



ASX:MEB

INVESTOR PRESENTATION

MAY 2022

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New US-based IR Firm Appointed



Medibio Limited is pleased to announce that it has retained the United States based investor relations firm, RB Milestone Group LLC ("RBMG"), for an initial period of twelve months which started in February 2022. RBMG will be providing corporate communications advisory, non-deal roadshow advisory, market intelligence advisory, and business referrals to the management team of Medibio.

Medibio's Founder and Managing Director, Claude Solitario, stated:

"Medibio remains focused on advancing towards becoming one of the first companies with a regulated screening technology for mental health. RBMG has an excellent track record of targeting the right investor groups. We look forward to working with RBMG which will be leveraging its long-standing relationships to ensure that our narrative is communicated effectively to a US audience that is very receptive to mental health solutions, particularly considering the effects of COVID on mental health."

RBMG's Managing Director, Trevor Brucato, stated:

"We are very proud and protective of our track record with our clients globally. RBMG partners with companies that meet our criteria and appear to be set up for long-term success. My team and I look forward to working with the Medibio team and introducing the Company and its disruptive screening technology for mental health to a wider audience within the investment community throughout the United States."



medibio

Looking at Mental Health Objectively

Medibio is a mental health technology company pioneering the use of artificial intelligence, deep-learning algorithms and neural network methodology to identify biological markers and patterns to aid in early detection and screening of mental health conditions.



CHANGING SUBJECTIVITY TO OBJECTIVITY

THE PROBLEM

Current diagnostic tests rely on clinical interviews with subjective interpretation



OUR SOLUTION

Identify biomarkers and patterns based on the patient's own biological data to objectively aid in the early detection and screening of mental health conditions



WHY INVEST IN MEDIBIO?



FDA Program Underway

If successful, Medibio's solution will be one of the first objective screening test for depression approved by the FDA (De Novo Application)



Targeting a Large Market

In 2019, national spending on mental health services totaled \$225 billion and accounted for 5.5% of all health spending* 26% of the US population suffer from a mental health condition in a given year.



Patent-Protected, Innovative Technology

Assessing mental state using overnight heart rate
Cost effective to administer

Scalable



Early Revenue Potential

Early revenue opportunity in the non-clinical market with Medibio's **LUCA** and **ilumen** mobile apps



BUSINESS UNITS

Significant revenue potential in each market

CLINICAL



MEB-001 Sleep Clinics

Primary care physicians and other specialists referring patients to sleep clinics for sleep disturbance and depression screening



MEB-002 Homes

Depression Screening in Home Sleep Environment

NON-CLINICAL











FOR CORPORATES



CLINICAL

MEB-001
Depression Testing in Sleep Clinics

STATUS: FINAL TRIAL TO VALIDATE MEB-001

The FDA, during the pre-submission meeting on 2nd February 2022 and subsequent discussions, has been very supportive of Medibio's objectives and complimentary of the Company's MEB-001 validation trial design.

The FDA has acknowledged that MEB-001 is novel and will likely pave the way for other research and products.

Medibio and the FDA are currently in communication to agree on end-points that remained outstanding within the 60 minutes allocated for the meeting.

In anticipation of agreement during March, Medibio has 17 sleep clinics in various stages of preparedness to launch the validation trial.



Medibio's vision is that everyone referred for a sleep study will also be simultaneously screened for a Major Depressive Episode using MEB-001.



CLINICAL

MEB-002
Depression Testing in the Home

More and more sleep studies are being performed in the patient's own home as opposed to in sleep clinics due to technological advancements, reduced cost and convenience.

In recognition of the growing patient cohort moving from in-clinic sleep studies to the home environment, **planning has now begun for the development of MEB-002**, which will screen for depression in patients undertaking a sleep study in the home environment.

