



Item 8.01 Other Events.

On September 12, 2013, Raptor Pharmaceutical Corp. issued a press release announcing that the European Commission has approved PROCYSBI® gastro-resistant hard capsules of cysteamine (as mercaptamine bitartrate) as an orphan medicinal product for the treatment of proven nephropathic cystinosis for marketing in the European Union. A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference into this Item 8.01.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

Exhibit No. Exhibit Description

99.1 Press release dated September 12, 2013.

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SIGNATURES

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

Date: September 12, 2013 RAPTOR PHARMACEUTICAL CORP.

By: /s/ Georgia Erbez

Name: Georgia Erbez

Title: Chief Financial Officer, Secretary and Treasurer

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Exhibit Index

Exhibit No.	Exhibit Description
99.1	Press release dated September 12, 2013.