



November 28, 2016

Mylan Signs Sub-license Agreement with the Medicines Patent Pool to Increase Access to Hepatitis C Treatment in Developing Countries

HERTFORDSHIRE, England and BENGALURU, India, Nov. 28, 2016 /PRNewswire/ -- Mylan N.V. (NASDAQ, TASE: MYL) today announced that the company has signed an agreement with the Medicines Patent Pool (MPP) to expand access to chronic hepatitis C medicines in developing countries. The agreement licenses Mylan to produce and market a generic version of Bristol-Myers Squibb's DAKLINZA™ (daclatasvir) Tablets, 30 mg and 60 mg, for distribution in 112 low and middle income countries.



Daclatasvir Tablets, 30 mg and 60 mg, are indicated for use with sofosbuvir, with or without ribavirin, for the treatment of patients with chronic hepatitis C virus (HCV) genotype 1 or genotype 3 infection in the U.S. and genotype 1, 3 and 4 in Europe. The license allows Mylan to develop fixed-dose combinations that offer the potential to treat all of the six major genotypes of HCV. Earlier this year, the World Health Organization added several new hepatitis C treatments, including daclatasvir, to its essential medicines list, highlighting the urgent need to promote equitable access to innovative medicines¹.

Commenting on today's announcement, Mylan President Rajiv Malik said, "We're committed at Mylan to reducing the burden of hepatitis C on communities around the world by providing access to high quality medicines that treat the disease. We are pleased to work together with the MPP and Bristol-Myers Squibb to help make daclatasvir available to low and middle income countries at affordable prices."

Globally, 130 to 150 million people have chronic hepatitis C infection² and the vast majority live in low and middle income countries³.

Forward-Looking Statements

This press release includes statements that constitute "forward-looking statements," including with regard to future plans and product offerings. These statements are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Because such 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: any changes in or difficulties with Mylan's or its partners' ability to develop, manufacture, and commercialize products; any regulatory, legal, or other impediments to Mylan's or its partners' ability to bring products to market; Mylan's and its partners' ability to protect intellectual property and preserve intellectual property rights; the effect of any changes in Mylan's or its partners' customer and supplier relationships and customer purchasing patterns; other changes in third-party relationships; the impact of competition; changes in the economic and financial conditions of the businesses of Mylan or its partners; the scope, timing, and outcome of any ongoing legal proceedings and the impact of any such proceedings on Mylan's or its partners' business; actions and decisions of healthcare and pharmaceutical regulators, and changes in healthcare and pharmaceutical laws and regulations, in the United States and abroad; risks associated with international operations; other uncertainties and matters beyond the control of management; and the other risks detailed in Mylan's filings with the Securities and Exchange Commission. Mylan undertakes no obligation to update these statements for revisions or changes after the date of this release.

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 2,700 generic and branded pharmaceuticals, including antiretroviral therapies on which nearly 50% of people being treated for HIV/AIDS worldwide depend. We market our products in more than 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 40,000 strong workforce is dedicated to creating better health for a better world, one person at a time. Learn more at mylan.com

1. <http://www.who.int/mediacentre/news/releases/2015/new-essential-medicines-list/en/>
2. <http://www.who.int/mediacentre/factsheets/fs164/en/>
3. Global epidemiology and genotype distribution of the hepatitis C virus infection, Gower, Erin et al., Journal of Hepatology , Volume 61 , Issue 1 , S45 - S57

Logo - <http://photos.prnewswire.com/prnh/20140423/77793>

To view the original version on PR Newswire, visit:<http://www.prnewswire.com/news-releases/mylan-signs-sub-license-agreement-with-the-medicines-patent-pool-to-increase-access-to-hepatitis-c-treatment-in-developing-countries-300368162.html>

SOURCE Mylan N.V.

News Provided by Acquire Media