

April 8, 2021



NeuBase Therapeutics to Host a Virtual R&D Day on June 8th to Provide Updates on the Drug Development Pipeline Targeting Genetic Diseases

PITTSBURGH, April 08, 2021 (GLOBE NEWSWIRE) -- [NeuBase Therapeutics, Inc.](https://www.neubasetherapeutics.com) (Nasdaq: NBSE) ("NeuBase" or the "Company"), a biotechnology company accelerating the genetic revolution with a new class of precision genetic medicines, announced today that it will host a virtual R&D day for investors and analysts on Tuesday, June 8, 2021, from 12:30 p.m. to 2:30 p.m. EDT.

During the event, NeuBase will present new data, along with an in-depth review of the Company's pipeline of drug candidates – including in Huntington's disease, myotonic dystrophy type 1, and a new oncology program for a target that has previously been thought of as undruggable – as well as an introduction to the expanded management team.

Additional details will be made available prior to the event. The event will be webcast live on NeuBase's website at <https://ir.neubasetherapeutics.com/news-events/ir-calendar>. Following the live webcast, a replay will be available on the Company's website and archived for approximately 90 days.

About NeuBase Therapeutics, Inc.

NeuBase is accelerating the genetic revolution by developing a new class of precision genetic medicines which can be designed to increase, decrease, or change gene function, as appropriate, to resolve genetic defects that drive disease. NeuBase's targeted PATrOL™ therapies are centered around its proprietary drug scaffold to address genetic diseases at the DNA or RNA level by combining the highly targeted approach of traditional genetic therapies with the broad organ distribution capabilities of small molecules. With an initial focus on silencing disease-causing mutations in debilitating neurological, neuromuscular, and oncologic disorders, NeuBase is committed to redefining medicine for the millions of patients with both common and rare conditions. To learn more, visit www.neubasetherapeutics.com.

Use of Forward-Looking Statements

This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act. These forward-looking statements are distinguished by use of words such as "will," "would," "anticipate," "expect," "believe," "designed," "plan," or "intend," the negative of these terms, and similar references to future periods. Forward-looking statements include, among others, those related to our expectations regarding virtual R&D day and the information to be presented during the event. These views involve risks and uncertainties that are difficult to predict and, accordingly, our actual results may differ materially from the results discussed in our forward-looking statements. Our forward-looking

statements contained herein speak only as of the date of this press release. Factors or events that we cannot predict, including those risk factors contained in our filings with the U.S. Securities and Exchange Commission, may cause our actual results to differ from those expressed in forward-looking statements. The Company may not actually achieve the plans, carry out the intentions or meet the expectations or projections disclosed in the forward-looking statements, and you should not place undue reliance on these forward-looking statements. Because such statements deal with future events and are based on the Company's current expectations, they are subject to various risks and uncertainties, and actual results, performance or achievements of the Company could differ materially from those described in or implied by the statements in this press release, including: the Company's plans to develop and commercialize its product candidates; the timing of initiation of the Company's planned clinical trials; the timing of the availability of data from the Company's clinical trials; the timing of any planned investigational new drug application or new drug application; the Company's plans to research, develop and commercialize its current and future product candidates; the clinical utility, potential benefits and market acceptance of the Company's product candidates; the Company's commercialization, marketing and manufacturing capabilities and strategy; global health conditions, including the impact of COVID-19; the Company's ability to protect its intellectual property position; and the requirement for additional capital to continue to advance these product candidates, which may not be available on favorable terms or at all, as well as those risk factors contained in our filings with the U.S. Securities and Exchange Commission. Except as otherwise required by law, the Company disclaims any intention or obligation to update or revise any forward-looking statements, which speak only as of the date hereof, whether as a result of new information, future events or circumstances or otherwise.

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