



Concert Pharmaceuticals Selected for Late-Breaking Oral Presentation of CTP-543 Phase 2 Data in Alopecia Areata at the 2020 American Academy of Dermatology Annual Meeting

February 24, 2020

LEXINGTON, Mass.--(BUSINESS WIRE)--Feb. 24, 2020-- [Concert Pharmaceuticals, Inc.](#) (NASDAQ: CNCE) today announced that results from its CTP-543 Phase 2 dose-ranging trial in alopecia areata have been selected for an oral presentation at the Late-Breaking Research Program during the American Academy of Dermatology (AAD) Annual Meeting being held March 20-24, 2020, in Denver, CO.

The details of the presentation, which will include new analyses from the Company's completed Phase 2 dose-ranging study, are as follows:

- Title: Oral JAK1/JAK2 Inhibitor CTP-543 Achieves Primary Endpoint in Patients with Alopecia Areata
- Date and Time: Saturday, March 21, 2020, 9:00 AM – 12:00 PM
- Session: S027 - Late-Breaking Research: Clinical Trials
- Room: Bellco Theatre 2

About CTP-543 and Alopecia Areata

CTP-543 is an oral inhibitor of Janus kinases JAK1 and JAK2. The U.S. Food and Drug Administration (FDA) has granted Fast Track designation for CTP-543 for alopecia areata.

Alopecia areata is an autoimmune disease in which the immune system attacks hair follicles, resulting in partial or complete loss of hair on the scalp and body. Alopecia areata may affect up to 650,000 Americans at any given time¹. The scalp is the most commonly affected area, but any hair-bearing site can be affected alone or together with the scalp. Onset of the disease can occur throughout life and affects both women and men. Alopecia areata can be associated with serious psychological consequences, including anxiety and depression. There are currently no drugs approved by the FDA for the treatment of alopecia areata.

The FDA selected alopecia areata as one of eight new disease areas that it focused on under its Patient-Focused Drug Development Initiative (PFDDI) in 2016-2017. The goal of the PFDDI was to bring patient perspectives into an earlier stage of product development. Following the FDA's Patient-Focused Drug Development meeting held in September 2017 on alopecia areata, the FDA summarized the input shared by patients and patient representatives in a [Voice of the Patient](#) report. Additional information on the PFDDI is available [online](#).

About Concert

[Concert Pharmaceuticals](#) is a clinical stage biopharmaceutical company focused on applying its [DCE Platform®](#) (deuterated chemical entity platform) to create novel medicines designed to treat serious diseases and address unmet patient needs. The Company's approach starts with previously studied compounds, including approved drugs, in which deuterium substitution has the potential to enhance clinical safety, tolerability or efficacy. Concert's [pipeline](#) of innovative medicines targets autoimmune diseases and central nervous systems (CNS) disorders. For more information please visit www.concertpharma.com or follow us on Twitter at [@ConcertPharma](#) or on [LinkedIn](#).

Cautionary Note on Forward Looking Statements

Any statements in this press release about our future expectations, plans and prospects, including, among others, statements about our expectations on the clinical development of CTP-543, and any other statements containing the words "anticipate," "believe," "continue," "could," "estimate," "expect," "intend," "may," "plan," "potential," "predict," "project," "should," "target," "will," "would" and similar expressions, constitute forward-looking statements within the meaning of The Private Securities Litigation Reform Act of 1995. Actual results may differ materially from those indicated by such forward-looking statements as a result of various important factors, including: the uncertainties inherent in the initiation of future clinical trials, whether preliminary results from a clinical trial will be predictive of the final results of that trial or whether results of early clinical trials will be indicative of the results of later clinical trials, expectations for regulatory approvals and other factors discussed in the "Risk Factors" section of our most recent Quarterly Report on Form 10-Q filed with the Securities and Exchange Commission and in other filings that we make with the Securities and Exchange Commission. In addition, any forward-looking statements included in this press release represent our views only as of the date of this release and should not be relied upon as representing our views as of any subsequent date. We specifically disclaim any obligation to update any forward-looking statements included in this press release.

¹ Fricke M. Epidemiology and Burden of alopecia areata: a systemic review. *Clinical, Cosmetic and Investigational Dermatology*. 2015; Vol 8. 397-403.

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