



EYEPOINT®

EyePoint to Report Second Quarter 2025 Financial Results on August 6, 2025

Jul 30, 2025

WATERTOWN, Mass., July 30, 2025 (GLOBE NEWSWIRE) -- EyePoint Pharmaceuticals, Inc. (NASDAQ: EYPT), a company committed to developing and commercializing innovative therapeutics to improve the lives of patients with serious retinal diseases, today announced it will host a conference call and live webcast at 8:30 a.m. ET on Wednesday, August 6, 2025 to report its second quarter 2025 financial results and highlight recent corporate developments.

To access the live conference call, please register using the audio conference link: <https://register-conf.media-server.com/register/BI2f02d8b4966b40da83f2ef4135b2ba78>. A live audio webcast of the event can be accessed via the Investors section of the Company website at www.eyepointpharma.com. A webcast replay will also be available on the corporate website at the conclusion of the call.

About EyePoint

EyePoint Pharmaceuticals, Inc. (Nasdaq: EYPT) is a clinical-stage biopharmaceutical company committed to developing and commercializing innovative therapeutics to improve the lives of patients with serious retinal diseases. The Company's lead product candidate, DURAVYU™, is an innovative investigational sustained delivery treatment for VEGF-mediated retinal diseases combining vorolanib, a selective and patent-protected tyrosine kinase inhibitor (TKI), in next-generation bioerodible Durasert E™ technology. Supported by robust safety and efficacy data to date, DURAVYU is currently being evaluated in two Phase 3 pivotal trials for wet age-related macular degeneration (wet AMD) with topline data anticipated in 2026. DURAVYU also completed a positive Phase 2 clinical trial in diabetic macular edema (DME) with Phase 3 pivotal planning underway. Despite current therapies, patients with wet AMD and DME still tend to lose vision in the long term and wet AMD is the leading cause of vision loss among people 50 years of age and older in the United States.

The Company is committed to partnering with the retina community to improve patient lives while creating long-term value, with four approved drugs over three decades and tens of thousands of eyes treated with EyePoint innovation.

EyePoint is headquartered in Watertown, Massachusetts, with a commercial manufacturing facility in Northbridge, Massachusetts.

Vorolanib is licensed to EyePoint exclusively by Equinox Sciences, a Betta Pharmaceuticals affiliate, for the localized treatment of all ophthalmic diseases outside of China, Macao, Hong Kong and Taiwan.

DURAVYU™ has been conditionally accepted by the FDA as the proprietary name for EYP-1901. DURAVYU is an investigational product candidate; it has not been approved by the FDA. FDA approval and the timeline for potential approval is uncertain.

Investors:

Tanner Kaufman / Jenni Lu
FTI Consulting
Direct: 203-722-8743 / 667-321-6018
tanner.kaufman@fticonsulting.com / jenni.lu@fticonsulting.com

Media Contact:

Amy Phillips
Green Room Communications
Direct: 412-327-9499
aphillips@greenroompr.com



EYEPOINT®

Source: EyePoint Pharmaceuticals, Inc.