Urovant Sciences to Ring the Nasdaq Stock Market Opening Bell

January 17, 2020

IRVINE, Calif. & BASEL, Switzerland--(BUSINESS WIRE)--Jan. 17, 2020-- Urovant Sciences (Nasdaq: UROV) announced today that Chief Executive Officer Keith Katkin and other Urovant executives will ring the opening bell of the Nasdaq Stock Market on Tuesday, January 21, 2020, at 9:30 a.m. Eastern.

The ceremony will begin at 9:20 a.m. To view a live stream of the Nasdaq Opening Bell, please visit Nasdaq's MarketSite: https://www.nasdaq.com/marketsite/bell-ringing-ceremony.

"We are honored to be ringing the Nasdaq opening bell to commemorate our initial public offering just over one year ago," said Keith Katkin, Chief Executive Officer. "Since going public we have made significant progress towards our goal of becoming a leading specialty urology company. Our recently submitted New Drug Application to the U.S. Food and Drug Administration for vibegron in overactive bladder was another major milestone for Urovant. Vibegron, if approved this year, would be the first new branded prescription drug for the treatment of OAB in nearly a decade, potentially providing relief for the millions of patients suffering from OAB. We extend our gratitude to the employees, board members, clinical investigators and shareholders who have supported us in our mission and we look forward to potentially bringing a new treatment option to the market this year."

About Urovant Sciences

Urovant Sciences is a clinical-stage biopharmaceutical company focused on developing and commercializing innovative therapies for urologic conditions. The Company's lead product candidate, vibegron, an oral, once-daily small molecule beta-3 agonist is being evaluated for overactive bladder (OAB). Urovant Sciences reported positive data from the vibegron 12-week, phase 3 pivotal EMPOWUR study and demonstrated favorable longer-term efficacy, safety, and tolerability in a 40-week extension study. The Company submitted a New Drug Application to the FDA seeking approval of vibegron for the treatment of patients with OAB in December 2019. Vibegron is also being evaluated for treatment of OAB in men with benign prostatic hyperplasia (OAB+BPH) and for abdominal pain associated with irritable bowel syndrome (IBS). Urovant's second product candidate, URO-902, is a novel gene therapy being developed for patients with OAB who have failed oral pharmacologic therapy. Urovant Sciences, a subsidiary of Sumitomo Dainippon Pharma Co., Ltd., intends to develop novel treatments for additional urologic diseases. Learn more about us at www.urovant.com.

Forward-Looking Statements

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements include all statements that are not historical statements of fact and statements regarding the Company's intent, belief or expectations and can be identified by words such as "anticipate," "believe," "can," "continue," "could," "estimate," "expect," "intend," "likely," "may," "might," "objective," "ongoing," "plan," "potential," "predict," "project," "should," "strive," "to be," "will," "would," or the negative or plural of these words or other similar expressions or variations, although not all forward-looking statements contain these identifying words. In this press release, forward-looking statements include, but are not limited to, statements regarding Urovant's plans to advance the clinical development of vibegron in patients with OAB and obtain FDA approval. Forward-looking statements are subject to risks and uncertainties that could cause actual results to differ materially and reported results should not be considered as an indication of future performance. These risks and uncertainties include, but are not limited to, risks associated with: the success, cost, and timing of Urovant's development activities, including the timing of the initiation and completion of clinical trials and the timing of expected regulatory filings; the clinical utility and potential attributes and benefits of vibegron, including reliance on collaboration partners and the ability to procure additional sources of financing; our intellectual property position, including the ability to identify and in-license or acquire third-party patents and licenses, and associated costs; and other risks and uncertainties listed in the Company's filings with the United States Securities and Exchange Commission (SEC), including under the heading "Risk Factors" in the Company's most recently filed Annual Report on Form 10-K and any subsequent Quarterly Reports on Form 10-Q filed with the SEC, as such risk factors may be amended, supplemented or superseded from time to time by other filings with the SEC. Given these risks and uncertainties, you should not place undue reliance on any forward-looking statements. These forward-looking statements are based on information available to Urovant as of the date of this press release and speak only as of the date of this release. Urovant disclaims any obligation to update these forward-looking statements, except as may be required by law.

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