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 of Report (Date of earliest event reported): August 14, 2023

Mind Medicine (MindMed) Inc.

(Exact name of Registrant as Specified in Its Charter)

New York (State or Other Jurisdiction of Incorporation)

One World Trade Center Suite 8500 New York, New York

(Address of Principal Executive Offices)

001-40360 (Commission File Number) 98-1582438 (IRS Employer Identification No.)

> 10007 (Zip Code)

Registrant's Telephone Number, Including Area Code: (212) 220-6633

Not Applicable

(Former Name or Former Address, if Changed Since Last Report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

□ Written communications pursuant to Rule 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 pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

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

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

	Trading	
Title of each class	Symbol(s)	Name of each exchange on which registered
Common Shares	MNMD	The Nasdaq Stock Market LLC

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).

Emerging growth company \boxtimes

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.

Item 7.01 Regulation FD Disclosure.

On Auguest 14, 2023, Mind Medicine (MindMed) Inc. (the "Company") posted an updated corporate presentation on the Company's website, which the Company may use from time to time in conversations with investors, analysts or other third parties.

In accordance with General Instruction B.2. of Form 8-K, the information in this Item 7.01 and Exhibit 99.1 hereto shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liability of that section, nor shall it be deemed incorporated by reference in any of the Company's filings under the Securities Act of 1933, as amended, or the Exchange Act, whether made before or after the date hereof, regardless of any incorporation language in such a filing, except as expressly set forth by specific reference in such a filing.

Item 8.01 Other Events.

As disclosed above, on August 14, 2023, the Company updated its corporate presentation, which is attached as Exhibit 99.1 hereto. The information in Exhibit 99.1 is incorporated by reference herein.

Item 9.01 Financial Statements and Exhibits.

Exhibit No.	Description
99.1	Investor presentation dated August 14, 2023
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURES

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

MIND MEDICINE (MINDMED) INC.

Date: August 14, 2023

By:/s/ Robert BarrowName:Robert BarrowTitle:Chief Executive Officer

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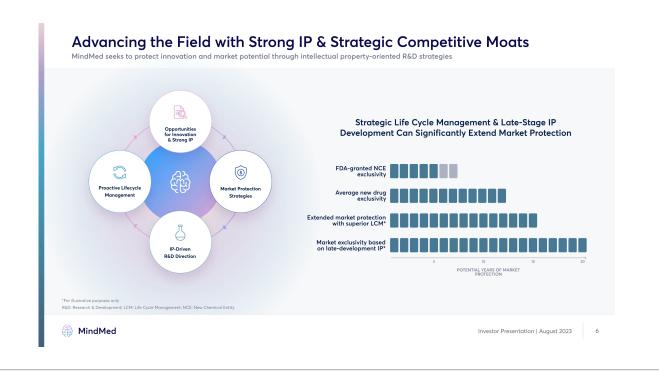
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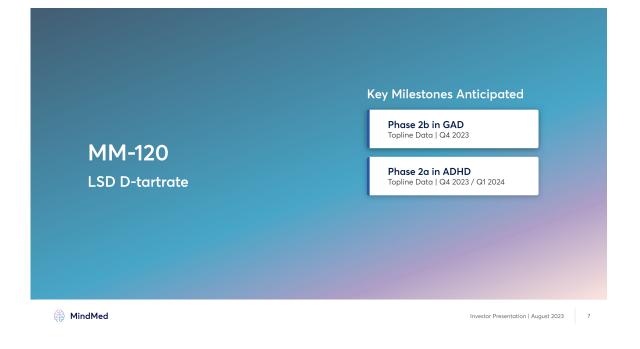
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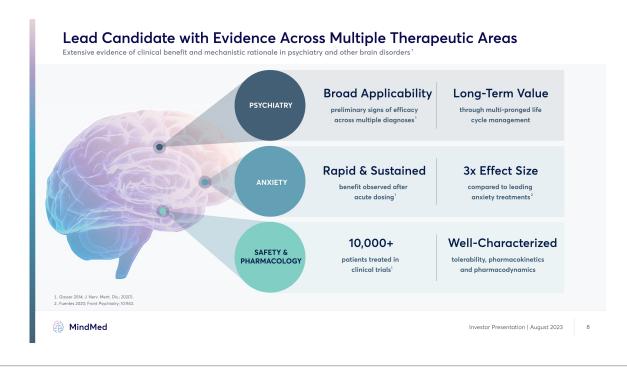
	GAD	10%	1-year prevalence of anxiety disorders in the US ¹
	ADHD	4.4%	estimated prevalence rate o ADHD among all US adults ²
1 in 4 U.S. Adults has a Diagnosable			
Mental Health Disorder	ASD	\$461B	economic cost of ASD in the US predicted by 2025 ³

