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

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER PURSUANT TO RULE 13a-16 OR 15d-16 UNDER THE SECURITIES EXCHANGE ACT OF 1934

For the month of November 2025

Commission File Number: 001-35165

BRAINSWAY LTD.

(Translation of registrant's name into English)

16 Hartum Street RAD Tower, 14th Floor Har HaHotzvim Jerusalem, 9777516, Israel (+972-2) 582-4030

(Address and telephone number of Registrant's principal executive office)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F. Form 20-F [X] Form 40-F []

This Form 6-K is incorporated by reference into the Company's Registration Statement on Form S-8 filed with the Securities and Exchange Commission on April 22, 2019 (Registration No. 333-230979) and the Company's Registration Statements on Form F-3 filed with the Securities and Exchange Commission on July 22, 2024 (Registration No. 333-280934) and on April 22, 2025 (Registration No. 333-286672).

EXHIBIT INDEX

Exhibit Title

99.1 <u>Investor Deck</u>

SIGNATURES

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

BRAINSWAY LTD.
(Registrant)

Date: November 12, 2025

/s/ Hadar Levy
Hadar Levy
Chief Executive Officer



Advanced Noninvasive Neurostimulation Treatments for Mental Health Disorders

Investor Deck November 2025

Nasdaq/TASE: BWAY



Safe Harbor and Non-GAAP/IFRS Financial Measures

This presentation does not constitute an offer or invitation to sell or issue, or any solicitation of an offer to subscribe for or acquire, any securities of the Company, nor to participate in any investment. This presentation shall not constitute advertising or be construed as commercial or promotional in nature. No representation or warranty is made as to the accuracy or completeness of this presentation. You must make your own investigation and assessment of the matters contained herein. In particular, no responsibility is assumed as to the achievement or reasonableness of any forecasts, estimates, or statements as to prospects contained or referred to in this presentation.

This presentation contains information that includes or is based on forward-looking statements within the meaning of the federal securities laws. Such statements are not guarantees of future performance. They reflect current expectations and are subject to risks and uncertainties that could cause actual results to differ materially from those expressed or implied. Factors include, but are not limited to: continued business impact from COVID-19; adverse economic conditions; changes in demand and pricing; manufacturing difficulties or delays; legislative and regulatory actions; changes in reimbursement; third-party payor decisions; product liability claims; U.S. healthcare reform; financial market conditions; competitive dynamics; failure to achieve sufficient market adoption; regulatory actions or delays; and our ability to realize expected operational and manufacturing efficiencies. Additional information about these and other risks is contained in our filings with the U.S. Securities and Exchange Commission. If one or more of these factors materialize, or if underlying assumptions prove incorrect, actual results may differ materially. This presentation also includes market and industry data from government and private publications and internal estimates and projections based on a number of assumptions. We undertake no obligation to update any forward-looking statement as a result of new information, future events, or otherwise.

Certain non-GAAP/IFRS financial measures are included in this presentation. These measures are presented to complement the financial information prepared in accordance with IFRS because management believes they are useful to investors. For example, Adjusted EBITDA, a non-IFRS measure, is widely used by investors and securities analysts to evaluate a company's operating performance without regard to one-time items (such as restructuring and litigation expenses) that can vary substantially from company to company. Management also uses Adjusted EBITDA with IFRS measures for planning purposes, including preparation of annual operating budgets, as a measure of operating performance. These non-GAAP/IFRS measures should be considered only as supplements to, and not superior to, financial measures prepared in accordance with IFRS. Other companies may calculate similarly titled non-GAAP/IFRS measures differently than the Company.



Investor Deck | November 2025

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BrainsWay: Global Leader in Transcranial Magnetic Stimulation (TMS)

Installed base of >1,600 systems

> 7M individual treatment

4 FDA-cleared indications

2013 - Major Depressive Disorder (MDD)

2018 - Obsessive-Compulsive Disorder (OCD)

2020 - Smoking Addiction

2021- Anxious Depression

September 16, 2025: FDA Clearance for Accelerated Deep TMS Protocol for MDD

Established reimbursement





Q3 2025 Highlights

Strong Growth & Profitability

• Revenue: \$13.5 M (+29% YoY)

· Gross Margin: 75%

Adjusted EBITDA *: \$2.0 M (+81%)

Net Profit: \$1.6 M (+137%)

• Cash: ~\$71 M as of Sept 30, 2025

"We're driving record adoption and sustained profitability, building longterm value through clinical innovation, strategic investments, and durable recurring revenue."

