

# G1 Therapeutics to Present Preclinical Data on CDK4/6 Inhibitors Trilaciclib and G1T38 at the 2018 American Association for Cancer Research (AACR) Annual Meeting

April 9, 2018

RESEARCH TRIANGLE PARK, N.C., April 09, 2018 (GLOBE NEWSWIRE) -- G1 Therapeutics, Inc. (Nasdaq:GTHX), a clinical-stage oncology company, today announced that preclinical data on both of the company's CDK4/6 inhibitors will be presented at the 2018 American Association for Cancer Research Annual Meeting (AACR 2018), being held April 14-18 in Chicago, IL.

Data on G1T38, a potential best-in-class oral CDK4/6 inhibitor being developed for use in combination with other targeted therapies in multiple oncology indications, were selected to be featured in the AACR 2018 Poster Discussion Session. Data on trilaciclib, a short-acting IV CDK4/6 inhibitor in development to preserve hematopoietic stem cells and enhance immune system function (myelopreservation) during chemotherapy, will also be presented.

Details on the presentations are as follows:

Title: The CDK4/6 inhibitor. G1T38. enhances response to targeted therapies in preclinical models of non-small cell lung cancer

Abstract Number: 1522 Session: Targeting the Cell Cycle: Mechanism and Therapy Poster Number: 1

Date / Time / Location: April 15, 4:10 p.m. – 4:15 p.m. CDT in McCormick Place South, Level 4, Room S402 (poster discussion); April 16, 8 a.m. – 12 p.m. CDT in McCormick Place South, Exhibit Hall A, Poster Section 23 (poster presentation) Presenters: Jessica Sorrentino, Daniel Freed, John Bisi, Jay Strum, Patrick Roberts (G1 Therapeutics)

Title: Transient exposure to trilaciclib, a CDK4/6 inhibitor, modulates gene expression in tumor immune infiltrates and promotes a pro-inflammatory tumor microenvironment

# Abstract Number: 1752

Session: Modifiers of the Tumor Microenvironment 2 Poster Number: 17 Date / Time / Location: April 16, 8 a.m. – 12 p.m. CDT in McCormick Place South, Exhibit Hall A, Poster Section 33 Presenters: Anne Lai, Jessica Sorrentino, Jay Strum, Patrick Roberts (G1 Therapeutics)

Additional information on the meeting can be found on the AACR website: www.aacr.org

# About Trilaciclib (G1T28)

Trilaciclib is a short-acting CDK4/6 inhibitor in development to preserve hematopoietic stem cells and enhance immune system function (myelopreservation) during chemotherapy. Trilaciclib is administered intravenously prior to chemotherapy and has the potential to significantly improve treatment outcomes.

Trilaciclib is being evaluated in four randomized Phase 2 clinical trials. The company recently announced positive Phase 2a topline data showing myelopreservation benefits in newly diagnosed, treatment-naive small cell lung cancer (SCLC) patients (<u>NCT02499770</u>). There are three additional ongoing trilaciclib trials: a trial in previously treated SCLC patients (<u>NCT02514447</u>), a trial in patients with triple-negative breast cancer (<u>NCT02978716</u>), and a trial in combination with Tecentriq® and chemotherapy in SCLC patients (<u>NCT03041311</u>).

# About G1T38

G1T38 is a potential best-in-class oral CDK4/6 inhibitor being developed for use in combination with other targeted therapies in multiple oncology indications. Preclinical data presented at the American Association for Cancer Research 2016 Annual Meeting and published in *Molecular Cancer Research* and *Oncotarget* demonstrated the compound's differentiation from other CDK4/6 inhibitors.

G1T38 is currently being evaluated in two Phase 1/2 clinical trials: a trial in combination with Faslodex® for people with estrogen receptor-positive, HER2-negative breast cancer (<u>NCT02983071</u>), and a trial in combination with Tagrisso<sup>®</sup> for people with EGFR-mutant non-small cell lung cancer (<u>NCT03455829</u>).

# **About G1 Therapeutics**

G1 Therapeutics, Inc., is a clinical-stage biopharmaceutical company focused on the discovery and development of novel therapeutics for the treatment of cancer. G1's two clinical assets, trilaciclib and G1T38, are CDK4/6 inhibitors, a validated and promising class of targets for anti-cancer therapeutics. Trilaciclib and G1T38 have broad therapeutic potential in many forms of cancer and may serve as the backbone of multiple combination regimens. In addition, G1 is advancing G1T48, a potential first- / best-in-class oral selective estrogen receptor degrader, or SERD, which is targeted for the treatment of ER+ breast cancer.

G1 is based in Research Triangle Park, NC. For additional information, please visit <u>www.g1therapeutics.com</u> and follow us on Twitter @G1Therapeutics.

# **Forward-Looking Statements**

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Words such as "may," "will," "expect," "plan," "anticipate," "estimate," "intend" and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances) are intended to identify forward-looking statements. Forward-looking statements in this news release include, but are not limited to, the therapeutic potential of trilaciclib, G1T38 and G1T48, and are based on G1 Therapeutics' expectations and assumptions as of

the date of this press release. Each of these forward-looking statements involves risks and uncertainties. Factors that may cause G1 Therapeutics' actual results to differ from those expressed or implied in the forward-looking statements in this press release are discussed in G1 Therapeutics' filings with the U.S. Securities and Exchange Commission, including the "Risk Factors" sections contained therein and include, but are not limited to, the inherent uncertainties associated with developing new products or technologies and operating as a development-stage company; G1 Therapeutics' ability to complete clinical trials for, obtain approvals for and commercialize any of its product candidates; G1 Therapeutics' ability to recruit and enroll patients in its studies; competition in the industry in which G1 Therapeutics operates; and market conditions. Except as required by law, G1 Therapeutics assumes no obligation to update any forward-looking statements contained herein to reflect any change in expectations, even as new information becomes available.

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