Cytokinetics Appoints Edward M. Kaye, M.D., to Board of Directors

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SOUTH SAN FRANCISCO, Calif., May 20, 2016 (GLOBE NEWSWIRE) -- Cytokinetics, Inc. (Nasdaq:CYTK) announced today that Edward M. Kaye, M.D., has been appointed to the company's Board of Directors. Dr. Kaye is the Interim Chief Executive Officer and Senior Vice President, Chief Medical Officer of Sarepta Therapeutics.

Dr. Kaye joins the Cytokinetics Board of Directors with substantial industry experience in rare diseases including neurological conditions. He has served as Interim Chief Executive Officer of Sarepta since 2015 and as the SVP, Chief Medical Officer since 2011. Prior to joining Sarepta, Dr. Kaye was employed by Genzyme Corporation for ten years, holding various senior management positions, the most recent of which was Group Vice President of Clinical Development, in which he supervised clinical research in lysosomal storage disease programs and genetic neurological disorders. Previously, Dr. Kaye served as Chief of Biochemical Genetics at Children's Hospital of Philadelphia and Associate Professor of Neurology and Pediatrics at the University Of Pennsylvania School Of Medicine. Dr. Kaye serves as a Neurological Consultant at the Children's Hospital of Boston and is on the editorial boards of a number of medical journals. He is also a member of several scientific advisory boards, including United Leukodystrophy Foundation, Spinal Muscular Atrophy Foundation, CureCMD, CureDuchenne and Prize4Life. Dr. Kaye received his medical education and pediatric training at Loyola University Stritch School of Medicine and University Hospital, child neurology training at Boston City Hospital, Boston University, and completed his training as a neurochemical research fellow at Bedford VA Hospital, Boston University.

"We are pleased to welcome Ed to our Board. His expertise in rare neuromuscular diseases is highly relevant as we advance our clinical programs in ALS and SMA into late-stage development," said Robert I. Blum, Cytokinetics' President and Chief Executive Officer. "Ed has overseen many clinical research and development programs, and we look forward to the benefit of his industry-leading experience."

About Cytokinetics

Cytokinetics is a late-stage biopharmaceutical company focused on discovering, developing and commercializing first-in-class muscle activators as potential treatments for debilitating diseases in which muscle performance is compromised and/or declining. As a leader in muscle biology and the mechanics of muscle performance, the company is developing small molecule drug candidates specifically engineered to increase muscle function and contractility. Cytokinetics' lead drug candidate is *tirasemtiv*, a fast skeletal muscle troponin activator, for the potential treatment of ALS. *Tirasemtiv* has been granted orphan drug designation and fast track status by the U.S. Food and Drug Administration and orphan medicinal product designation by the European Medicines Agency for the potential treatment of ALS. Cytokinetics retains the right to develop and commercialize *tirasemtiv*. Cytokinetics is collaborating with Amgen Inc. to develop *omecamtiv mecarbil*, a novel cardiac muscle activator, for the potential treatment of heart failure. Cytokinetics is collaborating with Astellas Pharma Inc. to develop CK-2127107, a fast skeletal muscle activator, for the potential treatment of spinal muscular atrophy and chronic obstructive pulmonary disease. Amgen holds an exclusive license worldwide to develop and commercialize *omecamtiv mecarbil* and Astellas holds an exclusive license worldwide to develop and commercialize CK-2127107. Both licenses are subject to Cytokinetics' specified development and commercialization participation rights. For additional information about Cytokinetics, visit http://www.cytokinetics.com/.

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 the potential advancement of its drug candidates in late-stage clinical trials. 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 approval and production of Cytokinetics' drug candidates and potential drug candidates that could slow or prevent clinical development or product approval, including risks that current and past results of clinical trials or preclinical studies may not be indicative of future clinical trials results and that Cytokinetics' drug candidates and potential drug candidates may have unexpected adverse side effects or inadequate therapeutic efficacy. 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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Cytokinetics, Inc