



## Rocket Pharmaceuticals to Participate in the Chardan Virtual 5th Annual Genetic Medicines Conference

September 30, 2021

CRANBURY, N.J.--(BUSINESS WIRE)--Sep. 30, 2021-- [Rocket Pharmaceuticals, Inc.](#) (NASDAQ: RCKT), a clinical-stage company advancing an integrated and sustainable pipeline of genetic therapies for rare childhood disorders, today announced that Gaurav Shah, M.D., chief executive officer, will participate in a fireside chat on Tuesday, October 5, 2021, at 10:30 am ET at the Chardan Virtual 5<sup>th</sup> Annual Genetic Medicines Conference. In addition, Jonathan Schwartz, M.D., chief medical officer, will be part of a panel discussion: “*Genetic Medicines: The Ongoing Emergence of the Heart as a Target Tissue*” on Monday, October 4, at 3:00 pm ET.

The live webcast of the fireside chat will be accessible via Rocket’s website on the [Events](#) page. An archived copy of the webcast will be available on the Rocket website for 30 days after the event.

### About Rocket Pharmaceuticals, Inc.

Rocket Pharmaceuticals, Inc. (NASDAQ: RCKT) (“Rocket”) is advancing an integrated and sustainable pipeline of genetic therapies that correct the root cause of complex and rare childhood disorders. The company’s platform-agnostic approach enables it to design the best therapy for each indication, creating potentially transformative options for patients afflicted with rare genetic diseases. Rocket’s clinical programs using lentiviral vector (LVV)-based gene therapy are for the treatment of Fanconi Anemia (FA), a difficult to treat genetic disease that leads to bone marrow failure and potentially cancer, Leukocyte Adhesion Deficiency-I (LAD-I), a severe pediatric genetic disorder that causes recurrent and life-threatening infections which are frequently fatal, Pyruvate Kinase Deficiency (PKD) a rare, monogenic red blood cell disorder resulting in increased red cell destruction and mild to life-threatening anemia and Infantile Malignant Osteopetrosis (IMO), a bone marrow-derived disorder. Rocket’s first clinical program using adeno-associated virus (AAV)-based gene therapy is for Danon disease, a devastating, pediatric heart failure condition. For more information about Rocket, please visit [www.rocketpharma.com](http://www.rocketpharma.com).

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Source: Rocket Pharmaceuticals, Inc.