Momentum Across the Business

· Remaining Performance Obligations: \$65 M (up 37% YoY)

• Shipments: 90 Deep TMS™ systems (+43%), installed base >1,600

~70% of new engagements as multi-year leases

• FDA Clearance: Accelerated protocol for Major Depressive Disorder (MDD)

 NIH Grant: \$2.5 M for Stanford/PAVIR study on Deep TMS for Alcohol Use Disorder

· Strategic Investments: 4 minority stakes



*Adjusted EBITDA is a non-IFRS measure. See slide 33 for operating income results, the closest IFRS measure, and the reconciliation table in the Company's earnings release



Going Forward, All About Execution



Strong Profitability

75% gross margins (Q3 25) +81% adjusted EBITDA growth * (Q3 25 vs. Q2 24) \$7 M
adjusted EBITDA *
(Q3 25 annualized)

10
consecutive quarters of positive free cash flow

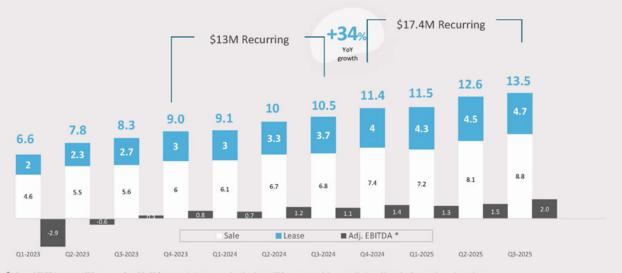




*Adjusted EBITDA is a non-IFRS measure. See slide 33 for operating income results, the closest IFRS measure, and the reconciliation table in the Company's earnings release.

The ARR Model

Recurring revenue growth (+34% YoY)
Exceeds Non-Recurring Revenue growth (+23% YoY)





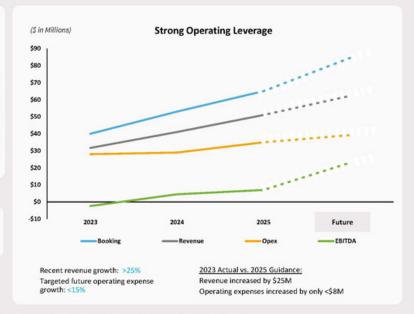
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Focused on ARR and Operating Leverage

Revenue Mix Shift Towards ARR

	2023	2024	2025 (est.)	% Growth
\$ Millions				
Recurring Revenue	\$10.0	\$14.0	\$18.4	+31%
Non-Recurring Revenue	\$21.7	\$27.0	\$33.7	+25%

Recurring Revenue Growth Exceeds Non-Recurring Revenue Growth





*Adjusted EBITDA is a non-IFRS measure. See slide 33 for operating income results, the closest IFRS measure, and the reconciliation table in the Company's earnings release.

------ Business Model - Leasing

We transformed to a winning business model

From:

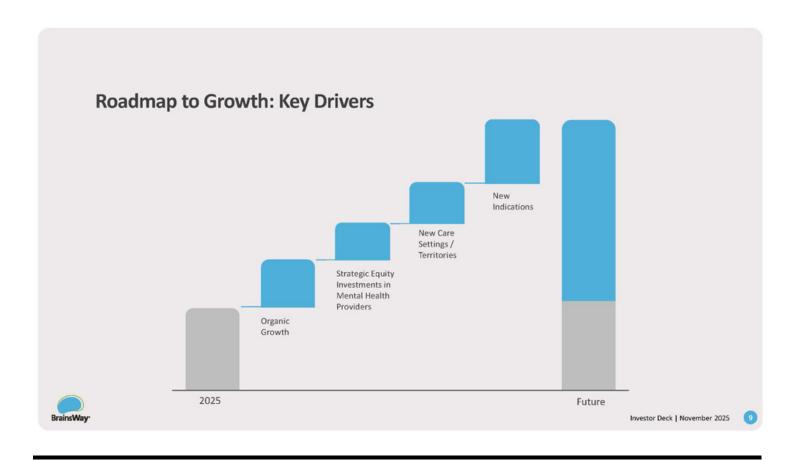
One time sale



To:

A high-margin ARR model





Fix Lease Model & Pay Per Use Driving \$65M RPO – U.S. Only

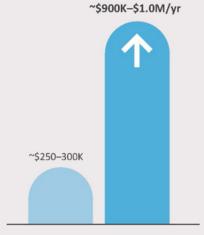
Removes barriers, boosts adoption

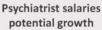
Reduces upfront capital commitment for physicians

