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Immtech Pharmaceuticals to Present at the BIO-Europe 2006 Conference

NEW YORK, November 7, 2006 – Immtech Pharmaceuticals, Inc. (AMEX: IMM) today announced that Daniel M. Schmitt, Vice President, Licensing & Commercial Development is scheduled to provide a corporate update at the upcoming BIO-Europe 2006 12th Annual International Partnering Conference at the Congress Center Sued, Stadthalle in Duesseldorf, Germany on November 8, 2006 at 9:15 am CET.

About Immtech Pharmaceuticals, Inc.

Immtech Pharmaceuticals, Inc. is focused on developing and commercializing drugs to treat infectious diseases, and the Company is expanding its targeted markets by applying its proprietary pharmaceutical platform to treat other disorders. Immtech has advanced clinical programs that include new oral treatments for Pneumocystis pneumonia (PCP), malaria, and trypanosomiasis (African Sleeping Sickness), and a well defined, expanding library of compounds targeting fungal infections, Hepatitis C and other serious diseases. Immtech holds the exclusive worldwide licenses to certain patents, patent applications and technology for products derived from a proprietary pharmaceutical platform. For additional information, please go to <http://www.immtechpharma.com>

“Safe Harbor” Statement under the Private Securities Reform Act of 1995: Statements in this press release regarding Immtech Pharmaceuticals, Inc.’s business which are not historical facts are “forward-looking statements” that involve risks and uncertainties. Actual results could differ materially from these forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to, those discussed under the headings "Management's Discussion and Analysis of Financial Condition and Results of Operations" and "Risk Factors" in the Immtech’s annual report on Form 10-K for the year ended March 31, 2006 and in its other SEC filings and include: (i) Immtech’s ability to develop commercially viable products; (ii) Immtech’s ability to achieve profitability; (iii) Immtech’s ability to retain key personnel; (iv) the ability of Immtech’s scientists and collaborators to discover new compounds; (v) the availability of additional research grants; (vi) Immtech’s ability to obtain regulatory approval of its drug candidate; (vii) the success of Immtech’s clinical trials; (viii) Immtech’s ability to manufacture or to have a third party manufacture its drug candidate at a reasonable cost; (ix) Immtech’s ability to protect its intellectual property; (x) competition and alternative technologies; (xi) Immtech’s ability to obtain reimbursement from third party payers for any product it commercializes; and (xii) potential exposure to significant product liability.