NEWS RELEASE

Sarepta Therapeutics to Present New Data from its Neuromuscular Portfolio at 2024 World Muscle Society Congress

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CAMBRIDGE, Mass.--(BUSINESS WIRE)-- Sarepta Therapeutics, Inc. (NASDAQ:SRPT), the leader in precision genetic medicine for rare diseases, will present at the 29 th Annual Congress of the World Muscle Society 2024 Congress (WMS 2024), taking place Oct. 8-12, 2024, in Prague, Czechia.

Among the data to be presented are new safety and efficacy results from several studies in the delandistrogene moxeparvovec clinical development program, including data from Studies 9001-101, 9001-103 (ENDEAVOR) and 9001-301 (EMBARK).

“New data to be presented at WMS from clinical studies of delandistrogene moxeparvovec include the first look at skeletal muscle and cardiac MRI outcomes from the EMBARK study, a late-breaking analysis of safety data from across all delandistrogene moxeparvovec clinical studies, and five-year functional results from Study SRP-9001-101, the longest-term data to date for a gene therapy in Duchenne,” said Louise Rodino-Klapac, Ph.D., executive vice president, chief scientific officer and head of research and development, Sarepta Therapeutics. “The results add to the substantial and growing body of data for delandistrogene moxeparvovec.”

Podium Presentation:

19O: Muscle MRI outcomes in patients with Duchenne Muscular Dystrophy treated with delandistrogene moxeparvovec: Findings from EMBARK Part 1	Oct. 12, 2024 1:45-2:00 a.m. EDT (07:45-08:00 CEST)

Late Breaking Poster:

726LBP: Long-term safety and tolerability of delandistrogene moxeparvovec in Duchenne muscular dystrophy: phase 1 to phase 3 clinical trials	Oct. 11, 2024 9:45-10:45 a.m. EDT (15:45-16:45 CEST)

Poster Presentations (*Denotes encore presentation)

391P: Agreement and accuracy of ambulatory definitions in Duchenne muscular dystrophy (DMD): a cross-sectional analysis*	Oct. 9, 2024 8:30-9:30 a.m. EDT (14:30-15:30 CEST)
94P: JOURNEY: a natural history study of Limb Girdle muscular dystrophies R3-R5: baseline characteristics of study cohort	Oct. 9, 2024 11:15 a.m.-12:15 p.m. EDT (17:15-18:15 CEST)
423P: Safety and efficacy of delandistrogene moxeparvovec versus placebo in Duchenne muscular dystrophy: Phase 3 EMBARK primary results*	Oct. 9, 2024 11:15 a.m.-12:15 p.m. EDT (17:15-18:15 CEST)
424P: Micro-dystrophin expression and safety with delandistrogene moxeparvovec gene therapy for DMD in a broad population: Phase 1B trial (ENDEAVOR)	Oct. 9, 2024 11:15 a.m.-12:15 p.m. EDT (17:15-18:15 CEST)
425P: Five-year outcomes with delandistrogene moxeparvovec in patients with Duchenne Muscular dystrophy (DMD): a phase 1/2a study	Oct. 9, 2024 11:15 a.m.-12:15 p.m. EDT (17:15-18:15 CEST)
428P: Cardiac MRI outcomes in patients with Duchenne Muscular Dystrophy treated with delandistrogene moxeparvovec: Findings from EMBARK Part 1	Oct. 9, 2024 11:15 a.m.-12:15 p.m. EDT (17:15-18:15 CEST)

The full WMS 2024 program is available at **https://www.wms2024.com/page/programme** .

About Sarepta Therapeutics

Sarepta is on an urgent mission: engineer precision genetic medicine for rare diseases that devastate lives and cut futures short. We hold leadership positions in Duchenne muscular dystrophy (DMD) and limb-girdle muscular dystrophies (LGMDs), and we currently have more than 40 programs in various stages of development. Our vast pipeline is driven by our multi-platform Precision Genetic Medicine Engine in gene therapy, RNA and gene editing. For more information, please visit **www.sarepta.com** or follow us on **LinkedIn** , **X (formerly Twitter)** , **Instagram** and **Facebook** .

Internet Posting of Information

We routinely post information that may be important to investors in the 'For Investors' section of our website at **www.sarepta.com** . We encourage investors and potential investors to consult our website regularly for important information about us.

Forward-Looking Statements

This press release contains forward-looking statements. Any statements contained in this press release that are not statements of historical fact may be deemed to be forward-looking statements. Words such as "believes," "anticipates," "plans," "expects," "will," "intends," "potential," "possible" and similar expressions are intended to identify forward-looking statements. These forward-looking statements include statements related to our research and development programs, technologies, scientific approaches, and product and product candidates, including potential benefits related to safety and efficacy, and our mission.

These forward-looking statements involve risks and uncertainties, many of which are beyond Sarepta's control. Known risk factors include, among others: Sarepta may not be able to execute on its business plans, including meeting its expected or planned regulatory milestones and timelines, clinical development plans, and bringing its products to markets for various reasons including possible limitations of Company financial and other resources, manufacturing limitations that may not be anticipated or resolved for in a timely manner, and regulatory, court or agency decisions, such as decisions by the United States Patent and Trademark Office with respect to patents that cover Sarepta's product candidates; the results of future research may not be consistent with past positive results or may fail to meet regulatory approval requirements for the safety and efficacy of product candidates; different methodologies, assumptions and applications we use to assess particular safety or efficacy parameters may yield different statistical results, and even if we believe the data collected from clinical trials of our product candidates are positive, these data may not be sufficient to support approval by the FDA or other global regulatory authorities; and those risks identified under the heading "Risk Factors" in Sarepta's most recent Annual Report on Form 10-K for the year ended December 31, 2023, and most recent Quarterly Report on Form 10-Q filed with the Securities and Exchange Commission (SEC) as well as other SEC filings made by the Company which you are encouraged to review.

Any of the foregoing risks could materially and adversely affect the Company's business, results of operations and the trading price of Sarepta's common stock. For a detailed description of risks and uncertainties Sarepta faces, you are encouraged to review the SEC filings made by Sarepta. We caution investors not to place considerable reliance on the forward-looking statements contained in this press release. Sarepta does not undertake any obligation to publicly update its forward-looking statements based on events or circumstances after the date hereof, except as required by law.

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