

IMMUNE CEO ANNOUNCES NEW OPERATIONAL LEADERSHIP TEAM Elliott Goldstein, MD, Joins as Chief Medical Officer; Eugene Williams Joins as Chief Operating Officer

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NEW YORK and HERZLIYA, Israel, April 2, 2014 -- Immune Pharmaceuticals Inc. (OTCQX IMNP and NASDAQ OMX First North Premier: IMNP) announced today the addition of two new senior members to its management team.

Elliot Goldstein, MD, joins as Chief Medical Officer. Dr. Goldstein brings a unique track record in the clinical, regulatory and commercial development of new pharmaceuticals. Dr. Goldstein began his career with Sandoz Pharmaceuticals (now Novartis), a fourteen-year period on drug development in France, Basel, Switzerland Global Headquarters, including as Head of Clinical R&D in the United States. He subsequently held positions as SVP of Strategic Product Development at SmithKline Beecham (now GSK), CEO of British Biotech (Oxford, UK), Chief Operating Officer and Chief Medical Officer of Maxygen, and President and CMO of a startup biotech devoted to development of biosimilar monoclonal antibodies. Dr. Goldstein holds an M.D. from the University Aix-Marseille II, Marseille, France, and a B.Sc. from McGill University, Montreal.

Eugene Williams joins as Chief Operating Officer. Mr. Williams is a former SVP at Genzyme, with senior roles integrating commercialization, drug development, and deal making. He is also an entrepreneur, as the founder and director of Adheris, which became the largest company in the patient adherence area. He was previously a strategy consultant at Bain and Corporate Decisions Inc. (a Bain Spin off, now part of Oliver Wyman), where he was co-Head of Healthcare and spent extensive time on speeding and improving the drug development process and on commercialization strategies. Mr. Williams was most recently the CEO of Dart Therapeutics, an Orphan Disease drug development company. Mr. Williams holds a B.A. from Harvard University and an M.B.A. from Harvard Business School.

"We are very happy to welcome Dr. Elliot Goldstein and Mr. Eugene Williams to our senior management team," said Immune Pharmaceuticals' CEO, Dr. Daniel Teper. "Following on our successful fund raising in March of this year, we are building a strong management team with the experience and vision to capitalize on our leading assets in the field of personalized medicine. Both Elliot and Eugene have a proven track record as innovators, and as senior leaders in very successful pharma and biotech companies. They bring significant experience in the disease areas which we have prioritized for our lead assets - rare diseases, autoimmune diseases, and oncology."

Immune Pharmaceuticals Inc. applies a personalized approach to treatment, developing novel, highly targeted antibody therapeutics to improve the lives of patients with inflammatory diseases and cancer. The Company's lead product candidate, bertilimumab, is in clinical development for moderate to severe ulcerative colitis and Crohn's Disease as well as bullous pemphigoid, an orphan auto-immune dermatological condition. Immune licensed worldwide rights for systemic indications of bertilimumab from iCo Therapeutics (TSX: ICO; OTCQX: ICOTF) in June 2011, while iCo retained rights to all ophthalmic indications. iCo originally licensed the exclusive world-wide rights to bertilimumab in 2006 from MedImmune, the Global Research and Development Arm of AstraZeneca. Immune's pipeline also includes NanomAbs®, antibody nanoparticle conjugates, for the targeted delivery of chemotherapeutics and Amiket™, a Neuropathic Pain drug candidate ready for Phase III. Amiket has received Fast Track designation for chemotherapy induced neuropathic pain and Orphan Drug Designation for Post Herpetic Neuralgia.

For more information, visit Immune's website at www.immunepharmaceuticals.com, the content of which is not a part of this press release.

Erik Penser Bankaktiebolag is engaged as Immune's Certified Adviser on NASDAQ OMX First North Premier.

Forward-Looking Statements

This news release and any oral statements made with respect to the information contained in this news release contain forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. You are urged to consider statements that include the words "may," "will," "would," "could," "should," "believes," "estimates," "projects," "potential," "expects," "plans," "anticipates," "intends," "continues," "forecast," "designed," "goal" or the negative of those words or other comparable words to be uncertain and forward-looking. Such forward-looking statements include statements that express plans, anticipation, intent, contingency, goals, targets, future development and are otherwise not statements of historical fact. These statements are based on our current expectations and are subject to risks and uncertainties that could cause actual results or developments to be materially different from historical results or from any future results expressed or implied by such forward-looking statements. Factors that may cause actual results or developments to differ materially include, but not limited to: the risks associated with the adequacy of our existing cash resources and our ability to continue as a going concern; the risks associated with our ability to continue to meet our obligations under our existing debt agreements; the risk that we will not be able to find a partner to help conduct the Phase III trials for AmiKet™ on attractive terms, a timely basis or at all; the risk that we will not obtain approval to market and commercialize any of our product candidates; the risks associated with dependence upon key personnel; the risks associated with reliance on collaborative partners and others for further clinical trials, development, manufacturing and commercialization of our product candidates; the cost, delays and uncertainties associated with our scientific research, product development, clinical trials and regulatory approval process; our history of operating losses since our inception; the highly competitive nature of our business; risks associated with litigation; risks associated with our ability to protect our intellectual

property; risks associate with our ability to raise additional funds; and our liquidity. These factors and other material risks are more fully discussed in our periodic reports, including our reports on Forms 8-K, 10-Q and 10-K and other filings with the U.S. Securities and Exchange Commission. You are urged to carefully review and consider the disclosures found in our filings which are available at www.sec.gov or at www.immunepharmaceuticals.com. You are cautioned not to place undue reliance on any forward-looking statements, any of which could turn out to be wrong due to inaccurate assumptions, unknown risks or uncertainties or other risk factors. We expressly disclaim any obligation to publicly update any forward-looking statements contained herein, whether as a result of new information, future events or otherwise, except as required by law.

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