NEWS RELEASE

IMMUNOMEDICS EXTENDS PATENT LIFE ON THERAPEUTIC ANTIBODIES

-US Patent Issues Covering Novel Formulation for Subcutaneous Injections-

Morris Plains, NJ, February 26, 2014 --- Immunomedics, Inc. (Nasdaq: IMMU), a biopharmaceutical company primarily focused on the development of monoclonal antibody-based products for the targeted treatment of cancer, autoimmune and other serious diseases, today announced the issuance of U.S. Patent No. 8,658,773, which protects a method of concentrating antibodies for subcutaneous, intramuscular or transdermal administration. The patent covers all of the Company's therapeutic antibodies, and can also be applied to most other antibodies, permitting a concentrated formulation to be given in low volume (3 ml or less) and not requiring a special device. The patent has an expiration date of May 1, 2032. It is intended for use with naked, unconjugated antibodies, particularly where subcutaneous administrations are desired in patients with cancer, autoimmune, neurological, and metabolic diseases.

The Company's President and CEO, Cynthia L. Sullivan, commented: "This is a very important extension of patent coverage for all of our therapeutic antibodies that are desirable for subcutaneous, intramuscular or transdermal administration in very low volumes, thus being a simple, fast, and convenient procedure for use in the doctor's office or even at home." "Importantly, this patent extends the coverage of all of our antibodies to 2032, including the anti-CD20, anti-CD74, and anti-CD22 antibodies that are in clinical studies for both oncology and autoimmune disease indications," she clarified.

About Immunomedics

Immunomedics is a New Jersey-based biopharmaceutical company primarily focused on the development of monoclonal antibody-based products for the targeted treatment of cancer, autoimmune and other serious diseases. We have developed a number of advanced proprietary technologies that allow us to create humanized antibodies that can be used either alone in unlabeled or "naked" form, or conjugated with radioactive isotopes, chemotherapeutics, cytokines or toxins, in each case to create highly targeted agents. Using these technologies, we have built a pipeline of therapeutic product candidates that utilize several different mechanisms of action. Our lead product candidate, epratuzumab, is currently in two Phase III clinical trials in lupus. In oncology, clivatuzumab tetraxetan labeled with a radioisotope is in a Phase III pivotal trial in advanced pancreatic cancer patients. Other solid tumor therapeutics in Phase II clinical development include 2 antibody-drug conjugates, IMMU-132 (anti-TROP-2-SN-38) and IMMU-130 (anti-CEACAM5-SN-38). We also have a majority ownership in IBC Pharmaceuticals, Inc., which is developing a novel DOCK-AND-LOCK™ (DNL™) method with us for making fusion proteins and multifunctional antibodies. DNL™ is being used particularly to make bispecific antibodies targeting cancers and infectious diseases as a T-cell redirecting immunotherapy, as well as bispecific antibodies for next-generation cancer and autoimmune disease therapies. We believe that our portfolio of intellectual property, which includes approximately 245 active patents in the United States and more than 400 foreign patents, protects our product candidates and technologies. Our strength in intellectual property has resulted in the top-4 ranking in the January 2014 Patent Board scorecard in the Biotechnology industry. For additional information

~ more ~

on us, please visit our website at www.immunomedics.com. The information on our website does not, however, form a part of this press release.

This release, in addition to historical information, may contain forward-looking statements made pursuant to the Private Securities Litigation Reform Act of 1995. Such statements, including statements regarding clinical trials (including the funding therefor, outcomes, timing or associated costs), out-licensing arrangements (including the timing and amount of contingent payments), forecasts of future operating results, potential collaborations, and capital raising activities, involve significant risks and uncertainties and actual results could differ materially from those expressed or implied herein. Factors that could cause such differences include, but are not limited to, risks associated with any cash payment that the Company might receive in connection with a sublicense involving a third party and UCB, which is not within the Company's control, new product development (including clinical trials outcome and regulatory requirements/actions), our dependence on UCB for the further development of epratuzumab for non-cancer indications, competitive risks to marketed products and availability of required financing and other sources of funds on acceptable terms, if at all, as well as the risks discussed in the Company's filings with the Securities and Exchange Commission. The Company is not under any obligation, and the Company expressly disclaims any obligation, to update or alter any forward-looking statements, whether as a result of new information, future events or otherwise.

For More Information:
Dr. Chau Cheng
Senior Director, Investor Relations & Grant Management
(973) 605-8200, extension 123
ccheng@immunomedics.com

###