



BioDelivery Sciences to Present at the 6th Annual Credit Suisse Small & Mid Cap Conference and the Ladenburg Thalmann 2015 Healthcare Conference

September 10, 2015

RALEIGH, N.C., Sept. 10, 2015 /PRNewswire/ -- BioDelivery Sciences International, Inc. (NASDAQ: BDSI) announced that the company will present at two upcoming healthcare conferences:

6th Annual Credit Suisse Small & Mid Cap Conference

Date: Wednesday, September 16

Location: Waldorf Astoria, New York City

Presentation Time: 10:10 AM Eastern Time

Ladenburg Thalmann 2015 Healthcare Conference

Date: Tuesday, September 29

Location: Sofitel New York, New York City

Presentation Time: 10:00 AM Eastern Time

BDSI will be discussing progress with the commercial launch of BUNAVAIL[®] (buprenorphine and naloxone buccal film), including the recently secured exclusive preferred formulary status with Tennessee Medicaid. Also discussed will be BELBUCA[™] (Buprenorphine HCL buccal film), which is under review by the U.S. Food and Drug Administration (FDA) with a PDUFA date of October 23, 2015 for use in patients with pain severe enough to require daily, around-the-clock, long-term opioid treatment for which alternative treatment options are inadequate. The presentation will also provide an update on the status of Clonidine Topical Gel being studied for the treatment of painful diabetic neuropathy and Buprenorphine Depot Injection, which is being developed for both chronic pain and opioid dependence.

The Ladenburg Thalmann presentation will be webcast live and can be accessed at www.bdsi.com. For those who are not available to listen to the live broadcast, a replay of the webcast will be available on the BDSI website.

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.

For more information, please visit or follow us:

Internet: www.bdsi.com

Facebook: [Facebook.com/BioDeliverySI](https://www.facebook.com/BioDeliverySI)

Twitter: @BioDeliverySI

Cautionary Note on Forward-Looking Statements

This press release, the presentations described herein, and any statements of employees, representatives and partners of BioDelivery Sciences International, Inc. (the "Company") related thereto (including, without limitation, at the presentations 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 the commercial launch of BUNAVAIL and the Company's clinical trials for, and FDA 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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For further information: Investors: Matthew P. Duffy, Managing Director, LifeSci Advisors, LLC, 212-915-0685, matthew@lifesciadvisors.com; Al Medwar, Vice President, Marketing and Corporate Development, BioDelivery Sciences International, Inc., 919-582-9050, amedwar@bdsi.com; Media: Susan Forman/Laura Radocaj, Dian Griesel Int'l., 212-825-3210, sforman@dgicomm.com, lradocaj@dgicomm.com