CELGENE CORP /DE/
Form 8-K
February 07, 2018

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): February 6, 2018

CELGENE CORPORATION

(Exact name of registrant as specified in its charter)

Delaware	001-34912	22-2711928
(State or other jurisdiction of incorporation)	(Commission File Number)	(IRS Employer Identification No.)

86 Morris Avenue, Summit, New Jersey 07901
(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (908) 673-9000

(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))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ¨

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ¨

ITEM 8.01 OTHER EVENTS

On February 6, 2018, Celgene Corporation issued a press release announcing that the Phase III, randomized, open-label, international clinical study, OPTIMISMM (evaluating the efficacy and safety of POMALYST/IMNOVID (pomalidomide) plus bortezomib and low-dose dexamethasone (PVd) versus bortezomib and low-dose dexamethasone in patients with relapsed/refractory multiple myeloma), achieved its primary endpoint, showing a statistically significant and clinically meaningful improvement in progression-free survival (PFS) for the pomalidomide arm versus the comparator arm.

Attached hereto and incorporated herein by reference as Exhibit 99.1 is the press release announcement.

ITEM 9.01 FINANCIAL STATEMENTS AND EXHIBITS.

Exhibit 99.1 – Press Release dated February 6, 2018

SIGNATURES

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

CELGENE CORPORATION

Date: February 6, 2018 By: /s/ Peter N. Kellogg
Peter N. Kellogg
Executive Vice President and
Chief Financial Officer
(principal financial and accounting officer)

EXHIBIT INDEX

Exhibit No.	Description
99.1	Press Release dated February 6, 2018