CytomX Therapeutics, Inc. Logo

CytomX Therapeutics Announces Teleconference and Webcast to Provide Corporate Update

March 19, 2017 at 7:01 PM EDT

Teleconference Scheduled for Monday, March 20, 2017 at 8:00 a.m. ET

SOUTH SAN FRANCISCO, Calif., March 19, 2017 (GLOBE NEWSWIRE) -- CytomX Therapeutics, Inc. (Nasdaq:CTMX), a biopharmaceutical company developing investigational Probody™ therapeutics for the treatment of cancer, today announced that members of the management team will host a teleconference on Monday, March 20, 2017, at 8:00 a.m. ET to provide a corporate update.

Conference Call/Webcast Information
Interested parties may access the live audio webcast of the teleconference through the Investor and News page of CytomX's website at http://ir.cytomx.com or by dialing 877-809-6037 and using the passcode 90993451. A replay will be available on the CytomX website or by dialing 855-859-2056 and using the passcode 90993451. The replay will be available from March 20, 2017, at 11:00 a.m. ET until March 27, 2017, at 11:00 a.m. ET.

About CytomX Therapeutics
CytomX is a clinical-stage, oncology-focused biopharmaceutical company pioneering a novel class of investigational antibody therapeutics based on its Probody technology platform. The company utilizes the platform to create proprietary cancer immunotherapies against clinically-validated targets, such as PD-L1, and first-in-class cancer therapeutics against novel targets, such as CD166, that are difficult to drug without causing damage to healthy tissues. Probody therapeutics are designed to take advantage of unique conditions in the tumor microenvironment to enhance the tumor-targeting features of an antibody and reduce drug activity in healthy tissues. The company's lead program, CX-072, a wholly-owned PD-L1-targeting Probody therapeutic, is being evaluated in a Phase 1/2 study. The Investigational New Drug filing for CX-2009 is planned for the first half of 2017. CX-2009 is a first-in-class Probody drug conjugate targeting the highly expressed tumor antigen, CD166. CX-072 is part of PROCLAIM (Probody Clinical Assessment In Man), an international umbrella clinical trial program that provides clinical trial sites with access to the company's novel therapies under one central protocol. In addition to its proprietary programs, CytomX is collaborating with strategic partners including AbbVie, Bristol-Myers Squibb Company, Pfizer Inc., MD Anderson Cancer Center and ImmunoGen, Inc. For more information, visit www.cytomx.com or follow us on Twitter.

CytomX Therapeutics Forward-Looking Statements
This press release includes forward-looking statements. Such forward-looking statements involve known and unknown risks, uncertainties and other important factors that are difficult to predict, may be beyond our control, and may cause the actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied in such statements. Accordingly, you should not rely on any of these forward-looking statements. Our Probody platform is beginning clinical development, and the process by which clinical development could potentially lead to an approved product is long and subject to significant risks and uncertainties. Projected net cash utilization and capital resources are subject to substantial risk of variance based on a wide variety of factors that can be difficult to predict. Applicable risks and uncertainties include those relating to our preclinical research and development, clinical development, and other risks identified under the heading "Risk Factors" included in our filings with the SEC. The forward-looking statements contained in this press release are based on information currently available to CytomX and speak only as of the date on which they are made. CytomX does not undertake and specifically disclaims any obligation to update any forward-looking statements, whether as a result of any new information, future events, changed circumstances or otherwise.

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