

December 3, 2019

**BAUDAX BIO**<sup>®</sup>

# **Baudax Bio to Present at the Piper Jaffray 31st Annual Healthcare Conference**

MALVERN, Pa., Dec. 03, 2019 (GLOBE NEWSWIRE) -- Baudax Bio, Inc., (NASDAQ:BXRX) a pharmaceutical company focused on advancing non-opioid analgesics and other products for the hospital and other acute care settings, today announced that Gerri Henwood, the Company's President and Chief Executive Officer, will present at the Piper Jaffray 31<sup>st</sup> Annual Healthcare Conference on Thursday, December 5, 2019 at 12:00 p.m. ET at the Lotte New York Palace.

A live and archived webcast of the Piper Jaffray presentation will be available on the Events page of the company's website at <https://www.baudaxbio.com/>. The webcast will be archived for a period of 30 days following the conclusion of the live event.

## **About Baudax Bio**

Baudax Bio is a pharmaceutical company focused on therapeutics for acute care settings. The Company's lead product candidate is a proprietary intravenous (IV) form of meloxicam, a non-opioid, long-acting preferential COX-2 inhibitor. IV meloxicam has successfully completed two pivotal Phase III clinical efficacy trials, a large double-blind placebo-controlled Phase III safety trial, four Phase II clinical efficacy trials, as well as other safety studies. Upon IV meloxicam approval, ANJESO<sup>™</sup>, will be a novel non-opioid option for the management of moderate to severe pain. As a non-opioid, IV meloxicam has the potential to overcome many of the issues associated with commonly prescribed opioid therapeutics, including respiratory depression, constipation, excessive nausea and vomiting, as well as having no addictive potential while maintaining meaningful analgesic effects for relief of pain. For more information please visit [www.baudaxbio.com](http://www.baudaxbio.com).

## **Cautionary Statement Regarding Forward Looking Statements**

This press release contains forward-looking statements that involve risks and uncertainties. Such forward-looking statements reflect Baudax Bio's expectations about its future performance and opportunities that involve substantial risks and uncertainties. When used herein, the words "anticipate," "believe," "estimate," "may," "upcoming," "plan," "target," "intend" and "expect" and similar expressions, as they relate to Baudax Bio or its management, are intended to identify such forward-looking statements. These forward-looking statements are based on information available to Baudax Bio as of the date of this press release and are subject to a number of risks, uncertainties, and other factors that could cause Baudax Bio's performance to differ materially from those expressed in, or implied by, these forward-looking statements. Baudax Bio assumes no obligation to update any such forward-looking statements. Factors that could cause Baudax Bio's actual performance to materially differ from those expressed in the forward-looking statements set forth in this press release include, without limitation: the Company's ability to execute its strategy for further development and commercialization of IV meloxicam, the Company's ability to execute its strategic initiatives, the Company's ability to adequately resolve the

outstanding labeling issues with the FDA for IV meloxicam, and the time frame associated with any such resolution; the Company's ability to raise future financing for continued product development and IV meloxicam commercialization; with regard to the Company's clinical trial results, whether there may be changes in the interpretation by the FDA of the data of the Company's clinical trials and the length, cost and uncertain results and timing of our ongoing clinical trials; with regard to the potential commercial opportunity of IV meloxicam, whether any FDA approval of IV meloxicam will include labeling restrictions and the potential that IV meloxicam does not receive regulatory approval or does not receive reimbursement by third party payors, that IV meloxicam is not accepted by the medical community, including physicians, patients, health care providers and hospital formularies or that a commercial market for IV meloxicam does not develop; the Company's ability to manage costs and execute on its operational and budget plans; the Company's ability to achieve its financial goals; the Company's ability to obtain, maintain and successfully enforce adequate patent and other intellectual property protection; the Company's lack of operating history as a standalone business; risks relating to the separation from Recro, including, among others, failure to achieve the anticipated benefits from the separation, reliance on Recro and other third parties to provide certain services post-separation, and the Company's ability to satisfy liabilities and potential indemnification obligations in connection with the separation. The forward-looking statements in this press release should be considered together with the risks and uncertainties that may affect Baudax Bio's business and future results included in Baudax Bio's filings with the Securities and Exchange Commission at [www.sec.gov](http://www.sec.gov).

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