
**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): April 26, 2022

MIND MEDICINE (MINDMED) INC.

(Exact Name of Registrant as Specified in Charter)

British Columbia, Canada
(State or Other Jurisdiction
of Incorporation)

001-40360
(Commission
File Number)

98-1582438
(IRS Employer
Identification No.)

One World Trade Center, Suite 8500
New York, New York
(Address of Principal Executive Offices)

10007
(Zip Code)

Registrant's telephone number, including area code: (650) 208-2454

Not applicable
(Former Name or Former Address, if Changed Since Last Report)

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

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

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 (see General Instruction A.2. below):

- 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))

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

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 April 26, 2022, MindMedicine (MindMed) Inc. (the “Company”) posted a corporate presentation titled Corporate Overview to the “Investor Relations” page of its website at <https://mindmed.co/investor-resources/>, which the Company may use from time to time in communications or conferences. A copy of the corporate presentation is attached as Exhibit 99.1 to this Current Report on 8-K (the “Report”).

The information in this Report and Exhibit 99.1 attached hereto is being furnished to the Securities and Exchange Commission and shall not be deemed to be “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”), or otherwise subject to the liabilities of that Section, nor shall it be deemed incorporated by reference into any filing under the Securities Act of 1933, as amended, or the Exchange Act, regardless of any general incorporation language in such filing, except as expressly set forth by specific reference in such filing. The Company’s submission of this Report shall not be deemed an admission as to the materiality of any information required to be disclosed solely to satisfy the requirements of Regulation FD.

This Report and Exhibit 99.1 hereto contain forward-looking statements within the meaning of the federal securities laws. These forward-looking statements are based on current expectations and are not guarantees of future performance. Further, the forward-looking statements are subject to limitations listed in Exhibit 99.1 and in the other reports of the Company filed with the Securities and Exchange Commission, including that actual events or results may differ materially from those in the forward-looking statements.

Item 9.01 Financial Statements and Exhibits.

| Exhibit No. | Description |
|--------------------|---|
| 99.1 | MindMed Corporate Overview, dated April 2022 |
| 104 | Cover Page Interactive Data File (embedded within the Inline XBRL document) |

SIGNATURE

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.

Date: April 26, 2022

MIND MEDICINE (MINDMED) INC.

By: /s/ Cynthia Hu
Name: Cynthia Hu
Title: Chief Legal Officer & Secretary



MindMed

Corporate Overview

April 2022

Disclaimer

This presentation (the "Presentation") has been prepared by Mind Medicine (MindMed) Inc. ("MindMed" or the "Company") solely for informational purposes. None of MindMed, its affiliates or any of their respective employees, directors, officers, contractors, advisors, members, successors, representatives or agents makes any representation or warranty as to the accuracy or completeness of any information contained in this Presentation and shall have no liability for any representations (expressed or implied) contained in, or for any omissions from, this Presentation. This presentation shall not constitute an offer, nor a solicitation of an offer, of the sale or purchase of securities. This Presentation does not constitute an offering of securities of MindMed and under no circumstances is it to be construed as a prospectus or advertisement or public offering of securities.

Any trademarks included herein are the property of the owners thereof and are used for reference purposes only. Such use should not be construed as an endorsement of the products or services of MindMed.

Any amounts are in USD unless otherwise noted. MindMed's securities have not been approved or disapproved by the SEC or by any state, provincial or other securities regulatory authority, nor has the SEC or any state, provincial or other securities regulatory authority passed on the accuracy or adequacy of this Presentation. Any representation to the contrary is a criminal offense.

Cautionary Note Regarding Forward-Looking Statements

This Presentation contains, and our officers and representatives may from time to time make, "forward-looking statements" within the meaning of the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995 and other applicable securities laws. Forward-looking statements can often, but not always, be identified by words such as "plans", "expects", "is expected", "budget", "scheduled", "estimates", "forecasts", "intends", "anticipates", "will", "projects", or "believes" or variations (including negative variations) of such words and phrases, or statements that certain actions, events, results or conditions "may", "could", "would", "might" or "will" be taken, occur or be achieved, and similar references to future periods. Except for statements of historical fact, examples of forward-looking statements include, among others, statements pertaining to the development and commercialization of any medicine or treatment, or the efficacy of either of the foregoing, the success and timing of our development activities, the success and timing of our planned clinical trials, our ability to meet the milestones set forth herein, the likelihood of success of any clinical trials or of obtaining FDA or other regulatory approvals, the likelihood of obtaining patents or the efficacy of such patents once granted, and the potential for the markets that MindMed is anticipating to access.

Forward-looking statements are neither historical facts nor assurances of future performance. Instead, they are based only on our current beliefs, expectations and assumptions regarding the future of our business, future plans and strategies, projections, anticipated events and trends, the economy and other future conditions as of the date of this Presentation. While we consider these assumptions to be reasonable, the assumptions are inherently subject to significant business, social, economic, political, regulatory, competitive and other risks and uncertainties that are difficult to predict and many of which are outside of our control, and our actual results and financial condition may differ materially from those indicated in the forward-looking statements. Therefore, you should not rely on any of these forward-looking statements. Important factors that could cause our actual results and financial condition to differ materially from those indicated in the forward-looking statements include, among others, the following: our ability to raise capital to complete its plans and fund its studies; the medical and commercial viability of the contemplated medicines and treatments being developed; our ability to raise additional capital in the future as we continue to develop our products; our history of negative cash flow; our limited operating history; incurrence of future losses; availability of additional capital; lack of revenue; compliance with laws and regulations; difficulty associated with research and development; risks associated with clinical trials or studies; heightened regulatory scrutiny; early stage product development; clinical trial risks; regulatory approval processes; novelty of the psychedelic inspired medicines industry; as well as those risk factors discussed or referred to throughout the "Risk Factors" sections of our most recently filed Annual Report on Form 10-K filed with the Securities and Exchange Commission (the "SEC") and in other filings we make in the future with the SEC and the securities regulatory authorities in all provinces and territories of Canada, available under the Company's profile on SEDAR at www.sedar.com.