Clinics pay only when they treat: faster adoption & utilization

Provides 4-5 years contracts

Built for today's market: solves high-rate CapEx hesitation and staffing constraints

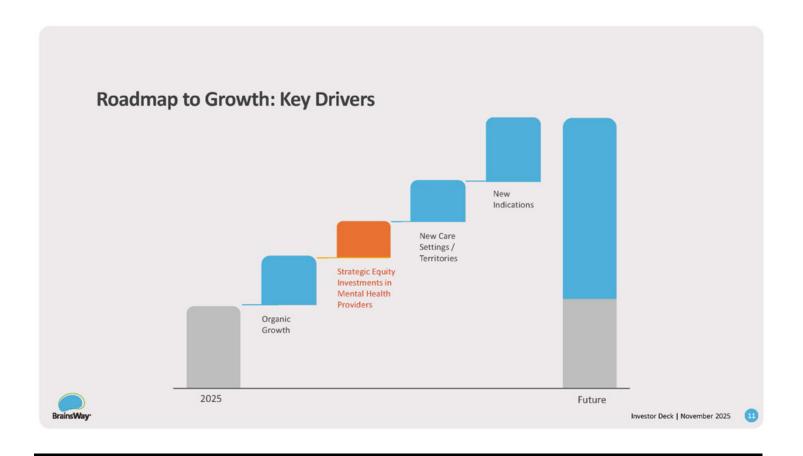












Strategic Partnership With Valor Equity Partners

Valor, a leading \$20B equity firm behind market leaders, is now BrainsWay's largest investor and an active partner in our scale strategy





















Strategic Equity Investments in Mental Health Providers

New Strategy to Accelerate Company Growth



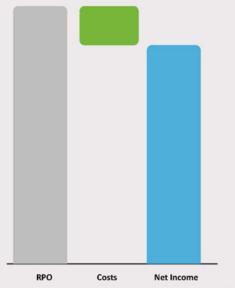
Strategic Initiative Led by Valor Fund 200+

Identified Mental Health Clinics as Potential Partners

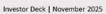
No Added OPEX

Each deal can potentially add 10-15 Clinics Annually

Each deal can deliver Significant incremental gross margin









Access to Data



Access to Patients



Best-in-class Infrastructure for Expansion

Deals signed in 2025:

\$5M investment



20+ dinics
 30,000 patients treated to date

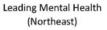
How do we partner?



- \$2.3M in initial investment
 \$2.5
- Additional \$1M milestonebased investment
- Clinics in Colorado



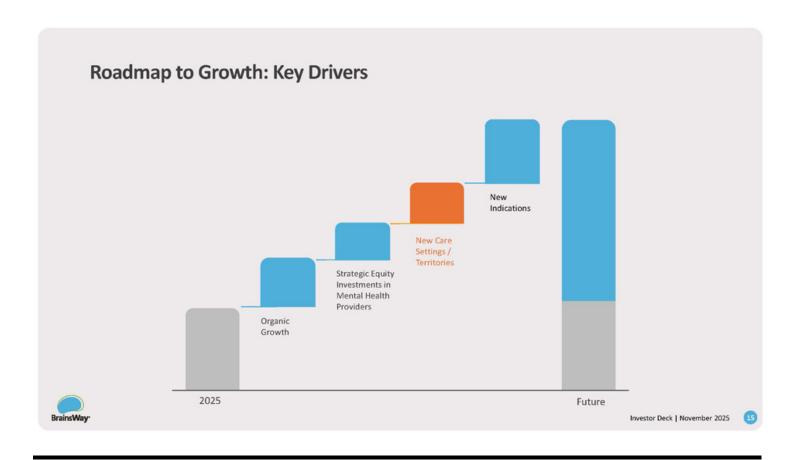
- \$2.5M in initial investment
- Additional \$1.5M milestonebased investment
- · Clinics in Texas



· Clinics in NY, Boston, Florida, PA

?





