

Outlook Therapeutics, Inc. Logo

## Outlook Therapeutics Announces Inclusion in the Russell 2000® Index

June 14, 2021

ISELIN, N.J., June 14, 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 for use in retinal indications, today announced that as part of the annual reconstitution of the Russell stock indexes, Outlook Therapeutics has been selected to be added to the Russell 2000® Index effective June 25, 2021, after the close of the U.S. equity markets.

"We are very pleased to have been selected for inclusion in the Russell 2000® Index. Our placement in this widely used performance benchmark for small-cap companies reflects the hard work of the Outlook Therapeutics team and the value we have created over the past year. With our inclusion in the Russell 2000® we believe we are well-positioned to continue driving value with this increased market exposure," said Lawrence A. Kenyon, President, CEO and CFO, Outlook Therapeutics.

The Russell 2000® Index measures the performance of the small-cap segment of the US equity market. The Russell 2000® Index is a subset of the Russell 3000® Index representing approximately 10% of the total market capitalization of that index. It includes approximately 2,000 of the smallest securities based on a combination of their market cap and current index membership.

Russell indexes are widely used by investment managers and institutional investors for index funds and as benchmarks for active investment strategies. Approximately \$9 trillion in assets are benchmarked against Russell's U.S. indexes which are part of FTSE Russell, a global index leader that provides innovative benchmarking, analytics and data solutions for investors worldwide.

For more information on the Russell Indexes, please visit the FTS Russell website at [www.ftserussell.com](http://www.ftserussell.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 for use in retinal indications, including wet AMD, DME and BRVO. 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. Outlook Therapeutics expects to file ONS-5010 ophthalmic bevacizumab with the U.S. FDA as a new BLA under the PHSA 351(a) regulatory pathway. For more information, please visit [www.outlooktherapeutics.com](http://www.outlooktherapeutics.com).

### Forward-Looking Statements

This press release contains 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 "may," "might," "will," "should," "expect," "plan," "anticipate," "project," "believe," "estimate," "predict," "potential," "intend" or "continue," the negative of terms like these or other comparable terminology, and other words or terms of similar meaning. 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 pharmaceutical product candidates, risks of conducting clinical trials and risks in obtaining necessary regulatory approvals, as well as those risks detailed in Outlook Therapeutics' filings with the Securities and Exchange Commission, including its Annual Report on Form 10-K for the fiscal year ended September 30, 2020, as amended, and subsequent Quarterly Reports on Form 10-Q, which include the uncertainty of future impacts related to the ongoing COVID-19 pandemic. 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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