

January 2, 2013

Theravance to Present at the 31st Annual J.P. Morgan Healthcare Conference

SOUTH SAN FRANCISCO, CA -- (Marketwire) -- 01/02/13 -- Theravance, Inc. (NASDAQ: THRX) announced today that that Rick E Winningham, Theravance's Chief Executive Officer, is scheduled to present at the 31st Annual J.P. Morgan Healthcare Conference on Wednesday, January 9, 2013, at 10:30 a.m. PST. The conference will be held from January 7-10, 2013 at the Westin St. Francis Hotel in San Francisco, CA.

The presentation will be webcast live and can be reached from Theravance's web site at www.theravance.com. Listeners are encouraged to visit the site at least 15 minutes prior to the scheduled presentation to register, download and install any necessary audio software.

About Theravance

Theravance is a biopharmaceutical company with a pipeline of internally discovered product candidates and strategic collaborations with pharmaceutical companies. Theravance is focused on the discovery, development and commercialization of small molecule medicines across a number of therapeutic areas including respiratory disease, bacterial infections, and central nervous system (CNS)/pain. Theravance's key programs include: RELVAR™ or BREO™ (FF/VI), ANORO™ (UMEC/VI) and MABA (Bifunctional Muscarinic Antagonist-Beta2 Agonist), each partnered with GlaxoSmithKline plc, and its oral Peripheral Mu Opioid Receptor Antagonist program. By leveraging its proprietary insight of multivalency to drug discovery, Theravance is pursuing a best-in-class strategy designed to discover superior medicines in areas of significant unmet medical need. For more information, please visit Theravance's web site at www.theravance.com.

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RELVAR™ or BREO™ (FF/VI) and ANORO™ (UMEC/VI) are investigational medicines and are not currently approved anywhere in the world. RELVAR™, BREO™ and ANORA™ are trademarks of the GlaxoSmithKline group of companies. T use of these brand names has not yet been approved by any regulatory authority.

VIBATIV® is a registered trademark of Theravance, Inc.

Theravance's presentation will contain certain "forward-looking" statements as that term is defined in the Private Securities Litigation Reform Act of 1995 regarding, among other things, statements relating to goals, plans, objectives and future events. Theravance intends such forward-looking statements to be covered by the safe harbor provisions for forward-looking statements contained in Section 21E of the Securities Exchange Act of 1934 and the Private Securities Litigation Reform Act of 1995. Examples of such statements include statements relating to the status and timing of clinical studies, data analysis and communication of results, statements regarding the potential benefits and mechanisms of action of drug candidates, statements concerning the timing of seeking regulatory approval of our product candidates (including with respect to VIBATIV® statements regarding any expectation that we will be able to respond fully or adequately to FDA's requests using currently existing clinical data and any expectation that the FDA will approve the VIBATIV® nosocomial pneumonia NDA on the basis of existing preclinical and clinical data or at all), statements concerning the enabling capabilities of Theravance's approach to drug discovery and its proprietary insights, statements concerning expectations for the discovery, development and commercialization of product candidates and projections of revenue, expenses and other financial items. These statements are based on the current estimates and assumptions of the management of Theravance as of the date of the presentation and are subject to risks, uncertainties, changes in circumstances, assumptions and other factors that may cause the actual results of Theravance to be materially different from those reflected in its forward-looking statements. Important factors that could cause actual results to differ materially from those indicated by such forward-looking statements include, among others, risks related to delays or difficulties in commencing or completing clinical and non-clinical studies, the potential that results of clinical or non-clinical studies indicate product candidates are unsafe or ineffective, our dependence on third parties in the conduct of our clinical studies, delays or failure to achieve regulatory approvals for product candidates, risks of relying on third-party manufacturers for the supply of our product and product candidates and risks of collaborating with third parties to discover, develop and commercialize products. These and other risks are described in greater detail under the heading "Risk Factors" contained in Theravance's Quarterly Report on Form 10-Q filed with the Securities and Exchange Commission (SEC) on October 31, 2012 and the risks discussed in our other period filings with SEC. Given these uncertainties, you should not place undue reliance on these forward-looking statements. Theravance assumes no obligation to update its forward-looking statements.

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Contact Information:

Michael W. Aguiar

Senior Vice President and Chief Financial Officer

650-808-4100

investor.relations@theravance.com

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