Endo Launches Unscripted Video Series With Real Dupuytren's Contracture Patients

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Real Patients Talk About the Hand Condition and Treatments

- *Six short, unscripted "Coffee & Cords" videos explore various topics, including the tabletop test, finding a hand specialist and discussing treatment options*
- *Videos can be seen online, with digital and social advertising to come later this year*

DUBLIN, Oct. 5, 2022 /PRNewswire/ -- Endo International plc (OTC: ENDPQ) announced today the launch of its new branded video series, *Coffee & Cords*, which aims to educate people about Dupuytren's contracture (DC), a progressive hand condition that results in the formation of a collagen rope-like cord that pulls fingers toward the palm and affects an estimated 14 million Americans.

The *Coffee & Cords* videos feature five real patients with DC in a coffee shop discussing their experiences with the finger contracture condition, including the impact on their daily activities and treatment options. The conversations are unscripted and present the patients' actual experiences and views.

"Those who best understand and relate to DC patients are other patients—they've faced similar physical challenges, feelings about the condition and hesitations or confusion around treatment," said Justin Mattice, Vice President & General Manager, Medical Therapeutics at Endo. "We believe these authentic voices can empower others who think they may have DC to talk to a hand surgeon about treatment options, including XIAFLEX® (collagenase clostridium histolyticum), the only FDA-approved nonsurgical treatment for adults with DC with a palpable cord."

Do not receive XIAFLEX if you have had an allergic reaction to collagenase clostridium histolyticum or any of the ingredients in XIAFLEX, or to any other collagenase product.

The six *Coffee & Cords* videos range from one to four minutes in length. The short episodes focus on specific topics so that people can easily find what they're interested in, or they can watch the entire series to get a fuller view of the patient experience—dealing with the effects of the finger contraction condition, understanding DC, doing the tabletop test, finding a hand specialist and discussing treatment options.

The videos are viewable on Endo's newly redesigned consumer website for XIAFLEX, found at XIAFLEX.com. Later this year, they will be used in digital and social advertising.

About Dupuytren's Contracture

DC is a lifelong condition that may get worse over time. It's caused by a buildup of collagen in the hand, which forms a rope-like cord that pulls fingers toward the palm so they can't be straightened. As DC progresses, it may become difficult for individuals to use their hand(s) for daily tasks and activities.

WHAT IS XIAFLEX®?

XIAFLEX is a prescription medicine used to treat adults with Dupuytren's contracture when a "cord" can be felt. It is not known if XIAFLEX is safe and effective in children under the age of 18.

IMPORTANT SAFETY INFORMATION FOR XIAFLEX

Do not receive XIAFLEX if you have had an allergic reaction to collagenase clostridium histolyticum or any of the ingredients in XIAFLEX, or to any other collagenase product. See the end of the Medication Guide for a complete list of ingredients in XIAFLEX.

XIAFLEX can cause serious side effects, including:

- **Tendon rupture or ligament damage.** Receiving an injection of XIAFLEX may cause damage to a tendon or ligament in your hand and cause it to break or weaken. This could require surgery to fix the damaged tendon or ligament. Call your healthcare provider right away if you have trouble bending your injected finger (towards the wrist) after the swelling goes down or you have problems using your treated hand after your follow-up visit

- **Nerve injury or other serious injury of the hand.** After finger procedures, some people developed tears in the skin (lacerations), and local skin and soft-tissue necrosis (death of skin cells). Some lacerations and necrosis required skin grafting, or other surgery including amputation. **Call your healthcare provider right away** if you get numbness, tingling, increased pain, or tears in the skin (laceration) in your treated finger or hand after your injection or after your follow-up visit
- **Hypersensitivity reactions, including anaphylaxis.** Severe allergic reactions can happen in people who receive XIAFLEX because it contains foreign proteins. **Call your healthcare provider right away if you have any of these symptoms of an allergic reaction after an injection of XIAFLEX:**
    - hives
    - swollen face
    - breathing trouble
    - chest pain
    - low blood pressure
    - dizziness or fainting
- **Fainting.** Fainting (passing out) or near fainting can happen in people who receive XIAFLEX, especially following finger procedures
    - **If you have dizziness or feel faint after receiving XIAFLEX, lie down until the symptoms go away.**
- **Increased chance of bleeding.** Bleeding or bruising at the injection site can happen in people who receive XIAFLEX. Talk to your healthcare provider if you have a problem with your blood clotting. XIAFLEX may not be right for you.

Before receiving XIAFLEX, tell your healthcare provider if you have had an allergic reaction to a previous XIAFLEX injection, or have a bleeding problem or any other medical conditions. Tell your healthcare provider about all the medicines you take, including prescription and non-prescription medicines, vitamins, and herbal supplements. Using XIAFLEX with certain other medicines can cause serious side effects. Especially tell your healthcare provider if you take medicines to thin your blood (anticoagulants). If you are told to stop taking a blood thinner before your XIAFLEX injection, your healthcare provider should tell you when to restart the blood thinner. Ask your healthcare provider or pharmacist for a list of these medicines if you are unsure.

The most common side effects with XIAFLEX for the treatment of Dupuytren's contracture include:

- swelling of the injection site or the hand
- bruising or bleeding at the injection site
- pain or tenderness of the injection site or the hand
- swelling of the lymph nodes (glands) in the elbow or armpit
- itching
- breaks in the skin
- redness or warmth of the skin
- pain in the armpit

Tell your healthcare provider if you have any side effect that bothers you or does not go away. These are not all of the possible side effects with XIAFLEX. For more information, ask your healthcare provider or pharmacist.

Click for full Prescribing Information, including Medication Guide.

