EyePoint Announces Participation at Upcoming Investor Conferences

May 13, 2025

WATERTOWN, Mass., May 13, 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 that company management will participate at the following upcoming investor conferences:

- **RBC Capital Markets Global Healthcare Conference**
  Forum: Fireside Chat
  Date: Tuesday, May 20, 2025
  Time: 10:00 a.m. ET

- **Mizuho Neuro & Ophthalmology Summit 2025**
  Forum: 1x1 Meetings
  Date: Wednesday, May 21, 2025

- **Stifel 2025 Virtual Ophthalmology Forum**
  Forum: Fireside Chat
  Date: Tuesday, May 27, 2025
  Time: 8:30 a.m. ET

A live webcast and subsequent archived replay of each presentation may be accessed via the Investors section of the Company website at www.eyepointpharma.com.

About EyePoint

EyePoint Pharmaceuticals, Inc. (Nasdaq: EYPT) is a clinical-stage biopharmaceutical company committed to developing and commercializing innovative therapeutics to help improve the lives of patients with serious retinal diseases. The Company's pipeline leverages its proprietary bioerodible Durasert E™ technology for sustained intraocular drug delivery. The Company's lead product candidate, DURAVYU™ is an investigational sustained delivery treatment for VEGF-mediated retinal diseases combining vorolanib, a selective and patent-protected tyrosine kinase inhibitor with bioerodible Durasert E ™. Supported by robust safety and efficacy data to date, DURAVYU is presently in Phase 3 global, pivotal clinical trials for wet age-related macular degeneration (wet AMD), the leading cause of vision loss among people 50 years of age and older in the United States and recently completed a Phase 2 clinical trial in diabetic macular edema (DME).

Pipeline programs include EYP-2301, a TIE-2 agonist, razuprotafib, formulated in Durasert E ™ to potentially improve outcomes in serious retinal diseases. The proven Durasert® drug delivery technology has been safely administered to thousands of patient eyes across four U.S. FDA approved products in multiple disease indications. EyePoint is headquartered in Watertown, Massachusetts, and operates a commercial-ready 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.

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