Extending to Home-Use Neuromodulation

August 21, 2025: Announced Structured M&A Option





Explore extending deep TMS treatments with Athome* therapy - start in clinic, continue at home





Structured deal: \$5M convertible debt; milestone tranches; call option to acquire





Clean P&L: no consolidation, no added OPEX





Bigger reach: taps \sim 50% telemedicine activity, clinic-prescribed home device grows TAM without extra marketing and feeds in-clinic use. Patients can go directly to their primary care physician (PCP)



*Neurolief's at-home therapy is not yet FDA-cleared

Investor Deck | September 2025





Upcoming Planned Milestones and Catalysts*

Near-Term

12-36

Mid-Term

Long-Term



- FDA accelerated protocol clearance
- New AUD trial launch
- FDA adolescent MDD clearance
- First 5 enterprise deployments
- FDA Comorbid PTSD application
- Neurolief Home use device launch**
- Full enterprise rollout (10+ accounts)
- Broader coil adoption
- New payer coverage wins

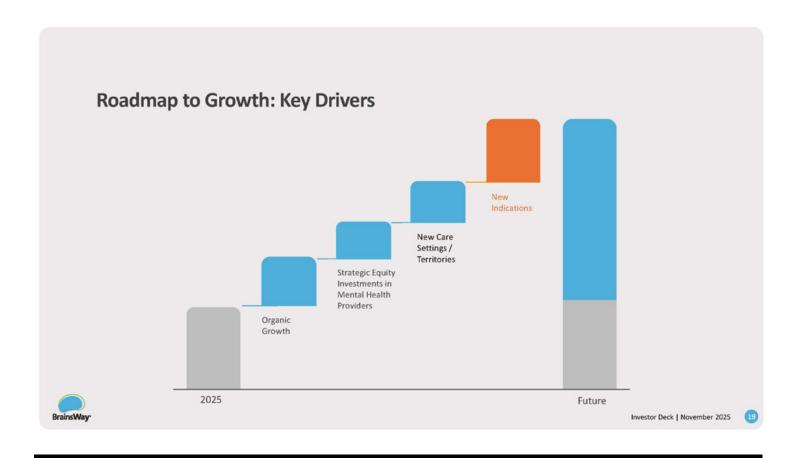
- Neurology portfolio
- BrainsWay 360 global launch
- Selective global expansion



*Milestones on this slide are forward looking in nature. See cover slide.

**Neurolief's at-home therapy is not yet FDA-cleared



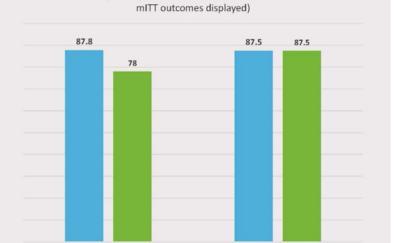


From 6 weeks



To 6 days*

- Accelerated Deep TMS protocol:
 1,800 iTBS pulses at 110% of MT using the H1 coil
- 30 total sessions
- Flexible scheduling days can be nonconsecutive



Response (mITT, PP) Remission (PP)

Accel H1

Accelerated Trial Outcomes (n=104 patients enrolled, 89 completers



*Acute phase, before required maintenance sessions

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Standard H1

Smoking Addiction

28.3_M



U.S Adult Smokers

\$2B/year
nearly spent by smokers on Quitting



85% of them don't succeed

~1 out of 3

patients quit for 4 weeks

~67%

of 6-week completers remained non-smokers for +3 months

1st and Only TMS Addiction Clearance

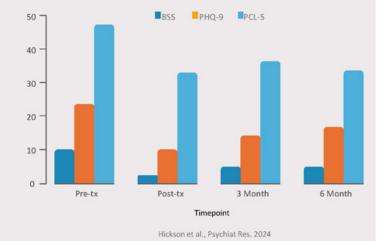
With Brainsway:



PTSD New Article (2024)*

Real-world evidence showing significant reductions in comorbid PTSD symptoms in MDD patients for up to 6 months after treatment

- · 99 patients in VA clinic
- · H1-coil, FDA-approved depression protocol
- · 3-week taper (maintenance)
- Significant reductions in PTSD symptoms, depression, and suicidal thoughts observed immediately after treatment, at 3 months, and at 6 months
- · Impressive outcomes of response and remission





*Deep TMS is not FDA-cleared for PTSD. Data collection efforts underway.



