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

January 13, 2014
Date of Report (Date of earliest event reported)

JAZZ PHARMACEUTICALS PUBLIC LIMITED COMPANY
(Exact name of registrant as specified in its charter)

Ireland	**001-33500**	**98-1032470**
(State or other jurisdiction of incorporation)	**(Commission File No.)**	**(IRS Employer Identification No.)**

Fourth Floor, Connaught House,
1 Burlington Road, Dublin 4, Ireland
(Address of principal executive offices, including zip code)

011-353-1-634-7800
(Registrant's telephone number, including area code)

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))

Item 2.02 Results of Operations and Financial Condition.

On January 13, 2014, at the J.P. Morgan Healthcare Conference in San Francisco, California, Jazz Pharmaceuticals plc (the "Company") presented a corporate overview and financial update, which presentation included a confirmation of the Company's current expectations with respect to certain of the financial guidance for the year ended December 31, 2013 that the Company previously provided on November 5, 2013. The presentation was announced by a widely disseminated press release and was made available to the public via audio webcast, and the slides that accompanied the presentation were available to the public at the time of the webcast through the Company's website. A transcript of the relevant portion of the presentation relating to the confirmation of certain of its aforementioned financial guidance is attached hereto as Exhibit 99.1, along with a copy of the relevant slides containing such information.

The information contained in this Item 2.02 and in the accompanying Exhibit 99.1 to this current report shall not be deemed to be "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, or otherwise subject to the liabilities of that Section or Sections 11 and 12(a)(2) of the Securities Act of 1933, as amended. The information contained in this Item 2.02 and in the accompanying Exhibit 99.1 to this current report shall not be incorporated by reference into any filing with the U.S. Securities and Exchange Commission made by the Company, whether made before or after the date hereof, regardless of any general incorporation language in such filing.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit Number	Description
99.1	Portion of slides and related transcript of presentation by Jazz Pharmaceuticals plc on January 13, 2014

SIGNATURES

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.

JAZZ PHARMACEUTICALS PUBLIC LIMITED COMPANY

By: /s/ Suzanne Sawochka Hooper
Name: Suzanne Sawochka Hooper
Title: Executive Vice President and General Counsel

Date: January 14, 2014

EXHIBIT INDEX

Exhibit Number	Description
99.1	Portion of slides and related transcript of presentation by Jazz Pharmaceuticals plc on January 13, 2014

Exhibit 99.1

Relevant portion of the slides presented by Jazz Pharmaceuticals plc at the J.P. Morgan Healthcare Conference in San Francisco, California on January 13, 2014:

