May 7, 2025

Cellectar Biosciences to Report First Quarter Financial Results and Host a Conference Call on Tuesday, May 13, 2025

FLORHAM PARK, N.J., May 07, 2025 (GLOBE NEWSWIRE) -- Cellectar Biosciences, Inc. (NASDAQ: CLRB), a late-stage clinical biopharmaceutical company focused on the discovery, development and commercialization of drugs for the treatment of cancer, today announced that the Company will report financial results for the first quarter ended March 31, 2025, and provide a corporate update on May 13, 2025, at 8:30 a.m. Eastern Time.

Conference Call & Webcast Details:	
Date:	Tuesday, May 13, 2025
Time:	8:30 am Eastern Time
Toll Free:	1-800-717-1738
Conference ID:	53983
Webcast:	Click HERE

A replay of the corporate presentation will be available on the Events section of the Company's Investor Relations website.

About Cellectar Biosciences, Inc.
Cellectar Biosciences is a late-stage clinical biopharmaceutical company focused on the discovery and development of proprietary drugs for the treatment of cancer, independently and through research and development collaborations. The company's core objective is to leverage its proprietary Phospholipid Drug Conjugate™ (PDC) delivery platform to develop the next-generation of cancer cell-targeting treatments, delivering improved efficacy and better safety as a result of fewer off-target effects.

The company's product pipeline includes its lead assets: iopofosine I 131, a PDC designed to provide targeted delivery of iodine-131 (radioisotope); CLR 225, an actinium-225 based program being targeted to several solid tumors with significant unmet need, such as pancreatic cancer; and CLR 125, an iodine-125 Auger-emitting program targeted in solid tumors, such as triple negative breast, lung and colorectal, as well as proprietary preclinical PDC chemotherapeutic programs and multiple partnered PDC assets.

In addition, iopofosine I 131 has been studied in Phase 2b trials for relapsed or refractory multiple myeloma (MM) and central nervous system (CNS) lymphoma, and the CLOVER-2 Phase 1b study, targeting pediatric patients with high-grade gliomas, for which Cellectar is eligible to receive a Pediatric Review Voucher from the FDA. The FDA has also granted iopofosine I 131 six Orphan Drug, four Rare Pediatric Drug, and two Fast Track designations for various cancer indications. The EMA (European Medicines Agency) has granted iopofosine I 131 two Orphan Drug designations and PRIME designation for WM.

For more information, please visit www.cellectar.com or join the conversation by liking and following us on the company's social media channels: X, LinkedIn, and Facebook.

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Source: Cellectar Biosciences, Inc.