MannKind Announces FDA Acknowledgement of Resubmission of New Drug Application for AFREZZA

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VALENCIA, Calif.--(BUSINESS WIRE)--Oct. 30, 2013-- **MannKind Corporation (Nasdaq: MNKD)** today announced that the U.S. Food and Drug Administration (FDA) has acknowledged the resubmission of a New Drug Application (NDA) for AFREZZA® (insulin human [rDNA origin]) Inhalation Powder. The FDA considered the updated NDA to be a complete class 2 response to its Complete Response Letter issued in January 2011 and assigned a user fee goal date of April 15, 2014.

About MannKind Corporation

MannKind Corporation (Nasdaq: MNKD) focuses on the discovery, development and commercialization of therapeutic products for patients with diseases such as diabetes. Its lead product candidate, AFREZZA®, has completed Phase 3 clinical trials. MannKind maintains a website at www.mannkindcorp.com to which MannKind regularly posts copies of its press releases as well as additional information about MannKind. Interested persons can subscribe on the MannKind website to e-mail alerts that are sent automatically when MannKind issues press releases, files its reports with the Securities and Exchange Commission or posts certain other information to the website.

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