Palvella Therapeutics to Host Third Quarter 2026 Financial Results and Corporate Update Conference Call on November 5, 2026

October 29, 2026

WAYNE, Pa., Oct. 29, 2026 (GLOBE NEWSWIRE) -- Palvella Therapeutics, Inc. (Palvella or the "Company") (Nasdaq: PVLA), a clinical-stage biopharmaceutical company focused on developing and commercializing novel therapies for serious, rare skin diseases and vascular anomalies for which there are no U.S. Food and Drug Administration (FDA)-approved therapies, today announced that it will report its third quarter 2026 financial results before market open on Thursday, November 5, 2026. Palvella management will host a conference call for investors at 8:30 a.m. ET on that same day to discuss the results and provide a corporate update.

To access the live webcast, including presentation slides, please click here or visit the "Events & Presentations" section of Palvella's website. To access the conference call by phone, register using this link, and you will be provided with dial-in details. A replay of the webcast will be available approximately two hours after the conclusion of the call and will remain archived for 90 days under the "Events & Presentations" section of the Company's website at www.palvellatx.com.

About Palvella Therapeutics

Founded and led by rare disease biotech veterans, Palvella Therapeutics, Inc. (Nasdaq: PVLA) is a clinical-stage biopharmaceutical company focused on developing and commercializing novel therapies to treat patients living with serious, rare skin diseases and vascular anomalies for which there are no FDA-approved therapies. Palvella is developing a broad pipeline of product candidates based on its patented QTORIN™ platform, with an initial focus on serious, rare skin diseases and vascular anomalies, many of which are lifelong in nature. Palvella's lead product candidate, QTORIN™ 3.9% rapamycin anhydrous gel (QTORIN™ rapamycin), is currently being developed for the treatment of microcystic lymphatic malformations, cutaneous venous malformations, and clinically significant angiokeratomas. Palvella's second product candidate, QTORIN™ pitavastatin, is currently being developed for the treatment of disseminated superficial actinic porokeratosis. For more information, please visit www.palvellatx.com or follow Palvella on LinkedIn or X (formerly known as Twitter).

QTORIN™ rapamycin and QTORIN™ pitavastatin are for investigational use only and neither has been approved by the FDA o by any other regulatory agency for any indication.

Contact Information

Investors

Wesley H. Kaupinen
Founder and CEO
Palvella Therapeutics
wes.kaupinen@palvellatx.com

Media

Marcy Nanus
Vice President, Investor Relations and Corporate Affairs
Palvella Therapeutics
marcy.nanus@palvellatx.com