

REGENERON PHARMACEUTICALS INC

Form 8-K

November 21, 2011

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): November 18, 2011 (November 18, 2011)

REGENERON PHARMACEUTICALS, INC.  
(Exact Name of Registrant as Specified in Charter)

New York  
(State or other jurisdiction of  
Incorporation)

000-19034  
(Commission File No.)

13-3444607  
(IRS Employer Identification No.)

777 Old Saw Mill River Road, Tarrytown, New York 10591-6707  
(Address of principal executive offices, including zip code)

(914) 347-7000  
(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))
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### Item 8.01 Other Events.

On November 18, 2011, Regeneron Pharmaceuticals, Inc. issued a press release announcing that the U.S. Food and Drug Administration approved EYLEA™ (aflibercept) Injection for the treatment of patients with neovascular (wet) Age-related Macular Degeneration (AMD).

A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K and incorporated by reference into this Item.

### Item 9.01 Financial Statements and Exhibits.

#### (d) Exhibits

99.1 Press Release Announcing FDA Approval of EYLEA™ (aflibercept) Injection for the Treatment of Wet Age-Related Macular Degeneration, dated November 18, 2011.

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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.

Date: November 18, 2011

REGENERON PHARMACEUTICALS, INC.

By: /s/ Joseph J. LaRosa

Name: Joseph J. LaRosa

Title: Senior Vice President, General Counsel and Secretary

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Exhibit Index

Number	Description
99.1	Press Release Announcing FDA Approval of EYLEA™ (aflibercept) Injection for the Treatment of Wet Age-Related Macular Degeneration, dated November 18, 2011.

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