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FOR IMMEDIATE RELEASE

LYOPHILIZATION SERVICES OF NEW ENGLAND MANUFACTURES COMMERCIAL STERILE DRUG PRODUCT

Bedford, NH – Lyophilization Services of New England (LSNE), has been notified by one of its clients that their ANDA for a commercial sterile product, listing LSNE as the sole drug product manufacturing site, has recently been approved by the FDA. This product sparked the most recent FDA inspection (PAI) of the 25 Commerce Drive manufacturing site in April of 2015. While this is not the first commercial approval for the 25 Commerce Drive site, it does mark the first commercial aseptic product approved to be manufactured at the site. Additionally, this is the first approved drug which referenced LSNE's TYPE V DMF.

About LSNE Contract Manufacturing

LSNE Contract Manufacturing is a privately held company with three GMP facilities located in New England. LSNE has been providing contract lyophilization services to the pharmaceutical, biotechnology and medical device industries since 1997, specializing in a wide range of services including cycle development, cGMP fill/finish, and lyophilization. Through the thoughtful integration of three processing facilities, qualified staffing, and an extensive manufacturing history, LSNE is strategically positioned to provide uninterrupted manufacturing for clinical through commercial supply for medical device and pharmaceuticals to a multi-national market.

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