Any forward-looking statement made by us in this Presentation is based only on information currently available to us and speaks only as of the date on which it is made. MindMed undertakes no obligation to publicly update any forward-looking statement, whether written or oral, that may be made from time to time, whether as a result of new information, future developments or otherwise.

Cautionary Note Regarding Regulatory Matters

The United States federal government regulates drugs through the Controlled Substances Act. The Company works with a non-hallucinogenic synthetic derivative of the psychedelic substance ibogaine, known as "18-MC", which is a synthetic organic molecule designed around a common coramidine chemical backbone. 18-MC is not a Schedule I substance in the United States and the Company does not foresee it becoming a Schedule I substance due to its non-hallucinogenic properties. While the Company is focused on programs using psychedelic inspired compounds and classic psychedelics, the Company does not have any direct or indirect involvement with the illegal selling, production or distribution of any substances in the jurisdictions in which it operates. The Company is a neuro-pharmaceutical drug development company and does not deal with psychedelic substances except within laboratory and clinical trial settings conducted within approved regulatory frameworks. The Company's products will not be commercialized prior to applicable regulatory approval, which will only be granted if clinical evidence of safety and efficacy for the intended uses is successfully developed.

Market and Industry Data

This Presentation includes market and industry data that has been obtained from third party sources, including industry publications. MindMed believes that the industry data is accurate and that the estimates and assumptions are reasonable, but there is no assurance as to the accuracy or completeness of this data. Third party sources generally state that the information contained therein has been obtained from sources believed to be reliable, but there is no assurance as to the accuracy or completeness of included information. Although the data is believed to be reliable, MindMed has not independently verified any of the data from third party sources referred to in this Presentation or ascertained the underlying economic assumptions relied upon by such sources. References in this Presentation to research reports or to articles and publications should be not construed as depicting the complete findings of the entire referenced report or article. MindMed does not make any representation as to the accuracy of such information.

Business Highlights

Our mission is to deliver on the therapeutic potential of psychedelics and other novel targets to treat brain health disorders

- **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** to maximize market exclusivity and protection
- **Leveraging decades of research** on clinical and preclinical potential of product candidates
- **Industry-leading expertise** in drug and digital medicine development and commercialization
- **Fully funded** through key clinical readouts and into 2024

There is an Urgent Need for Better Treatments

Substantial opportunities exist to advance novel treatments for a wide range of brain health disorders



1 in 5 U.S. Adults is Diagnosed with a Mental Health Disorder¹

1. NIMH 2020; Mental Illness.
2. Bandelow 2015; Dialogues Clin. Neurosci; 17(3).
3. Zelaya 2019; NCHS Data Brief; 2020; (390).
4. NIDA 2022; Overdose Death Rates.
5. Leigh & Du 2015; J. Autism Dev. Disord.; 45(12).

ANXIETY

21%

1-year prevalence of anxiety disorders in the US

PAIN

50M

US adults live with chronic pain²

OUD

68,630

US deaths from opioid overdose in 2020⁴

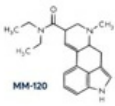

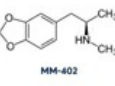
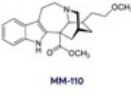
ASD

\$461B

economic cost of ASD in the US predicted by 2025⁵

Advancing Multiple Generations of Drug Candidates

Our strategy is to deliver on well-characterized psychedelic candidates and next generation candidates with enhanced drug profiles

| | CONCEPT | MINDMED PRODUCT CANDIDATES | PIPELINE EXPANSION OPPORTUNITIES |
|-----------------------------------|---|---|---|
| CLASSIC PSYCHEDELICS | <ul style="list-style-type: none"> Clinical evidence of efficacy¹ Well-characterized pharmacology Accelerated development potential |  <p>MM-120</p> | <ul style="list-style-type: none"> Expanded clinical indications Psychedelics with distinct PK/PD  |
| 2ND GENERATION / OPTIMIZED | <ul style="list-style-type: none"> Enhanced pharmacology Overcome safety liabilities Increased IP potential |  <p>MM-402</p> | <ul style="list-style-type: none"> Advanced drug delivery Novel treatment models Novel treatment regimen |
| 3RD GENERATION / NCEs | <ul style="list-style-type: none"> Analogues of classic psychedelics Require full development program Strongest IP potential |  <p>MM-110</p> | <ul style="list-style-type: none"> Novel tryptamines Novel phenethylamines Non-hallucinogenic analogues |

1. Gasser 2014; J Nerv Ment Dis; 202(7)

IP: intellectual property; DMT: N,N-dimethyltryptamine; NCE: new chemical entity; PD: pharmacodynamics; PK: pharmacokinetics

Research & Development Pipeline

Our pipeline diversification offers potential opportunities across therapeutic areas and mechanisms of action



ADHD: Attention-Deficit/Hyperactivity Disorder

Upcoming Portfolio Milestones

MindMed's clinical research portfolio creates multiple near-term and intermediate catalysts



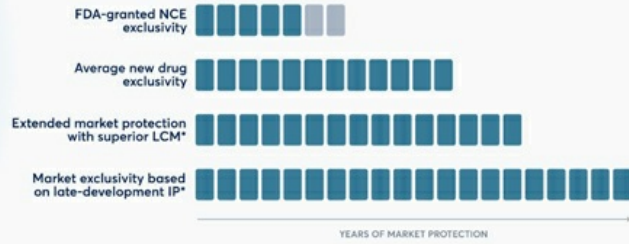
ADHD: Attention-Deficit/Hyperactivity Disorder; IT: Investigator-Initiated Trial; R&D: research & development; ESDE: early sign of efficacy

