



Spyre Therapeutics Announces Poster Presentations at United European Gastroenterology Week (UEGW) 2025

October 5, 2025

WALTHAM, Mass., Oct. 05, 2025 (GLOBE NEWSWIRE) -- Spyre Therapeutics, Inc. (NASDAQ: SYRE), a clinical-stage biotechnology company advancing best-in-class antibody engineering, dose optimization, and rational therapeutic combinations for the treatment of Inflammatory Bowel Disease ("IBD") and other immune-mediated diseases, today announced scientific presentations at the UEGW Congress.

"We are excited to share follow-up data out to six months from our Phase 1 study of SPY002, our potential best-in-class anti-TL1A agent in development for IBD. The data continue to show SPY002 is well tolerated, has a differentiated PK profile supporting quarterly or twice-yearly dosing, and suppresses free TL1A through 24 weeks," said Josh Friedman, M.D., Ph.D., SVP of Clinical Development at Spyre. "Additionally, we are pleased to share new preclinical data demonstrating that each of our combination programs ($\alpha 4\beta 7$ + TL1A, $\alpha 4\beta 7$ + IL-23, and TL1A + IL-23) exhibit superior efficacy relative to constituent monotherapies in rodent TNBS-induced colitis models, providing additional validation for our ongoing SKYLINE-UC Phase 2 study."

The poster will be available for viewing during the UEGW Congress, and details are as follows:

Title: Interim Phase 1 Results for SPY002, a Novel Half-Life Extended Monoclonal Antibody Targeting TL1A, Suggests a Potential for Q3M or Q6M Maintenance Dosing for Inflammatory Bowel Disease

Authors: Y. Vugmeyster, S. Sloan, JD Lu, K. Hew, P. Patel, C. Sheldon, D. Nguyen, R. McLean, M. Huyghe, B. Connolly, B. Wang, M. Kennedy, M. Rose, E. Svejnoha, J. Friedman

Title: Combined Inhibition of Integrin $\beta 7$ and TL1A, Integrin $\beta 7$ and IL-23, or TL1A and IL-23 Are Superior to Their Constituent Monotherapies in Mouse TNBS-Induced Colitis

Authors: M. Siegel, J. Friedman, E. Lewis, D. Giles, A. Spencer

Full session details can be accessed via the UEGW program.

About Spyre Therapeutics

Spyre Therapeutics is a clinical-stage biotechnology company that aims to create next-generation inflammatory bowel disease (IBD) and other immune-mediated disease products by combining best-in-class antibody engineering, dose optimization, and rational therapeutic combinations. Spyre's pipeline includes investigational extended half-life antibodies targeting $\alpha 4\beta 7$, TL1A, and IL-23.

For more information, please visit <http://spyre.com>.

Forward-Looking Statements

Certain statements in this press release, other than purely historical information, may constitute "forward-looking statements" within the meaning of the federal securities laws, including for purposes of the safe harbor provisions under the United States Private Securities Litigation Reform Act of 1995, concerning Spyre and other matters. These forward-looking statements include, but are not limited to, express or implied statements relating to Spyre's management team's expectations, hopes, beliefs, intentions or strategies regarding the future including, without limitation, Spyre's ability to achieve the expected benefits or opportunities with respect to its pipeline of product candidates such as the potential efficacy, tolerability, convenience, commercial viability, dosing regimen and safety profile of SPY002 in humans, including the potential for a quarterly or twice yearly dosing profile; the potential for SPY002 to become a best-in-class therapy for IBD; the potential consistency of the SPY002 Phase 2 trial final data readouts with interim Phase 1 results; and the potential therapeutic benefits of Spyre's product candidates as monotherapies or in combinations and their extended half-life, including the expected duration of half-life in comparison to competitor products and the potential potency, efficacy and convenience compared to today's standard of care. In addition, any statements that refer to projections, forecasts or other characterizations of future events or circumstances, including any underlying assumptions, are forward-looking statements. The words "opportunity," "potential," "milestones," "pipeline," "can," "goal," "aim," "strategy," "target," "seek," "anticipate," "achieve," "believe," "contemplate," "continue," "could," "estimate," "expect," "intends," "may," "might," "plan," "possible," "predict," "project," "should," "will," "would," and similar expressions (including the negatives of these terms or variations of them) may identify forward-looking statements, but the absence of these words does not mean that a statement is not forward-looking. These forward-looking statements are based on current expectations and beliefs concerning future developments and their potential effects. There can be no assurance that future developments affecting Spyre will be those that have been anticipated. These forward-looking statements involve a number of risks, uncertainties (some of which are beyond Spyre's control) or other assumptions that may cause actual results or performance to be materially different from those expressed or implied by these forward-looking statements. These risks and uncertainties include, but are not limited, uncertainties and risks arising from regulatory feedback, including potential disagreement by regulatory authorities with the Company's interpretation of data and the Company's clinical trials for its product candidates, including our plans for and timing of

cohort initiation for combination therapy arms for the ongoing SKYLINE-UC Phase 2 platform trial across different jurisdictions; the potential for final data not being consistent with or different than the interim data reported for our programs; the potential impact of Trump Administration policies and changes in law on our business; and those uncertainties and factors described under the heading "Risk Factors," "Risk Factor Summary" and "Note about Forward-Looking Statements" in Spyre's most recent Annual Report on Form 10-K, as supplemented and updated by subsequent Quarterly Reports on Form 10-Q and Current Reports on Form 8-K that the Company has filed or will file with the SEC, as well as discussions of potential risks, uncertainties, and other important factors included in other filings by Spyre from time to time. Should one or more of these risks or uncertainties materialize, or should any of Spyre's assumptions prove incorrect, actual results may vary in material respects from those projected in these forward-looking statements. Nothing in this press release should be regarded as a representation by any person that the forward-looking statements set forth therein will be achieved or that any of the contemplated results of such forward-looking statements will be achieved. You should not place undue reliance on forward-looking statements in this press release, which speak only as of the date they are made and are qualified in their entirety by reference to the cautionary statements herein. Spyre does not undertake or accept any duty to make any updates or revisions to any forward-looking statements. This press release does not purport to summarize all of the conditions, risks and other attributes of an investment in Spyre.

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