

Immunic, Inc. to Participate in Scientific and Investor Conferences in September

SAN DIEGO, September 3, 2019 – Immunic, Inc. (Nasdaq: IMUX), a clinical-stage biopharmaceutical company focused on developing potentially best-in-class, oral therapies for the treatment of chronic inflammatory and autoimmune diseases, today announced management's participation in the following scientific and investor conferences in September:

- September 11-13: **Congress of the European Committee for Treatment and Research in Multiple Sclerosis (ECTRIMS) 2019**: Andreas Muehler, M.D., Chief Medical Officer of Immunic, will present a poster summarizing the profile of lead oral compound, IMU-838, in phase 2 development for the treatment of relapsing-remitting multiple sclerosis, as compared to the approved DHODH inhibitor, teriflunomide, at the ECTRIMS 2019 Congress in Stockholm, Sweden.
 - Abstract Number: A-1026-0031-00242
 - Title: *The DHODH Inhibitor IMU-838/Vidofludimus Calcium Shows a Superior Compound Profile as Compared to the Approved DHODH Inhibitor, Teriflunomide*
 - Session Title: Poster Session 1
 - Session Date: Wednesday, September 11, 2019
 - Presenting Time: 5:15 pm – 7:15 pm CEST

- September 24: **Ladenburg Thalmann 2019 Healthcare Conference**: Daniel Vitt, Ph.D., Chief Executive Officer and President of Immunic, will present a company overview at the Ladenburg Thalmann 2019 Healthcare Conference in New York on Tuesday, September 24, at 3:30 pm EDT. A live audio webcast of the presentation will be available on the “Events and Presentations” section of Immunic’s website at: ir.immunic-therapeutics.com. An archived replay will be available on the company’s website for a period of 90 days after the conference.

About Immunic, Inc.

Immunic, Inc. (Nasdaq: IMUX) is a clinical-stage biopharmaceutical company developing a pipeline of selective oral immunology therapies aimed at treating chronic inflammatory and autoimmune diseases, including relapsing-remitting multiple sclerosis, ulcerative colitis, Crohn’s disease, and psoriasis. The company is developing three small molecule products: IMU-838 is a selective immune modulator that inhibits the intracellular metabolism of activated immune cells by blocking the enzyme DHODH; IMU-935 is an inverse agonist of ROR γ t; and IMU-856 targets the restoration of the intestinal barrier function. Immunic’s lead development program, IMU-838, is in phase 2 clinical development for relapsing-remitting multiple sclerosis and ulcerative colitis, with an additional phase 2 trial in Crohn’s disease planned for the second half of 2019. An investigator-sponsored proof-of-concept clinical trial for IMU-838 in primary sclerosing cholangitis is ongoing at the Mayo Clinic. For further information, please visit: www.immunic-therapeutics.com.



Cautionary Statement Regarding Forward-Looking Statements

This press release contains “forward-looking statements” that involve substantial risks and uncertainties for purposes of the safe harbor provided by the Private Securities Litigation Reform Act of 1995. All statements, other than statements of historical facts, included in this press release regarding strategy, future operations, future financial position, future revenue, projected expenses, prospects, plans and objectives of management are forward-looking statements. Examples of such statements include, but are not limited to, statements relating to management’s participation in scientific and investor conferences. Immunic may not actually achieve the plans, carry out the intentions or meet the expectations or projections disclosed in the forward-looking statements and you should not place undue reliance on these forward-looking statements. Such statements are based on management’s current expectations and involve risks and uncertainties. Actual results and performance could differ materially from those projected in the forward-looking statements as a result of many factors, including, without limitation, risks and uncertainties associated with the ability to project future cash utilization and reserves needed for contingent future liabilities and business operations, the availability of sufficient resources to meet business objectives and operational requirements, the fact that the results of earlier studies and trials may not be predictive of future clinical trial results, the protection and market exclusivity provided by Immunic’s intellectual property, risks related to the drug development and the regulatory approval process and the impact of competitive products and technological changes. Immunic disclaims any intent or obligation to update these forward-looking statements to reflect events or circumstances that exist after the date on which they were made.

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