

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date of Report (Date of Earliest Event Reported): January 11, 2018

ENDO INTERNATIONAL PLC

(Exact Name of Registrant as Specified in Its Charter)

Ireland
(State or other jurisdiction
of incorporation)

001-36326
(Commission File Number)

68-0683755
(IRS Employer
Identification No.)

**First Floor, Minerva House, Simmonscourt Road,
Ballsbridge, Dublin 4, Ireland**
(Address of principal executive offices)

Not Applicable
(Zip Code)

Registrant's telephone number, including area code 011-353-1-268-2000

Not Applicable
Former name or former address, if changed since last report

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

- Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 8.01. Other Events.

On January 11, 2018, Endo International plc (“Endo”) issued a press release announcing that its subsidiary, Endo Pharmaceuticals Inc. (“EPI”), has received a grand jury subpoena from the United States Attorney’s Office for the Southern District of Florida seeking documents and information relating to products containing oxymorphone. A copy of the press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

To Endo’s knowledge, no proceedings have currently been initiated against EPI as a result of the subpoena, although Endo cannot predict whether or when proceedings might be initiated. Endo and EPI intend to be responsive to the subpoena and cooperate with any related government investigation. Neither Endo nor EPI can make assurances as to the time or resources that will need to be devoted to this inquiry or its final outcome, or the impact of this inquiry or any proceedings on Endo’s consolidated financial condition, results of operations and cash flows.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits.

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press Release of Endo International plc, dated January 11, 2018

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, hereunto duly authorized.

ENDO INTERNATIONAL PLC

By: /s/ Matthew J. Maletta
Name: Matthew J. Maletta
Title: Executive Vice President,
Chief Legal Officer

Dated: January 11, 2018



Endo Comments on United States Attorney’s Office Grand Jury Subpoena Relating to Products Containing Oxymorphone

DUBLIN, Jan. 11, 2018 -- Endo International plc (NASDAQ: ENDP) (“Endo”) today announced that its subsidiary, Endo Pharmaceuticals Inc. (“EPI”), has received a grand jury subpoena from the United States Attorney’s Office for the Southern District of Florida seeking documents and information relating to products containing oxymorphone.

The subpoena broadly requests documents including, among others, those produced in past or pending lawsuits and those relating to product safety and efficacy, overdoses, diversion, thefts, overprescribing, abuse/misuse, dependency or tolerance, withdrawal, addictiveness, adverse events and manipulation. The subpoena also requests distribution and other third party agreements, together with sales and marketing, training, financial, compensation and corporate information, as well as documents relating to interactions with various government agencies, including the U.S. Food and Drug Administration, Drug Enforcement Administration, Veterans Administration, Federal Trade Commission, Department of Health & Human Services, Medicare and Medicaid. Endo and EPI intend to be responsive to the subpoena and cooperate with any related government investigation.

In all circumstances, it is Endo’s policy to comply with applicable laws, rules, regulations and industry guidance governing the sale and marketing of pharmaceutical products.

About Endo International plc

Endo International plc (NASDAQ: ENDP) is a highly focused generics and specialty branded pharmaceutical company delivering quality medicines to patients in need through excellence in development, manufacturing and commercialization. Endo has global headquarters in Dublin, Ireland, and U.S. headquarters in Malvern, PA. Learn more at www.endo.com.

Cautionary Note Regarding Forward-Looking Statements

This press release contains “forward-looking statements,” including statements regarding a federal grand jury subpoena. All forward-looking statements in this press release reflect the Company’s current analysis of information and represent the Company’s judgment only as of the date of this press release. If underlying assumptions prove inaccurate or unknown risks or uncertainties materialize, actual results could vary materially from the Company’s expectations. Risks and uncertainties include, among other things, general industry and market conditions; technological advances and patents attained by competitors; challenges inherent in the research and development and regulatory processes, including regulatory decisions, product recalls, withdrawals and other unusual items; challenges related to product marketing, such as the unpredictability of market acceptance for new products and/or the acceptance of new indications for such products; inconsistency of treatment results among patients; potential difficulties in manufacturing; the outcome of litigation, settlement discussions or other adverse proceedings; general economic conditions; and governmental laws and regulations affecting domestic and foreign operations. The Company expressly disclaims any intent or obligation to update these forward-looking statements except as required by law. Additional information concerning these and other risk factors can be found in the Company’s periodic reports filed with the U.S. Securities and Exchange Commission and in Canada on the System for Electronic Data Analysis and Retrieval (“SEDAR”), including current reports on Form 8-K, quarterly reports on Form 10-Q and annual reports on Form 10-K.

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