UNITED STATES SECURITIES AND EXCHANGE COMMISSION Washington, D.C. 20549

FORM 8-K

CURRENT REPORT Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934

Date o	f Report (Date of earliest event reported): July	15, 2025
	CNS Pharmaceuticals, Inc. (Exact name of registrant as specified in its charter)	r)
Nevada (State or other jurisdiction of incorporation or organization)	001-39126 (Commission File Number)	82-2318545 (I.R.S. Employer Identification No.)
	2100 West Loop South, Suite 900 Houston, Texas 77027 (Address of principal executive offices) (Zip Code	e)
Registrar	nt's telephone number, including area code: (800	946-9185
(Forme	Not Applicable er Name or Former Address, if Changed Since Last	Report)
k the appropriate box below if the Form provisions (<i>see</i> General Instruction A.2. b		e filing obligation of the registrant under any of th
Written communications pursuant to Ru	le 425 under the Securities Act (17 CFR 230.425)	
Soliciting material pursuant to Rule 14a-	-12 under the Exchange Act (17 CFR 240.14a-12)	
Pre-commencement communications pu	rsuant to Rule 14d-2(b) under the Exchange Act (1	7 CFR 240.14d-2(b))

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Emerging growth company □

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. \Box

Securities registered pursuant to Section 12(b) of the Act:

following

Title of each class	Trading Symbols(s)	Name of each exchange on which registered
Common stock, par value \$0.001 per share	CNSP	The NASDAQ Stock Market LLC

Item 7.01. Regulation FD Disclosure.

On July 15, 2025, CNS Pharmaceuticals, Inc. (the "Company") posted the investor presentation set forth in Exhibit 99.1 on its website.

The information contained in Item 7.01 of this Current Report on Form 8-K is being furnished and shall not be "filed" for the purpose of the Securities Exchange Act of 1934, as amended ("Exchange Act"), nor shall it be incorporated by reference in any filing under the Exchange Act or the Securities Act of 1933, as amended ("Securities Act"), unless specifically identified therein as being incorporated by reference.

Item 9.01. Financial Statements and Exhibits.

No.	Description
99.1	Presentation dated July 2025
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)
	2

Signature

Pursuant to the requirements of the	e Securities Exchange	e Act of 1934,	the registrant	has duly cause	d this report to	be signed on its	s behalf by the
undersigned hereunto duly authorized.							

CNS Pharmaceuticals, Inc.

By: /s/ Chris Downs
Chris Downs
Chief Financial Officer

Dated: July 15, 2025



Forward Looking Statements

This presentation incorporates information from materials filed with the SEC and contains forward-looking statements. All statements contained herein other than statements of historical fact, including statements regarding our future results of operations and financial position, our business strategy and plans, and our objectives for future operations, are forward-looking statements. The words "believe," "may," "will," "estimate," "continue," "anticipate," "intend," "expect," and similar expressions are intended to identify forward looking statements. We have based these forward-looking statements largely on our current expectations and projections about future events and trends that we believe may affect our financial condition, results of operations, business strategy, short-term and long-term business operations and objectives, and financial needs.

These forward-looking statements are subject to a number of risks, uncertainties and assumptions, including those described in the "Risk Factors" section of most recent Form 10-K as updated by any subsequent Form 10-Q filings. It is not possible for our management to predict all risks, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statements we may make. In light of these risks, uncertainties and assumptions, the future events and trends discussed in this presentation may not occur and actual results could differ materially and adversely from those anticipated or implied in the forward looking statements.