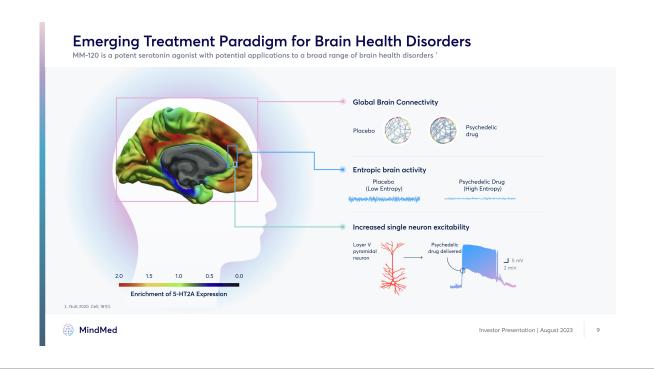
Research & Development Pipeline Our pipeline diversification offers potential opportunities across therapeutic areas and mechanisms of action

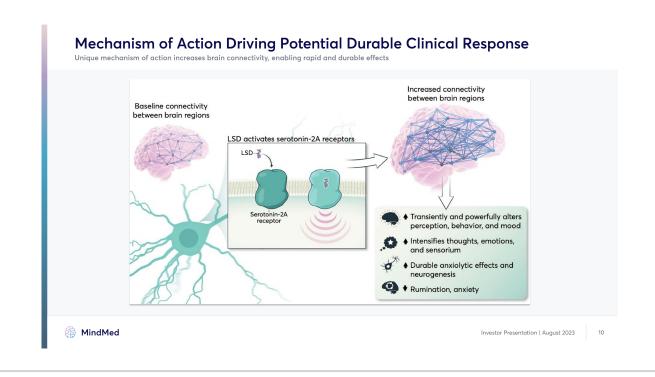
PRODUCT CANDIDATE	INDICATION	PRE-CLINICAL	PHASE 1	PHASE 2	PHASE 3	REGISTRATION
PSYCHIATRY						
MM-120 (LSD D-tartrate)	Generalized Anxiety Disorder					
	ADHD					
MM-402 (R(-)-MDMA)	Autism Spectrum Disorder					
OTHER PROGRAMS						
MM-110 (zolunicant HCI)*	Opioid Withdrawal					
DISCOVERY & EARLY DEVELOPM	ENT					
Novel tryptamines	undisclosed					
Novel phenethylamines	undisclosed					
Advanced drug delivery	undisclosed					
INVESTIGATOR-INITIATED TRIALS						
Lysergic Acid Diethylamide (LSD)	Major Depressive Disorder					
Lysergic Acid Diethylamide (LSD)	Cluster Headache					
PK/PD of MDMA enantiomers	Healthy Subjects					
Continued development of MM-110 is cu Full trial details and clinical trials.gov li						
Attention-Deficit/Hyperactivity Disorder.	LSD: lysergic acid diethylamide; MDN	1A: 3,4-methylenedioxymethamphetar	nine			











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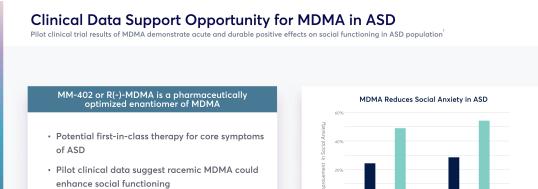
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Translational preclinical data suggest that R(-)-MDMA may have:	R(-)-MDMA Maintains Prosocial Effects with Reduced Stimulant Activity
 Strong prosocial effects Less stimulant activity compared to MDMA Plan to develop standard, at-home dose regimen 	200% 100% R R/S S R R/S S Interaction Locomotor Activity
Source: [1][2]	Source: [2]