MEB-002 algorithms will build on the success of MEB-001





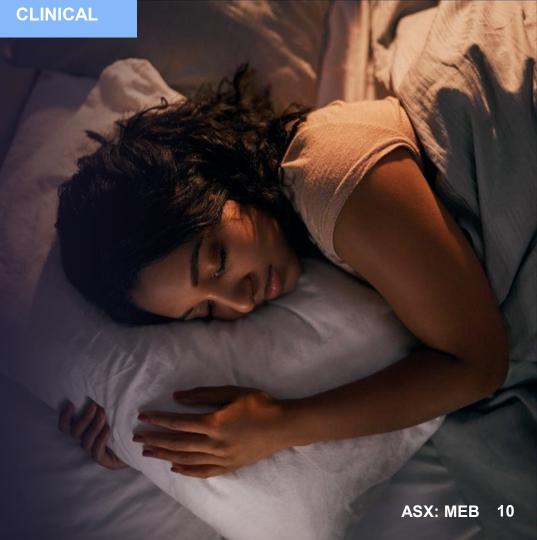
MEB-001 CLINICAL TRIAL PARTNER



MedBridge is the leading provider of sleep laboratory management services in the United States, operating 140 sleep disorder diagnostic centres.

"The objective identification of depression in patients that suffer certain sleep disorders is an <u>unmet need</u> and we are pleased to be working with Medibio to fast-track patient recruitment in the validation trial."

John Mathias - Chief Development Officer of MedBridge





NON-CLINICAL

CONSUMER HEALTH LUCA Mobile App



Biometric-driven stress monitoring with psychologistdeveloped tools to help users understand and track stress to avoid more serious conditions.





BIOMETRICS

Tracking three components of physiological stress: Cardiac (HR), Activity, and Sleep along with an Overall Stress score





PERCEIVED STRESS ASSESSMENT

A short series of questions to determine a user's level of perceived stress



ODYSSEY

Tools and exercises to confront negative thinking patterns and identify stressors

NON-CLINICAL

LUCA + NORTHERN MICHIGAN UNIVERSITY (NMU) TRIAL PARTNER

- Partnership: Medibio and The Department of Neuroscience at NMU
- Purpose: Raise awareness of the LUCA app prevention tool amongst the student body to potentially help students avoid getting into the red zone and/or taking more drastic options such as dialing 911
- **Trial:** Medibio will engage the student body through a call-to-action event with impactful head-line spokespersons.
 - Based upon the success of this program, Medibio will look to scale this program across the US - a potential market of 20 million students and 4,000 schools and universities.
 - Medibio will collaborate with commercial partners and wearable device companies to participate in this worthwhile initiative.



Principal Investigators:

- Professor Adam Prus (Distinguished Professor & Head of Psychological Science NMU)
- Professor Joshua Carlson
- · Assistant Professor Dr. Lin Fang



NON-CLINICAL

CORPORATE HEALTH ilumen Mobile App



Provides the employer a dashboard of de-identified, aggregated data to better manage the mental wellbeing of their most important asset......its workforce



CLOSES THE LOOP

Employers can now assess the effectiveness of employee wellness programs



REVENUE MODELS



MEB-001 & MEB-002

Licensing + Royalty Revenue



Consumer Health App

Free download + paid subscription model



Corporate Health App

Enterprise license based on a corporation's total workforce



LOOKING FORWARD 2022 DEVELOPMENT PROGRAM

		2021	H1 2022	H2 2022
CLINICAL	MEB-001 Depression Screening in Sleep Clinics	Development ✓ Sleep Algorithms ✓ Heart Rate Variability Algorithms ✓ Depression Analysis Algorithms	Feasibility Trials	Complete FDA Validation Trial and submit the De Novo application.
	MEB-002 Depression Screening in the Home	✓ Algorithm development✓ Trial planning and design	Feasibility Trials Trial will compare wearable devices to FDA approved medical devices	
NON-CLINICAL	ilumen App Corporate Market	✓ Launched in Australia	Market Feasibility	Potential Launch in USA
	LUCA App Consumer Market	✓ App development✓ Soft launch in October 2021	Planning for launch in college marketPilot study for launch in college market	Expand college market initiative to an initial 3 to 4 colleges across USA



THE TEAM

EXECUTIVE MANAGEMENT



CLAUDE SOLITARIO Managing Director & Founder



ARCHIE DEFILLO
Chief Medical Officer



MASSIMILIANO GRASSI Head of Artificial Intelligence



ZORIAN FEDORYGAHead of Software Development



MARK WALINSKI Business Development (North America)

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