Advancing the Field with Strong IP & Strategic Competitive Moats

Our approach is to protect innovation and market potential through intellectual property-oriented R&D strategies



Strategic Life Cycle Management & Late-Stage IP Development Can Significantly Extend Market Protection



*For illustrative purposes only
R&D: Research & Development; LCM: Life Cycle Management

MM-120

LSD D-tartrate

Key Milestones

LSD-Anxiety Topline Readout
Q2 2022 | Phase 2 (IIT)

GAD First Patient Dosing
Q2-Q3 2022 | Phase 2b

Chronic Pain Study Initiation
Q4 2022 | Phase 2 ESOE

ESOE: early sign of efficacy

MM-120 | Lead Candidate with Evidence Across Multiple Therapeutic Areas

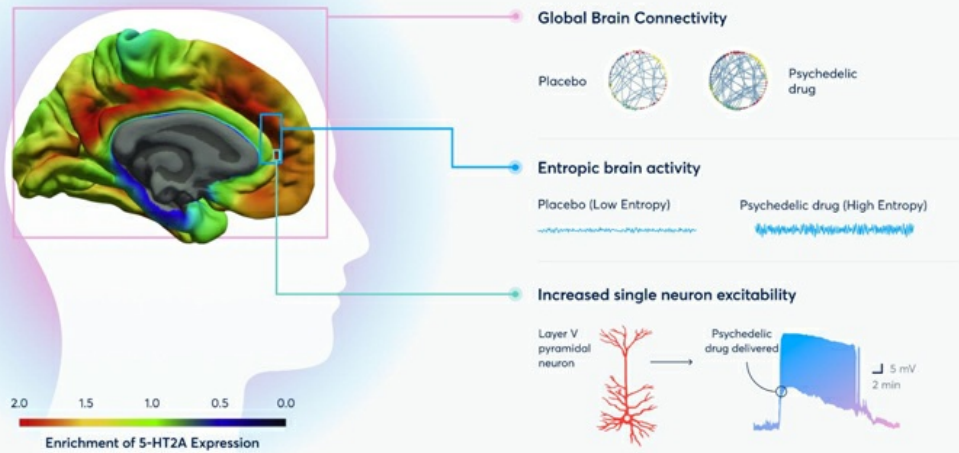
Extensive evidence of clinical benefit and mechanistic rationale in psychiatry, pain and substance use disorders ¹



1. Gasser 2014; J. Nerv. Ment. Dis.; 2020;
2. Fuentes 2020; Front Psychiatry; 10:943.
MOA: mechanism of action

MM-120 | Emerging Treatment Paradigm for Brain Health Disorders

MM-120 is a potent serotonin agonist with potential applications to a broad range of brain health disorders



Source: Nutt 2020, Cell; 181(1).

MM-120 | Legacy of LSD Clinical Research in Psychiatric Disorders

Building on decades of clinical research on LSD in anxiety and depression

| STUDIES | INDICATION(S) | SAMPLE SIZE | KEY FINDINGS |
|---------------------------------------|----------------------------------|--------------|---|
| 21 STUDIES PRIOR TO 1974 ¹ | Anxiety, depression & 'neuroses' | 512 patients | Up to 95% reduction in symptoms |
| GASSER 2014 ² | Anxiety in terminal illness | 12 patients | Effect size of 1.1 with durable reductions in anxiety at 1 year |
| LSD-ASSIST STUDY | Anxiety | 41 patients | Topline results expected in Q2 2022 |



1. Rucker 2016, J. Psychopharmacol; 30(12).
2. Gasser 2014, J. Nerv. Ment. Dis.; 202(7).

MM-120 | Phase 2b Generalized Anxiety Disorder (GAD)

Study design seeks to demonstrate dose-responsive effects and identify optimal dose for pivotal clinical trials

PSYCHIATRY | MM-120 (LSD D-tartrate) | Indication: GAD | PHASE 2B



Study MMED008 | MM-120 for GAD

A Phase 2b Dose Optimization Study of a Single Dose of MM-120 in Generalized Anxiety Disorder

KEY ENTRY CRITERIA

- Men and Women
- Ages 18-74
- Diagnosis of GAD
- HAM-A \geq 20

ADDITIONAL ENDPOINTS

- MADRS
- CGI-S / I
- PGI-S / C
- SDS
- EQ-5D-5L
- PSQI
- ASEX

Source: MindMed internal study documents

µg: microgram; HAM-A: Hamilton Anxiety Rating Scale; MADRS: Montgomery-Åsberg Depression Rating Scale; CGI-S: Clinical Global Impression - Severity; PGI-S: Patient Global Impression - Severity; SDS: Sheehan Disability Scale; EQ-5D-5L: EuroQol-5 Dimension; PSQI: Pittsburgh Sleep Quality Index; ASEX: Antasma Sexual Experiences Scale

MM-120 | Phase 2a Attention Deficit Disorder (ADHD)

Proof of concept study design seeks to explore potential clinical response in ADHD

PSYCHIATRY | MM-120 (LSD D-tartrate) | Indication: ADHD | PHASE 2A



Study MMED007 | MM-120 for ADHD

A Phase 2a Proof of Concept Study of Repeated Low Doses of MM-120 for the Treatment of ADHD in Adults

KEY ENTRY CRITERIA

- Men and Women
- Ages 18-65
- Diagnosis of ADHD
- AISRS \geq 26
- CGI-S \geq 4

ADDITIONAL ENDPOINTS

- AISRS @ 1 week
- CGI-S
- ASRS
- CAARS
- Sleep Diary

Source: MindMed internal study documents

AISRS: Adult ADHD Investigator Symptom Rating Scale; ASRS: Adult ADHD Self-Report Scale; CAAR: Conners' Adult ADHD Rating Scales; CGI-S: Clinical Global Impression - Severity

