Plus Therapeutics to Present at the 2023 SNO/ASCO CNS Cancer Conference

July 27, 2023

AUSTIN, Texas, July 27, 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 that the Company will share three poster presentations at the Society for Neuro Oncology (SNO)/American Society of Clinical Oncology (ASCO) Central Nervous System (CNS) Cancer Conference taking place August 10-12, 2023 in San Francisco, California.

The presentations will describe new data from the ReSPECT-LM and ReSPECT-GBM clinical studies evaluating the Company's lead radiotherapeutic, rhenium (^{186}Re) obisbemeda, for the treatment of leptomeningeal metastases (LM) and recurrent glioblastoma (rGBM), respectively, as well as details around the planned ReSPECT-PBC trial to evaluate rhenium (^{186}Re) obisbemeda for the treatment of recurrent, refractory, or progressive childhood ependymoma and high-grade glioma (HGG).

All presentations will be showcased during the Poster Reception on Thursday, August 10, 2023, beginning at 5:30 p.m. PT/8:30 p.m. ET.

Details of presentations:

Title	[LMAP-21] Preliminary Clinical Data in the Phase 1/2a Dose Escalation Trial of Rhenium (186Re) Obisbemeda (186RNL) in Leptomeningeal Metastases (LM): The ReSPECT-LM Trial
Title	[TIPS-23] Safety and Feasibility Results from a Phase 1/2 Clinical Trial of Rhenium (186Re) Obisbemeda (186RNL) in Recurrent Glioma: The ReSPECT-GBM Trial
Title	[TIPS-22] A two-part, Phase 1 study of Rhenium (186Re) Obisbemeda (186RNL) delivered by convection enhanced delivery (CED) for recurrent, refractory, or progressive childhood ependymoma and high-grade glioma (HGG)

A copy of the presentations will be made available under the Presentations tab of the Investors section of the Company's website following the meeting at https://ir.plustherapeutics.com.

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/.

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}Re including the ability of ^{186}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}Re; the continued evaluation of ^{186}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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