# BioDelivery Sciences to Webcast Analyst & Investor Day on February 25th

February 18, 2016

RALEIGH, N.C., Feb. 18, 2016 /PRNewswire/ -- BioDelivery Sciences International, Inc. (NASDAQ: BDSI) announced that it will webcast the company's Analyst and Investor Day, which will take place on Thursday, February 25, 2016 from 12:00 PM - 2:30 PM Eastern Time in New York City.

The event will focus on both of BDSI's marketed products, BUNAVAIL® (buprenorphine and naloxone) buccal film (CIII), and BELBUCA ™ (buprenorphine HCl) buccal film (CIII), which is licensed worldwide to Endo Pharmaceuticals. An update on BDSI's development portfolio, which includes Clonidine Topical Gel for painful diabetic neuropathy, and Buprenorphine Depot Injection for both opioid dependence and chronic pain, will also be provided.

The event will feature Dr. Richard Rauck, President, Carolinas Pain Institute and investigator in the BELBUCA Phase 3 clinical program, and Dr. Michael Frost, who is President of Frost Medical Group and specializes in the treatment of opioid dependence, as well as several members of the BDSI executive management team.

A live webcast of the event will be available at http://lifesci.rampard.com/20160225/reg.jspand on the BDSI website at www.bdsi.com. If you are an institutional investor or sell-side analyst, and would like to attend the event in person, please contact Mac MacDonald at 212-915-2567 or via e-mail at mac@lifesciadvisors.com for further information.

***About BioDelivery Sciences International***

BioDelivery Sciences International, Inc. (NASDAQ:BDSI) is a specialty pharmaceutical company with a focus in the areas of pain management and addiction medicine. BDSI is utilizing its novel and proprietary BioErodible MucoAdhesive (BEMA®) technology and other drug delivery technologies to develop and commercialize, either on its own or in partnership with third parties, new applications of proven therapies aimed at addressing important unmet medical needs.

BDSI's development strategy focuses on utilization of the FDA's 505(b)(2) approval process. This regulatory pathway creates the potential for more timely and efficient approval of new formulations of previously approved therapeutics.

BDSI's particular area of focus is the development and commercialization of products in the areas of pain management and addiction. These are areas where BDSI believes its drug delivery technologies and products can best be applied to address critical unmet medical needs. BDSI's marketed products and those in development address serious and debilitating conditions such as breakthrough cancer pain, chronic pain, painful diabetic neuropathy and opioid dependence. BDSI's headquarters is located in Raleigh, North Carolina.

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| Internet: | www.bdsi.com |
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**BDSI markets BUNAVAIL® (buprenorphine and naloxone) buccal film (CIII) and *ONSOLIS*®(fentanyl buccal soluble film) (CII). BELBUCA™ (buprenorphine) buccal film (CIII) will be commercialized in the U.S. by Endo Pharmaceuticals pursuant to the worldwide licensing and development agreement between BDSI and Endo. For full prescribing information and important safety information on BDSI products, including BOXED WARNINGS for ONSOLIS and BELBUCA, please visit www.bdsi.com where the Company promptly posts press releases, SEC filings and other important information or contact the Company at (800) 469-0261.**

**Cautionary Note on Forward-Looking Statements**

This press release, the presentation described herein, and any statements of employees, representatives and partners of BioDelivery Sciences International, Inc. (the "Company") related thereto (including, without limitation, at the presentation described herein) contain, or may contain, among other things, certain "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995. Such forward-looking statements involve significant risks and uncertainties. Such statements may include, without limitation, statements with respect to the Company's plans, objectives, projections, expectations and intentions and other statements identified by words such as "projects," "may," "will," "could," "would," "should," "believes," "expects," "anticipates," "estimates," "intends," "plans," "potential" or similar expressions. These statements are based upon the current beliefs and expectations of the Company's management and are subject to significant risks and uncertainties, including those detailed in the Company's filings with the Securities and Exchange Commission. Actual results (including, without limitation, the results of commercialization efforts for the Company's approved products and the clinical trials for, and regulatory review of, the Company's products in development) may differ significantly from those set forth in the forward-looking statements. These forward-looking statements involve certain risks and uncertainties that are subject to change based on various factors (many of which are beyond the Company's control). The Company undertakes no obligation to publicly update any forward-looking statements, whether as a result of new information, future presentations or otherwise, except as required by applicable

law.



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