Stemline Therapeutics to Present at the 2017 Wedbush PacGrow Healthcare Conference

August 14, 2017

NEW YORK, Aug. 14, 2017 (GLOBE NEWSWIRE) -- Stemline Therapeutics, Inc. (Nasdaq:STML), a clinical-stage biopharmaceutical company developing novel therapeutics for difficult to treat cancers, announced today that Ivan Bergstein, M.D., Stemline's CEO, will present at the 2017 Wedbush PacGrow Healthcare Conference on Tuesday, August 15, 2017 at 8:00 AM ET. The conference is being held at the Le Parker Meridien in New York, NY. A webcast of the presentation can be viewed on the company's website at www.stemline.com.

About Stemline Therapeutics
Stemline Therapeutics, Inc. is a clinical stage biopharmaceutical company developing novel therapeutics for difficult to treat cancers. SL-401 is a targeted therapy directed to the interleukin-3 receptor (CD123), a cell surface receptor overexpressed on BPDCN and a variety of other hematologic cancers. SL-401 was granted Breakthrough Therapy Designation (BTD) for the treatment of patients with BPDCN. A pivotal Phase 2 trial with SL-401 in BPDCN has completed enrollment in Stages 1, 2 and 3 of the trial; to ensure ongoing patient access to SL-401, an additional cohort, Stage 4, is enrolling. Additional Phase 1/2 trials are assessing SL-401 in patients with other malignancies including myeloproliferative neoplasms (MPN) (focused on chronic myelomonocytic leukemia [CMML] and myelofibrosis [MF]), acute myeloid leukemia (AML), and multiple myeloma, as a single agent or in combination with other agents. SL-801, a novel oral small molecule reversible inhibitor of XPO1, is enrolling patients with advanced solid tumors in a Phase 1 trial. SL-701, an immunotherapeutic, has completed dosing of patients with second-line glioblastoma in a Phase 2 trial and patients are being followed for outcomes including survival.

Forward-Looking Statements
Some of the statements included in this press release may be forward-looking statements that involve a number of risks and uncertainties. For those statements, we claim the protection of the safe harbor for forward-looking statements contained in the Private Securities Litigation Reform Act of 1995. The factors that could cause our actual results to differ materially include: the success and timing of our clinical trials and preclinical studies for our product candidates, including site initiation, institutional review board approval, scientific review committee approval, patient accrual, safety, tolerability and efficacy data observed, and input from regulatory authorities including the risk that the FDA or other ex-U.S. national drug authority ultimately does not approve any of our product candidates; our plans to develop and commercialize our product candidates; market acceptance of our products; reimbursement available for our products; our available cash and investments; our ability to obtain and maintain intellectual property protection for our product candidates; our ability to manufacture; the performance of third-party manufacturers, clinical research organizations, clinical trial sponsors and clinical trial investigators; and other risk factors identified from time to time in our reports filed with the Securities and Exchange Commission. Any forward-looking statements set forth in this press release speak only as of the date of this press release. We do not intend to update any of these forward-looking statements to reflect events or circumstances that occur after the date hereof.

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