MM-120 | Novel Applications in Chronic Pain

Preclinical and early clinical evidence provide support for unique mechanism of action and potential clinical activity

| SELECT STUDIES | INDICATION(S) | SAMPLE SIZE | KEY FINDINGS |
|--------------------------------|---|--------------|--|
| KAST 1967 ¹ | Terminal cancer pain | 128 patients | 100 µg reduced cumulative pain scores for at least 12 hours post-treatment |
| FANCIULLACCI 1977 ² | Phantom limb pain | 7 patients | 50 µg (qd) reduced pain in 5 of 7 patients (full analgesia in 2 of 7) |
| RAMAEKERS 2021 ³ | Experimental pain in healthy volunteers | 24 patients | 20 µg increased pain tolerance and reduced cold pressor test painfulness |

Study MM-120C201 - Phase 2 ESOE in Chronic Pain

| | |
|-------------------------|---|
| Study Design | To be announced |
| Dosing Regimen | Repeat administration |
| Indication | Chronic Pain |
| Primary Endpoint | Change in Daily Pain on 11-point Numerical Rating Scale |

1. Kast 1967. *Psych Quor* 41, 646-657.
2. Fanciullacci 1977. *The Journal of Head and Face Pain*, 17: 118-119.
3. Ramaekers 2021. *Journal of Psychopharmacology*, 35(4).

MM-110

Zolunicant HCl

Key Milestones

Phase 1 Topline Data Readout
Q2 2022 | Phase 1

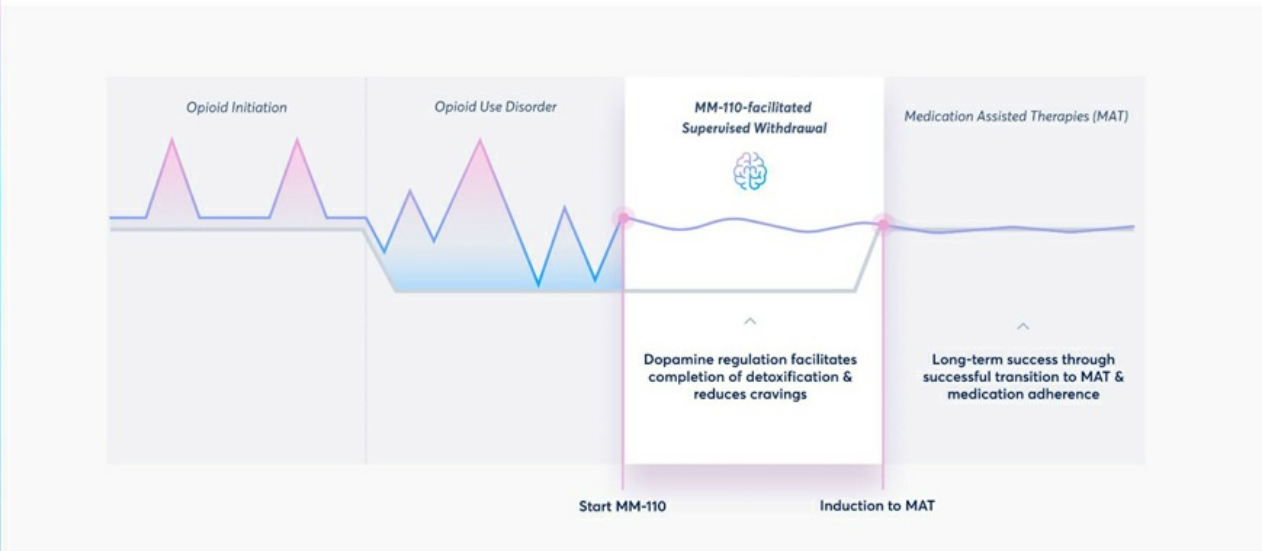
Opioid W/D Study Initiation
Q2 2022 | Phase 2a

Opioid W/D ESOE Readout
Q1 2023 | Phase 2a (Part A)

ESOE: early sign of efficacy; W/D: withdrawal

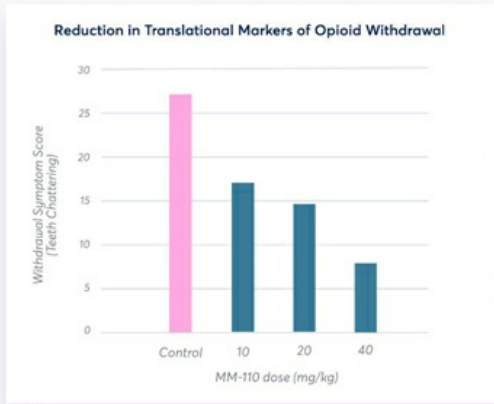
MM-110 | Novel Mechanism to Address a Critical Gap in OUD Treatment

Mechanism of action and target product profile complement standard-of-care and address a critical gap in available treatment landscape

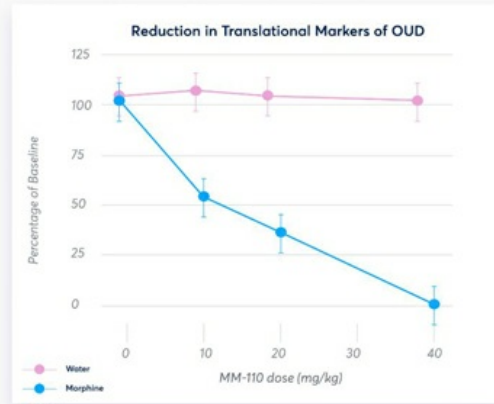


MM-110 | Strong Preclinical Efficacy on Key Translational Outcomes

A single dose of MM-110 mitigates withdrawal symptoms and opioid self-administration in preclinical models^{1,2}



