Syndax Pharmaceuticals Announces Presentations at the 2018 American Society of Clinical Oncology Annual Meeting

WALTHAM, Mass., April 25, 2018 (PRNEWSWIRE) -- Syndax Pharmaceuticals, Inc. ("Syndax," the "Company" or "we") (Nasdaq: SNDX), a clinical stage biopharmaceutical company developing an innovative pipeline of cancer therapies, today announced five poster presentations at the upcoming American Society of Clinical Oncology (ASCO) Annual Meeting being held June 1-5, 2018 in Chicago, Illinois.

Presentation Details

Title: ADVL1513: Results of a phase 1 trial of entinostat, an oral histone deacetylase inhibitor, in pediatric patients with recurrent or refractory solid tumors
First Author: Suman Malempati, MD, Oregon Health and Science University
Abstract Number: 10556
Poster Session: Pediatric Oncology
Poster Board: 229
Date and Time: Saturday, June 2, 2018, 8:00-11:30 AM CT, Hall A

Title: ENCORE 601: A phase 2 study of entinostat in combination with pembrolizumab in patients with microsatellite stable metastatic colorectal cancer
First Author: Nilofer Saba Azad, MD, Sidney Kimmel Cancer Center at Johns Hopkins University
Abstract Number: 3557
Poster Session: Gastrointestinal (Colorectal) Cancer
Poster Board: 50
Date and Time: Sunday, June 3, 2018, 8:00-11:30 AM CT, Hall A

Title: Efficacy and safety of entinostat (ENT) and pembrolizumab (PEMBRO) in patients with non-small cell lung cancer (NSCLC) previously treated with anti-PD-(L)1 therapy
First Author: Leena Gandhi, MD, PhD, NYU Perlmutter Cancer Center
Abstract Number: 9036
Poster Session: Lung Cancer—Non-Small Cell Metastatic
Poster Board: 359
Date and Time: Sunday, June 3, 2018, 8:00-11:30 AM CT, Hall A

Title: Entinostat in combination with nivolumab for patients with advanced cholangiocarcinoma and pancreatic adenocarcinoma
First Author: Marina Baretti, MD, Sidney Kimmel Cancer Center at Johns Hopkins University
Abstract Number: TPS4151
Poster Session: Gastrointestinal (Noncolorectal) Cancer
Poster Board: 330a
Date and Time: Sunday, June 3, 2018, 8:00-11:30 AM CT, Hall A

Title: Efficacy and safety of entinostat (ENT) and pembrolizumab (PEMBRO) in patients with melanoma progressing on or after a PD-1/L1 blocking antibody
First Author: Sanjiv S. Agarwala, MD, St. Luke's Hospital
Abstract Number: 9530

Poster Session: Melanoma/Skin Cancers
Poster Board: 357
Date and Time: Monday, June 4, 2018, 1:15-4:45 PM CT, Hall A

About Syndax Pharmaceuticals, Inc.

Syndax Pharmaceuticals is a clinical stage biopharmaceutical company developing an innovative pipeline of cancer therapies. The Company is developing its lead product candidate, entinostat, a once-weekly, oral, small molecule, class I HDAC inhibitor, in combination with exemestane and several approved PD-1/PD-L1 antagonists. The Company's pipeline also includes SNDX-6352, a monoclonal antibody that blocks the colony stimulating factor 1 (CSF-1) receptor, as well as a portfolio of potent and selective inhibitors targeting the binding interaction of Menin with MLLr. For more information, please visit www.syndax.com or follow the Company on Twitter and LinkedIn.

Syndax's Cautionary Note on Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Words such as "may," "will," "expect," "plan," "anticipate," "estimate," "intend," "believe" and similar expressions (as well as other words or expressions referencing future events, conditions or circumstances) are intended to identify forward-looking statements. These forward-looking statements are based on Syndax's expectations and assumptions as of the date of this press release. Each of these forward-looking statements involves risks and uncertainties. Actual results may differ materially from these forward-looking statements. Forward-looking statements contained in this press release include, but are not limited to, statements about the progress, timing, clinical development and scope of clinical trials and the reporting of clinical data for Syndax's product candidates, and the potential use of our product candidates to treat various cancer indications. Many factors may cause differences between current expectations and actual results including unexpected safety or efficacy data observed during preclinical or clinical studies, clinical trial site activation or enrollment rates that are lower than expected, changes in expected or existing competition, changes in the regulatory environment, failure of Syndax's collaborators to support or advance collaborations or product candidates and unexpected litigation or other disputes. Other factors that may cause Syndax's actual results to differ from those expressed or implied in the forward-looking statements in this press release are discussed in Syndax's filings with the U.S. Securities and Exchange Commission, including the "Risk Factors" sections contained therein. Except as required by law, Syndax assumes no obligation to update any forward-looking statements contained herein to reflect any change in expectations, even as new information becomes available.

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