

Matinas BioPharma to Present at the 8th Annual Partnership Opportunities in Drug Delivery Conference

BEDMINSTER, N.J., Oct. 16, 2018 (GLOBE NEWSWIRE) -- <u>Matinas BioPharma Holdings</u>, <u>Inc.</u> (NYSE AMER: MTNB), a clinical-stage biopharmaceutical company, announced today that <u>Raphael Mannino</u>, <u>Ph.D.</u>, <u>Chief Scientific Officer</u> of Matinas, will present at the <u>8th Annual Partnership Opportunities In Drug Delivery (PODD) Conference</u> on Wednesday, October 17, 2018 at 2:15 PM in Boston, MA.

Dr. Mannino will give an oral presentation titled, "Enabling the Intracellular Delivery of Life-Changing Medicines," as part of the conference's Drug Delivery Technology Presentations Session: Non-Injectable Technologies: Oral, discussing controlled release, modified-release, colon-targeted, and orally-disintegrating delivery systems, as well as bioavailability & solubility enhancing technologies. As part of his presentation, Dr. Mannino will discuss the Company's differentiated lipid nano-crystal (LNC) platform technology which offers an intracellular drug delivery solution with potential advantages for a range of therapeutics, including demonstrated use with small molecules, nucleic acid polymers, proteins, peptides, and vaccines, as well as the potential for efficient and non-immunogenic delivery of gene editing technologies.

In addition to the presentation, Dr. Mannino will also be available to participate in one-onone meetings with registered attendees at the conference.

About Matinas BioPharma

Matinas BioPharma is a clinical-stage biopharmaceutical company focused on developing innovative medicines using its lipid nano-crystal (LNC) platform delivery technology. The Company's proprietary, disruptive technology utilizes lipid-crystal nano-particle cochleates to nano-encapsulate small molecules, oligonucleotides, vaccines and other medicines potentially making them safer, more tolerable, less toxic and orally bioavailable.

The Company's lead anti-fungal product candidate, MAT2203, positions Matinas BioPharma to become a leader in the safe and effective delivery of anti-infective therapies utilizing its proprietary LNC formulation technology.

In addition, the Company is exploring development and partnership options for MAT9001, a prescription-only omega-3 fatty acid-based composition under development for hypertriglyceridemia, which has shown superiority versus Vascepa[®] (icosapent ethyl) in reducing serum triglycerides, Total- and Non-HDL-Cholesterol, apolipoproteins and PCSK9 levels.

For more information, please visit <u>www.matinasbiopharma.com</u> and connect with the Company on Twitter, LinkedIn and Facebook.

Forward Looking Statements: This release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995, including those relating to the Company's anticipated capital and liquidity needs, strategic focus and the future development of its product candidates, including MAT2203, the anticipated timing of regulatory submissions, the anticipated timing of clinical studies, the anticipated timing of regulatory interactions, the Company's ability to identify and pursue development and partnership opportunities for its products or platform delivery technology on favorable terms, if at all, and the ability to obtain required regulatory approval and other statements that are predictive in nature, that depend upon or refer to future events or conditions. All statements other than statements of historical fact are statements that could be forward-looking statements. Forward-looking statements include words such as "expects," "anticipates," "intends," "plans," "could," "believes," "estimates" and similar expressions. These statements involve known and unknown risks, uncertainties and other factors which may cause actual results to be materially different from any future results expressed or implied by the forwardlooking statements. Forward-looking statements are subject to a number of risks and uncertainties, including, but not limited to, our ability to obtain additional capital to meet our liquidity needs on acceptable terms, or at all, including the additional capital which will be necessary to complete the clinical trials of our product candidates; our ability to successfully complete research and further development and commercialization of our product candidates; the uncertainties inherent in clinical testing; the timing, cost and uncertainty of obtaining regulatory approvals; our ability to maintain and derive benefit from the Qualified Infectious Disease Product (QIDP), Orphan and/or Fast Track designations for MAT2203, which does not change the standards for regulatory approval or guarantee regulatory approval on an expedited basis, or at all; our ability to protect the Company's intellectual property; the loss of any executive officers or key personnel or consultants; competition; changes in the regulatory landscape or the imposition of regulations that affect the Company's products; and the other factors listed under "Risk Factors" in our filings with the SEC, including Forms 10-K, 10-Q and 8-K. Investors are cautioned not to place undue reliance on such forward-looking statements, which speak only as of the date of this release. Except as may be required by law, the Company does not undertake any obligation to release publicly any revisions to such forward-looking statements to reflect events or circumstances after the date hereof or to reflect the occurrence of unanticipated events. Matinas BioPharma's product candidates are all in a development stage and are not available for sale or use.

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