Source: [1]



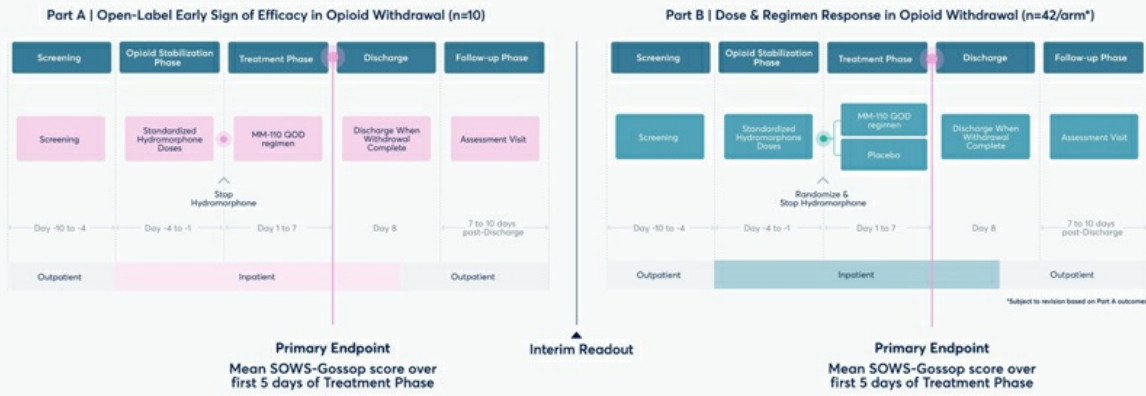
Source: [2]

1. Rho & Glick 1998; NeuroReport; 9.
2. Maisonneuve & Glick 2003; Pharmacol Biochem Behav; 75.

MM-110 | Phase 2a Supervised Withdrawal in Opioid Use Disorder

Gated two-part study design provides opportunity for early signs of efficacy (ESOE) and informs randomized proof of concept design

SUBSTANCE USE DISORDERS | MM-110 (zincuricant HC; 18-MC) | Indication: Opioid Withdrawal | PHASE 2A



Source: MindMed internal study documents
 ESOE: early sign of efficacy; QOD: Every Other Day (dosage timing); SOWS: Subjective Opiate Withdrawal Scale

MM-402

R(-)-MDMA

Key Milestones

PK/PD Study Initiation
Q3 2022 | Phase 1 IIT

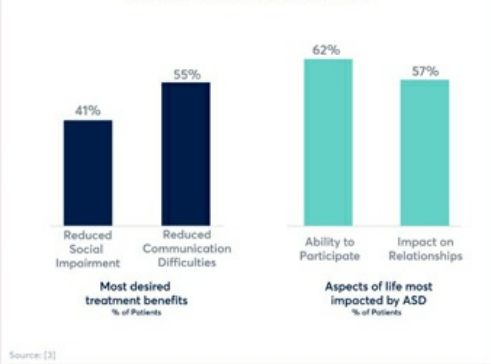
Phase 1 Study Initiation
2023 | Phase 1

No Approved Drugs for Core Symptoms of Autism Spectrum Disorder (ASD)

Growing prevalence and impact of ASD yields an urgent need for novel therapies that target core symptoms and align with patient preferences



MM-402 Activity Aligns with Reported Needs and Desired Benefits for Individuals with ASD



Source: [1]

1. Leigh & Du 2015; J Autism Dev Disord. 45(12).
2. Kogan, Pediatrics 2018; 142(6).
3. FDA Patient Focused Drug Development workshop on Autism Spectrum Disorder (2017)

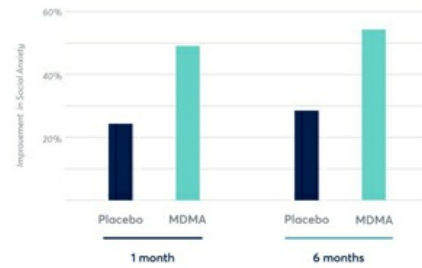
MM-402 | Clinical Data Support Opportunity in ASD

Pilot clinical trial results of MDMA demonstrate acute and durable positive effects on social functioning in ASD population¹

MM-402 or R(-)-MDMA is a pharmacologically optimized enantiomer of MDMA

- Potential first in class therapy for core symptoms of ASD
- Pilot clinical data suggest MDMA could enhance social functioning
- Pharmacological profile aligns with patient-desired treatment benefits

MDMA Reduces Social Anxiety in ASD



Source: [1]

1. Danforth 2018; Psychopharmacology; 235.
MDMA; 3,4-methylenedioxyamphetamines

MM-402 | Preclinical Data Indicate Potential Enhanced Benefit/Risk Profile

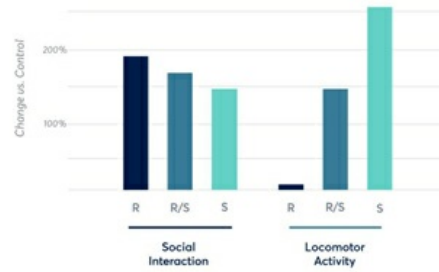
Preclinical data suggest the R-enantiomer of MDMA has enhanced prosocial effects with an improved safety profile

Translational preclinical data suggest that R(-)-MDMA may have:

- Strong prosocial effects
- Less stimulant activity compared to MDMA
- Superior safety and tolerability profile
- Potential to be administered in standard dosing regimen

Source: [1][2]

R(-)-MDMA Maintains Prosocial Effects with Reduced Stimulant Activity



Source: [2]

