

TEVA PHARMACEUTICAL INDUSTRIES LTD
Form 6-K
March 03, 2008

FORM 6-K

SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

Report of Foreign Private Issuer

**Pursuant to Rule 13a-16 or 15d-16
under the Securities Exchange Act of 1934**

For the month of February 2008

Commission File Number 0-16174

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Teva Pharmaceutical Industries Limited

(Translation of registrant's name into English)

5 Basel Street, P.O. Box 3190

Petach Tikva 49131 Israel

(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F:

Form 20-F X

Form 40-F

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Indicate by check mark whether by furnishing the information contained in this Form, the registrant is also hereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes

No X

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g(3)-2(b):
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Teva Pharmaceutical Industries Ltd.

Web Site: www.tevapharm.com

For Immediate Release

TEVA PROVIDES UPDATE ON GENERIC ACTONEL[®] LITIGATION

Jerusalem, Israel, February 29, 2008 - Teva Pharmaceutical Industries Ltd. (NASDAQ: TEVA) announced today that the U.S. District Court for the District of Delaware has issued a decision in its litigation over the Company's Abbreviated New Drug Application (ANDA) to market its generic version of Procter & Gamble's Actonel[®] (Risedronate Sodium) Tablets, 5mg, 30mg and 35 mg. The Court found Procter & Gamble's U.S. Patent No. 5,538,122 to be valid. Teva intends to appeal this decision.

About Teva

Teva Pharmaceutical Industries Ltd., headquartered in Israel, is among the top 20 pharmaceutical companies in the world and is the leading generic pharmaceutical company. The company develops, manufactures and markets generic and innovative human pharmaceuticals and active pharmaceutical ingredients. Over 80 percent of Teva's sales are in North America and Europe.

Safe Harbor Statement under the U. S. Private Securities Litigation Reform Act of 1995:

This release contains forward-looking statements, which express the current beliefs and expectations of management. Such statements are based on management's current beliefs and expectations and involve a number of known and unknown risks and uncertainties that could cause Teva's future results, performance or achievements to differ significantly from the results, performance or achievements expressed or implied by such forward-looking statements. Important factors that could cause or contribute to such differences include risks relating to: Teva's ability to rapidly integrate CoGenesys' operations with its own operations, the diversion of management time on merger-related issues, and Teva and CoGenesys' ability to successfully develop and commercialize biopharmaceutical products, Teva's ability to accurately predict future market conditions, potential liability for sales of generic products prior to a final

resolution of outstanding patent litigation, including that relating to the generic versions of Allegra®
Neurontin®, Lotrel® Famvir®, and Protonix®, Teva's ability to successfully develop and commercialize
additional pharmaceutical products, the introduction of competing generic equivalents, the extent to which Teva may
obtain U.S. market exclusivity for certain of its new generic products and regulatory changes that may prevent Teva
from utilizing exclusivity periods, competition from brand-name companies that are under increased pressure to
counter generic products, or competitors that seek to delay the introduction of generic products, the impact of
consolidation of our distributors and customers, the effects of competition on our innovative products, especially
Copaxone® sales, the impact of pharmaceutical industry regulation and pending legislation that could affect the
pharmaceutical industry, the difficulty of predicting U.S. Food and Drug Administration, European Medicines Agency
and other regulatory authority approvals, the regulatory environment and changes in the health policies and structures
of various countries, our ability to achieve expected results through our innovative R&D efforts, Teva's ability to
successfully identify, consummate and integrate acquisitions, potential exposure to product liability claims to the
extent not covered by insurance, dependence on the effectiveness of our patents and other protections for innovative
products, significant operations worldwide that may be adversely affected by terrorism, political or economical
instability or major hostilities, supply interruptions or delays that could result from the complex manufacturing of our
products and our global supply chain, environmental risks, fluctuations in currency, exchange and interest rates, and
other factors that are discussed in Teva's Annual Report on Form 20-F and its other filings with the U.S. Securities
and Exchange Commission. Forward-looking statements speak only as of the date on which they are made and the
Company undertakes no obligation to update or revise any forward-looking statement, whether as a result of new
information, future events or otherwise.

Teva Pharmaceutical Industries Ltd.

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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, thereunto duly authorized.

TEVA PHARMACEUTICAL INDUSTRIES LIMITED

(Registrant)

By: /s/ Dan Suesskind

Name: Dan Suesskind
Title: Chief Financial Officer

Date: February 29 , 2008

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