

Cadrenal Therapeutics to Participate in the Lytham Partners Fall 2025 Investor Conference on September 30, 2025

PONTE VEDRA, Fla., Sept. 29, 2025 (GLOBE NEWSWIRE) — Cadrenal Therapeutics, Inc. (Nasdaq: CVKD), a biopharmaceutical company developing transformative therapeutics to overcome the gaps in anticoagulation therapy, today announced that it will participate in a webcast presentation and host one-on-one meetings with investors at the Lytham Partners Fall 2025 Investor Conference, taking place virtually on Tuesday, September 30, 2025.

Company Webcast

The webcast presentation will take place at 3:30 p.m. ET on Tuesday, September 30, 2025. The webcast can be accessed by visiting the conference home page at <https://lythampartners.com/fall2025/> or directly at <https://app.webinar.net/QG1g6kJWEM0>. The webcast will also be available for replay following the event.

1×1 Meetings

Management will be participating in virtual one-on-one meetings throughout the event. To arrange a meeting with management, please contact Lytham Partners at 1x1@lythampartners.com or register for the event at <https://lythampartners.com/fall2025invreg/>.

About Cadrenal Therapeutics, Inc.

Cadrenal Therapeutics, Inc. is a biopharmaceutical company developing transformative therapeutics to overcome the gaps in anticoagulation therapy. Cadrenal's lead investigational product is tecarfarin, a novel oral Vitamin K antagonist anticoagulant that is designed to address unmet needs in anticoagulation therapy. Tecarfarin is a reversible anticoagulant (blood thinner) designed to prevent heart attacks, strokes, and deaths due to blood clots in patients requiring chronic anticoagulation. Although warfarin is widely used, extensive clinical and real-world data have shown it can have significant, serious side effects. With tecarfarin, Cadrenal aims to reduce the clinical complexities of managing Vitamin K antagonists, particularly where direct-acting oral anticoagulants (DOACs) remain inadequate or unproven.

Tecarfarin received Orphan Drug Designation (ODD) and fast-track designation for the prevention of systemic thromboembolism (blood clots) of cardiac origin in patients with end-stage kidney disease and atrial fibrillation (ESKD+AFib). The Company also received ODD for the prevention of thromboembolism and thrombosis in patients with implanted mechanical circulatory support devices, including Left Ventricular Assist Devices (LVADs).

For more information, visit <https://www.cadrenal.com/> and connect with the Company on LinkedIn.

For more information, please contact:

Cadrenal Therapeutics:

Matthew Szot, CFO

press@cadrenal.com

Investors:

Lytham Partners, LLC

Robert Blum, Managing Partner

602-889-9700

CVKD@lythampartners.com

