

Cytokinetics Announces Presentations Relating to Tirasemtiv at the 25th International Symposium on ALS/MND

December 1, 2014 9:00 PM EST

South San Francisco, CA, December 1, 2014 - Cytokinetics, (Nasdaq: CYTK) announced today that presentations relating to *tirasemtiv* are scheduled during the 25th International Symposium on ALS/MND to be held December 5-7, 2014 at the Square Brussels Meeting Centre in Brussels, Belgium. *Tirasemtiv* is the lead drug candidate from Cytokinetics' skeletal muscle contractility program. *Tirasemtiv* selectively activates the fast skeletal muscle troponin complex by increasing its sensitivity to calcium and, in preclinical studies and early clinical trials, demonstrated increases in skeletal muscle force in response to neuronal input and delays in the onset and reductions in the degree of muscle fatigue. *Tirasemtiv* was the subject of BENEFIT-ALS (Blinded Evaluation of Neuromuscular Effects and Functional Improvement with *T*irasemtiv ALS), a Phase IIb, multi-national, double-blind, randomized, placebo-controlled, clinical trial designed to evaluate the safety, tolerability and efficacy of *tirasemtiv* in patients with ALS.

Oral Presentation at the 25th International Symposium on ALS/MND

 Date:
 Friday, December 5, 2014

 Location:
 Square Brussels Meeting Centre, Copper Hall

 Presentation Time:
 2:40 PM-3:00 PM (Central European Time)

 Session:
 3B - Trials and Trial Design

 Title:
 The Effects of *Tirasemtiv* on Measures of Respiratory Function in Amyotrophic Lateral Sclerosis

 Presenter:
 Jinsy Andrews, MD, MSc, Director, Clinical Research and Development, Cytokinetics

Poster Presentations at the 25th International Symposium on ALS/MND

 Date:
 Friday, December 5, 2014

 Location:
 Square Brussels Meeting Centre, Grand Hall 1

 Presentation Time:
 5:30 PM - 7:30 PM

 Session:
 Poster Session A

 Theme:
 6 - Epidemiology

 Poster Number:
 P133

 Title:
 Profile of Medical Care Costs in Patients with Amyotrophic Lateral Sclerosis in Medicare Program and Under Commercial Insurance

 Presenter:
 Lisa Meng, PhD, Director, Biometrics, Cytokinetics

 Poster Moderated:
 6:00 PM - 6:30 PM (Central European Time)

Date: Saturday, December 6, 2014 Location: Square Brussels Meeting Centre, Grand Hall 1 Presentation Time: 5:30 PM - 7:30 PM Session: Poster Session B Theme: 4 - Respiratory and Nutritional Management Poster Number: P82 Title: Effect of *Tirasemtiv* on Submaximal Rodent Diaphragm Strength and Respiratory Function Presenter: Darren Hwee, PhD, Scientist, Cytokinetics Poster Moderated: 5:45 PM - 6:20 PM (Central European Time)

 Date:
 Saturday, December 6, 2014

 Location:
 Square Brussels Meeting Centre, Grand Hall 1

 Presentation Time:
 5:30 PM - 7:30 PM

 Session:
 Poster Session B

 Theme:
 11 - Therapeutic Strategies

 Poster Number:
 P283

 Title:
 Relationships Between *Riluzole* and *Tirasemtiv* Levels on Outcomes in the BENEFIT-ALS Trial

 Presenter:
 Jeremy Shefner, MD, PhD, Professor & Chair of Neurology, Barrow Neurological Institute

 Poster Moderated:
 6:20 PM - 6:55 PM (Central European Time)

Date: Saturday, December 6, 2014 Location: Square Brussels Meeting Centre, Grand Hall 1 Presentation Time: 5:30 PM - 7:30 PM Session: Poster Session B Theme: 11 - Therapeutic Strategies Poster Number: P286 Title: Fast Skeletal Muscle Troponin Activator *Tirasemtiv* Increases Muscle Function and Performance in Mouse Models of Spinal Muscular Atrophy Presenter: Darren Hwee, PhD, Scientist, Cytokinetics Poster Moderated: 6:20 PM - 6:55 PM (Central European Time)

About Cytokinetics

Cytokinetics is a clinical-stage biopharmaceutical company focused on the discovery and development of novel small molecule therapeutics that modulate muscle function for the potential treatment of serious diseases and medical conditions. Cytokinetics' lead drug candidate from its cardiac muscle contractility program, *omecantiv mecarbil*, is in Phase II clinical development for the potential treatment of heart failure. Amgen Inc. holds an exclusive license worldwide to develop and commercialize *omecantiv mecarbil* and related compounds, subject to Cytokinetics' specified development and commercialization participation rights. Cytokinetics is independently developing *tirasemtiv*, a fast skeletal muscle activator, as a potential treatment for diseases and medical conditions associated with neuromuscular dysfunction. *Tirasemtiv* is the subject of a Phase II clinical trials program and 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 amyotrophic lateral sclerosis (ALS). Cytokinetics is collaborating

with Astellas Pharma Inc. to develop CK-2127107, a skeletal muscle activator structurally distinct from *tirasemtiv*, for non-neuromuscular indications. All of these drug candidates have arisen from Cytokinetics' muscle biology focused research activities and are directed towards the cytoskeleton. The cytoskeleton is a complex biological infrastructure that plays a fundamental role within every human cell. Additional information about Cytokinetics can be obtained at www.cytokinetics.com.

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 Cytokinetics' research and development activities, including the potential significance and utility of the results from preclinical studies and clinical trials of tirasemtiv; planned further analyses of the results from BENEFIT-ALS and the potential outcomes of such analyses; potential further development of tirasemtiv; planned presentations; and the properties and potential benefits of skeletal muscle activators and of tirasemtiv and 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 approval and production of Cytokinetics' drug candidates and potential drug candidates and potential development, testing results results and that Cytokinetics' drug candidates and potential drug candidates 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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