Cytokinetics Announces Three Upcoming Presentations at the European Society of Cardiology Heart Failure 2022 Congress

May 16, 2022 8:00 PM EDT

SOUTH SAN FRANCISCO, Calif., May 16, 2022 (GLOBE NEWSWIRE) -- Cytokinetics, Incorporated (Nasdaq: CYTK) today announced three Late-Breaking Science presentations at Heart Failure 2022, an International Congress of the European Society of Cardiology taking place online and in Madrid, Spain from May 21, 2022 – May 24, 2022. The presentations will include interim data from the open-label extension study of *aficamten*, REDWOOD-HCM OLE (**R**andomized **E**valuation of **D**osing **W**ith CK-274 in **O**bstructive **O**utflow **D**isease in **HCM**) and two additional analyses from GALACTIC-HF (**G**lobal **A**pproach to **L**owering **A**dverse **C**ardiac Outcomes **T**hrough **I**mproving **C**ontractility in **H**eart **F**ailure), one on the effect of treatment with *omecamtiv mecarbil* in patients with heart failure with reduced ejection fraction (HFrEF) and low blood pressure, and the other on the impact of tricuspid regurgitation on clinical outcomes in patients with HFrEF.

Title: GALACTIC-HF - *Omecamtiv Mecarbil* in HFrEF and Low Blood Pressure
Presenter: Marco Metra, M.D., Professor of Cardiology & Director of the Institute of Cardiology, Department of Medical & Surgical Specialties, Radiological Sciences & Public Health, University & Civil Hospitals of Brescia, Italy
Date: May 22, 2022
Topic: Chronic Heart Failure
Session Title: Late-Breaking Trials - Pharmacological Treatment I
Session Type: Late Breaking Science
Presentation Time: 12:13 PM CEST
Location: Room 1

Title: REDWOOD-HCM OLE: *Aficamten* in HCM
Presenter: Ahmad Masri, M.D., Assistant Professor of Medicine, Division of Cardiovascular Medicine, School of Medicine, Oregon Health & Science University
Date: May 23, 2022
Topic: Chronic Heart Failure
Session Title: Late-Breaking Trials – Registries and Trial Updates II
Session Type: Late Breaking Science
Presentation Time: 5:10 PM CEST
Location: Room 3

Title: GALACTIC-HF and Tricuspid Regurgitation
Presenter: Marianna Adamo, M.D., Interventional Cardiologist, University of Brescia, Italy
Date: May 23, 2022
Topic: Chronic Heart Failure
Session Title: Late-Breaking Trials – Registries and Trial Updates II
Session Type: Late Breaking Science
Presentation Time: 5:30 PM CEST
Location: Room 3

About Cytokinetics

Cytokinetics is a late-stage biopharmaceutical company focused on discovering, developing and commercializing first-in-class muscle activators and next-in-class muscle inhibitors as potential treatments for debilitating diseases in which 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 muscle function and contractility. Cytokinetics is readying for the potential commercialization of *omecamtiv mecarbil*, its cardiac muscle activator, following positive results from GALACTIC-HF, a large, international Phase 3 clinical trial in patients with heart failure. Cytokinetics is also developing *aficamten*, a next-generation cardiac myosin inhibitor, currently the subject of SEQUOIA-HCM, the Phase 3 clinical trial of *aficamten* in patients with symptomatic obstructive hypertrophic cardiomyopathy (HCM). *Aficamten* is also being evaluated in non-obstructive HCM in Cohort 4 of the Phase 2 clinical trial, REDWOOD-HCM. Cytokinetics is also developing *reldesemtiv*, an investigational fast skeletal muscle troponin activator, currently the subject of COURAGE-ALS, a Phase 3 clinical trial in patients with amyotrophic lateral sclerosis (ALS). Cytokinetics continues its over 20-year history of pioneering innovation in muscle biology and related pharmacology focused to diseases of muscle dysfunction and conditions of muscle weakness.

For additional information about Cytokinetics, visit www.cytokinetics.com and follow us on Twitter, 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 *omecamtiv mecarbil*, *aficamten*, 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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Contact:
Cytokinetics
Diane Weiser
Senior Vice President, Corporate Communications, Investor Relations
(650) 624-3071

Source: Cytokinetics, Incorporated