

Contacts

Investor Inquiries:

Mike Moyer
LifeSci Advisors, LLC
Managing Director
mmoyer@lifesciadvisors.com

Media:

KWM Communications
Kellie Walsh / Rachel Kessler
pelthos@kwmcommunications.com
(914) 315-6072