Upside Potential: Deep TMS 360™ Research*

Overview

- Potential Novel solution being explored for shorter treatment, better efficacy, and possible new indications
 - Enables activation of greater numbers of neurons in the brain than currently available forms of TMS

May be uniquely suited for older adults with neurodegenerative conditions and reduced neuroplasticity

Today's TMS: Single coil

- · Neurons aligned parallel to coil's electrical field are more likely to be stimulated
- · As a result, only a fraction of neurons in the targeted brain region are actually impacted
- . Deep TMS 360: Rotational Field system has 2 perpendicular orthogonal TMS coils
- 2 coils are operated with short time lag (milliseconds) in order to induce a circularly rotating electrical field
- · Results in uniform stimulation of neurons oriented across a wide range of directions

Deep TMS 360 Clinical Plans

- · Launched study on post-stroke rehabilitation in 1H 2024
- Launching alcohol use disorder trial in Q3 2025
- Plans to initiate feasibility studies on various neurology indications (dementia, Parkinsons, Alzheimer's) in [2H 2025]

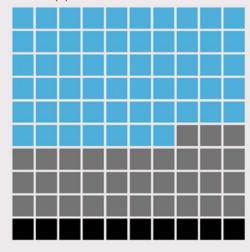


*Deep TMS 360 is being investigated for clinical trial research and is not commercially available. This slide is forward-looking only.



Expanding TAM in a Massive Market

U.S Patient population



Major Depressive Disorder (MDD) &
Anxious Depression,
OCD,
Smoking Addiction

FDA cleared indications

29м

Alcohol Use Disorder (AUD)

Future Pipeline Indications



Alzheimer Post-Stroke



BrainsWay: Note: Other than MDD/Anxious Depression, OCD, and Smoking Addiction, the above indications are currently investigational, not available in the U.S, and not cleared by the FDA.

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2

35 Years of Established Technology

With Demonstrated Safety and Efficacy

How it works:

- · An electromagnetic H-coil is placed on the scalp
- · A rapid current flows through the coil
- This generates a magnetic field that induces an electric field in the brain.
- The changing electric field depolarizes axons and triggers action potentials
- ~2.5× deeper penetration than standard TMS
- · Deeper, broader stimulation activates more neurons





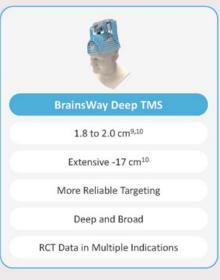






BrainsWay Deep TMS vs. Traditional TMS

Clear, compelling technological advantage



Features
Depth
Brain Volume Stimulated
Therapy Delivery
Structures Treated
Technology Platform



RCT Data Limited to Depression



All competitors in the commercial depression space use variations of the traditional "figure-8" TMS coil design Technological distinctions do not necessarily correlate with clinical outcomes

Depression Clinical Efficacy: Demonstrated Safety and Efficacy

After 30 sessions

81.6 %

65.3 %



Response

- Durability in TMS is >1 Year in ~50% of Responders
- A published abstract of 200 patients from a single BrainsWay site showed average durability of 860 days
- Remission
- ~2 in 3 patients achieved remission with Deep TMS
- 1,351 MDD patients who received at least 20 Deep TMS sessions
- · No systemic side effects



OCD Clinical Efficacy After 29 sessions

57.9 %

87.0 %



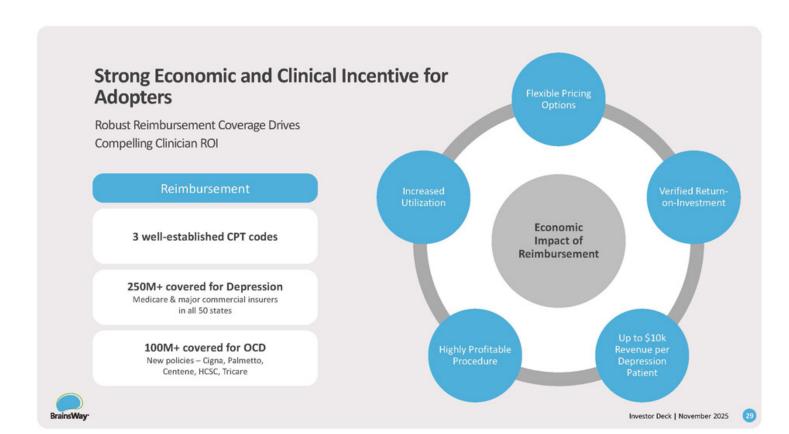
Response

- 219 patients across 22 centers
- Sustained response achieved in ~20 sessions
- No systemic side effects

Durability of 1+ Years

- n = 60 patients (from pivotal and post-market studies)
- Durability = time from end of Deep TMS course to change in ongoing treatment





Most Extensive and Broadest TMS Intellectual Property

Encompassing Core Technology and Applications

Patent Portfolio

30+ US 50+ OUS

Issued Patents or Allowed Applications

Key Portfolio Coverage Areas

Deep TMSTM Coils

Multi-Channel TMS

Rotational Field Deep TMS

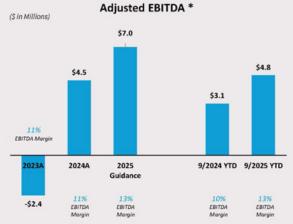
Closed Loop TMS/EEG





Strong Financial Momentum.







