



Celcuity To Participate in Upcoming Investor Conferences

February 25, 2026

MINNEAPOLIS, Feb. 25, 2026 (GLOBE NEWSWIRE) -- Celcuity Inc. (Nasdaq: CELC), a clinical-stage biotechnology company pursuing development of targeted therapies for oncology, today announced that Brian Sullivan, Chief Executive Officer, and Co-founder of Celcuity, will present and be available for one-on-one investor meetings at the following investor conferences:

- A fireside chat at the TD Cowen 46th Annual Healthcare Conference at 10:30 a.m. ET on Wednesday, March 4, 2026. A live webcast will be available using this weblink: <https://event.summitcast.com/view/9z5q2VrV6e6rbCqQgDRoHA/NYER3h287L9qyxak8BZiq2>
- A fireside chat at the Leerink Global Healthcare Conference at 2:20 p.m. ET on Tuesday, March 10, 2026. A live webcast will be available using this weblink: <https://event.summitcast.com/view/mT9pocHNDNthc6b89WqVf/VziaefXK3xAkvtByRaxLpT>
- Jefferies Biotech on the Beach Summit; Wednesday, March 11, 2026. Management will host one-on-one investor meetings only.

Alternatively, the live webcasts will be accessible from the Investors section of the company's website at <https://ir.celcuity.com/events-presentations/> with a replay available shortly after the live events.

About Celcuity

Celcuity is a clinical-stage biotechnology company pursuing the development of targeted therapies for the treatment of multiple solid tumor indications. The company's lead therapeutic candidate is gedatolisib, a potent, pan-PI3K and mTORC1/2 inhibitor that comprehensively blockades the PI3K/AKT/mTOR ("PAM") pathway. Its mechanism of action and pharmacokinetic properties are differentiated from other currently approved and investigational therapies that target PI3K α , AKT, or mTORC1 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 ("ABC"), has completed enrollment, and the company has reported detailed results for the *PIK3CA* wild-type cohort. 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- ABC, is ongoing. A Phase 1/2 clinical trial, CELC-G-201, evaluating gedatolisib in combination with darolutamide in patients with metastatic castration resistant prostate cancer, is ongoing. More detailed information about Celcuity's active clinical trials can be found at [ClinicalTrials.gov](https://clinicaltrials.gov). Celcuity is headquartered in Minneapolis. Further information about Celcuity can be found at www.celcuity.com. Follow us on [LinkedIn](#) and [X](#).

Contacts:

Celcuity Inc.
Brian Sullivan, bsullivan@celcuity.com
Vicky Hahne, vhahne@celcuity.com
(763) 392-0123
Jodi Sievers, jsievers@celcuity.com
(415) 494-9924