Eagle Pharmaceuticals Announces Acceptance of Two Abstracts at the Society for Academic Emergency Medicine (SAEM) Annual Meeting

May 11, 2023

Abstracts to be presented cover preclinical data on RYANODEX® (dantrolene sodium) for injectable suspension in Acute Radiation Syndrome and Traumatic Brain Injury

WOODCLIFF LAKE, N.J., May 11, 2023 (GLOBE NEWSWIRE) -- Eagle Pharmaceuticals, Inc. (Nasdaq: EGRX) ("Eagle" or the "Company") today announced that two abstracts were accepted for presentation at the Society for Academic Emergency Medicine (SAEM) Annual Meeting, being held May 16-19, 2023, in Austin, Texas.

"We are pleased to have the opportunity to present this investigative work at such a prestigious forum dedicated to original research in academic emergency medicine. Eagle is actively engaged with the scientific community, as we focus on developing innovative medicines to address unmet medical needs. RYANODEX plays an important role in the treatment of malignant hyperthermia, today," stated Scott Tarriff, President and Chief Executive Officer of Eagle Pharmaceuticals.

Details of the presentations are as follows:

Abstract: 207
Title: **Ryanodex-Dantrolene Sodium for Injectable Suspension Improves Survival in Mouse Model of Acute Radiation Syndrome**
Date: Wednesday May 17, 2023
Time: 4:16 PM-4:24 PM CDT
Location: Room 307: Level Three

Abstract: 359
Title: **Ryanodex® Reduces Persistent Hippocampal Effects of Single Mild Traumatic Brain Injury in Rats**
Date: Thursday May 18, 2023
Time: 2:38 PM-2:46 PM CDT
Location: Room 208: Level Two

About Eagle Pharmaceuticals, Inc.
Eagle is a fully integrated pharmaceutical company with research and development, clinical, manufacturing and commercial expertise. Eagle is committed to developing innovative medicines that result in meaningful improvements in patients' lives. Eagle's commercialized products include vasopressin, PEMFEXY®, RYANODEX®, BENDEKA®, BELRAPZO®, TREAKISYM® (Japan), and BYFAVO® and BARHEMSYS® through its wholly owned subsidiary Acacia Pharma Inc. Eagle's oncology and CNS/metabolic critical care pipeline includes product candidates with the potential to address underserved therapeutic areas across multiple disease states. Additional information is available on Eagle's website at www.eagleus.com.

Forward-Looking Statements
This press release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, as amended, and other securities law. Forward-looking statements are statements that are not historical facts. Words and phrases such as "anticipated," "forward," "will," "would," "could," "should," "may," "remain," "potential," "prepare," "expected," "believe," "plan," "future," "believe," "guidance," "project," "estimate," "intend," "advance," "continue" and similar expressions are intended to identify forward-looking statements. These statements include, but are not limited to, statements with respect to: the Company's ability to develop innovative medicines that address unmet medical needs; RYANODEX role in the treatment of malignant hyperthermia; and RYANODEX potential development for other indications. All of such statements are subject to certain risks and uncertainties, many of which are difficult to predict and generally beyond the Company's control, that could cause actual results to differ materially from those expressed in, or implied or projected by, the forward-looking information and statements. Such risks and uncertainties include, but are not limited to: the risk that the anticipated benefits of the Company's acquisition of Acacia are not realized; the ability of Enalare to achieve milestones and deliverables and achieve successful results in the development of ENA-001 and the Company's ability to exercise its option to acquire the remaining outstanding share capital of Enalare; the impacts of the continuing effects of the COVID-19 pandemic and geopolitical events such as the conflict in Ukraine, including disruption or impact in the sales of the Company's marketed products, interruptions or other adverse effects to clinical trials, delays in regulatory review, manufacturing and supply chain interruptions, adverse effects on healthcare systems, disruption in the operations of the Company's third party partners and disruption of the global economy or other events on the Company's business, financial condition and results of operations; macroeconomic conditions, including rising inflation and interest rates, uncertain credit and financial markets and recent and potential disruptions in banking systems; whether the Company will incur unforeseen expenses or liabilities or other market factors; whether the Company will successfully implement its development plan for its product candidates; delay in or failure to obtain regulatory approval of the Company's or its partners' product candidates; whether the Company can successfully market and commercialize its product candidates; the success of the Company's relationships with its partners; the availability and pricing of third party sourced products and materials; the outcome of litigation involving any of its products or that may have an impact on any of our products; successful compliance with the FDA and other governmental regulations applicable to product approvals, manufacturing facilities, products and/or businesses; general economic conditions, including the potential adverse effects of public health issues, including the COVID-19 pandemic and geopolitical events, on economic activity and the performance of the financial markets generally; the strength and enforceability of the Company's intellectual property rights or the rights of third parties; competition from other pharmaceutical and biotechnology companies and the potential for competition from generic entrants into the market; the risks inherent in the early stages of drug development and in conducting clinical trials; and any unanticipated factors in addition to the foregoing that may impact the Company's financial and business projections and guidance and may cause the Company's actual results and outcomes to materially differ from its projections and guidance;

and those risks and uncertainties identified in the “Risk Factors” sections of the Company's Annual Report on Form 10-K for the year ended December 31, 2022, filed with the Securities and Exchange Commission (the “SEC”) on March 23, 2023, and its other subsequent filings with the SEC, including the Company’s Quarterly Report on Form 10-Q for the quarter ended March 31, 2023, filed with the SEC on May 9, 2023. Readers are cautioned not to place undue reliance on these forward-looking statements. All forward-looking statements contained in this press release speak only as of the date on which they were made. Except to the extent required by law, the Company undertakes no obligation to update such statements to reflect events that occur or circumstances that exist after the date on which they were made.

Investor Relations for Eagle Pharmaceuticals, Inc.:
Lisa M. Wilson
In-Site Communications, Inc.
T: 212-452-2793
E: lwilson@insitecony.com