Overview

Developing Anti-Cancer Drug Candidates for the Treatment of Primary and Metastatic Brain Cancers

Strong Financial Position

Proven Clinical Development "Engine" with Global Trial Site Network in Place to Accelerate Complex CNS Focused Trials

Advancing Lead Product Candidate, TPI 287 for treatment of Glioblastoma Multiforme (GBM)

- · Late-stage, novel, blood brain-barrier permeable taxane-derivative (abeotaxane)
- Studies in over 350 patients to date, include clinical trials as monotherapy and combination with bevacizumab

Reported Primary Analysis of Berubicin Monotherapy in 2nd line GBM

· Ongoing analysis of outcomes ongoing to determine next steps



Did not a statistically significant difference in overall survival, the primary endpoint



A Focused and Targeted CNS Oncology Pipeline

Highlights
Studied in over 350 patients to date
Plan to engage with regulators to design potential registration study in 2025
• P

A Much Bigger Story Beyond GBM

	Potential Fut	cure Indications	
Primary Brain	High Grade Gliomas	Brain Metastases - Combo	Primary CNS
Tumors	in Pediatrics	with Radiation Therapy	Lymphoma (PCSNL)
15,000	6,000	45,000	1,200
Patients	Patients	Patients	Patients



Established "Engine" to Execute Global CNS Clinical Trials

Key Learnings and Established Network From Berubicin Monotherapy Potentially Pivotal Trial





Successfully Built CNS Trial Network and Enrolled Patients in Record Time, All During a Global Pandemic

E

Proven Clinical Development Infrastructure Optimized for Brain Cancer Drug Development

Relationships

- Deliberate establishment of a global, CNS focused network
- · Commitment to work in this disease
- Deep understanding of the landscape of clinical trials in GBM

Program Development Infrastructure & Efficiencies

- Seamless transition to our next asset
- · Built to last
- · Set up for success



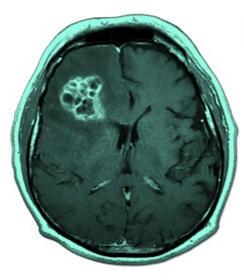


Glioblastoma Multiforme (GBM)

One of the most aggressive, deadly and treatment-resistant cancers that forms in the brain

Current standard of care ineffective in ~60% of patients

Can affect cognition, mood, behavior and organ function



12 - 18 MONTHS

Average Life Expectancy¹

>50,000

New Cases in the 8 Major Markets² Each Year³

>151,000

Forecast of Annual New Cases in the 8 Major Markets² by 2027³

~48%

Of All Primary Malignant Brain Tumors1

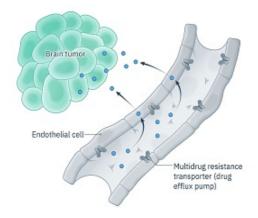


1: https://braintumor.org/take-action/about-gbm/

8 Major Markets includes USA, France, Germany, Italy, Spain, UK, Japan and urban China
 Global Data, "Glioblastoma Multiforme (GBM): Opportunity Analysis and Forecasts to 2027" (2017)

The Blood Brain Barrier (BBB)

Highly Selective, Semi-Permeable Barrier that Separates the Circulating Blood from the Brain

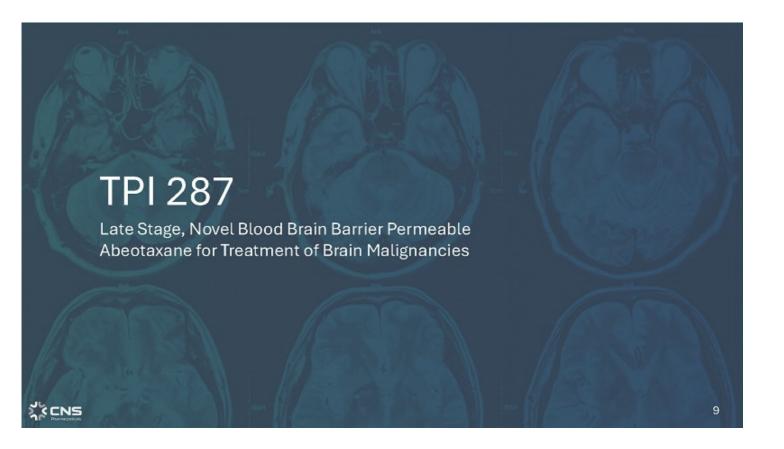


Key Functions

- · Protection:
 - Blocks toxins, pathogens and potentially harmful molecules from entering the brain by transporters that bind to these substances and deliver them back to the bloodstream
- · Selective Permeability:
 - Allows essential nutrients like glucose and amino acids to pass through while restricting larger or harmful molecules
- · Maintaining Homeostasis:
 - Ensures a controlled environment for proper neuronal function



