



Cidara Therapeutics to Present New Data for Rezafungin at TIMM Congress

September 29, 2021

SAN DIEGO, Sept. 29, 2021 (GLOBE NEWSWIRE) -- Cidara Therapeutics, Inc. (NASDAQ: CDTX), a biotechnology company developing long-acting therapeutics designed to improve the standard of care for patients facing serious diseases, today announced that it will present new clinical and non-clinical data in two poster presentations at the 10th Congress on Trends in Medical Mycology (TIMM) taking place in-person in Aberdeen, Scotland and virtually from October 8-11, 2021.

Rezafungin is a novel once-weekly echinocandin currently being studied in Phase 3 trials for the treatment and prevention of serious fungal infections, including in critically ill patients. Cidara expects to announce top-line data from its Phase 3 ReSTORE trial by the end of 2021 and anticipates filing a New Drug Application (NDA) in the U.S. and similar regulatory filings outside the U.S. in mid-2022.

Presentation details are summarized below.

Title: Rezafungin Safety and Pharmacokinetics in Subjects with Moderate or Severe Hepatic Impairment

Lead Author: Taylor Sandison, M.D., M.P.H., Cidara Therapeutics

Poster Number: P335

Date and Time: Oct. 9, 2021 at 11:00 a.m. – 12:00 p.m. CEST

Title: Evaluation of the Post-Antifungal Effect of Rezafungin and Micafungin against *Candida albicans*, *Candida parapsilosis*, and *Candida glabrata*

Lead Author: Jeffrey B. Locke, Ph.D., Cidara Therapeutics

Poster Number: P338

Date and Time: Oct. 10, 2021 at 11:00 a.m. – 12:00 p.m. CEST

Additional details can be found on the TIMM website [here](#). Copies of the presentations will be made available on the Publications section of Cidara's website.

About Rezafungin

Rezafungin is a novel once-weekly echinocandin being developed for both the treatment and prevention of serious fungal infections, such as candidemia and invasive candidiasis. The structure and properties of rezafungin are specifically designed to improve upon a clinically validated mechanism intended to enhance its efficacy and safety potential for patients. Cidara is currently conducting a Phase 3 clinical trial with rezafungin for the first-line treatment of candidemia and/or invasive candidiasis (ReSTORE trial) and a second Phase 3 clinical trial of rezafungin for the prevention of invasive fungal disease in patients undergoing allogeneic blood and marrow transplantation (ReSPECT trial).

About Cidara Therapeutics

Cidara is developing long-acting therapeutics designed to improve the standard of care for patients facing serious diseases. The Company's portfolio is comprised of new approaches aimed at transforming existing treatment and prevention paradigms, first with its lead Phase 3 antifungal candidate, rezafungin, in addition to drug-Fc conjugates (DFCs) targeting viral and oncology diseases from Cidara's proprietary Cloudbreak[®] platform. Cidara is headquartered in San Diego, California. For more information, please visit www.cidara.com.

Forward-Looking Statements

This release contains "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, and such forward-looking statements are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. "Forward-looking statements" describe future expectations, plans, results, or strategies and are generally preceded by words such as "anticipates," "expect," "may," "plan" or "will". Forward-looking statements in this release include, but are not limited to, statements related to our expectations for the timing of receiving top-line data for the Phase 3 ReSTORE trial and potentially filing a NDA for rezafungin. Such statements are subject to a multitude of risks and uncertainties that could cause future circumstances, events, or results to differ materially from those projected in the forward-looking statements, such as unanticipated delays in or negative results from Cidara's pre-clinical or clinical trials, delays in action by regulatory authorities due to limitations on inspections and other COVID-19-related effects, and impacts of the COVID-19 pandemic or other obstacles on the enrollment of patients or other aspects of rezafungin development. These and other risks are identified under the caption "Risk Factors" in Cidara's most recent Quarterly Report on Form 10-Q and other filings subsequently made with the Securities and Exchange Commission. All forward-looking statements contained in this press release speak only as of the date on which they were made and are based on management's assumptions and estimates as of such date. Cidara does not undertake any obligation to publicly update any forward-looking statements, whether as a result of the receipt of new information, the occurrence of future events or otherwise.

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