

ALEXION PHARMACEUTICALS INC  
Form 8-K  
October 06, 2006

---

**UNITED STATES**  
**SECURITIES AND EXCHANGE COMMISSION**  
**WASHINGTON, D.C. 20549**

---

**FORM 8-K**

---

**CURRENT REPORT**

**PURSUANT TO SECTION 13 OR 15(D) OF THE**  
**THE SECURITIES EXCHANGE ACT OF 1934**

**Date of report (Date of earliest event reported): September 20, 2006**

---

**ALEXION PHARMACEUTICALS, INC.**

(Exact name of registrant as specified in its charter)

---

**Delaware**  
(State or other jurisdiction of  
incorporation or organization)

**000-27756**  
(Commission File Number)

**13-3648318**  
(I.R.S. Employer

Identification No.)

**352 Knotter Drive, Cheshire, Connecticut 06410**

(Address of Principal Executive Offices) (Zip Code)

**Registrant's telephone number, including area code: (203) 272-2596**

---

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):

Edgar Filing: ALEXION PHARMACEUTICALS INC - Form 8-K

- .. 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 September 20, 2006, Alexion Pharmaceuticals, Inc. (the Company) issued a press release announcing that the Company has submitted a Biologics License Application with the U.S. Food and Drug Administration for its lead product candidate Soliris (eculizumab), as a treatment for patients diagnosed with paroxysmal nocturnal hemoglobinuria, a rare life-threatening genetic blood disorder (PNH). A copy of that press release is furnished as Exhibit 99.1 to this Form 8-K.

On September 26, 2006, the Company issued a press release announcing that Alexion Europe, a wholly-owned subsidiary of the Company, has submitted a Market Authorization Application to the European Medicines Evaluation Agency for Soliris (eculizumab) as a treatment for patients diagnosed with PNH. A copy of that press release is furnished as Exhibit 99.2 to this

Form 8-K.

**Item 9.01 Financial Statements and Exhibits.**

(c) Exhibits

99.1 Press Release issued by Alexion Pharmaceuticals, Inc. on September 20, 2006.

99.2 Press Release issued by Alexion Pharmaceuticals, Inc. on September 26, 2006.

**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.

ALEXION PHARMACEUTICALS, INC.

Date: October 5, 2006

By: /s/ Thomas I. H. Dubin

Name: Thomas I. H. Dubin

Title: Senior Vice President and General Counsel

**Index to Exhibits**

<b>Exhibit No.</b>	<b>Description</b>
99.1	Press Release issued by Alexion Pharmaceuticals, Inc. on September 20, 2006.
99.2	Press Release issued by Alexion Pharmaceuticals, Inc. on September 26, 2006.