April 20, 2017

TG Therapeutics, Inc. Announces Clinical Data Presentations at the Upcoming 53rd Annual Meeting of the American Society of Clinical Oncology

Positive results from the GENUINE Phase 3 study to be featured in an oral presentation

NEW YORK, April 20, 2017 (GLOBE NEWSWIRE) -- TG Therapeutics, Inc. (NASDAQ:TGTX), today announced that clinical abstracts featuring TG-1101 and TGR-1202 have been selected for presentation at the upcoming 53rd Annual Meeting of the American Society of Clinical Oncology (ASCO), to be held from June 2 - 6, 2017, at McCormick Place in Chicago, Illinois. Details of the data presentations are outlined below.

Oral Presentation:

- Title: Ublituximab and ibrutinib for previously treated genetically high-risk chronic lymphocytic leukemia: Results of the GENUINE Phase 3 study
  - Abstract Number: 7504
  - Presentation Date & Time: Saturday, June 3, 2017 3:00 PM - 6:00 PM CT
  - Session Title: Hematologic Malignancies—Lymphoma and Chronic Lymphocytic Leukemia
  - Presenter: Jeffrey P. Sharman, MD

Poster Discussion Presentation:

- Title: Tolerability and activity of chemo-free triplet combination of TGR-1202, ublituximab, and ibrutinib in patients with advanced CLL and NHL
  - Abstract Number: 7511
  - Presentation Date & Time: Monday, June 5, 2017 8:00 AM-11:30 AM CT (Poster Viewing); 1:15 PM-2:30 PM CT (Poster Discussion)
  - Session Title: Poster Discussion Session, Hematologic Malignancies—Lymphoma and Chronic Lymphocytic Leukemia
  - Presenter: Loretta Nastoupil, MD

Trials in Progress Poster Presentation:

- Title: KI intolerance study: A phase 2 study to assess the safety and efficacy of TGR-1202 in pts with chronic lymphocytic leukemia (CLL) who are intolerant to prior BTK or PI3K-delta inhibitor therapy
  - Abstract Number: TPS7569
  - Presentation Date & Time: Monday, June 5, 2017 8:00 AM-11:30 AM CT
  - Session Title: Poster Session, Hematologic Malignancies—Lymphoma and Chronic Lymphocytic Leukemia
  - Presenter: Colleen Dorsey, BSN, RN

The above abstracts will be released publicly on May 17, 2017 through the ASCO meeting website at www.asco.org. Following each presentation, the data presented will be available on the Publications page, located within the Pipeline section, of the Company's website at www.tgtherapeutics.com.

TG THERAPEUTICS INVESTOR & ANALYST EVENT

TG Therapeutics will also host a reception on Monday, June 5, 2017 beginning at 7:00pm CT, with featured presentations beginning promptly at 7:15pm CT. The event will take place at the Peninsula Chicago Hotel in the Avenues Ballroom. This event will be webcast live and will be available on the Events page, located within the Investors & Media section of the Company's website at www.tgtherapeutics.com, as well as archived for future review. This event will also be broadcast via conference call. To access the conference line, please call 1-877-407-8029 (U.S.), 1-201-689-8029 (outside the U.S.), and reference Conference Title: TG Therapeutics June 2017 Investor & Analyst Event.

ABOUT TG THERAPEUTICS, INC.

TG Therapeutics is a biopharmaceutical company focused on the acquisition, development and commercialization of novel treatments for B-cell malignancies and autoimmune diseases. Currently, the company is developing two therapies targeting hematological malignancies and autoimmune diseases. TG-1101 (ublituximab) is a novel, glycoengineered monoclonal antibody that targets a specific and unique epitope on the CD20 antigen found on mature B-lymphocytes. TG Therapeutics is also developing TGR-1202, an orally available PI3K delta inhibitor. The delta isoform of PI3K is strongly expressed in cells of hematopoietic origin and is believed to be important in the proliferation and survival of B-lymphocytes. Both TG-1101 and TGR-1202 are in clinical development for patients with hematologic malignancies, with TG-1101 also in clinical development for autoimmune disorders. The Company also has pre-clinical programs to develop IRAK4 inhibitors, BET inhibitors, and anti-PD-L1 and anti-GITR antibodies. TG Therapeutics is headquartered in New York City.

Cautionary Statement

Some of the statements included in this press release, particularly those with respect to anticipating future clinical trials, the timing of commencing or completing such trials and business prospects for TG-1101, TGR-1202, the IRAK4 inhibitor program, the BET inhibitor program, and the anti-PD-L1 and anti-GITR antibodies may be forward-looking statements that involve a number of risks and uncertainties. For those statements, we claim the protection of the safe harbor for forward-looking statements contained in the Private Securities Litigation Reform Act of 1995. Among the factors that could cause our actual results to differ materially are the following: our ability to successfully and cost-effectively complete preclinical and clinical trials for TG-1101, TGR-1202, the IRAK4 inhibitor program, the BET inhibitor program, and the anti-PD-L1 and anti-GITR antibodies; the risk that early preclinical and clinical results that supported our decision to move forward with TG-1101, TGR-1202, the IRAK4 inhibitor program, the BET inhibitor program, and the anti-PD-L1 and anti-GITR antibodies will not be reproduced in additional patients or in future studies; the risk that the pending GENUINE confirmation scans may have a negative effect on the topline data presented; the risk that the FDA will not grant us a pre-BLA meeting to discuss the results of the GENUINE study; the risk of the willingness of the FDA to review the data for approval and any likelihood of FDA approval or disapproval and the timing of filing of a BLA for TG-1101; the risk that trends observed which underlie certain assumptions of future performance of TGR-1202 will not continue; the risk that TGR-1202 will not produce satisfactory safety and efficacy results to warrant further development following the completion of the current Phase 1 study; the risk that the combination of TG-1101 and TGR-1202, referred to as TG-1303, will not prove to be a safe and efficacious backbone for triple and quad combination therapies; the risk that the data (both safety and efficacy) from future clinical trials will not coincide with the data produced from prior preclinical and clinical trials; the risk that trials will take longer to enroll than expected; our ability to achieve the milestones we project over the next year; our ability to manage our cash in line with our projections, and other risk factors identified from time to time in our reports filed with the Securities and Exchange Commission. Any forward-looking statements set forth in this press release speak only as of the date of this press release. We do not undertake to update any of these forward-looking statements to reflect events or circumstances that occur after the date hereof. This press release and prior releases are available at www.tgtherapeutics.com. The information found on our website is not incorporated by reference into this press release and is included for reference purposes only.

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