

Outlook Therapeutics, Inc. Logo

Outlook Therapeutics® to Present at the H.C. Wainwright 24th Annual Global Investment Conference

September 7, 2022

ISELIN, N.J., Sept. 07, 2022 (GLOBE NEWSWIRE) -- [Outlook Therapeutics, Inc.](#) (Nasdaq: OTLK), a pre-commercial biopharmaceutical company working to develop and launch the first FDA-approved ophthalmic formulation of bevacizumab for use in retinal indications, today announced that [Russ Trenary, President and Chief Executive Officer](#) of Outlook Therapeutics, will present at the [H.C. Wainwright 24th Annual Global Investment Conference](#) being held in New York, NY and virtually on September 12-14, 2022.

In addition to the presentation, management will be available to participate in virtual one-on-one meetings with qualified members of the investor community who are registered to attend the conference. For more information about the conference, please visit the [conference website](#).

A [video webcast](#) of the presentation will be accessible for viewing on-demand beginning on Monday, September 12 at 7:00 AM ET for those registered for the event and will be available on the [Events](#) page in the [Investors](#) section of the Company's website ([outlooktherapeutics.com](#)). The webcast replay will be archived for 90 days following the event.

About Outlook Therapeutics, Inc.

Outlook Therapeutics is a biopharmaceutical company working to develop and launch ONS-5010/ LYTENAVA™ (bevacizumab-vikg), an investigational therapy, as the first FDA-approved ophthalmic formulation of bevacizumab for use in retinal indications, including wet AMD, DME and BRVO. Outlook Therapeutics has submitted its Biologics License Application (BLA) to the U.S. Food and Drug Administration (FDA) for ONS-5010 to treat wet AMD. The submission is supported by Outlook Therapeutics' wet AMD registration clinical program, which consists of three clinical trials: NORSE ONE, NORSE TWO, and NORSE THREE. If ONS-5010 ophthalmic bevacizumab is approved, Outlook Therapeutics expects to commercialize it as the first and only FDA-approved ophthalmic formulation of bevacizumab for use in treating retinal diseases in the United States, United Kingdom, Europe, Japan and other markets. For more information, please visit [www.outlooktherapeutics.com](#).

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Source: Outlook Therapeutics, Inc.