32nd Annual JP Morgan Healthcare Conference

Bruce Cozadd, Chairman and CEO

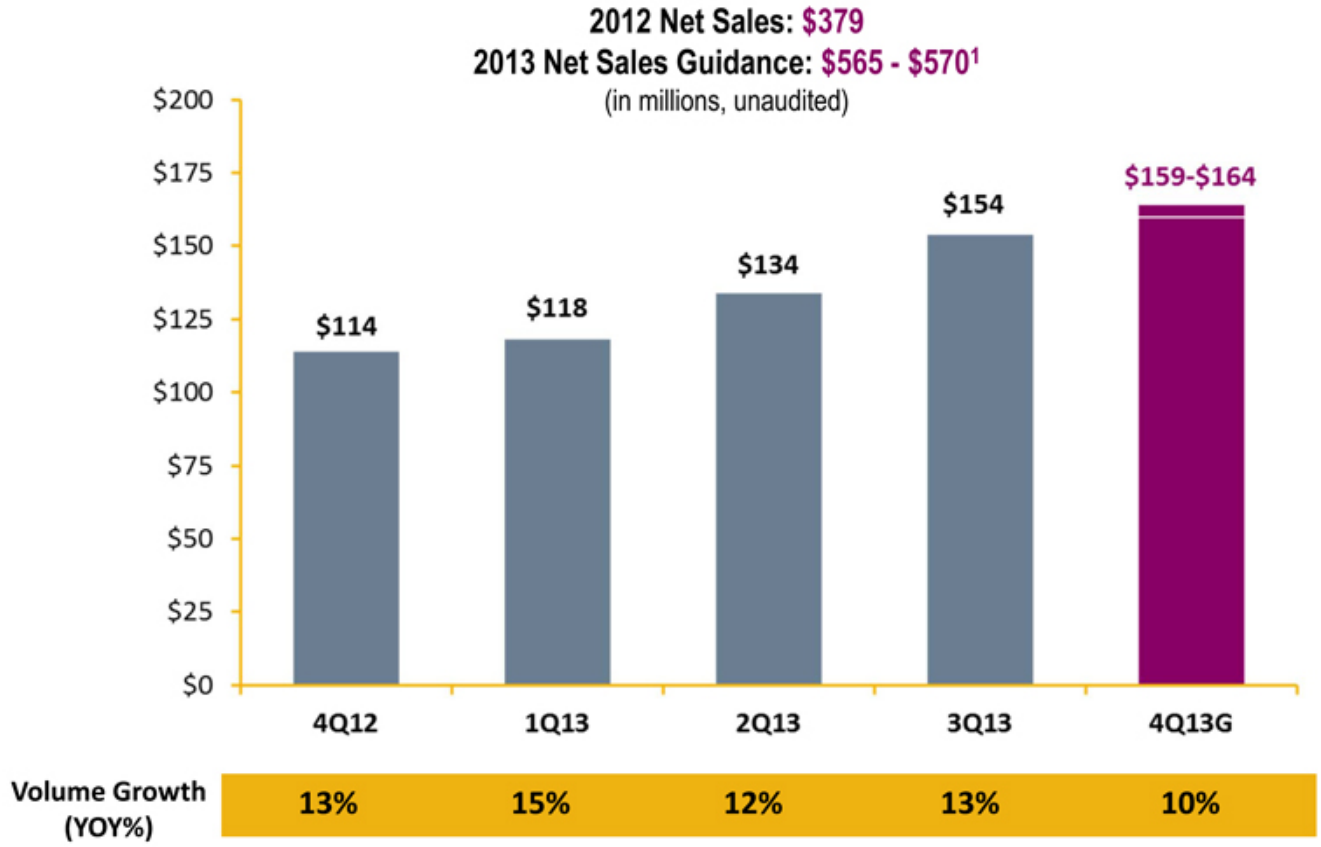
January 13, 2014

Forward-Looking Statements

"Safe Harbor" Statement Under the Private Securities Litigation Reform Act of 1995