1. Pitts 2018; Psychopharmacology; 235.
2. Curry 2018; Neuropharmacology; 126.

Collaborations & Early R&D

External Collaborations Accelerate Discovery & Development

Leveraging key partnerships and collaborations to accelerate drug discovery and de-risk clinical development

NEW CHEMICAL ENTITY DISCOVERY ENGINE



DISCOVERY &
LEAD OPTIMIZATION

MindShift
Compounds

ADVANCED DRUG DELIVERY



BRAIN TARGETED LIPOSOMES
(BTLs)

nextage
GENERIC INNOVATION CENTER

EFFICIENT CLINICAL PROVING GROUND

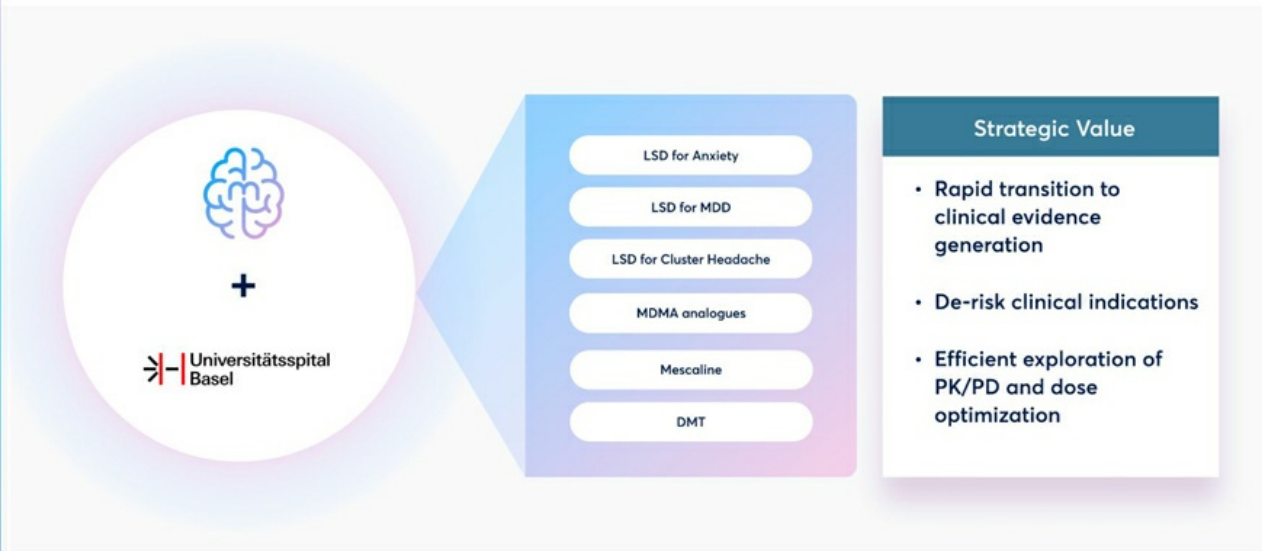


RAPID DATA GENERATION &
CLINICAL CONCEPT TESTING

Universitätsspital
Basel

Exclusive Collaboration with Leading Researchers

MindMed's exclusive collaboration with the Liechti Lab at UHB enables efficient evidence generation to support R&D strategy



Digital Medicine

Digital Unlocks Potential Opportunities Throughout the Product Lifecycle

Generating data, insights, models, and tools from early development through market management

Preclinical Research

IND & Phases 1 - 3

Drug Launch

Enhancement and Lifecycle Management

Clinical Development Tools



- Deep Digital Diagnoses
- Decentralized Trials
- Advanced Analytics

Companion Products



- Decision Support
- Predictive Intervention
- Patient Engagement

Post Approval Research



- Surveillance & Registries
- Remote Management
- HEOR

Combination Products

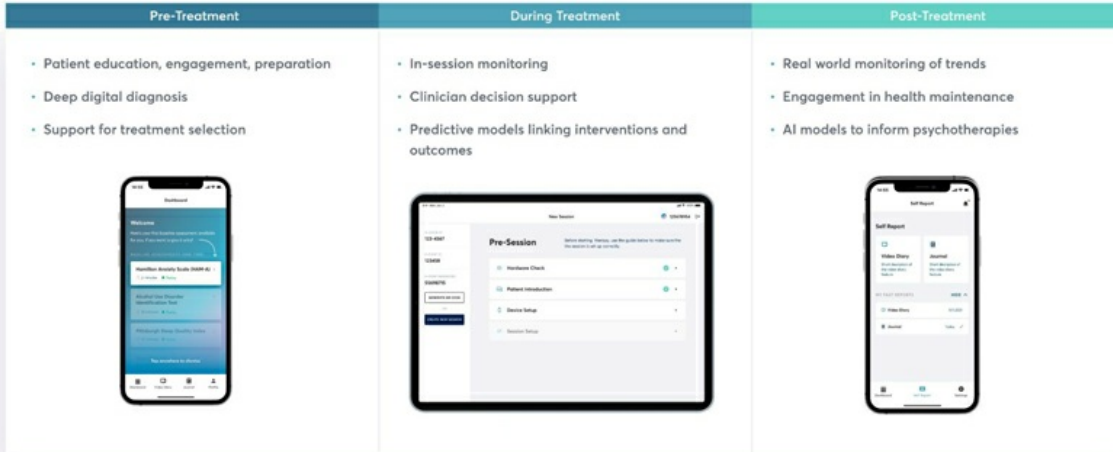


- Drug-Device Combinations
- Lifecycle Enhancement
- Efficient Phase 4 Research

HEOR: health economics and outcomes research

Digital Platform Will Add Value Through the Patient Journey

Developing a scalable delivery platform to enable adoption leveraging the existing treatment ecosystem



