November 5, 2014

ADMA Biologics Approved for Trading on NASDAQ

RAMSEY, N.J., Nov. 5, 2014 (GLOBE NEWSWIRE) -- ADMA Biologics, Inc. (OTCQB:ADMA), a late-stage biopharmaceutical company that develops, manufactures, and intends to market specialty plasma-based biologics for the treatment and prevention of certain infectious diseases, today announced that the NASDAQ stock market has approved ADMA's common stock for listing on the NASDAQ Capital Market and is expected to begin trading on November 10, 2014 under the symbol "ADMA."

"We are very excited to announce the uplisting of our common stock to NASDAQ. This approval from NASDAQ is a significant milestone for ADMA, as we believe trading on NASDAQ will enhance our visibility as a late-stage biotech company, along with providing increased liquidity for our shareholders," stated Brian Lenz, Vice President and Chief Financial Officer of ADMA Biologics, Inc. "The uplist comes at an opportune time, as we expect to provide top-line, preliminary results from our Phase III clinical study of RI-002 by the end of this year," stated Adam Grossman, President and Chief Executive Officer.

The Company's common stock will continue to trade on the over-the-counter market, under its existing symbol "ADMA," until the market close on November 7, 2014.

About ADMA Biologics, Inc.

ADMA is a late stage biopharmaceutical company that develops, manufactures, and intends to market specialty plasma-based biologics for the treatment and prevention of certain infectious diseases. ADMA's mission is to develop and commercialize plasma-derived, human immune globulins targeted to niche patient populations for the treatment and prevention of certain infectious diseases. The target patient populations include immune-compromised individuals who suffer from an underlying immune deficiency disease or who may be immune-compromised for medical reasons. ADMA also operates ADMA BioCenters a source plasma collection operation which provides ADMA with a portion of its blood plasma for the manufacture of RI-002 as well as generates revenues from the sale of source plasma to third parties. For more information, please visit the Company's website at www.admabiologics.com.

About ADMA's lead product candidate RI-002

ADMA's lead product candidate, RI-002 is a specialty plasma-derived, polyclonal, Intravenous Immune Globulin, or IGIV, derived from human plasma containing naturally occurring polyclonal antibodies (e.g., Streptococcus pneumoniae, H. influenza type B, Cytomegalovirus (CMV), measles, tetanus, etc.) as well as standardized, high levels of antibodies targeted to respiratory syncytial virus (RSV). ADMA is pursuing an indication for the use of this specialty IGIV product for treatment of patients diagnosed with primary immune deficiency diseases, or PIDD. Polyclonal antibodies are the primary component of IGIV products. Polyclonal antibodies are proteins that are used by the body's immune system to neutralize microbes, such as bacteria and viruses. The polyclonal antibodies that are present in RI-002 are expected to prevent infections in immune-compromised patients. The product is currently being evaluated in a Phase III trial in the United States.

Cautionary Statement Regarding Forward-Looking Information

This press release contains "forward looking statements." Forward-looking statements include, without limitation, any statement that may predict, forecast, indicate, or imply future results, performance or achievements, and may contain the words "estimate," "project," "intend," "forecast," "target," "anticipate," "plan," "planning," "expect," "believe," "will," "will likely," "should," "could," "would," "may" or, in each case, their negative, or words or expressions of similar meaning. These forward-looking statements include, but are not limited to, statements concerning the timing, progress and results of the clinical development, the availability of preliminary data, the reporting of data, regulatory processes, potential clinical trial initiations, potential investigational new product applications, biologics license applications, expansion plans, the achievement of clinical and regulatory milestones, build out, opening and regulatory approval of plasma facilities, commercialization efforts of the Company's product candidate(s), trends relating to demand for source plasma and our ability to sustain the listing of our common stock on the NASDAQ Capital Market. Forward-looking statements are subject to many risks and uncertainties that could cause our actual results and the timing of certain events to differ materially from any future results expressed or implied by the forward-looking statements, including, but not limited to, the risks listed under the heading "Risk Factors" in our Annual Report on Form 10-K for the year ended December 31, 2013, as filed with the U.S. Securities and Exchange Commission on March 28, 2014 and our other filings with the U.S. Securities and Exchange Commission including, among other things, risks as to whether any preliminary data will, if and when available, be encouraging, positive or will otherwise lead to an effective or approved product, whether we will be able to demonstrate efficacy or gain necessary approvals to market and commercialize any product, whether we will meet any of our clinical or regulatory milestones, open any new facilities, changes in supply and demand for source plasma, whether we will be able to sustain the listing of our common stock on the NASDAQ Capital Market and whether we will meet any timing targets expressed by the Company. Therefore, current and prospective security holders are cautioned that there also can be no assurance that the forward-looking statements included in this press release will prove to be accurate. In light of the significant uncertainties inherent to the forward-looking statements included herein, the inclusion of

such information should not be regarded as a representation or warranty by ADMA or any other person that the objectives and plans of ADMA will be achieved in any specified time frame, if at all. Except to the extent required by applicable laws or rules, ADMA does not undertake any obligation to update any forward looking statements or to announce revisions to any of the forward-looking statements.

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Source: ADMA Biologics, Inc.