

April 17, 2019



# Syros to Host Key Opinion Leader Symposium on CDK7 Inhibition on April 24, 2019

CAMBRIDGE, Mass.--(BUSINESS WIRE)-- Syros Pharmaceuticals (NASDAQ:SYRS), a leader in the development of medicines that control the expression of genes, today announced that it will host a key opinion leader (KOL) breakfast symposium focused on the unmet need, treatment landscape and opportunities for selective cyclin-dependent kinase 7 (CDK7) inhibition in ovarian and breast cancers on Wednesday, April 24, 2019 from 8:30 – 10:30 a.m. ET in New York City. Syros is currently evaluating SY-1365, a first-in-class selective CDK7 inhibitor, in a Phase 1 clinical trial as a single agent and in combination with other therapies in multiple ovarian and breast cancer patient populations.

The event will feature presentations from Dejan Juric, M.D., Medical Oncologist and Director at the Termeer Center for Targeted Therapies, Massachusetts General Hospital and Charles A. Leath, III, M.D., M.S.P.H., Professor in the Division of Gynecologic Oncology and Ellen Gregg Shook Culverhouse Chair in Gynecologic Oncology at the University of Alabama Medical Center. Additionally, members of the Syros management team will provide an overview of SY-1365, including preclinical and clinical data supporting its ongoing development in ovarian and breast cancer patients, and SY-5609, its oral selective CDK7 inhibitor, which is expected to enter a Phase 1 oncology study in early 2020.

A live webcast of the event will be available on the Investors & Media section of the Syros website at [www.syros.com](http://www.syros.com). An archived replay of the webcast will be available for approximately 30 days following each presentation.

## About Syros Pharmaceuticals

Syros is pioneering the understanding of the non-coding regulatory region of the genome to advance a new wave of medicines that control the expression of genes. Syros has built a proprietary platform that is designed to systematically and efficiently analyze this unexploited region of DNA to identify and drug novel targets linked to genomically defined patient populations. Because gene expression is fundamental to the function of all cells, Syros' gene control platform has broad potential to create medicines that achieve profound and durable benefit across a range of diseases. Syros is currently focused on cancer and monogenic diseases and is advancing a growing pipeline of gene control medicines. Syros' lead drug candidates are SY-1425, a selective RAR $\alpha$  agonist in a Phase 2 clinical trial for genomically defined subsets of patients with acute myeloid leukemia, and SY-1365, a selective CDK7 inhibitor in a Phase 1 clinical trial focused on patients with ovarian and breast cancers. Syros is also developing a deep preclinical and discovery pipeline, including SY-5609, an oral CDK7 inhibitor, as well as programs in immuno-oncology and sickle cell disease. Led by a team with deep experience in drug discovery, development and commercialization, Syros is located in Cambridge, Mass.

## Cautionary Note Regarding Forward-Looking Statements

This press release contains forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995, including without limitation statements regarding the ability to complete IND-enabling preclinical studies and begin clinical development of SY-5609; and the benefits of Syros' gene control platform and product development pipeline . The words "anticipate," "believe," "continue," "could," "estimate," "expect," "hope," "intend," "may," "plan," "potential," "predict," "project," "target," "should," "would," and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words. Actual results or events could differ materially from the plans, intentions and expectations disclosed in these forward-looking statements as a result of various important factors, including Syros' ability to: advance the development of its programs, including SY-5609, under the timelines it projects in current and future clinical trials; demonstrate in any current and future clinical trials the requisite safety, efficacy and combinability of its drug candidates; successfully progress SY-5609 through IND-enabling preclinical and toxicology studies; replicate scientific and non-clinical data in clinical trials; obtain and maintain patent protection for its drug candidates and the freedom to operate under third party intellectual property; obtain and maintain necessary regulatory approvals; identify, enter into and maintain collaboration agreements with third parties, including its ability to perform under the collaboration agreement with Incyte; manage competition; manage expenses; raise the substantial additional capital needed to achieve its business objectives; attract and retain qualified personnel; and successfully execute on its business strategies; risks described under the caption "Risk Factors" in Syros' Annual Report on Form 10-K for the year ended December 31, 2018, which is on file with the Securities and Exchange Commission; and risks described in other filings that Syros makes with the Securities and Exchange Commission in the future. Any forward-looking statements contained in this press release speak only as of the date hereof, and Syros expressly disclaims any obligation to update any forward-looking statements, whether because of new information, future events or otherwise.

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