This presentation contains forward-looking statements, including, but not limited to, statements related to Jazz Pharmaceuticals' estimated financial results and future growth potential, including expectations and estimates regarding 2013 sales of Xyrem® (sodium oxybate) oral solution and Erwinaze® (asparaginase *Erwinia chrysanthemi*), other financial and operating results and financial guidance, the company's growth and acquisition strategy and its 2014 goals (financial and otherwise), the therapeutic and commercial potential of the company's product candidates, potential future clinical trials and other development of the company's product candidates and the indications the company plans to pursue, potential approval and commercialization of the company's product candidates, expected patent protection for ADX-N05 and the potential extension of that patent protection, the anticipated consummation of the tender offer for Gentium S.p.A. ordinary shares and American Depositary Shares and the timing and benefits thereof, the plan to launch Defitelio™ (defibrotide) and the timing thereof, future commercial opportunities and potential expansion of European commercial operations after the expected completion of the Gentium acquisition, the potential to develop Defitelio for approval in other conditions, anticipated pipeline opportunities, future clinical development and regulatory matters, the expected launch of Versacloz™ (clozapine, USP) oral suspension and other statements that are not historical facts. These forward-looking statements are based on Jazz Pharmaceuticals' current expectations and inherently involve significant risks and uncertainties. Actual results and the timing of events could differ materially from those anticipated in such forward-looking statements as a result of these risks and uncertainties, which include, without limitation, risks and uncertainties associated with maintaining and increasing sales of and revenue from Xyrem, such as the potential introduction of generic competition and changed or increased regulatory restrictions on or requirements with respect to Xyrem, as well as similar risks related to effectively commercializing the company's other marketed products, including Erwinaze and Prialt® (ziconotide) intrathecal infusion; protecting and expanding the company's intellectual property rights; obtaining appropriate pricing and reimbursement for the company's products in an increasingly challenging environment; ongoing regulation and oversight by U.S. and non-U.S. regulatory agencies; dependence on key customers and sole source suppliers; the difficulty and uncertainty of pharmaceutical product development, including the timing thereof, the uncertainty of clinical success, such as the risk that results from early clinical trials may not be predictive of results obtained in later and larger clinical trials planned or anticipated to be conducted for the company's product candidates, and the uncertainty of regulatory approval; the company's ability to successfully manage the risks associated with integrating ADX-N05 and other acquired products or product candidates into the company's product portfolio, including the availability of funding to complete the development of, obtain regulatory approval for and commercialize acquired product candidates; the satisfaction of closing conditions and the availability and terms of the financing for the proposed Gentium acquisition; risks associated with business combination or product acquisition transactions, such as the risk that the acquired business or the products or product candidates acquired will not be integrated successfully or that such integration may be more difficult, time-consuming or costly than expected; risks related to future opportunities and plans for the combined company following the proposed Gentium acquisition, including uncertainty of the expected financial performance and results of the combined company following completion of the proposed acquisition; disruption from the proposed Gentium acquisition, making it more difficult to conduct business as usual or maintain relationships with customers, employees or suppliers; the possibility that if Jazz Pharmaceuticals does not achieve the perceived or anticipated benefits of the proposed acquisition of Gentium or its acquisition of ADX-N05 as rapidly or to the extent anticipated by financial analysts or investors, the market price of Jazz Pharmaceuticals' ordinary shares could decline; the company's ability to identify and acquire, in-license or develop additional products or product candidates to grow its business; and possible restrictions on the company's ability and flexibility to pursue certain future opportunities as a result of its substantial outstanding debt obligations; as well as risks related to future opportunities and plans, including the uncertainty of expected future financial performance and results, and those other risks detailed from time-to-time under the caption "Risk Factors" and elsewhere in Jazz Pharmaceuticals plc's Securities and Exchange Commission filings and reports (Commission File No. 001-33500), including in the Quarterly Report on Form 10-Q for the quarter ended September 30, 2013 and future filings and reports by the company. Jazz Pharmaceuticals undertakes no duty or obligation to update any forward-looking statements contained in this presentation as a result of new information, future events or changes in its expectations.

