January 2, 2019

Tonix Pharmaceuticals to Present at Upcoming Investor Conferences and Participate in Featured Panel Discussion

NEW YORK, Jan. 02, 2019 (GLOBE NEWSWIRE) -- Tonix Pharmaceuticals Holding Corp. (Nasdaq: TNXP) (Tonix or the Company) announced today that Seth Lederman, M.D., President and Chief Executive Officer of Tonix, will present a company overview at two investor conferences in January 2019.

In addition, Dr. Lederman will participate in a panel during the 2nd Annual Neuroscience Innovation Forum titled, "*Advances in Neuropsychiatry and Pain Management*," scheduled from 11:10 a.m. – 11:45 a.m. ET, January 6, 2019, in Room Commandants.

Details of the Tonix Pharmaceuticals company presentations and webcasts are as follows:

Event:	2nd Annual Neuroscience Innovation Forum
Date:	Sunday, January 6, 2019
Time:	6:45 p.m. ET (3:45 p.m. PT)
Location:	Marines' Memorial Club & Hotel, San Francisco (Room Commandants)

Event:	Biotech Showcase
Date:	Tuesday, January 8, 2019
Time:	6:00 p.m. ET (3:00 p.m. PT)
Location:	Hilton San Francisco Union Square (Franciscan A, Ballroom Level)

About Tonix Pharmaceuticals Holding Corp.

A live webcast and subsequent archived recording of the Company presentations will be available under the IR Events tab of the Investor Relations section of the Tonix Pharmaceuticals website at www.tonixpharma.com.

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 Tonmya®*, which is in Phase 3 development and has been granted Breakthrough Therapy designation by the FDA, as a bedtime treatment for PTSD. Tonix is also developing TNX-102 SL as a bedtime treatment for agitation in Alzheimer's disease under a separate IND to support a Phase 2, potential pivotal, efficacy study and has been designated a Fast Track development program by the FDA for this indication. 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. Phase 1 clinical study of TNX-601 in healthy volunteers will be

conducted outside of the U.S. in 2019. 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 U.S. Food and Drug Administration (FDA) as the proposed trade name for TNX-102 SL (cyclobenzaprine HCl sublingual tablets) for the treatment of PTSD. TNX-102 SL is an investigational new drug and has not been approved for any indication.*

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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