About Endo
Endo (OTC: ENDPQ) is a specialty pharmaceutical company committed to helping everyone we serve live their best life through the delivery of quality, life-enhancing therapies. Our decades of proven success come from passionate team members around the globe collaborating to bring treatments forward. Together, we boldly transform insights into treatments benefiting those who need them, when they need them. Learn more at www.endo.com or connect with us on LinkedIn.

Cautionary Note Regarding Forward-Looking Statements
Certain information in this press release may be considered "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995 and any applicable Canadian securities legislation, including, but not limited to, the statements by Mr. Mattice and any statements relating to the video series, patient experiences, treatment options and XIAFLEX®, and any other statements that refer to expected, estimated or anticipated future results or that do not relate solely to historical facts. Statements including words or phrases such as "believe," "expect," "anticipate," "intend," "estimate," "plan," "will," "may," "look forward," "intend," "guidance," "future," "potential" or similar expressions are forward-looking statements. All forward-looking statements in this communication reflect the Company's current views as of the date of this communication about its plans, intentions, expectations, strategies and prospects, which are based on the information currently available to it and on assumptions it has made. Actual results may differ materially and adversely from current expectations based on a number of factors, including, among other things, the following: the outcome of the Company's contingency planning and restructuring activities; the timing, impact or results of any pending or future litigation, investigations, proceedings or claims, including opioid, tax and antitrust related matters; actual or contingent liabilities; settlement discussions or negotiations; the Company's liquidity, financial performance, cash position and operations; the Company's strategy; risks and uncertainties associated with Chapter 11 proceedings; the negative impacts on the Company's businesses as a result of filing for and operating under Chapter 11 protection; the time, terms and ability to confirm a sale of the Company's businesses under Section 363 of the U.S. Bankruptcy Code; the adequacy of the capital resources of the Company's businesses and the difficulty in forecasting the liquidity requirements of the operations of the Company's businesses; the unpredictability of the Company's financial results while in Chapter 11 proceedings; the Company's ability to discharge claims in Chapter 11 proceedings; negotiations with the holders of the Company's indebtedness and its trade creditors and other significant creditors; risks and uncertainties with performing under the terms of the restructuring support agreement and any other arrangement with lenders or creditors while in Chapter 11 proceedings; the Company's ability to conduct business as usual; the Company's ability to continue to serve customers, suppliers and

other business partners at the high level of service and performance they have come to expect from the Company; the Company's ability to continue to pay employees, suppliers and vendors; the ability to control costs during Chapter 11 proceedings; adverse litigation; the risk that the Company's Chapter 11 Cases may be converted to cases under Chapter 7 of the Bankruptcy Code; the Company's ability to secure operating capital; the Company's ability to take advantage of opportunities to acquire assets with upside potential; the Company's ability to execute on its strategic plan to pursue, evaluate and close an asset sale of the Company's businesses pursuant to Section 363 of the U.S. Bankruptcy Code; the impact of competition, including the loss of exclusivity and generic competition for VASOSTRICT®; our ability to satisfy judgments or settlements or pursue appeals including bonding requirements; our ability to adjust to changing market conditions; our ability to attract and retain key personnel; our inability to maintain compliance with financial covenants and operating obligations which would expose us to potential events of default under our outstanding indebtedness; our ability to incur additional debt or equity financing for working capital, capital expenditures, business development, debt service requirements, acquisitions or general corporate or other purposes; our ability to refinance our indebtedness; a significant reduction in our short-term or long-term revenues which could cause us to be unable to fund our operations and liquidity needs or repay indebtedness; supply chain interruptions or difficulties; changes in competitive or market conditions; changes in legislation or regulatory developments; our ability to obtain and maintain adequate protection for our intellectual property rights; the timing and uncertainty of the results of both the research and development and regulatory processes, including regulatory decisions, product recalls, withdrawals and other unusual items; domestic and foreign health care and cost containment reforms, including government pricing, tax and reimbursement policies; technological advances and patents obtained by competitors; the performance, including the approval, introduction, and consumer and physician acceptance of new products and the continuing acceptance of currently marketed products; our ability to integrate any newly acquired products into our portfolio and achieve any financial or commercial expectations; the impact that known and unknown side effects may have on market perception and consumer preference for our products; the effectiveness of advertising and other promotional campaigns; the timely and successful implementation of any strategic initiatives; unfavorable publicity regarding the misuse of opioids; the uncertainty associated with the identification of and successful consummation and execution of external corporate development initiatives and strategic partnering transactions; our ability to advance our strategic priorities, develop our product pipeline and continue to develop the market for QWO® and other products; and our ability to obtain and successfully manufacture, maintain and distribute a sufficient supply of products to meet market demand in a timely manner. In addition, U.S. and international economic conditions, including consumer confidence and debt levels, inflation, taxation, changes in interest and currency exchange rates, international relations, capital and credit availability, the status of financial markets and institutions, the impact of and response to the ongoing COVID-19 pandemic and the impact of continued economic volatility, can materially affect our results. Therefore, the reader is cautioned not to rely on these forward-looking statements. Endo expressly disclaims any intent or obligation to update these forward-looking statements, except as required to do so by law.

Additional information concerning risk factors, including those referenced above, can be found in press releases issued by Endo, as well as Endo's public periodic filings with the U.S. Securities and Exchange Commission and with securities regulators in Canada, including the discussion under the heading "Risk Factors" in Endo's most recent Annual Report on Form 10-K and any subsequent Quarterly Reports on Form 10-Q or other filings with the U.S. Securities and Exchange Commission. Copies of Endo's press releases and additional information about Endo are available at www.endo.com or you can contact the Endo Investor Relations Department at relations.investor@endo.com.

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