

Teva Pharmaceuticals And Perrigo Company Announce The U.S. Launch Of Generic Temozolomide

JERUSALEM and ALLEGAN, Mich., Aug. 12, 2013 [/PRNewswire/](#) -- Teva Pharmaceutical Industries Ltd. (NYSE: TEVA) and Perrigo Company (NYSE: PRGO;TASE) today announced the launch of the generic equivalent to Temodar® (temozolomide). Teva will manufacture, market and distribute the product in the U.S. and both companies will equally share in the cost and profitability of the product in the U.S. Teva was first to file, making the product eligible for 180 days of marketing exclusivity.

(Logo: <http://photos.prnewswire.com/prnh/20120301/DE62255LOGO>)

This product is the generic equivalent to Temodar® (temozolomide), indicated for the treatment of adult patients with newly diagnosed glioblastoma multiforme concomitantly with radiotherapy and then as maintenance treatment and refractory anaplastic astrocytoma patients who have experienced disease progression on a drug regimen containing nitrosourea and procarbazine. Temodar® had annual sales of approximately \$423 million in the United States, according to IMS data as of December 31, 2012. The launch of this product provides a quality alternative, making cancer therapy more cost effective for patients who suffer from this devastating cancer.

Perrigo's Chairman, President and CEO Joseph C. Papa stated, "This first-to-file launch with our partner Teva is another example of our focus to manufacture complex API's. We are pleased to offer this important product to patients in the United States."

"We are pleased to partner with Perrigo to offer patients a high-quality, less expensive alternative of this important medicine. This launch demonstrates Teva's commitment to continue to pursue first-to-market opportunities and enhance the value of our portfolio by concentrating on high-margin, low competition markets," stated Allan Oberman, President and CEO of Teva Americas Generics.

About Teva

Teva Pharmaceutical Industries Ltd. (NYSE: TEVA) is a leading global pharmaceutical company, committed to increasing access to high-quality healthcare by developing, producing and marketing affordable generic drugs as well as innovative and specialty pharmaceuticals and active pharmaceutical ingredients. Headquartered in Israel, Teva is the world's leading generic drug maker, with a global product portfolio of more than 1,000 molecules and a direct presence in about 60 countries. Teva's branded businesses focus on CNS, oncology, pain, respiratory and women's health therapeutic areas as well as biologics. Teva currently employs approximately 46,000 people around the world and reached \$20.3 billion in net revenues in 2012.

About Perrigo

From its beginnings as a packager of generic home remedies in 1887, Allegan, Michigan-based Perrigo Company has grown to become a leading global provider of quality, affordable healthcare products. Perrigo develops, manufactures and distributes over-the-counter (OTC) and generic prescription (Rx) pharmaceuticals, infant formulas, nutritional products, animal health, dietary supplements and active pharmaceutical ingredients (API). The Company is the world's largest manufacturer of OTC pharmaceutical products for the store brand market. The Company's primary markets and locations of logistics operations have evolved over the years to include the United States, Israel, Mexico, the United

Kingdom, India, China and Australia. Visit Perrigo on the Internet (<http://www.perrigo.com>).

Teva's 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 our 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: our ability to develop and commercialize additional pharmaceutical products, including our ability to develop, manufacture, market and sell biopharmaceutical products, competition for our innovative products, especially Copaxone® (including competition from innovative orally-administered alternatives, as well as from potential purported generic equivalents), competition for our generic products (including from other pharmaceutical companies and as a result of increased governmental pricing pressures), competition for our specialty pharmaceutical businesses, our ability to achieve expected results through our specialty, including innovative, R&D efforts, the effectiveness of our patents and other protections for innovative products, decreasing opportunities to obtain U.S. market exclusivity for significant new generic products, our ability to identify, consummate and successfully integrate acquisitions, the effects of increased leverage as a result of recent acquisitions, the extent to which any manufacturing or quality control problems damage our reputation for high quality production and require costly remediation, our potential exposure to product liability claims to the extent not covered by insurance, increased government scrutiny in both the U.S. and Europe of our agreements with brand companies, potential liability for sales of generic products prior to a final resolution of outstanding patent litigation. our exposure to currency fluctuations and restrictions as well as credit risks, the effects of reforms in healthcare regulation and pharmaceutical pricing and reimbursement, any failures to comply with complex Medicare and Medicaid reporting and payment obligations, governmental investigations into sales and marketing practices (particularly for our specialty pharmaceutical products), uncertainties surrounding the legislative and regulatory pathways for the registration and approval of biotechnology-based products, adverse effects of political or economical instability, corruption, major hostilities or acts of terrorism on our significant worldwide operations, interruptions in our supply chain or problems with our information technology systems that adversely affect our complex manufacturing processes, any failure to retain key personnel or to attract additional executive and managerial talent, the impact of continuing consolidation of our distributors and customers, variations in patent laws that may adversely affect our ability to manufacture our products in the most efficient manner, potentially significant impairments of intangible assets and goodwill, potential increases in tax liabilities, the termination or expiration of governmental programs or tax benefits, environmental risks and other factors that are discussed in our Annual Report on Form 20-F for the year ended December 31, 2012 and in our 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.

Perrigo's Safe Harbor Statement: Certain statements in this press release are forward-looking statements within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended, and are subject to the safe harbor created thereby. These statements relate to future events or the Company's future financial performance and involve known

and unknown risks, uncertainties and other factors that may cause the actual results, levels of activity, performance or achievements of the Company or its industry to be materially different from those expressed or implied by any forward-looking statements. In some cases, forward-looking statements can be identified by terminology such as "may," "will," "could," "would," "should," "expect," "plan," "anticipate," "intend," "believe," "estimate," "predict," "potential" or other comparable terminology. The Company has based these forward-looking statements on its current expectations, assumptions, estimates and projections. While the Company believes these expectations, assumptions, estimates and projections are reasonable, such forward-looking statements are only predictions and involve known and unknown risks and uncertainties, many of which are beyond the Company's control. These and other important factors, including those discussed under "Risk Factors" in the Company's Form 10-K for the year ended June 30, 2012, as well as the Company's subsequent filings with the Securities and Exchange Commission, may cause actual results, performance or achievements to differ materially from those expressed or implied by these forward-looking statements. The forward-looking statements in this press release are made only as of the date hereof, and unless otherwise required by applicable securities laws, the Company disclaims any intention or obligation to update or revise any forward-looking statements, whether as a result of new information, future events or otherwise.

SOURCE Perrigo Company

For further information: Teva, IR Contacts: Kevin C. Mannix, United States, (215) 591-8912, Ran Meir, United States, (215) 591-3033, Tomer Amitai, Israel, 972 (3) 926-7656, PR Contacts: Iris Beck Codner, Israel, 972 (3) 926-7246, Denise Bradley, United States, (215) 591-8974, Perrigo, IR Contacts: Arthur J. Shannon, (269) 686-1709, ajshannon@perrigo.com, Bradley Joseph, (269) 686-3373, bradley.joseph@perrigo.com

<https://investor.perrigo.com/2013-08-12-Teva-Pharmaceuticals-And-Perrigo-Company-Announce-The-U-S-Launch-Of-Generic-Temozolomide>