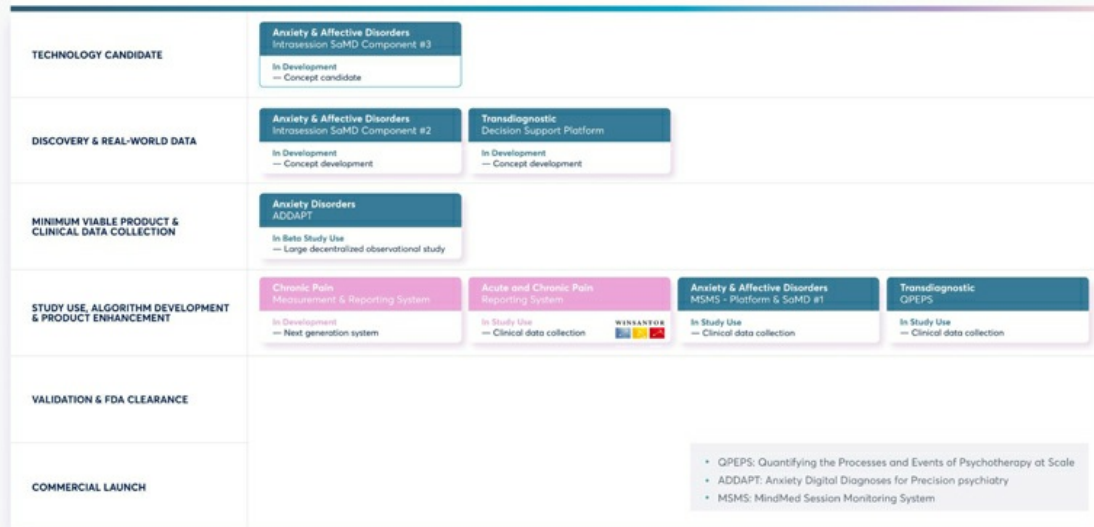
Digital Enables Alignment of Incentives for Broad Market Access

Complementary digital medicine products and studies for improved brain health outcomes



Digital Pipeline Progression Aligns with Drug Development

Executing across product categories with strong technical development and clinical research



Corporate Information

NASDAQ: MNMD // NEO: MMED

First Publicly Listed Company Developing Psychedelic Product Candidates

SHARE OWNERSHIP AS OF DECEMBER 31, 2021

| | | |
|-----------------------------------|-------------|-------|
| EXECUTIVE TEAM/DIRECTORS/INSIDERS | 44,796,490 | 9.2% |
| NON-INSIDER SHARES | 377,108,827 | 76.8% |
| EQUITY INCENTIVE PLAN (ISSUED) | 46,269,703 | 9.4% |
| OUTSTANDING WARRANTS | 22,539,931 | 4.6% |
| TOTAL (FULLY DILUTED) | 490,714,951 | 100% |



Market Capitalization: USD \$653 million | December 31, 2021 (\$1.38 per share)

Market Capitalization: C\$823 million | December 31, 2021 (C\$1.74 per share)

\$204 million

Raised since inception including warrants

\$133.5 million

Cash position as of December 31, 2021

Leadership: Leading Expertise in Innovative Drug & Digital Development



Robert Barrow
Chief Executive Officer & Board Director

Rob is an accomplished pharmaceutical executive and clinical pharmacologist with over a decade of experience leading drug development programs in a variety of disease areas. Mr. Barrow previously served as Director of Drug Development & Discovery at Unicon Institute, where he oversaw preclinical, clinical and regulatory development efforts for all of Unicon's development programs. Prior to joining Unicon, he served as Chief Operating Officer of Olistic Therapeutics where he oversaw the execution of numerous early- and late-stage clinical trials in the fields of analgesics, rheumatology, immunology and cardiovascular disease. Rob holds a Master's degree in Pharmacology from The Ohio State University and a Bachelor of Science degree from Wake Forest University, where he graduated summa cum laude.



Miri Halperin Wernli, PhD
Executive President & Board Director

Miri co-founded Creso Pharma, a cannabis company, and listed the company on the Australian Stock exchange (ASX) in October 2016. Prior to founding Creso Pharma Dr. Halperin Wernli worked in clinical psychiatry in Swiss academic hospital settings and then held various global senior leadership positions in the pharma and biotech industries in Switzerland and in the US (Merck, Sharp and Dohme, Roche and Actelion pharmaceuticals) covering Product Development, R&D, and Strategic Marketing. Her extensive pharmaceutical industry and biomed research and development experience covers the full spectrum of areas and activities from Preclinical to Clinical Development and Strategy, to Drug Registration and Launch, across several Therapeutic Areas.



Cynthia Hu, JD
Chief Legal Officer & Corporate Secretary

Cynthia joined in December 2021 as Chief Legal Officer & Corporate Secretary. Previously, from 2009-2021, she served as COO, General Counsel & Secretary at CASI Pharmaceuticals, Inc. and, from 2006 to 2009, as VP, General Counsel, of its predecessor, EnvoMed, Inc. Prior to that, she served as senior associate for the corporate and finance practice group at Powell Goldstein LLP in Washington, DC, where she advised clients on all corporate and financing matters, including complex public and private financings, mergers and acquisitions, SEC and regulatory compliance, and corporate governance and compliance. Before that, Ms. Hu was counsel for a NYSE-listed financial institution and prior to that was in private law practice at Klaw, Harrison, Harvey & Brantburg, LLP and Littmoss & Krooks, LLP focusing on corporate transactions and compliance with corporate and securities laws.



