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Imprimis Pharmaceuticals Sells Pennsylvania Facility and Sinus Assets

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SAN DIEGO, June 29, 2017 /PRNewswire/ -- Imprimis Pharmaceuticals, Inc. (NASDAQ: IMMY), an ophthalmology-focused pharmaceutical company, today announced it has entered into an agreement to sell its Folcroft, Pennsylvania facility and sinus related assets for \$450,000. The closing of this transaction is expected to occur on or around July 17, 2017, subject to customary terms and conditions.

Mark L. Baum, CEO of Imprimis said, "We are committed to focusing on our core growing ophthalmology business. The markets we serve with proprietary products in ophthalmology, including our cataract surgery portfolio, medications used in post-LASIK surgery care, our Simple Drops glaucoma program we recently launched and the upcoming launches we have planned to serve dry eye disease patients and other markets in ophthalmology should allow us to build a profitable and valuable company." Baum continued, "In furtherance of this focus, we entered into an agreement to sell our Folcroft facility and exit the sinus space. We believe this is an important step towards reducing non-strategic overhead expenses and focusing internal resources on strengthening our growing ophthalmology business."

About Imprimis Pharmaceuticals

Imprimis Pharmaceuticals, Inc. (NASDAQ: IMMY) is a pharmaceutical company dedicated to producing and dispensing high quality innovative medications in all 50 states. The company's unique business model increases patient access and affordability to many critical medicines. Headquartered in San Diego, California, Imprimis owns and operates three production and dispensing facilities located in California, New Jersey and Pennsylvania. For more information about Imprimis, please visit the corporate website at www.ImprimisRx.com.

Forward-Looking Statements

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. 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.

Other than drugs compounded at a registered outsourcing facility, all Imprimis compounded formulations may only be prescribed pursuant to a physician prescription for an individually identified patient consistent with federal and state laws.

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