Drug Delivery to the Brain is Challenging Due to the BBB's Selective Nature, Limiting the Access and Effectiveness of Cancer Therapies in the Brain



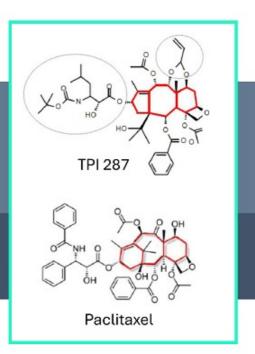
TPI 287: A Novel Taxane Derivative

Taxanes

- A class of chemotherapy that binds to microtubules and prevents them from functioning normally, which stops cancer cells from dividing
- A substrate for P-glycoprotein (Pgp), which is upregulated in cells that become taxane-resistant, and is part of the BBB

• TPI 287

- · A derivative of taxane (abeotaxane) that is not a substrate for Pgp
- · Effective in taxane-resistance and able to cross the BBB



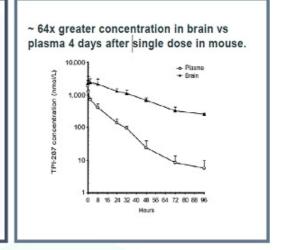


Readily Penetrates the Blood Brain Barrier in Animal Models

	COMPOUND	Blood ug*hr/ml	Brain ug*hr/g	Brain:Blood
	paclitaxel	3.2	3.2 1.6 8.7 2.5 16.8 65.9	0.5
Wild-type	docetaxel	8.7	2.5	0.3
	TPI 287	16.8	8.7 2.5 16.8 65.9 4.7 18.6	3.9
	paclitaxel	4.7	18.6	4.0
Pgp knock-out	docetaxel	9.0	15.4	1.7
	TPI 287	N/A	N/A	



pacifixed dosed 10 mg/kg AUC cal. 0.8 hr blood, 0.12 hr. brain (Clin Can Research, 9.2849, 2003). docetaxel dosed 33 mg/kg AUC cal. 0.8 hr blood, 0.12 hr. brain (Eur J Can. 40:1289, 2004). TPI 287 dosed 20 mg/kg AUC cal. 0.96 hr blood and brain (Mol Can Ther. 11:1959, 2012).



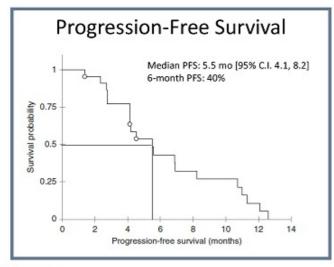


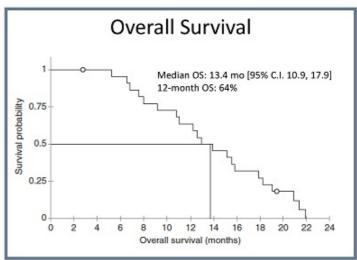
Clinical Trials with TPI 287





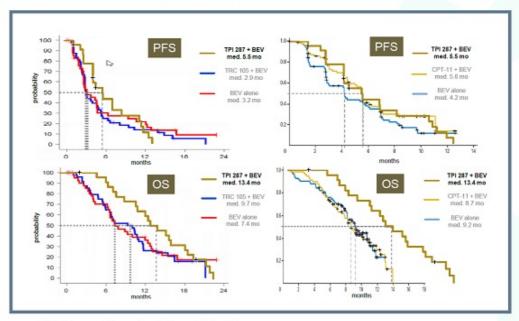
TPI 287 in Combination with Bevacizumab for the Treatment of Recurrent Glioblastoma