Daniel Karlin, MD, MA
Chief Medical Officer

Dan co-founded HealthMade in 2018 and served as its CEO until it was acquired by MindMed in 2021. Before that, he built and led Clinical, Informatics, and Regulatory Strategy for Pfizer Digital Medicine and Innovation Research Lab. He also served as a Global Clinical Lead for psychiatric clinical assets at Pfizer. Previously, he was the founder and Chief Medical Officer at Colum Health, a leading technology-enabled psychiatry and addiction practice. He is a founding board member of the Digital Medicine Society, and a strategic advisor to multiple big pharma and digital therapeutics companies. Dan is board Certified in Psychiatry, Addiction Medicine, and Clinical Informatics. He is an Assistant Professor of Psychiatry at Tufts University School of Medicine. He graduated with degrees in Neuroscience and Behavior (BA) and Clinical Informatics (MA) from Columbia University; and Medicine (MD) from the University of Colorado School of Medicine.



Carrie Liao, CPA
VP, Corporate Controller & Accounting Principal

Carrie Liao, CPA is an active Certified Public Accountant in the state of California with over 20 years of accounting and finance leadership experience in public and private companies. Starting at Deloitte, her career has focused on the life science industry from early development through commercialization and manufacturing. Recently, she has successfully supported multiple Initial Public Offering filings. She specializes in Sarbanes-Oxley Act compliance, resolution of complex accounting issues, process improvement, and U.S. Securities and Exchange Commission interim and annual filings.



Francois Lilienthal, MD, MBA
Chief Commercial Officer

Francois is a globally accomplished biopharmaceutical executive and a trained physician with extensive experience leading end-to-end development and commercialization of innovative medicines, driving significant growth across diversified portfolios through product launches, life cycle management and business development. Before joining MindMed, Francois was a Vice President at Merck's commercial division for 14 years. He built and led a new department focused on developing the commercial strategy for multiple products across several therapeutic areas, including neurology, psychiatry and pain. He previously drove double-digit growth of the Virology and Liver Diseases global business and oversaw global launches of innovative brands for the treatment of HIV and chronic hepatitis C.

Scientific Advisory Board



Robert Malenka, MD, PhD

Chairman of the Scientific Advisory Board, Nancy Friend Pitzer Professor in Psychiatry and Behavioral Sciences at Stanford University. Dr. Malenka is an elected member of the National Academy of Sciences and the National Academy of Medicine as well as an elected fellow of the American Academy of Arts and Sciences, the American Association for the Advancement of Science, and the American College of Neuropsychopharmacology. He has served on the National Advisory Council on Drug Abuse and as a Counselor for the Society for Neuroscience and the American College of Neuropsychopharmacology. He is known for his landmark contributions to understanding of brain plasticity mechanisms, and has extensive experience as an advisor to various pharmaceutical and biotechnology companies.



Maria A Oquendo, MD, PhD

Ruth Maltzer Professor and Chairman of Psychiatry at University of Pennsylvania, Psychiatrist-in-Chief at the Hospital of the University of Pennsylvania. Dr. Oquendo is a member of the National Academy of Medicine, one of the highest honors in medicine. She is Past President of the American Psychiatric Association (APA), the International Academy of Suicide Research and the American College of Neuropsychopharmacology (ACNP). She is President of the American Foundation for Suicide Prevention's Board of Directors, Vice President of the College of International Neuropsychopharmacology and has served on the National Institute of Mental Health's Advisory Council. Dr. Oquendo is a member of Tufts University's Board of Trustees, serves on its Executive Committee and chairs Tufts' Academic Affairs Committee.



Maurizio Fava, MD

Psychiatrist-in-Chief of the Massachusetts General Hospital (MGH), director, Division of Clinical Research of the MGH Research Institute, executive director of the Clinical Trials Network and Institute, (MGH), associate dean for clinical and translational research and the Slater Family Professor of Psychiatry at Harvard Medical School. Dr. Fava is a world leader in the field of depression. He has edited eight books and authored or co-authored more than 800 original articles published in medical journals with international circulation, articles which have been cited more than 80,000 times in the literature and with an h index of over 140. Dr. Fava is a world leader in the field of depression. He has edited eight books and authored or co-authored more than 800 original articles published in medical journals with international circulation, articles which have been cited more than 80,000 times in the literature and with an h index of over 140.



Robert H Dworkin, PhD

Professor of Anesthesiology and Perioperative Medicine, Neurology, and Psychiatry, and Professor in the Center for Health + Technology, at the University of Rochester School of Medicine and Dentistry. Dr. Dworkin has spent over 25 years conducting clinical research on pain. He is also Director of the Analgesic, Anesthetic, and Addiction Clinical Trial Translations, Innovations, Opportunities, and Networks (ACTTION) public-private partnership with the U.S. Food and Drug Administration (FDA). Dr. Dworkin received the American Pain Society's Wilbert E. Fordyce Clinical Investigator Award in 2005 and John and Emma Bonica Public Service Award in 2014, the American Academy of Neurology's Mitchell B. Max Award for Neuropathic Pain in 2015, and the International Association for the Study of Pain's John D. Loeser Award in 2020.



Peter Bergethon, MD

VP and Head of Quantitative & Clinical Technologies, Biogen, Inc., where he leads the effort to transform clinical trials and humanize drug discovery by encouraging the transition of clinical trial measures from a qualitative to a quantitative discipline. The Quantitative Medicine transformation has advanced Biogen's leadership in neuroscience therapeutics and personalized medicine. Dr. Bergethon came to Biogen in 2017 from Pfizer Worldwide Research and Development where he was Vice President and Head of the Pfizer Innovation Research Lab within the Early Clinical Development group. Before joining the biopharmaceutical industry in 2012, Dr. Bergethon spent 20 years in academic medicine as a Professor at Boston University and Tufts University in the Departments of Biochemistry, Neurology, Neurobiology & Anatomy, and Biomedical Engineering.

