

# WARNER CHILCOTT LTD

## FORM 8-K

(Current report filing)

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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION**  
Washington, D.C. 20549

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**Form 8-K**

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**Current Report Pursuant to Section 13 or 15(d) of the  
Securities Exchange Act of 1934**

Date of Report: March 17, 2009  
Date of earliest event reported: March 17, 2009

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**Warner Chilcott Limited**

(Exact name of registrant as specified in its charter)

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**Commission File Number: 1 – 33039**

**Bermuda**  
(State or other jurisdiction  
of incorporation)

**98-0496358**  
(IRS Employer  
Identification No.)

**Gibbons Building, 10 Queen Street, Suite 109 – First Floor**  
**Hamilton HM11, Bermuda**  
(Address of principal executive offices)

**(441) 292-0068**  
(Registrant's telephone number, including area code)

**Not Applicable**  
(Former name or former address, if changed since last report)

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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 March 17, 2009, Warner Chilcott Limited (the "Company") issued a press release announcing its receipt of a Paragraph IV Certification Notice from Impax Laboratories, Inc. advising that it has filed an Abbreviated New Drug Application ("ANDA") for a generic version of Doryx 150 mg delayed-release tablets. A copy of the Company's press release is filed as Exhibit 99.1 hereto and is incorporated by reference into this Item 8.01.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

<u>Exhibit Number</u>	<u>Description</u>
99.1	Press Release issued March 17, 2009 by Warner Chilcott Limited.

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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.

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By: /s/ Paul Herendeen

Name: Paul Herendeen

Title: Executive Vice President and Chief Financial  
Officer

Date: March 17, 2009

**WARNER CHILCOTT ANNOUNCES RECEIPT OF PARAGRAPH IV CERTIFICATION NOTICE**

HAMILTON, Bermuda, March 17, 2009 — Warner Chilcott Limited (Nasdaq: WCRX) announced today that one of its subsidiaries and Mayne Pharma International Pty. Ltd. (“Mayne”) have received a Paragraph IV Certification Notice from Impax Laboratories, Inc. (“Impax”) advising that Impax has filed an Abbreviated New Drug Application (ANDA) for a generic version of DORYX 150 mg delayed-release tablets.

DORYX, which Warner Chilcott markets and sells in 150, 100 and 75 mg strengths in the United States under a license agreement with Mayne, is a tetracycline-class oral antibiotic protected by Mayne’s Patent No. 6,958,161 (the “161 Patent”) which expires in 2022.

The Company and Mayne are currently reviewing the detail of the Paragraph IV Certification Notice from Impax and intend to vigorously defend the 161 patent and pursue their legal rights.

**About Warner Chilcott**

Warner Chilcott is a leading specialty pharmaceutical company currently focused on the women’s healthcare and dermatology segments of the U.S. pharmaceuticals market. The Company is a fully integrated company with internal resources dedicated to the development, manufacturing and promotion of its products. WCRX-G

Read more on <http://www.warnerchilcott.com>.

**Company Contact:** Rochelle Fuhrmann  
Investor Relations  
973-442-3281  
[rfuhrmann@wcrx.com](mailto:rfuhrmann@wcrx.com)

**Warner Chilcott’s Forward Looking Statements:**

This press release contains forward-looking statements, including statements concerning our operations, our anticipated economic performance and financial condition, and our business plans and growth strategy and product development efforts. These statements constitute forward-looking statements within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. The words “may,” “might,” “will,” “should,” “estimate,” “project,” “plan,” “anticipate,” “expect,” “intend,” “outlook,” “believe” and other similar expressions are

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intended to identify forward-looking statements. Readers are cautioned not to place undue reliance on these forward-looking statements, which speak only as of their dates. These forward-looking statements are based on estimates and assumptions by our management that, although we believe to be reasonable, are inherently uncertain and subject to a number of risks and uncertainties.

The following represent some, but not necessarily all, of the factors that could cause actual results to differ from historical results or those anticipated or predicted by our forward-looking statements: our substantial indebtedness; competitive factors in the industry in which we operate (including the approval and introduction of generic or branded products that compete with our products); our ability to protect our intellectual property; a delay in qualifying our manufacturing facility to produce our products or production or regulatory problems with either third party manufacturers upon whom we may rely for some of our products or our own manufacturing facilities; pricing pressures from reimbursement policies of private managed care organizations and other third party payors, government sponsored health systems, the continued consolidation of the distribution network through which we sell our products, including wholesale drug distributors and the growth of large retail drug store chains; the loss of key senior management or scientific staff; adverse outcomes in our outstanding litigation or an increase in the number of litigation matters to which we are subject; government regulation affecting the development, manufacture, marketing and sale of pharmaceutical products, including our ability and the ability of companies with whom we do business to obtain necessary regulatory approvals; our ability to manage the growth of our business by successfully identifying, developing, acquiring or licensing new products at favorable prices and marketing such new products; our ability to obtain regulatory approval and customer acceptance of new products, and continued customer acceptance of our existing products; changes in tax laws or interpretations that could increase our consolidated tax liabilities; the other risks identified in our Annual Report on Form 10-K for the year ended December 31, 2008; and other risks detailed from time-to-time in our public filings, financial statements and other investor communications.

We caution you that the foregoing list of important factors is not exclusive. In addition, in light of these risks and uncertainties, the matters referred to in our forward-looking statements may not occur. We undertake no obligation to publicly update or revise any forward-looking statement as a result of new information, future events or otherwise, except as may be required by law.