• Pharmacological profile aligns with patientdesired treatment benefits

MDMA Placebo 1 month Source: [1]

1. Danforth 2018; Psychopharmacology; 235. MDMA: 3,4-methylenedioxymethamphetamine; ASD

💮 MindMed

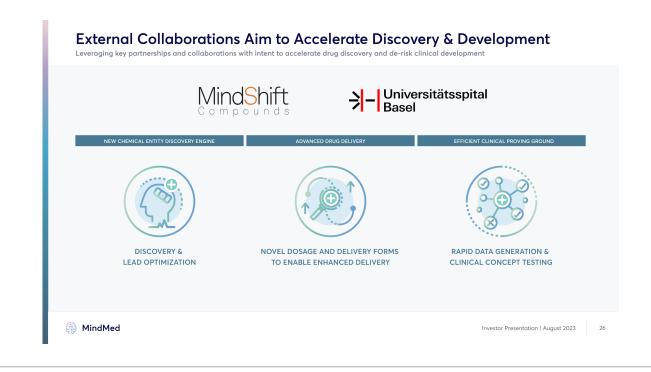
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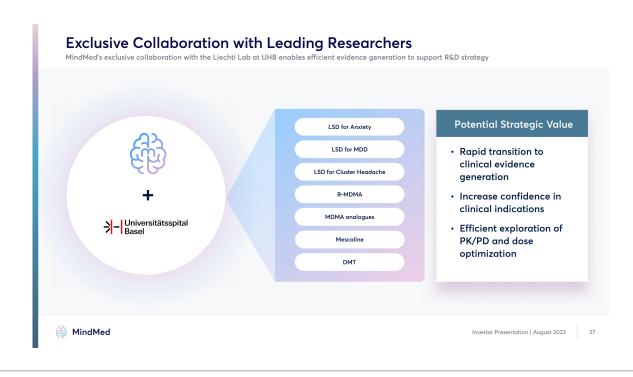
MDMA

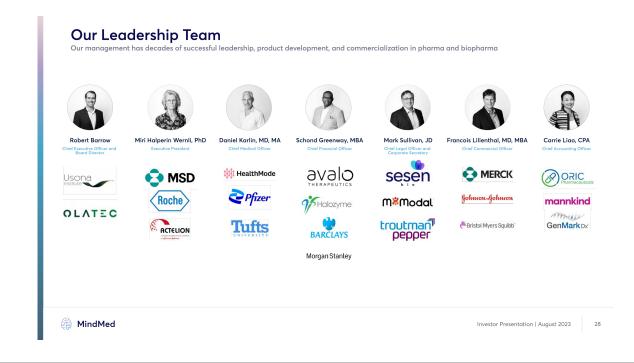
Placebo

6 months











CNS Products	Other Products
CROZECTEM rameliteon Brog subtra	Pifeltro Wing was to be a series with the series of the se
Selincro	
Subclassing vortioxetine smg=tomg=20mg tablets	
Emoningine tabless	
Suboxone Sublingual Ruperophine and reducing @ Film Anatoria - Anatoria - Stratoria	International Control

Business Highlights

- A leader in developing psychedelic product candidates to treat brain health disorders
- Diversified pipeline of clinical programs targeting significant unmet medical needs
- IP and R&D strategies intended to maximize market exclusivity and protection
- · Leveraging decades of research on clinical and preclinical potential of product candidates
- Expertise in drug and digital medicine development and commercialization
- Expected cash runway through key clinical readouts and into 2026¹
- MM-120 (LSD D-tartrate) for the treatment of GAD and ADHD
- Phase 2b dose-optimization study ongoing for the treatment of GAD; topline results expected in Q4 2023
- + Phase 2a study ongoing for the treatment of ADHD; topline results expected in Q4 2023 / Q1 2024
- MM-402 or R(-)-MDMA for the treatment of core symptoms of ASD
- IND-enabling studies ongoing; initiation of a Phase 1 clinical trial is planned in Q4 2023
- Phase 1 (UHB) investigator-initiated trial of R-, S- and R/S-MDMA in healthy volunteers ongoing; topline results expected H1 2024

1. The company's ending Q2 2023 cash and cash equivalents of \$116.9 million and committed credit facility are expected to fund operations into 2026, if certain milestones are achieved that unlock additional capital.

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