Tetraphase Pharmaceuticals to Present Data at the 29th European Congress of Clinical Microbiology and Infectious Diseases

April 8, 2019

WATERTOWN, Mass.--(BUSINESS WIRE)--Apr. 8, 2019-- Tetraphase Pharmaceuticals, Inc. (NASDAQ:TTPH), a biopharmaceutical company focused on developing and commercializing novel tetracyclines to treat serious and life-threatening conditions, today announced five data presentations at the 29thEuropean Congress of Clinical Microbiology and Infectious Diseases (ECCMID), taking place April 13-16 in Amsterdam, Netherlands. Presentations will include information about XERAVA^{TM} (eravacycline), a tetracycline antibiotic approved for the treatment of complicated intra-abdominal infections (cIAI), as well as data for the Company's clinical development candidates, TP-271 and TP-6076.

The details for the data presentations at ECCMID are as follows:

Eravacycline poster presentations

Poster title: Multicenter evaluation of eravacycline MIC results for *Esherichia coli* using MicroScan Dried Gram-negative MIC panels
Date and time: Monday, April 15 from 12:30 – 1:30 p.m. CEST (6:30 – 7:30 a.m. ET)
Location: Paper Poster Arena
Poster number: #P1735
Session information: PS098 – Diverse methodologies for MIC testing in Gram-negatives

Poster title: *In vitro* activity of eravacycline and comparators against Gram-positive bacteria collected from European hospitals in 2017
Date and time: Monday, April 15 from 1:30 – 2:30 p.m. CEST (7:30 – 8:30 a.m. ET)
Location: Paper Poster Arena
Poster number: #P1874
Session information: PS107 – *In vitro* activity of newer antimicrobial agents

Poster title: Surveillance of the *in vitro* activity of eravacycline and comparators against clinical isolates from Europe during 2017
Date and time: Monday, April 15 from 1:30 – 2:30 p.m. CEST (7:30 – 8:30 a.m. ET)
Location: Paper Poster Arena
Poster number: #P1875
Session information: PS107 – *In vitro* activity of newer antimicrobial agents

TP-271 poster presentation

Poster title: Safety, tolerability and pharmacokinetics of multiple doses of TP-271, a novel fluorocycline, in normal healthy subjects
Date and time: Monday, April 15 from 1:30 – 2:30 p.m. CEST (7:30 – 8:30 a.m. ET)
Location: Paper Poster Arena
Poster number: #P2012
Session information: PS116 – Safety of antibacterial agents in the clinic

TP-6076 poster presentation

Poster title:*In vivo* efficacy of TP-6076 in murine thigh and lung infection models challenged with *Acinetobacter baumannii*
Date and time: Monday, April 15 from 1:30 – 2:30 p.m. CEST (7:30 – 8:30 a.m. ET)
Location: Paper Poster Arena
Poster number: #P1994
Session information: PS115 – Evaluation of diverse antimicrobials *in vitro* and experimental models

Full abstracts can be found on the ECCMID website at www.eccmid.org. Additional questions or follow up should be directed to poster authors.

About Tetraphase Pharmaceuticals, Inc.

Tetraphase Pharmaceuticals, Inc., is a biopharmaceutical company using its proprietary chemistry technology to create novel tetracyclines for serious and life-threatening conditions, including infections caused by many of the multidrug-resistant bacteria highlighted as urgent public health threats by the World Health Organization and the Centers for Disease Control and Prevention. The Company has created more than 3,000 novel tetracycline compounds using its proprietary technology platform. Tetraphase's lead product XERAVA^{TM} is approved for the treatment of complicated intra-abdominal infections by the U.S. Food and Drug Administration and the European Medicines Agency. The Company's pipeline also includes TP-271 and TP-6076, which are in Phase 1 clinical trials, and TP-2846, which is in preclinical testing for acute myeloid leukemia. Please visit www.tphase.com for more company information.

Forward-Looking Statements

Any statements in this press release about our future expectations, plans and prospects, including statements regarding our strategy, future operations, prospects, plans and objectives, including our key milestones for 2019 and our anticipated cash runway, and other statements containing the words "anticipates," "believes," "expects," "plans," "will" 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 whether our commercial launch of XERAVA in the U.S. will be successful; our cash resources and the expected revenue will be sufficient to fund our operations for the period anticipated; our product candidates will succeed in clinical trials and even if the

clinical trials are successful, we may never achieve regulatory approval of such product candidates and other clinical, regulatory and commercial risk factors discussed in the "Risk Factors" section of our annual report on Form 10-K for the period ended December 31, 2018, filed with the Securities and Exchange Commission on March 15, 2019. In addition, the forward-looking statements included in this press release represent our views as of April 8, 2019.We anticipate that subsequent events and developments will cause our views to change. However, while we may elect to update these forward-looking statements at some point in the future, we specifically disclaim any obligation to do so.

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