



## Plus Therapeutics Announces Key Opinion Leader Roundtable on New Clinical Trial Data Being Presented at the 2023 SNO/ASCO CNS Cancer Conference

August 8, 2023

*Trial investigators and company management to discuss latest data from the ReSPECT-LM leptomeningeal metastases clinical trial*

*Webinar scheduled for Friday, August 11, 2023 at 8:00 a.m. ET*

AUSTIN, Texas, Aug. 08, 2023 (GLOBE NEWSWIRE) -- [Plus Therapeutics, Inc.](#) (Nasdaq: [PSTV](#)) (the "Company"), a clinical-stage pharmaceutical company developing targeted radiotherapeutics with advanced platform technologies for central nervous system cancers, today announced a key opinion leader roundtable discussion to be held on Friday, August 11, 2023, 8:00 a.m. – 9:00 a.m. ET to discuss [the latest data from the ReSPECT-LM clinical trial](#) of rhenium ( $^{186}\text{Re}$ ) obisbameda that will be presented at the Society for Neuro Oncology (SNO)/American Society of Clinical Oncology (ASCO) Central Nervous System (CNS) Cancer Conference on August 10, 2023.

The webinar will feature a comprehensive discussion about the ongoing ReSPECT-LM Phase 1/2a dose escalation clinical trial, including key safety, tolerability, dosing, feasibility and efficacy data. Speakers include:

- Andrew J. Brenner, M.D., Ph.D., Professor of Medicine, Neurology, and Neurosurgery at The University of Texas, Health Services Center at San Antonio and principal investigator of the ReSPECT-GBM trial and co-principal investigator of the ReSPECT-LM trial.
- Priya Kumthekar, M.D., Associate Professor of Neurology and Medicine (Hematology and Oncology) at Northwestern University's Feinberg School of Medicine and co-principal investigator of the ReSPECT-LM trial.
- Marc H. Hedrick, M.D., President and Chief Executive Officer of Plus Therapeutics
- Justin Walsh, Ph.D., Senior Healthcare Analyst, Jones Research will moderate the roundtable and in addition will provide an overview of the radiotherapeutic market.

### Webcast Details

A webinar with accompanying slides will be available in the [Events](#) page of the Investor Relations section of the Plus Therapeutics website beginning Friday, August 11, 2023 at 8:00 a.m. ET. Participants may also pre-register any time before the call [here](#). Please access the webinar 15 minutes prior to the start time.

The webinar will be available on the Company's website under the ['For Investor'](#) section. The webcast will be available on the Company's website for 90 days following the live call.

### About Plus Therapeutics

Plus Therapeutics, Inc. is a clinical-stage pharmaceutical company developing targeted radiotherapeutics for difficult-to-treat cancers of the central nervous system with the potential to enhance clinical outcomes for patients. Combining image-guided local beta radiation and targeted drug delivery approaches, the Company is advancing a pipeline of product candidates with lead programs in recurrent glioblastoma (GBM) and leptomeningeal metastases (LM). The Company has built a robust supply chain through strategic partnerships that enable the development, manufacturing and future potential commercialization of its products. Plus Therapeutics is led by an experienced and dedicated leadership team and has operations in key cancer clinical development hubs including Austin and San Antonio, Texas. For more information, visit <https://plustherapeutics.com/> and <https://respect-trials.com>.

### Cautionary Statement Regarding Forward-Looking Statements

This press release contains statements that may be deemed "forward-looking statements" within the meaning of U.S. securities laws. All statements in this press release other than statements of historical fact are forward-looking statements. These forward-looking statements may be identified by future verbs, as well as terms such as "designed to," "will," "can," "potential," "focus," "preparing," "next steps," "possibly," and similar expressions or the negatives thereof. Such statements are based upon certain assumptions and assessments made by management in light of their experience and their perception of historical trends, current conditions, expected future developments and other factors they believe to be appropriate. These statements include, without limitation, statements regarding the following: the potential promise of  $^{186}\text{Re}$  including the ability of  $^{186}\text{Re}$  to safely and effectively deliver radiation directly to the tumor at high doses; expectations as to the Company's future performance including the next steps in developing the Company's current assets; the Company's clinical trials including statements regarding the timing and characteristics of the ReSPECT-GBM and ReSPECT-LM clinical trials; possible negative effects of  $^{186}\text{Re}$ ; the continued evaluation of  $^{186}\text{Re}$  including through evaluations in additional patient cohorts; and the intended functions of the Company's platform and expected benefits from such functions.

The forward-looking statements included in this press release are subject to a number of risks and uncertainties that may cause actual results to differ materially from those discussed in such forward-looking statements. These risks and uncertainties include, but are not limited to: the Company's actual results may differ, including materially, from those anticipated in these forward-looking statements as a result of various factors, including, but not limited to, the following: the early stage of the Company's product candidates and therapies, the results of the Company's research and development activities, including uncertainties relating to the clinical trials of its product candidates and therapies; the Company's liquidity and capital resources and its ability to raise additional cash, the outcome of the Company's partnering/licensing efforts, risks associated with laws or regulatory requirements applicable to it, market conditions, product performance, litigation or potential litigation, and competition within the cancer diagnostics and therapeutics field, among others; and additional risks described under the heading "Risk Factors" in the Company's Securities and Exchange Commission filings, including in the Company's annual and quarterly reports. There may be events in the future that the Company is unable to predict, or over which it has

no control, and its business, financial condition, results of operations and prospects may change in the future. The Company assumes no responsibility to update or revise any forward-looking statements to reflect events, trends or circumstances after the date they are made unless the Company has an obligation under U.S. federal securities laws to do so.

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