

Protalix BioTherapeutics, Inc.  
Form 8-K  
August 29, 2014

---

**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): August 28, 2014**

---

**Protalix BioTherapeutics, Inc.**

**(Exact name of registrant as specified in its charter)**

<b>Florida</b>	<b>001-33357</b>	<b>65-0643773</b>
<b>(State or other jurisdiction</b>	<b>(Commission File Number)</b>	<b>(IRS Employer</b>
<b>of incorporation)</b>		<b>Identification No.)</b>

**2 Snunit Street  
Science Park, POB 455**

**20100**

**Carmiel, Israel**  
**(Address of principal executive offices) (Zip Code)**

**Registrant's telephone number, including area code +972-4-988-9488**

**(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 (see General Instruction A.2. below):

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 8.01. Other Events**

On August 28, 2014, Protalix BioTherapeutics, Inc. (the “Company”) and Pfizer Inc. (“Pfizer”) issued a joint press release announcing that the U.S. Food and Drug Administration (FDA) approved ELELYSO™ (taliglucerase alfa) for injection for pediatric patients. ELELYSO is therefore now indicated for long-term enzyme replacement therapy (ERT) for adult and pediatric patients with a confirmed diagnosis of Type 1 Gaucher disease. Also on August 28, 2014, the Company issued a second press release announcing that the Company will host a conference call on Wednesday, September 3, 2014 at 8:30am ET to discuss the approval of ELELYSO for pediatric patients described herein. In addition, the Company’s management will provide an update on the additional ongoing clinical programs, PRX-112 and PRX-102, and hold a Q&A session. Copies of the press releases are attached hereto as Exhibits 99.1 and 99.2.

**Item 9.01. Financial Statements and Exhibits**

**(d) Exhibits**

99.1 Press release dated August 28, 2014.

99.2 Press release dated August 28, 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.

**PROTALIX  
BIOTHERAPEUTICS, INC.**

Date: August 28, 2014 By: /s/ David Aviezer  
Name: David Aviezer, Ph.D.  
Title: President and  
Chief Executive Officer