



## Imprimis Pharmaceuticals to Announce Third Quarter 2018 Financial Results on November 13, 2018

November 6, 2018

SAN DIEGO, Nov. 06, 2018 (GLOBE NEWSWIRE) -- Imprimis Pharmaceuticals, Inc. (NASDAQ: IMMY) today announced it will release third quarter 2018 financial results after the close of trading on Tuesday, November 13, 2018. The company will host a conference call at 4:30 p.m. Eastern Time / 1:30 p.m. Pacific Time on the same day to discuss the financial results and recent business developments.

To participate in the call, please dial (877) 407-8031 for domestic callers or (201) 689-8031 for international callers. To listen to the webcast, please click [here](#) or visit the investor relations section of the Imprimis website by [clicking here](#). A dial in replay of the call will be available until December 13, 2018. To access the replay, dial (877) 481-4010 domestically or (919) 882-2331 internationally and reference Replay ID: 40047. The webcast replay will be available until February 13, 2019.

### About Imprimis Pharmaceuticals

Imprimis Pharmaceuticals, Inc. (NASDAQ: IMMY) is a commercial-stage pharmaceutical company based in San Diego, California. In addition to owning the nation's leading ophthalmology pharmaceutical compounding business, ImprimisRx, the Company holds large equity positions in Eton Pharmaceuticals, Surface Pharmaceuticals and Melt Pharmaceuticals, companies originally founded as subsidiaries of Imprimis. The Company also owns royalty rights in certain 505(b)(2) drug candidates being developed by Eton, Surface and Melt. For more information about Imprimis, please visit the Investor Relations section of the corporate website by [clicking here](#).

### SAFE HARBOR

This press release contains forward-looking statements within the meaning of the U.S. Private Securities Litigation Reform Act of 1995. Any statements in this release that are not historical facts may be considered such "forward-looking statements." Forward-looking statements are based on management's current expectations and are subject to risks and uncertainties which may cause results to differ materially and adversely from the statements contained herein. Some of the potential risks and uncertainties that could cause actual results to differ from those predicted include our ability to make commercially available our compounded formulations and technologies in a timely manner or at all; physician interest in prescribing our formulations; risks related to our compounding pharmacy operations; our ability to enter into other strategic alliances, including arrangements with pharmacies, physicians and healthcare organizations for the development and distribution of our formulations; our ability to obtain intellectual property protection for our assets; our ability to accurately estimate our expenses and cash burn, and raise additional funds when necessary; risks related to research and development activities; the projected size of the potential market for our technologies and formulations; unexpected new data, safety and technical issues; regulatory and market developments impacting compounding pharmacies, outsourcing facilities and the pharmaceutical industry; competition; and market conditions. These and additional risks and uncertainties are more fully described in Imprimis' filings with the Securities and Exchange Commission, including its Annual Report on Form 10-K and its Quarterly Reports on Form 10-Q. Such documents may be read free of charge on the SEC's web site at [www.sec.gov](http://www.sec.gov). Undue reliance should not be placed on forward-looking statements, which speak only as of the date they are made. Except as required by law, Imprimis undertakes no obligation to update any forward-looking statements to reflect new information, events or circumstances after the date they are made, or to reflect the occurrence of unanticipated events.

*No Imprimis compounded formulation is FDA-approved. Other than drugs compounded at a registered outsourcing facility, all Imprimis compounded formulations require a prescription for an individually identified patient consistent with federal and state laws.*

### CONTACTS

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