



One North End Avenue \_ Suite 1111 \_ New York \_ New York 10282  
Telephone: (877) 898-8038 (212) 791-2911 Fax: 212-791-2917

MEDIA CONTACT: Bill Berry, Berry & Co.: 212-253-8881

INVESTOR RELATIONS CONTACT: Sean Collins, CCG – 310-477-9800, ext. 202

## **IMMTECH ANNOUNCES POSTING OF ANNUAL SHAREHOLDERS MEETING PRESENTATION**

New York, NY, November 29, 2007 - Immtech Pharmaceuticals, Inc. (AMEX:IMM) announced that the presentation to be made during the annual shareholders meeting will be posted on the Company's website, <http://www.immtechpharma.com>, prior to the meeting.

### **About Immtech Pharmaceuticals, Inc.**

Immtech Pharmaceuticals, Inc. is focused on developing and commercializing drugs to treat infectious diseases. Immtech has advanced clinical programs that include new treatments for Pneumocystis pneumonia (PCP), malaria, and trypanosomiasis (African sleeping sickness), and a well defined, expanding library of compounds targeting Hepatitis C, fungal infections, and bacterial infections. Immtech holds 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, including the future prospects for PCP, 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 Immtech's annual report on Form 10-K for the year ended March 31, 2007 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, including PCP; (vii) the success of Immtech's clinical trials; (viii) dependence upon and contractual relationship with partners; (ix) Immtech's ability to manufacture or to have a third party manufacture its drug candidate at a reasonable cost; (x) Immtech's ability to protect its intellectual property; (xi) competition and alternative technologies; (xii) Immtech's ability to obtain reimbursement from third party payers for any product it commercializes; and (xiii) potential exposure to significant product liability.

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