



Stoke Therapeutics to Present at Upcoming Investor Conferences in June

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BEDFORD, Mass.--(BUSINESS WIRE)--May 28, 2026-- [Stoke Therapeutics, Inc.](https://www.stoketherapeutics.com) (Nasdaq: STOK) is a biotechnology company dedicated to restoring protein expression by harnessing the body's potential with RNA medicine and has a lead investigational medicine, zorevunersen, in development as a first-in-class potential disease-modifying treatment for Dravet syndrome. Today, the Company announced that Chief Executive Officer Ian F. Smith and Chief Patient Officer Jason Hoitt will present at the following upcoming investor conferences:

2026 Jefferies Global Healthcare Conference

Date: Thursday, June 4, 2026

Time: 9:20 a.m. ET

Location: New York, NY

Goldman Sachs 47th Annual Global Healthcare Conference 2026

Date: Monday, June 8, 2026

Time: 10:40 a.m. ET

Location: Miami, FL

Live webcasts of each event, along with archived replays, will be available in the Investors & News section of Stoke's website at <https://investor.stoketherapeutics.com/>.

About Stoke Therapeutics

Stoke Therapeutics (Nasdaq: STOK), is a biotechnology company dedicated to restoring protein expression by harnessing the body's potential with RNA medicine. Using Stoke's proprietary TANGO (Targeted Augmentation of Nuclear Gene Output) approach, Stoke is developing antisense oligonucleotides (ASOs) to selectively restore naturally-occurring protein levels. Stoke's first medicine in development, zorevunersen, has demonstrated the potential for disease modification in patients with Dravet syndrome and is currently being evaluated in a Phase 3 study. Stoke's initial focus are diseases of the central nervous system and the eye that are caused by a loss of ~50% of normal protein levels (haploinsufficiency). Proof of concept has been demonstrated in other organs, tissues, and systems, supporting broad potential for Stoke's proprietary approach. Stoke is headquartered in Bedford, Massachusetts. For more information, visit <https://www.stoketherapeutics.com/>.

About Zorevunersen

Zorevunersen is an investigational antisense oligonucleotide that is designed to treat the underlying cause of Dravet syndrome by increasing functional NaV1.1 protein production in brain cells from the unaffected (wild-type) copy of the *SCN1A* gene. This highly differentiated mechanism of action aims to reduce seizure frequency beyond what has been achieved with anti-seizure medicines and to improve neurodevelopment, cognition and behavior. Zorevunersen has demonstrated the potential for disease modification and has been granted orphan drug designation by the FDA and the EMA. The FDA has also granted zorevunersen rare pediatric disease designation and Breakthrough Therapy Designation for the treatment of Dravet syndrome with a confirmed mutation not associated with gain-of-function, in the *SCN1A* gene. Stoke has a strategic collaboration with Biogen (Nasdaq: BIIB) to develop and commercialize zorevunersen for Dravet syndrome. Under the collaboration, Stoke retains exclusive rights for zorevunersen in the United States, Canada, and Mexico; Biogen receives exclusive rest of world commercialization rights. Zorevunersen is currently in clinical development, and its safety and efficacy have not been evaluated by any regulatory authority.

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Stoke Media & Investor Contacts:

Susan Willson

Vice President, Corporate Communications

swillson@stoketherapeutics.com

415-509-8202

Doug Snow

Director, Communications & Investor Relations

IR@stoketherapeutics.com

508-642-6485

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