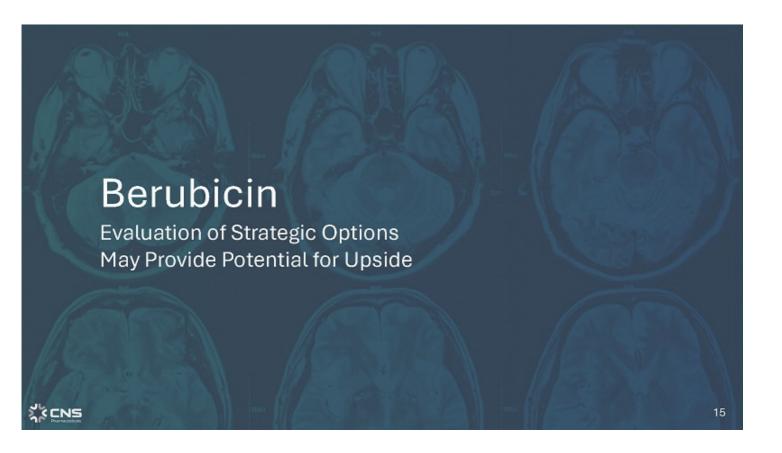


Improved GBM Survival in Combination with Bevacizumab





* Graphs represent aggregate data from multiple studies



Berubicin

Reported Primary Analysis of Berubicin in 2nd line GBM

Ongoing analysts of outcomes ongoing to determine next steps



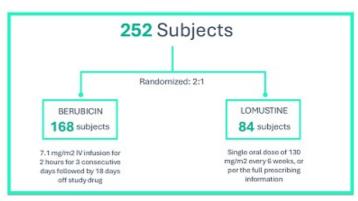
Did not a statistically significant difference in overall survival, the primary endorsin

Summary of Primary Analysis

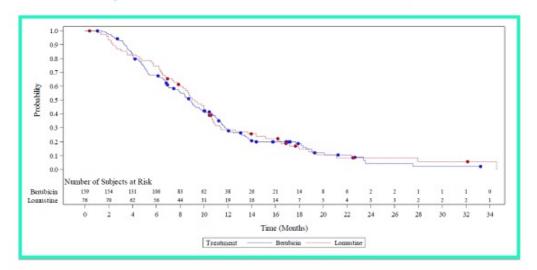
Showed clinically relevant outcomes comparable to Lomustine across multiple endpoints¹

Safety profile continues to be favorable, including the absence of anthracycline related cardiotoxicity

Analysis of outcomes are ongoing, including advanced imaging review, PK, and clinical endpoints



Berubicin Demonstrated Comparable Overall Survival Compared to Standard of Care, Lomustine





Financial Snapshot NASDAQ: CNSP

Cash Expected to Fund Operations Into the Second Half of 2026

\$17.5 Million

Cash*

As of March 31, 2025

~1.7M

Volume

Average 3 months

~\$4.4 Million

Market Cap

As of July 1, 2025

~\$4.5 Million

*Cash Position Includes Net Proceeds from Raise - \$13.0 Million with \$4.5 Million Raised After Quarter-End



Management Team



John M. Climaco, Esq PRESIDENT & CHIEF EXECUTIVE OFFICER

Twenty-one years experience managing the operations, strategies and finances of public and private lifescience companies.















Christopher S. Downs, CPA CHIEF FINANCIAL OFFICER

Nearly 20 years of finance and investment banking experience primarily in the healthcare industry









Sandra L. Silberman, MD, PhD CHIEF MEDICAL OFFICER

Board certified hematologist/medical oncologist with extensive experience in clinical development of novel therapies for the treatment of cancer. Frm Head Global Clinical Development at Novartis.















Donald Picker, PhD CHIEF SCIENTIFIC OFFICER

Over 35 years of drug development experience and responsible for the development of Carboplatin, one of the world's leading cancer drugs, acquired by Bristol-Myers Squibb and with annual sales of over \$500 million.





