

CytomX Therapeutics, Inc. Logo

## CytomX Therapeutics to Report First Quarter 2024 Results and Provide an Initial CX-904 Phase 1a Clinical Data Update on May 8, 2024

May 1, 2024 at 8:00 AM EDT

- CX-904 (masked EGFRxCD3 Probody® T-cell engager) preliminary Phase 1a results to be presented from ongoing dose escalation study -

- Management to hold conference call at 5 p.m. EDT / 2 p.m. PDT on May 8<sup>th</sup>-

SOUTH SAN FRANCISCO, Calif., May 01, 2024 (GLOBE NEWSWIRE) -- CytomX Therapeutics, Inc. (Nasdaq: CTMX), a leader in the field of masked, conditionally activated biologic therapeutics, today announced that it will report first quarter 2024 results and provide an initial CX-904 Phase 1a dose escalation update on Wednesday, May 8, 2024, after the close of U.S. markets. Following the announcement, the Company will host a conference call and webcast at 5:00 p.m. ET / 2:00 p.m. PT.

Participants may access the live webcast of the conference call from the Events and Presentations page of CytomX's website at <https://ir.cytomx.com/events-and-presentations>. Participants may register for the conference call [here](#) and are advised to do so at least 10 minutes prior to joining the call. An archived replay of the webcast will be available on the Company's website.

### About CytomX Therapeutics

CytomX is a clinical-stage, oncology-focused biopharmaceutical company focused on developing novel conditionally activated, masked biologics designed to be localized to the tumor microenvironment. By pioneering a novel pipeline of localized biologics, powered by its PROBODY® therapeutic platform, CytomX's vision is to create safer, more effective therapies for the treatment of cancer. CytomX's robust and differentiated pipeline comprises therapeutic candidates across multiple treatment modalities including antibody-drug conjugates ("ADCs"), T-cell engagers, and immune modulators such as cytokines. CytomX's clinical-stage pipeline includes CX-904, CX-2051 and CX-801. CX-904 is a masked, conditionally activated T-cell-engaging bispecific antibody targeting the epidermal growth factor receptor (EGFR) on tumor cells and the CD3 receptor on T cells. CX-904 is partnered with Amgen in a global co-development alliance. CX-2051 is a masked, conditionally activated ADC directed toward epithelial cell adhesion molecule, EpCAM, with potential applicability across multiple EpCAM-expressing epithelial cancers. CX-2051 was discovered in collaboration with Immunogen, now part of AbbVie. CX-801 is a masked interferon alpha-2b PROBODY® cytokine with broad potential applicability in traditionally immuno-oncology sensitive as well as insensitive (cold) tumors. CytomX has established strategic collaborations with multiple leaders in oncology, including Amgen, Astellas, Bristol Myers Squibb, Regeneron and Moderna. For more information about CytomX and how it is working to make conditionally activated treatments the new standard-of-care in the fight against cancer, visit [www.cytomx.com](http://www.cytomx.com) and follow us on [LinkedIn](#) and [X \(formerly Twitter\)](#).

PROBODY is a U.S. registered trademark of CytomX Therapeutics, Inc.

### Company Contact:

Chris Ogden  
SVP, Finance and Accounting  
[cogden@cytomx.com](mailto:cogden@cytomx.com)

### Investor Contact:

Stern Investor Relations  
Stephanie Ascher  
[stephanie.ascher@sternir.com](mailto:stephanie.ascher@sternir.com)

### Media Contact:

Redhouse Communications  
Teri Dahlman  
[teri@redhousecomms.com](mailto:teri@redhousecomms.com)



Source: CytomX Therapeutics Inc.