

Outlook Therapeutics, Inc. Logo

## Outlook Therapeutics to Present at the H.C. Wainwright Global Life Sciences Conference

March 4, 2021

MONMOUTH JUNCTION, N.J., March 04, 2021 (GLOBE NEWSWIRE) -- [Outlook Therapeutics, Inc.](#) (Nasdaq: OTLK), a late clinical-stage biopharmaceutical company working to develop the first FDA-approved ophthalmic formulation of bevacizumab-vikg for use in retinal indications, today announced that Lawrence A. Kenyon, President, CEO and CFO of Outlook Therapeutics, will participate in a fireside chat at the virtual [H.C. Wainwright Global Life Sciences Conference](#) taking place March 9-10, 2021.

In addition to the fireside chat, 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 event, please visit the conference website [here](#).

A [video webcast](#) of the fireside chat will be available for viewing on-demand beginning Tuesday, March 9, 2021, at 7:00 AM ET for those registered for the conference and will be accessible on the [Events](#) page of the [Investors](#) section of the Outlook Therapeutics website, [outlooktherapeutics.com](#).

### About Outlook Therapeutics, Inc.

Outlook Therapeutics is a late clinical-stage biopharmaceutical company working to develop ONS-5010/LYTENAVA™ (bevacizumab-vikg) as the first FDA-approved ophthalmic formulation of bevacizumab-vikg for use in retinal indications, including wet AMD, DME and BRVO. If ONS-5010 is approved, Outlook Therapeutics expects to commercialize it as the first and only FDA-approved ophthalmic formulation of bevacizumab-vikg for use in treating a range of retinal diseases in the United States, United Kingdom, Europe, Japan, China and other markets. Outlook Therapeutics expects to file ONS-5010 with the U.S. FDA as a new BLA under the PHS 351(a) regulatory pathway, initially for wet AMD. For more information, please visit [www.outlooktherapeutics.com](#).

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Source: Outlook Therapeutics, Inc.