August 15, 2018

Tonix Pharmaceuticals will Present Phase 3 HONOR and Phase 2 AtEase Study Results in a Poster Presentation at the 2018 Military Health System Research Symposium

NEW YORK, Aug. 15, 2018 (GLOBE NEWSWIRE) -- Tonix Pharmaceuticals Holding Corp. (Nasdaq: TNXP) (Tonix) announced today that it will be presenting findings and retrospective analyses from the Phase 3 HONOR study and Phase 2 AtEase Study in a poster, at the 2018 Military Health System Research Symposium (MHSRS) being held August 20 – 23, 2018, in Kissimmee, Fla.

Details of poster presentation at MHSRS

Title:	Differential Treatment Effects of a Sublingual Formulation of Cyclobenzaprine (TNX-102 SL*) on Dissociative Symptoms of Derealization and Depersonalization in a Military-Related PTSD Population: Retrospective Analysis of a Double-Blind Randomized Study
Poster Session:	Poster Session 1
Poster Discussion Session:	Innovations in PTSD Treatment Research
Date and Time:	Tuesday, August 21, 2018, 10:00 a.m. -12:00 p.m. ET
Abstract Number:	MHSRS-18-1970

Presenter: Gregory M. Sullivan, M.D., Chief Medical Officer, Tonix Pharmaceuticals Holdings Corp.

*TNX-102 SL is an investigational New Drug and has not been approved for any indications.

About Tonix Pharmaceuticals Holding Corp.

Tonix is a clinical-stage biopharmaceutical company focused on discovering and developing

pharmaceutical products to treat serious neuropsychiatric conditions and biological products to improve biodefense through potential medical counter-measures. Tonix is developing TNX-102 SL, or Tonmya®[#], which has been granted Breakthrough Therapy designation as a bedtime treatment for PTSD. TNX-102 SL is also being developed as a bedtime treatment for agitation in Alzheimer's disease under a separate IND to support a Phase 2, potential pivotal efficacy study. TNX-102 SL for agitation in Alzheimer's disease is a Fast Track development program. TNX-601 (tianeptine oxalate) is in the pre-IND application stage, also for the treatment of PTSD but by a unique mechanism and designed for daytime dosing. Tonix's lead biologic candidate, TNX-801, is a potential smallpox-preventing vaccine based on a live synthetic version of horsepox virus, currently in the pre-IND application stage.

[#]*Tonmya has been conditionally accepted by the FDA as the proposed trade name for TNX-102 SL (cyclobenzaprine HCl sublingual tablets) for PTSD.*

This press release and further information about Tonix can be found at www.tonixpharma.com.

Forward Looking Statements

Certain statements in this press release are forward-looking within the meaning of the Private Securities Litigation Reform Act of 1995. These statements may be identified by the use of forward-looking words such as "anticipate," "believe," "forecast," "estimate," "expect," and "intend," among others. These forward-looking statements are based on Tonix's current expectations and actual results could differ materially. There are a number of factors that could cause actual events to differ materially from those indicated by such forward-looking statements. These factors include, but are not limited to, risks related to failure to obtain FDA clearances or approvals and noncompliance with FDA regulations; our need for additional financing; uncertainties of patent protection and litigation; uncertainties of government or third party payor reimbursement; limited research and development efforts and dependence upon third parties; and substantial competition. As with any pharmaceutical under development, there are significant risks in the development, regulatory approval and commercialization of new products. Tonix does not undertake an obligation to update or revise any forward-looking statement. Investors should read the risk factors set forth in the Annual Report on Form 10-K for the year ended December 31, 2017, as filed with the Securities and Exchange Commission (the "SEC") on March 9, 2018, and periodic reports filed with the SEC on or after the date thereof. All of Tonix's forward-looking statements are expressly qualified by all such risk factors and other cautionary statements. The information set forth herein speaks only as of the date thereof.

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