



Rocket Pharmaceuticals to Join NASDAQ Biotechnology Index

December 21, 2018

NEW YORK--(BUSINESS WIRE)--Dec. 21, 2018-- [Rocket Pharmaceuticals, Inc.](#) (Nasdaq: RCKT) ("Rocket"), a leading U.S.-based multi-platform gene therapy company, today announces that the Company has been selected for addition to the NASDAQ Biotechnology Index® (NASDAQ: NBI). Rocket's addition to the NBI will become effective prior to market open on Monday, December 24, 2018.

The NASDAQ Biotechnology Index tracks the performance of a set of securities listed on The NASDAQ Stock Market® (NASDAQ®) that are classified as either biotechnology or pharmaceutical according to the Industry Classification Benchmark (ICB). The NASDAQ Biotechnology Index is calculated under a modified capitalization-weighted methodology and ranked on an annual basis. All securities in the NASDAQ Biotechnology Index are listed on the NASDAQ Global Market or the NASDAQ Global Select Market and meet minimum market value and share volume requirements among other criteria.

In addition, the NASDAQ Biotechnology Index is the basis for the iShares NASDAQ Biotechnology Index (SM) Fund. Options based on the iShares NASDAQ Biotechnology Index Fund trade on various exchanges. For more information about the NASDAQ Biotechnology Index visit <https://indexes.nasdaqomx.com>.

About Rocket Pharmaceuticals, Inc.

Rocket Pharmaceuticals, Inc. (NASDAQ:RCKT) ("Rocket") is an emerging, clinical-stage biotechnology company focused on developing first-in-class gene therapy treatment options for rare, devastating diseases. Rocket's multi-platform development approach applies the well-established lentiviral vector (LVV) and adeno-associated viral vector (AAV) gene therapy platforms. Rocket's lead clinical program is a LVV-based gene therapy for the treatment of Fanconi Anemia (FA), a difficult to treat genetic disease that leads to bone marrow failure and potentially cancer. Preclinical studies of additional bone marrow-derived disorders are ongoing and target Pyruvate Kinase Deficiency (PKD), Leukocyte Adhesion Deficiency-I (LAD-I) and Infantile Malignant Osteopetrosis (IMO). Rocket's first AAV-based gene therapy program targets Danon disease, a rare neuromuscular and cardiovascular disease. For more information about Rocket, please visit www.rocketpharma.com.

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