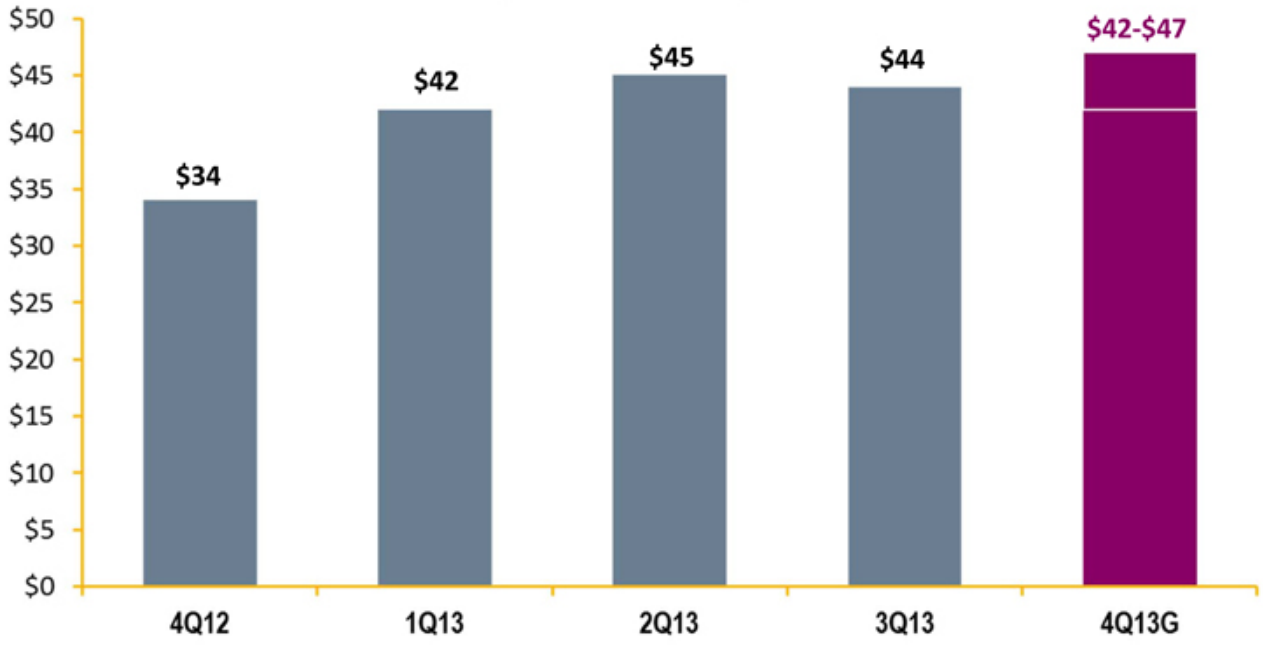
Xyrem: Strong Sales Growth



[1] G = Guidance. Guidance provided by Jazz Pharmaceuticals plc on and as of November 5, 2013. Jazz Pharmaceuticals plc currently expects that, for the year ended December 31, 2013, reported Xyrem net sales will meet the guidance range provided on November 5, 2013. The company has not finalized its financial results for the quarter and year ended December 31, 2013, and actual results may differ.

Erwinaze/Erwinase: Building for Success

2012 Pro Forma Net Sales: $132[1]
2013 Net Sales Guidance: $173 - $178[2]
(in millions, unaudited)



[1] Pro forma 2012 net sales of Erwinaze/Erwinase include net sales of $60 million from the historic EUSA Pharma business from January 1, 2012 through June 12, 2012, the closing date of the EUSA Pharma acquisition.
[2] G=Guidance. Guidance provided by Jazz Pharmaceuticals plc on and as of November 5, 2013. Jazz Pharmaceuticals plc currently expects that, for the year ended December 31, 2013, reported Erwinaze/Erwinase net sales will meet the guidance range provided on November 5, 2013. The company has not finalized its financial results for the quarter and year ended December 31, 2013, and actual results may differ.

Relevant portion of the transcript of the oral presentation by Jazz Pharmaceuticals plc at the J.P. Morgan Healthcare Conference in San Francisco, California on January 13, 2014:

Bruce C. Cozadd, Chairman & CEO, Jazz Pharmaceuticals plc

. . .

During my presentation today, I will make forward-looking statements. Actual results and timing may differ due to risks and uncertainties described here but also in our SEC filings.

. . .

Xyrem sales growth remains strong with annualized revenues north of $600 million. I can tell you today that we do expect to meet our Xyrem top-line guidance for 2013, which would represent growth of 50 percent over 2012. And volume growth remained strong. We saw 10 percent volume growth in the fourth quarter. This continues a string of double digit year-over-year volume growth for this product and gives us 12 percent volume growth for the year. We think that growth will continue. We are seeing growth in prescriptions by our lower and mid decile physicians, and we've recently expanded our sales force by 25 percent, to allow us to make more calls on a broader physician universe.

. . .

And, again, today, we can confirm that we will meet prior guidance, giving us sales for the year of $173 to $178 million, up 30 to 35 percent from pro forma 2012 sales. I can also tell you that, based on this, we expect that we will trigger the $50 million milestone payment to the EUSA shareholders.

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