Outlook Therapeutics, Inc. Logo

Outlook Therapeutics CEO, Bob Jahr, Participates in the Virtual Investor “Why Now” On-Demand Conference

August 27, 2026

Video webcast now available here

ISELIN, N.J., Aug. 27, 2026 (GLOBE NEWSWIRE) -- Outlook Therapeutics, Inc. (Nasdaq: OTLK), a biopharmaceutical company focused on the development and commercialization of LYTENAVA™ (bevacizumab-vikg, bevacizumab gamma) for the treatment of retinal diseases, today announced that Bob Jahr, Chief Executive Officer, recently participated in the Virtual Investor “Why Now” on-demand conference .

During the segment, Mr. Jahr highlighted why he believes Outlook Therapeutics is entering a transformative period following the FDA's approval of LYTENAVA™ on July 24, 2026, making it the only FDA-approved ophthalmic formulation of bevacizumab for the treatment of wet AMD in the United States. He discussed Outlook's transition from a development-stage to a commercial-stage company and the significant opportunity to bring an FDA-approved ophthalmic bevacizumab to an established market where bevacizumab is already widely used off label. Mr. Jahr also highlighted LYTENAVA's differentiated development and regulatory pathway, Outlook Therapeutics' commercial launch preparations in the U.S., its existing commercial presence in Europe, and the potential for LYTENAVA™ to redefine the standard of care by providing physicians with an FDA-approved ophthalmic bevacizumab specifically developed for retinal use. With regulatory risk now substantially behind the Company and commercial execution taking center stage, Mr. Jahr outlined why he believes this represents a pivotal moment in Outlook Therapeutics' evolution.

The video webcast is now accessible for on-demand viewing here.

JTC Team and Virtual Investor Co. are paid consultants to Outlook Therapeutics, Inc. JTC Team and Virtual Investor Co. are investor relations and corporate communications firms. Any content included in this release shall not be construed as an offer to purchase securities of Outlook Therapeutics, Inc. Interested parties are responsible for conducting their own due diligence and are encouraged to review the Company's website and the SEC website for the latest information and filings on the Company.

About Outlook Therapeutics, Inc.

Outlook Therapeutics is a biopharmaceutical company focused on the development and commercialization of LYTENAVA (bevacizumab-vikg (U.S.), bevacizumab gamma (E.U.)). LYTENAVA is the only ophthalmic formulation of bevacizumab to receive U.S. FDA approval and European Commission and MHRA Marketing Authorization for the treatment of wet AMD. Outlook Therapeutics commenced commercial launch of LYTENAVA™ (bevacizumab gamma) in Germany, Austria, and the UK as a treatment for wet AMD.

Forward-Looking Statements

This press release contains statements that may or are considered “forward-looking statements”. All statements other than statements of historical facts are “forward-looking statements,” including those relating to future events. In some cases, you can identify forward-looking statements by terminology such as “anticipate,” “believe,” “can,” “could,” “continue,” “expect,” “may,” “on track,” “plan,” “potential,” “target,” “will,” or “would”, the negative of terms like these or other comparable terminology, and other words or terms of similar meaning. These include, among others, express or implied discussions regarding the Company's planned launch of LYTENAVA in the United States and other jurisdictions and the timing thereof; expectations concerning potential revenue generation from sales of LYTENAVA; expectations surrounding market adoption of LYTENAVA; potential additional marketing approvals; expectations regarding the potential impact of LYTENAVA in the retina community; new indications or labeling for LYTENAVA; Outlook Therapeutics development or future revenue plans for LYTENAVA generally; and other statements that are not historical fact. Although Outlook Therapeutics believes that it has a reasonable basis for the forward-looking statements contained herein, they are based on current expectations about future events affecting Outlook Therapeutics and are subject to risks, uncertainties, and factors relating to its operations and business environment, all of which are difficult to predict and many of which are beyond its control. These risk factors include those risks associated with developing and commercializing pharmaceutical product candidates, risks in obtaining necessary regulatory approvals, the content and timing of decisions by regulatory bodies, as well as those risks detailed in Outlook Therapeutics' filings with the Securities and Exchange Commission (the SEC), including the Current Report on Form 8-K filed with the SEC on August 12, 2026, and future reports Outlook Therapeutics files with the SEC, which include uncertainty of market conditions and future impacts related to macroeconomic factors, including as a result of the ongoing overseas conflicts, tariffs, and trade tensions, fluctuations in interest rates and inflation, and potential future bank failures on the global business environment. These risks may cause actual results to differ materially from those expressed or implied by forward-looking statements in this press release. All forward-looking statements included in this press release are expressly qualified in their entirety by the foregoing cautionary statements. You are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date hereof. Outlook Therapeutics does not undertake any obligation to update, amend, or clarify these forward-looking statements, whether as a result of new information, future events, or otherwise, except as may be required under applicable securities law.

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