vTv Therapeutics Announces Inducement Grants under Nasdaq Listing Rule 5635(c)(4)

June 2, 2026

HIGH POINT, N.C., June 02, 2026 (GLOBE NEWSWIRE) -- vTv Therapeutics Inc. (Nasdaq: VTVT), a late-stage biopharmaceutical company focused on the development of *cadisegliatin*, a novel, potential first-in-class oral adjunctive therapy to insulin being investigated for the treatment of type 1 diabetes (T1D), today announced that it granted 2,500 stock options to purchase shares of common stock to *a* non-executive employee as a material inducement to employment in accordance with Nasdaq Listing Rule 5635(c)(4).

The stock options that were granted have an exercise price of $34.30 per share, which is equal to the closing price of the Company's common stock on June 1, 2026. Each option will vest over a 4-year period, with 25% of the shares underlying the employee's option vesting on the one-year anniversary of the applicable vesting commencement date and the remaining shares thereafter vesting quarterly over the following 36 months, subject to the employee's continued employment with vTv on such vesting dates. The options have a term of 10 years and are subject to the terms and conditions of the 2026 Inducement Plan and the stock option agreement covering the grant.

About vTv Therapeutics
vTv Therapeutics is a late-stage biopharmaceutical company focused on developing oral, small molecule drug candidates intended to help treat people living with diabetes and other chronic diseases. vTv's clinical pipeline is led by *cadisegliatin*, currently in a US Phase 3 trial, a potential first-in-class oral glucokinase activator being investigated for the treatment of type 1 diabetes. vTv and its development partners are investigating multiple molecules across different indications for chronic diseases. Learn more at vtvtherapeutics.com or follow the company on LinkedIn or X.

About Cadisegliatin
Cadisegliatin (TTP399) is a novel, oral small molecule, liver-selective glucokinase activator being investigated in the U.S. as a potential first-in-class oral adjunctive treatment for type 1 diabetes (T1D). In non-clinical studies, cadisegliatin acted selectively on the liver and increased glucokinase activity independently of insulin. These findings support clinical investigation of whether cadisegliatin can improve glycemic control through hepatic glucose uptake and glycogen storage. Cadisegliatin has been granted Breakthrough Therapy designation by the U.S. Food and Drug Administration (FDA).

Cadisegliatin is under investigation, and the safety and efficacy have not been established. There is no guarantee that this product will receive health authority approval or become commercially available for the use being investigated.

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