EyePoint Pharmaceuticals Announces First Quarter Fiscal Year 2019 Financial Results Release Date and Conference Call Information

October 30, 2018

WATERTOWN, Mass., Oct. 30, 2018 (GLOBE NEWSWIRE) -- EyePoint Pharmaceuticals, Inc. (NASDAQ:EYPT), a specialty biopharmaceutical company committed to developing and commercializing innovative ophthalmic products, will report results for its first quarter of fiscal year 2019 on Tuesday, November 6. Management will host a conference call to review the results at 8:00 AM ET on the same day.

The conference call may be accessed by dialing (877) 312-7507 from the U.S. and Canada, or (631) 813-4828 from international locations. The conference ID is 3098959. A live webcast will be available on the Investor Relations section of the corporate website at http://www.eyepointpharma.com. A webcast replay will also be available on the corporate website at the conclusion of the call.

About EyePoint Pharmaceuticals
EyePoint Pharmaceuticals, Inc. (formerly pSivida Corp.) (www.eyepointpharma.com), headquartered in Watertown, MA, is a specialty biopharmaceutical company committed to developing and commercializing innovative ophthalmic products in indications with high unmet medical need to help improve the lives of patients with serious eye disorders. With the approval of YUTIQTM, the Company has developed five of only six FDA-approved sustained-release treatments for eye diseases. In addition, DEXYCU™ was approved by the FDA on February 9, 2018. DEXYCU, administered as a single intraocular dose at the end of ocular surgery for the treatment of postoperative inflammation, is the first and only FDA-approved intraocular product with this indication. DEXYCU employs the Verisome® extended-release drug delivery technology, which encompasses a broad number of related, but distinct drug delivery systems capable of incorporating an extensive range of active agents, including small molecules, proteins and monoclonal antibodies. ILUVIEN® (fluocinolone acetonide intravitreal implant), a micro-insert for diabetic macular edema, licensed to Alimera Sciences, Inc., is currently sold directly in the U.S. and several EU countries. Retisert® (fluocinolone acetonide intravitreal implant), for posterior uveitis, is licensed to and sold by Bausch & Lomb, Inc. YUTIQ three-year treatment of chronic non-infectious uveitis affecting the posterior segment of the eye was approved by the FDA on October 12, 2018. The Company's pre-clinical development program is focused on using its core Durasert™ and the Verisome platform technologies to deliver drugs to treat wet age-related macular degeneration, glaucoma, and other diseases. To learn more about the Company, please visit www.eyepointpharma.com and connect on Twitter, LinkedIn, Facebook and Google+.

Contacts

Investors:

Argot Partners
Kimberly Minarovich
(646) 368-8014
kimberly@argotpartners.com

Joseph Rayne
(617) 340-6075
joseph@argotpartners.com

Media:
Thomas Gibson
(201) 476-0322
tom@tomgibsoncommunications.com

Source: EyePoint Pharmaceuticals, Inc.