

June 20, 2016

Attorney General Rejects VEB's Request Regarding Mylan Shareholder Approval of Meda Transaction

HERTFORDSHIRE, England, and PITTSBURGH, June 20, 2016 /PRNewswire/ -- Mylan N.V. (NASDAQ, TASE: MYL) today confirmed that the Attorney General of the Netherlands has rejected the request of the Dutch group VEB to initiate enquiry proceedings at the Enterprise Chamber of the Amsterdam Court of Appeal in respect of VEB's opinion that Mylan's shareholders should approve its recently announced public offer for all shares in Meda Aktiebolag (publ.). Mylan has consistently communicated to the VEB and to its shareholders that under applicable law such shareholder approval is not required.



Mylan N.V. launched its recommended public offer to the shareholders of Meda on June 17, 2016 to tender all their shares in Meda to Mylan.

Mylan Executive Chairman Robert J. Coury commented, "At Mylan, we are passionate about growing our company, while at the same time we have always and will always be fully committed to complying with all applicable laws and regulations. The Attorney General's decision supports our position that our acquisition of Meda is not subject to approval by Mylan shareholders. The acquisition of Meda will allow us to accelerate and deliver on the clear and compelling vision and long-term strategy we have continuously communicated to our shareholders, and once again deliver a transaction that will create significant value."

Forward-Looking Statements

This communication contains "forward-looking statements." Such forward-looking statements may include, without limitation, statements about the proposed acquisition of Meda by Mylan (the "Meda Transaction"), the Offer, the benefits and synergies of the Meda Transaction, future opportunities for Mylan, Meda, or the combined company and products and any other statements regarding Mylan's, Meda's or the combined company's future operations, anticipated business levels, future earnings, planned activities, anticipated growth, market opportunities, strategies, competition, and other expectations and targets for future periods. These may often be identified by the use of words such as "will," "may," "could," "should," "would," "project," "believe," "anticipate," "expect," "plan," "estimate," "forecast," "potential," "intend," "continue," "target" and variations of these words or comparable words. Because forward-looking statements inherently involve risks and uncertainties, actual future results may differ materially from those expressed or implied by such forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to: uncertainties related to the Meda Transaction, including as to the timing of the Meda Transaction, uncertainties as to whether Mylan will be able to complete the Meda Transaction, the possibility that competing offers will be made, the possibility that certain conditions to the completion of the Offer will not be satisfied, and the possibility that Mylan will be unable to obtain regulatory approvals for the Meda Transaction or be required, as a condition to obtaining regulatory approvals, to accept conditions that could reduce the anticipated benefits of the Meda Transaction; the ability to meet expectations regarding the accounting and tax treatments of Mylan's acquisition (the "EPD Transaction") of Mylan Inc. and Abbott Laboratories' non-U.S. developed markets specialty and branded generics business (the "EPD Business") and the Meda Transaction; changes in relevant tax and other laws, including but not limited to changes in the U.S. tax code and healthcare and pharmaceutical laws and regulations in the U.S. and abroad; the integration of the EPD Business and Meda being more difficult, time-consuming, or costly than expected; operating costs, customer loss, and business disruption (including, without limitation, difficulties in maintaining relationships with employees, customers, clients, or suppliers) being greater than expected following the EPD Transaction and the Meda Transaction; the retention of certain key employees of the EPD Business and Meda being difficult; the possibility that Mylan may be unable to achieve expected synergies and operating efficiencies in connection with the EPD Transaction and the Meda Transaction within the expected time-frames or at all and to successfully integrate the EPD Business and Meda; expected or targeted future financial and operating performance and results; the capacity to bring new products to market, including but not limited to where Mylan uses its business judgment and decides to manufacture, market, and/or sell products, directly or through third parties, notwithstanding the fact that allegations of patent infringement (s) have not been finally resolved by the courts (i.e., an "at-risk launch"); any regulatory, legal, or other impediments to Mylan's ability to bring new products to market; success of clinical trials and Mylan's ability to execute on new product opportunities; any changes in or difficulties with our inventory of, and our ability to manufacture and distribute, the EpiPen® Auto-Injector to meet anticipated demand; the scope, timing, and outcome of any ongoing legal proceedings and the impact of any such proceedings on financial condition, results of operations, and/or cash flows; the ability to protect intellectual property and preserve intellectual property rights; the effect of any changes in customer and supplier relationships and customer purchasing patterns; the ability to attract and retain key personnel; changes in third-party relationships; the impact of competition; changes in the economic and financial conditions of the businesses of Mylan, Meda or the combined company; the inherent challenges, risks, and costs in identifying, acquiring, and integrating complementary or strategic acquisitions of other companies, products or assets and in achieving anticipated synergies; uncertainties and matters beyond the control of management; and inherent uncertainties involved in the estimates and judgments used in the preparation of financial statements, and the providing of estimates of financial measures, in accordance with accounting principles generally accepted in the United States and related standards or on an adjusted basis. For more detailed information on the risks and uncertainties associated with Mylan's business activities, see the risks described in Mylan's Annual Report on Form 10-K for the year ended December 31, 2015, as amended, its Quarterly Report on Form 10-Q for the three months ended March 31, 2016 and its other filings with the SEC. These risks and uncertainties also include those risks and uncertainties that are discussed in the Offer Document that was published on June 16, 2016, the Registration Statement which was declared effective on June 16, 2016 and the EU Prospectus that was published on June 16, 2016. You can access Mylan's filings with the SEC through the SEC website at www.sec.gov, and Mylan strongly encourages you to do so. Mylan undertakes no obligation to update any statements herein for revisions or changes after the date of this communication, except as required by law.

About Mylan

Mylan is a global pharmaceutical company committed to setting new standards in healthcare. Working together around the world to provide 7 billion people access to high quality medicine, we innovate to satisfy unmet needs; make reliability and service excellence a habit; do what's right, not what's easy; and impact the future through passionate global leadership. We offer a growing portfolio of more than 1,400 generic and branded pharmaceuticals, including antiretroviral therapies on which approximately 50% of people being treated for HIV/AIDS in the developing world depend. We market our products in approximately 165 countries and territories. Our global R&D and manufacturing platform includes more than 50 facilities, and we are one of the world's largest producers of active pharmaceutical ingredients. Every member of our more than 35,000-strong workforce is dedicated to creating better health for a better world, one person at a time. Learn more at mylan.com.

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To view the original version on PR Newswire, visit: http://www.prnewswire.com/news-releases/attorney-general-rejects-vebs-request-regarding-mylan-shareholder-approval-of-meda-transaction-300287035.html

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