

May 23, 2018



## **Tonix Pharmaceuticals to Present at the 2018 American Society of Clinical Psychopharmacology**

NEW YORK, May 23, 2018 (GLOBE NEWSWIRE) -- Tonix Pharmaceuticals Holding Corp. (Nasdaq:TNXP) (Tonix) announced today that Gregory Sullivan, M.D., Chief Medical Officer of Tonix, will present at the 2018 American Society of Clinical Psychopharmacology (ASCP) Annual Meeting being held May 29 – June 1, 2018, in Miami Beach, Fla. The presentation is titled, *Including Suicidal Individuals in Treatment Trials: Treatment of Military-Related PTSD with TNX-102 SL, a Novel Formulation of Cyclobenzaprine Hypothesized to Address PTSD through Improvement in Sleep Quality*.

### *Tonix Pharmaceuticals Presentation Details*

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|----------------|------------------------------------------------------------------------|
| Event:         | 2018 American Society of Clinical Psychopharmacology                   |
| ASCP Workshop: | Inclusion of Suicidal Individuals in Treatment Trials: Now is the Time |
| Date:          | Wednesday, May 30 <sup>th</sup> , 2018                                 |
| Time:          | 2:45 – 4:45 p.m.                                                       |
| Workshop Room: | Poinciana 1-2, Loews Miami Beach                                       |

### **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's lead product candidate, Tonmya, or TNX-102 SL, is in Phase 3 development as a bedtime treatment for PTSD. Tonix is also developing TNX-102 SL as a bedtime treatment for agitation in Alzheimer's disease. A Phase 2 IND (Investigational New Drug) application was submitted in March 2018 after completion of a successful pre-IND meeting with the FDA. TNX-601 (tianeptine oxalate) is in the pre-IND application stage, also for the treatment of PTSD but 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.

This press release and further information about Tonix can be found at [www.tonixpharma.com](http://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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