*Adjusted EBITDA is a non-IFRS measure. See slide 33 for operating income results, the closest IFRS measure, and the reconciliation table in the Company's earnings release.

YTD 2025 VS. YTD 2024

	Actual	Actual	Variance	
USD in thousands	YTD 2025	YTD 2024	s	%
Revenues	37,680	29,602	8,078	27%
Cost of Revenues	9,412	7,532	1,880	25%
Gross profit	28,268	22,070	6,198	28%
Gross Margin	75.02%	74.56%		
Research and Development expenses	7,072	5,146	1,926	37%
Selling and Marketing expenses	13,831	11,731	2,100	18%
General and Administrative expenses	4,958	4,233	725	17%
Total Operating expenses	25,861	21,110	4,751	23%
Operating Profit	2,407	960	1,447	151%
% Operating Profit	6%	3%		
Finance Income, net	2,762	763	1,999	262%
Income Tax expense	469	350	119	34%
Net Income	4,700	1,373	3,327	242%
Adjusted EBITDA	4,767	3,080	1,687	55%
% EBITDA	13%	10%		
Basic Net Income Per Share	\$ 0.12	\$ 0.04	0.08	211%
Diluted Net Income Per Share	\$ 0.11	\$ 0.04	0.07	175%



Q3 2025 Balance Sheet

USD in thousands	As of September 30,	As of June 30,	As of December 31,	Change vs. 06/25		Change vs. 12/24	
	2025	2025	2024	s	%	s	%
ASSETS	- Table 1		-				
Cash and cash equivalents	70,458	67,912	69,345	2,546	4%	1,113	2%
Restricted cash	251	251	271	-	0%	(20)	-7%
Short-term deposits		10,087	-	(10,087)	-100%		100%
Trade receivables, net	5,129	3,871	4,596	1,258	32%	533	12%
Inventory	4,418	4,190	4,426	228	5%	(8)	0%
Other current financial assets	1,079	949	-	130	14%	1,079	#DIV/0!
Other current assets	1,598	1,771	1,032	(173)	-10%	566	55%
Total Current Assets	82,933	89,031	79,670	(6,098)	-7%	3,263	4%
Non-Current Assets	30,129	22,533	14,646	7,596	34%	15,483	106%
Total Assets	113,062	111,564	94,316	1,498	1%	18,746	20%
LIABILITIES AND EQUITY							
Current Liabilities	25,993	25,508	15,346	485	2%	10,647	69%
Non-Current Liabilities	17,004	17,924	16,657	(920)	-5%	347	2%
Equity	70,065	68,132	62,313	1,933	3%	7,752	12%
Total Liabilities and Equity	113,062	111,564	94,316	1,498	1%	18,746	20%



Successful, Experienced Medical Device Professionals - Decades of Results



Hadar Levy Chief Executive Officer 25+ Years Med Device



Ido Marom Chief Financial Officer 20+ Years Finance



Dr. Gilead Moiseyev Chief Technology Officer 20+ Years Med Device Dev



Dr. Richard Bermudes Chief Medical Officer 15+ Years TMS Leadership



Michael Cohen Vice President - US Sales 15+ Years Med Device Sales



Moria Ben Soussan Vice President - R&D 15+ Years Med Device Dev



Dor HagaiVice President – Operations
10+ Years Supply Chain & Ops



Dr. Colleen Hanlon Vice President - Medical Affairs 15+ Years Brain Stim Research



Summary Highlights



Category Leader - Only Deep TMS platform with 4 FDA cleared indications



Expanding TAM - New indications, Neurolief at-home entry, international scale



Recurring Revenue Model – ~70% new deals are multi-year rentals



High Visibility - \$65M Remaining Performance Obligations (RPO)



Strong Profitability – 10 straight quarters of positive free cash flow



Backed to Scale - \$71M cash; strategic partner Valor supports continued expansion



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