MEDICIS PHARMACEUTICAL CORP
Form 8-K
December 15, 2010

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
December 13, 2010
Date of Report (Date of earliest event reported)
Medicis Pharmaceutical Corporation
(Exact name of registrant as specified in its charter)

Delaware	**001-14471**	**52-1574808**
(State of Incorporation)	(Commission File Number)	(IRS Employer Identification Number)

7720 North Dobson Road
Scottsdale, Arizona 85256
(Address of principal executive offices) (Zip Code)
(602) 808-8800
(Registrant s telephone number, including area code)
N/A
(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:

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 8.01 Other Events.

The Company Receives a Paragraph IV Patent Certification from Ranbaxy Laboratories Limited

On December 13, 2010, Medicis Pharmaceutical Corporation (the Company) received a Paragraph IV Patent Certification from Ranbaxy Laboratories Limited (Ranbaxy) advising that Ranbaxy has filed a supplement or amendment to its earlier filed Abbreviated New Drug Application (ANDA) assigned ANDA number 91-118 (ANDA Supplement/Amendment) with the U.S. Food and Drug Administration (FDA) for generic versions of SOLODYN® (minocycline HCl, USP) Extended Release Tablets in 55mg and 105mg strengths. Ranbaxy has not advised the Company as to the timing or status of the FDA s review of its filing, or whether Ranbaxy has complied with FDA requirements for proving bioequivalence. Ranbaxy s Paragraph IV Certification alleges that the Company s U.S. Patent Nos. 5,908,838 (the 838 Patent) and 7,790,705 (the 705 Patent) will not be infringed by Ranbaxy s manufacture, importation, use, sale and/or offer for sale of the products for which the ANDA Supplement/Amendment was submitted because Ranbaxy has a licensing agreement with the Company. The expiration date for the 838 Patent is in 2018 and the expiration date for the 705 Patent is in 2025 or later.

SIGNATURES

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

Medicis Pharmaceutical Corporation

Date: December 15, 2010

By: /s/ Seth L. Rodner
Seth L. Rodner
Senior Vice President, General Counsel
and Corporate Secretary