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CELGENE CORP /DE/ Form 8-K June 20, 2007

Delaware

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): June 19, 2007

CELGENE CORPORATION
(Exact name of registrant as specified in its charter)

0-16132

22-2711928

| | te or other jurisdiction (Commission File Number) of incorporation) | (IRS Employer Identification No. |
|--|---|----------------------------------|
| 86 | Morris Avenue, Summit, New Jersey | 07901 |
| (Add | ress 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 (17 CFR 230.425) | Securities Act |
| [] | Soliciting material pursuant to Rule $14a-12$ under the (17 CFR $240.14a-12$) | Exchange Act |
| [] | Pre-commencement communications pursuant to Rule 14d-2 Exchange Act (17 CFR 240.14d-2(b)) | (b) under the |
| [] | Pre-commencement communications pursuant to Rule 13e-4 Exchange Act (17 CFR 240.13e-4(c)) | (c) under the |

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ITEM 8.01 OTHER EVENTS

On June 19, 2007, Celgene Corporation announced that REVLIMID(R) (lenalidomide) has been granted full marketing authorization by the European Commission for use in combination with dexamethasone as a treatment for patients with multiple myeloma who have received at least one prior therapy. Multiple myeloma is the second most commonly diagnosed blood cancer. According to the International Myeloma Foundation, there are an estimated 750,000 people with multiple myeloma worldwide. There are more than 85,000 men and women in Europe currently undergoing treatment for multiple myeloma, and 25,000 people are expected to die from this blood cancer in 2007.

The Marketing Authorization Application (MAA) for REVLIMID(R) was based upon the safety and efficacy results of two large, randomized pivotal Phase III special protocol assessment trials, North American Trial MM-009 and International Trial MM-010, evaluating REVLIMID(R) plus dexamethasone in multiple myeloma patients that have received at least one prior therapy.

Attached hereto and incorporated herein by reference as Exhibit 99.1 is the Press Release announcing such information.

ITEM 9.01 FINANCIAL STATEMENTS AND EXHIBITS.

(d) Exhibit 99.1 - Press Release dated June 19, 2007

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: June 19, 2007 By: /s/ Robert J. Hugin

Name: Robert J. Hugin
Title: President and

Chief Operating Officer