Cytokinetics Announces Two Presentations at the 2024 American College of Clinical Pharmacology Annual Meeting

September 5, 2024 8:00 PM EDT

Full Data from Phase 1 Study of CK-4021586 to be Presented on September 8, 2024

SOUTH SAN FRANCISCO, Calif., Sept. 05, 2024 (GLOBE NEWSWIRE) -- Cytokinetics, Incorporated (Nasdaq: CYTK) today announced two presentations at the 2024 American College of Clinical Pharmacology (ACCP) Annual Meeting in Bethesda, MD from September 8, 2024 – September 10, 2024.

Title: A First-in-Human, Single & Multiple Ascending Dose Study of CK-4021586, A Novel Cardiac Myosin Inhibitor
Presenter: Justin Lutz, Pharm. D., Ph.D., Senior Director, Clinical Pharmacology, Cytokinetics
Date: September 8, 2024
Session Title: Opening Reception, Exhibits & Poster Session 1
Topic: Pharmacokinetics (Including PopPK, ADME, Biopharmaceutics)
Poster Number: 074
Session Time: 5:00-7:00 PM ET
Location: Grand Ballroom E-H

Title: Clinical Evaluation of the Effect of *Aficamten* on QT/QTc Interval in Healthy Participants
Presenter: Polina German, Pharm. D., Executive Director, Clinical Pharmacology, Cytokinetics
Date: September 8, 2024
Session Title: Opening Reception, Exhibits & Poster Session 1
Topic: General Drug Development Strategy & Practice
Poster Number: 022
Session Time: 5:00-7:00 PM ET
Location: Grand Ballroom E-H

About Cytokinetics

Cytokinetics is a late-stage, specialty cardiovascular biopharmaceutical company focused on discovering, developing and commercializing muscle biology-directed drug candidates as potential treatments for debilitating diseases in which cardiac muscle performance is compromised. As a leader in muscle biology and the mechanics of muscle performance, the company is developing small molecule drug candidates specifically engineered to impact myocardial muscle function and contractility. Cytokinetics is preparing for regulatory submissions for *aficamten*, its next-in-class cardiac myosin inhibitor, following positive results from SEQUOIA-HCM, the pivotal Phase 3 clinical trial in obstructive hypertrophic cardiomyopathy which were published in the *New England Journal of Medicine*. *Aficamten* is also currently being evaluated in MAPLE-HCM, a Phase 3 clinical trial of *aficamten* as monotherapy compared to metoprolol as monotherapy in patients with obstructive HCM, ACACIA-HCM, a Phase 3 clinical trial of *aficamten* in patients with non-obstructive HCM, CEDAR-HCM, a clinical trial of *aficamten* in a pediatric population with obstructive HCM, and FOREST-HCM, an open-label extension clinical study of *aficamten* in patients with HCM. Cytokinetics is also developing *omecamtiv mecarbil*, a cardiac muscle activator, in patients with heart failure. Additionally, Cytokinetics is developing CK-586, a cardiac myosin inhibitor with a mechanism of action distinct from *aficamten* for the potential treatment of HFpEF.

For additional information about Cytokinetics, visit www.cytokinetics.com and follow us on X, LinkedIn, Facebook and YouTube.

Forward-Looking Statements

This press release contains forward-looking statements for purposes of the Private Securities Litigation Reform Act of 1995 (the "Act"). Cytokinetics disclaims any intent or obligation to update these forward-looking statements and claims the protection of the Act's Safe Harbor for forward-looking statements. Examples of such statements include, but are not limited to, statements relating to any of our other clinical trials, statements relating to the potential benefits of *aficamten*, CK-586 or any of our other drug candidates. Cytokinetics' research and development activities; the design, timing, results, significance and utility of preclinical and clinical results; and the properties and potential benefits of Cytokinetics' other drug candidates. Such statements are based on management's current expectations, but actual results may differ materially due to various risks and uncertainties, including, but not limited to, potential difficulties or delays in the development, testing, regulatory approvals for trial commencement, progression or product sale or manufacturing, or production of Cytokinetics' drug candidates that could slow or prevent clinical development or product approval; Cytokinetics' drug candidates may have adverse side effects or inadequate therapeutic efficacy; the FDA or foreign regulatory agencies may delay or limit Cytokinetics' ability to conduct clinical trials; Cytokinetics may be unable to obtain or maintain patent or trade secret protection for its intellectual property; standards of care may change, rendering Cytokinetics' drug candidates obsolete; and competitive products or alternative therapies may be developed by others for the treatment of indications Cytokinetics' drug candidates and potential drug candidates may target. For further information regarding these and other risks related to Cytokinetics' business, investors should consult Cytokinetics' filings with the Securities and Exchange Commission.

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