BioDelivery Sciences to Host Conference Call and Webcast Reporting First Quarter 2017 Financial Results on Monday, May 15

May 9, 2017

RALEIGH, N.C., May 9, 2017 /PRNewswire/ -- BioDelivery Sciences International, Inc. (NASDAQ: BDSI) announced that it will report its first quarter 2017 financial results and host a conference call and webcast at 4:30 PM Eastern Time on Monday, May 15, 2017.

Conference Call & Webcast	
Monday, May 15 @ 4:30 PM Eastern Time	
Domestic:	888-437-9364
International:	719-325-2133
Passcode:	7698100
Webcast:	http://public.viavid.com/index.php?id=124144
Replays available through May 29th:	
Domestic:	844-512-2921
International:	412-317-6671
Conference ID:	7698100

BDSI will also be filing a Form 12b-25 with the U.S. Securities and Exchange Commission (SEC) on May 10, 2017 in order to provide BDSI with five extra days to file its first quarter 2017 Form 10-Q. The extra five days will allow BDSI's and Endo Pharmaceuticals' auditors and the company the time to complete the required valuation, audit and statement preparation work related to BDSI's January 2017 license termination transaction with Endo related to BELBUCA®. As previously reported, such transaction (in which BDSI reacquired the rights to BELBUCA) technically requires (under generally accepted accounting principles and SEC reporting rules) the preparation of separate audited and pro forma information related specifically to BELBUCA. This is an SEC informational requirement only and has no bearing on BDSI's first quarter 2017 results of operations.

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 the 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 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 in Raleigh, North Carolina.

For more information, please visit or follow us:

Internet:	www.bdsi.com

Facebook:	Facebook.com/BioDeliverySI
Twitter:	@BioDeliverySI

BUNAVAIL® (buprenorphine and naloxone) buccal film (CIII) and BELBUCA® (buprenorphine) buccal film (CIII) are marketed in the U.S. by BioDelivery Sciences. *ONSOLIS*® (fentanyl buccal soluble film) (CII) is licensed in the U.S. to Collegium Pharmaceutical pursuant to the U.S. licensing and development agreement between BDSI and Collegium. For full prescribing information and important safety information on BDSI products, including BOXED WARNINGS for ONSOLIS, 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. For full prescribing and safety information on BELBUCA, please visit www.belbuca.com and for full prescribing and safety information on BUNAVAIL, please visit www.bunavail.com.

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 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 Company's financial results and the results of the Company's commercialization and strategic initiatives) 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.

SOURCE BioDelivery Sciences International, Inc.

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