REGENERON PHARMACEUTICALS INC
Form 8-K
March 16, 2015

# 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): **March 16, 2015 (March 14, 2015)**

# REGENERON PHARMACEUTICALS, INC.

(Exact name of registrant as specified in its charter)

**New York**

(State or other jurisdiction of incorporation)

| **000-19034** | **13-3444607** |
|---|---|
| (Commission File Number) | (I.R.S. Employer Identification No.) |
| **777 Old Saw Mill River Road, Tarrytown, New York** | **10591-6707** |
| (Address of principal executive offices) | (Zip Code) |

Registrant s telephone number, including area code: **(914) 847-7000**

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 Instructions A.2. below):

o Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

o Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

o Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

o Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

**Item 7.01. Regulation FD Disclosure.**

As previously announced, on March 14, 2015, positive results from the ODYSSEY CHOICE I and CHOICE II trials, which evaluated monthly dosing of PRALUENT (alirocumab) 300 mg and PRALUENT 150 mg, were presented at the American College of Cardiology s 64th Annual Scientific Sessions & Expo ( ACC 15 ) in San Diego, California. A copy of the poster presentation is furnished as Exhibit 99.1 to this Current Report on Form 8-K and is incorporated by reference herein.

On March 16, 2015, a pooled analysis of adverse events from four Phase 2 and five Phase 3 double-blind, placebo-controlled trials exploring multiple PRALUENT doses and regimens will be presented at ACC 15. A copy of the presentation slides is furnished as Exhibit 99.2 to this Current Report on Form 8-K and is incorporated by reference herein.

**Item 9.01. Financial Statements and Exhibits.**

(d) Exhibits

| | |
|---|---|
| 99.1 | Poster presentation entitled Efficacy and safety of alirocumab 150 mg and 300 mg every 4 weeks in patients with poorly controlled hypercholesterolemia: the ODYSSEY CHOICE I and CHOICE II studies. |
| 99.2 | Presentation slides entitled Pooled Safety and Adverse Events in Nine Randomized, Placebo-controlled, Phase 2 and 3 Clinical Trials of Alirocumab. |

**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 thereunto duly authorized.

**REGENERON PHARMACEUTICALS, INC.**

/s/ Joseph J. LaRosa
Joseph J. LaRosa
Senior Vice President, General Counsel and Secretary

Date: March 16, 2015

**EXHIBIT INDEX**

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