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Abeona Therapeutics to Present at Jefferies 2015 Global Healthcare Conference

NEW YORK, NY -- (Marketwired) -- 11/18/15 --

Abeona Therapeutics, Inc. (NASDAQ: ABEO), a biopharmaceutical company focused on developing and delivering gene therapy and plasma-based products for severe and life-threatening rare diseases, today announced that Tim Miller, Ph.D., President and CEO, will be presenting for the company at the Jefferies Autumn 2015 Global Healthcare Conference in London, UK on Thursday, November 19th, 2015 at 4pm GMT on Track 2 at The Mayfair in London, UK. A live webcast of this presentation or replay the webcast of this presentation by visiting ***http://wsw.com/webcast/jeff92/abeo***.

About Jefferies: Jefferies, the global investment banking firm focused on serving clients for over 50 years, is a leader in providing insight, expertise and execution to investors, companies and governments. The firm provides a full range of investment banking, sales, trading, research and strategy across the spectrum of equities, fixed income and foreign exchange, as well as wealth management, in the Americas, Europe and Asia. Jefferies Group LLC is a wholly-owned subsidiary of Leucadia National Corporation, a diversified holding company.

The Jefferies Global Healthcare Conference in London is the premier international healthcare conference and the largest healthcare-dedicated conference in Europe. This global gathering of leading healthcare executives and institutional, private equity and venture capital investors, will address near and long-term investment opportunities and discuss the drivers of growth in the healthcare sector.

About Abeona: Abeona Therapeutics, Inc. develops and delivers gene therapy and plasma-based products for severe and life-threatening rare diseases. Abeona's lead programs are AB0-101 (AAV9 NAGLU) and ABO-102 (scAAV9 SGSH), adeno-associated virus (AAV)-based gene therapies for Sanfilippo syndrome (MPS IIIB and IIIA) in collaboration with patient advocate groups, researchers and clinicians. We are also developing ABO-201 (scAAV9 CLN3) gene therapy for juvenile Batten disease (JBD); and ABO-301 (AAV FANCC) for Fanconi anemia (FA) disorder using a novel CRISPR/Cas9-based gene editing approach to gene therapy program for rare blood diseases. In addition, we are also developing rare plasma protein therapies including SDF Alpha™ (alpha-1 protease inhibitor) for inherited COPD using our proprietary SDF™ (Salt Diafiltration) ethanol-free process. For more information, visit www.abeonatherapeutics.com.

This press release contains certain statements that are forward-looking within the meaning of Section 27a of the Securities Act of 1933, as amended, and that involve risks

and uncertainties. These statements include, without limitation, our plans for the use of proceeds of the financing, our plans to begin enrolling patients in clinical trials for the treatment of Sanfilippo syndrome, development and internationalization of other clinical programs, management plans for the Company, the anticipated closing of the transaction, and general business outlook. These statements are subject to numerous risks and uncertainties, including but not limited to continued interest in our rare disease portfolio, our ability to enroll patients in clinical trials, the impact of competition; the ability to develop our products and technologies; the ability to achieve or obtain necessary regulatory approvals; the impact of changes in the financial markets and global economic conditions; and other risks as may be detailed from time to time in the Company's Annual Reports on Form 10-K and other reports filed by the Company with the Securities and Exchange Commission. The Company undertakes no obligations to make any revisions to the forward-looking statements contained in this release or to update them to reflect events or circumstances occurring after the date of this release, whether as a result of new information, future developments or otherwise.

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