



## Rocket Pharmaceuticals Announces Participation at Upcoming Conferences

February 18, 2021

CRANBURY, N.J.--(BUSINESS WIRE)--Feb. 18, 2021-- [Rocket Pharmaceuticals, Inc.](https://www.rocketpharma.com) (NASDAQ: RCKT), a clinical-stage company advancing an integrated and sustainable pipeline of genetic therapies for rare childhood disorders announces participation at the following upcoming conferences:

- **10<sup>th</sup> Annual SVB Leerink Global Healthcare Conference**
  - Gaurav Shah, M.D., Chief Executive Officer of Rocket, is scheduled to participate in a fireside chat on Thursday, February 25, 2021, at 2:20 p.m. Eastern Time.
- **Cowen's 41<sup>st</sup> Annual Health Care Conference**
  - Tuesday, March 2, 2021

A live audio webcast of the presentation will be available on the Investors section of the company's website, [www.rocketpharma.com](http://www.rocketpharma.com). A replay of the presentation will be archived on the Rocket website following the conference.

### 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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Claudine Prowse, Ph.D.  
SVP, Strategy & Corporate Development  
[investors@rocketpharma.com](mailto:investors@rocketpharma.com)

Source: Rocket Pharmaceuticals, Inc.