



## Celcuity Inc. Schedules Release of Fourth Quarter and Full Year 2024 Financial Results and Webcast/Conference Call

March 24, 2025

MINNEAPOLIS, March 24, 2025 (GLOBE NEWSWIRE) -- Celcuity Inc. (Nasdaq: CELC), a clinical-stage biotechnology company pursuing development of targeted therapies for oncology, today announced that it will release its financial results for the fourth quarter and full year 2024 after the market closes on Monday, March 31, 2025. Management will host a webcast/teleconference the same day at 4:30 p.m. Eastern Time to discuss the results and provide a corporate update.

### Webcast and Conference Call Information

To participate in the teleconference, domestic callers should dial 1-800-717-1738 and international callers should dial 1-646-307-1865. A live webcast presentation can also be accessed using this weblink: [https://viaid.webcasts.com/starthere.jsp?ei=1704367&tp\\_key=f99d4186f3](https://viaid.webcasts.com/starthere.jsp?ei=1704367&tp_key=f99d4186f3). A replay of the webcast will be available on the Celcuity website following the live event.

### About Celcuity

Celcuity is a clinical-stage biotechnology company focused on development of targeted therapies for treatment of multiple solid tumor indications. The company's lead therapeutic candidate is gedatolisib, a potent, pan-PI3K and mTOR inhibitor. Its mechanism of action and pharmacokinetic properties are differentiated from other currently approved and investigational therapies that target PI3K or mTOR alone or together. A Phase 3 clinical trial, VIKTORIA-1, evaluating gedatolisib in combination with fulvestrant with or without palbociclib in patients with HR+/HER2- advanced breast cancer is currently enrolling patients. More detailed information about the VIKTORIA-1 study can be found at [ClinicalTrials.gov](https://clinicaltrials.gov). A Phase 1b/2 clinical trial, CELC-G-201, evaluating gedatolisib in combination with darolutamide in patients with metastatic castration resistant prostate cancer, is ongoing. A Phase 3 clinical trial, VIKTORIA-2, evaluating gedatolisib plus a CDK4/6 inhibitor and fulvestrant as first-line treatment for patients with HR+/HER2- advanced breast cancer is expected to begin enrolling patients in the second quarter of 2025. Celcuity is headquartered in Minneapolis. Further information about Celcuity can be found at [www.celcuity.com](http://www.celcuity.com). Follow us on [LinkedIn](#) and [Twitter](#).

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### Contacts:

Celcuity Inc.  
Brian Sullivan, [bsullivan@celcuity.com](mailto:bsullivan@celcuity.com)  
Vicky Hahne, [vhahne@celcuity.com](mailto:vhahne@celcuity.com)  
(763) 392-0123

ICR Healthcare  
Patti Bank, [patti.bank@icrhealthcare.com](mailto:patti.bank@icrhealthcare.com)  
(415) 513-1284



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