

ONMRD MEDICAL

Investor Webcast Q3 2024 Business Update

November 26, 2024



















Forward Looking Statements

This Presentation may include statements, including the Company's financial and operational medium-term objectives, that may be deemed to be "forward-looking statements". These forward-looking statements may be identified by the use of forward-looking terminology, including the terms "believes", "aims", "forecasts", "continues", "estimates", "plans", "projects", anticipates", "expects", "intends", "may", "or "or, in each case, their negative or other variations or comparable terminology, or by discussions of strategy, plans, objectives, goals, future events or intentions. Forward-looking statements may and often do differ materially from actual results. Any forward-looking statements reflect the Company's current view concerning future events and are subject to risks relating to future events and other risks, uncertainties, and assumptions relating to the Company's business, results of operations, financial position, liquidity, prospects, growth, or strategies. Forward-looking statements speak only as of the date they are made.

Speaking Today



Dave Marver CEO



Jocelyne Bloch, MD
Neurosurgeon, Co-founder and
Medical Advisor



Amori Fraser Finance Director



Grégoire Courtine, PhDNeuroscientist, Co-founder and Scientific Advisor

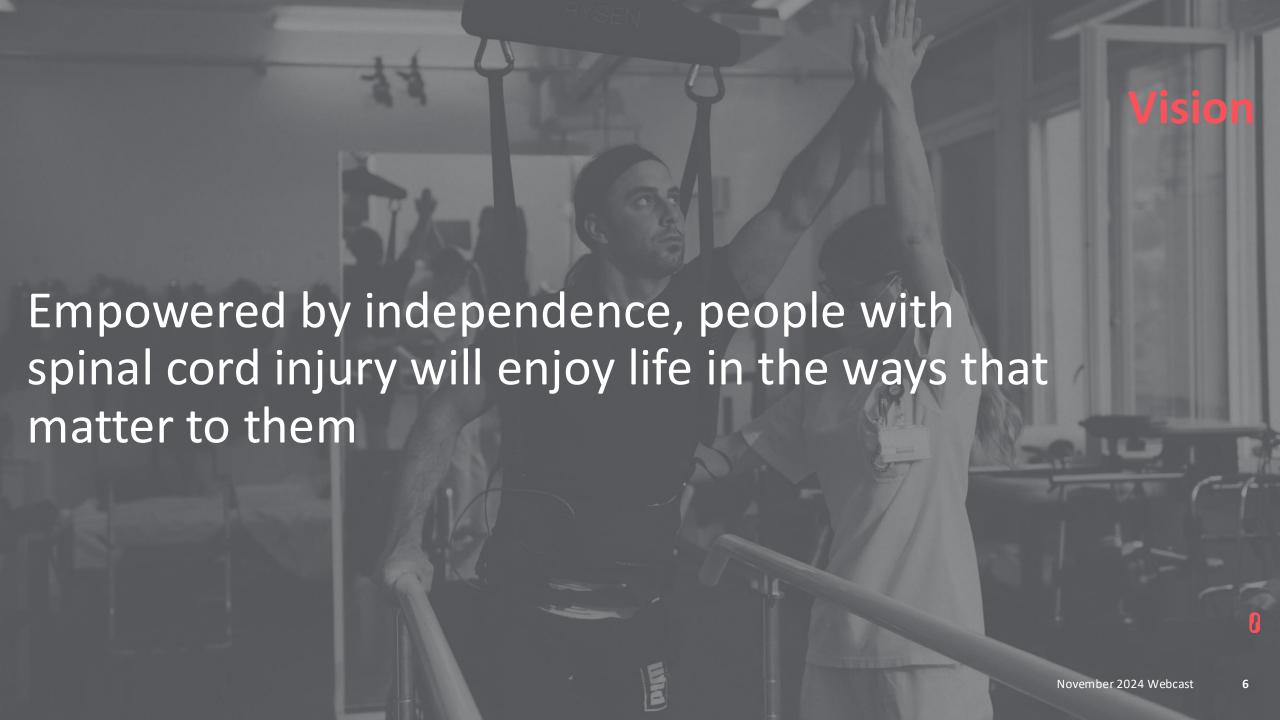


ONWARD® Medical at a Glance

Key Facts

- o Founded in 2015
- o ~100 FTEs
- HQ in the Netherlands
- Science and Engineering Center in Switzerland
- Growing US presence
- IPO 2021; listed on Euronext Brussels, Amsterdam, and Paris
- Research coverage from Stifel, Bryan, Garnier & Co, KBC Securities, Degroof Petercam and Kepler Cheuvreux

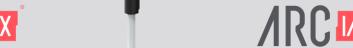
- Technology: 3 purpose-built neuromodulation platforms
 - ARC^{EX®} delivers ARC Therapy[™] externally through the skin
 - o ARC^{IM®} delivers ARC Therapy via a fully implanted system
 - ARC^{BCI™} pairs ARC^{IM} with an implanted brain-computer interface to restore thought-driven movement via our wireless DigitalBridge[™]
 - Innovation: 10 FDA Breakthrough Device Designations; 270+ issued patents¹
 - **Clinical Success:**
 - Safety and effectiveness of ARC^{EX} Therapy for upper limb mobility demonstrated in Up-LIFT clinical trial;
 results published in *Nature Medicine*, May 2024
 - Positive interim results for ARC^{IM} Therapy to improve blood pressure regulation
 - Market Opportunity: \$20B+ / €19B+ total addressable market with limited competition
 - Commercialization: FDA authorization and US launch anticipated 2H 2024



Targeted, programmed electrical stimulation of the spinal cord to restore movement, function, and independence in people with spinal cord injury



Note: Investigational devices, not available for commercial use.



















Reach commercial stage by year-end, then expand labeling and platforms

Company Focus

Short Term 2024

Medium Term 2026/2027

Long Term 2026/2027+

Commercialize external platform (ARCEX)

First indication: Upper Limb

Population: SCI

Generate revenue and develop market for ARCIM



Commercialize implantable platform (ARCIM)

First indication: Blood Pressure

Population: SCI

Enter traditional medtech NASDAQ IPO/M&A window



Expand labeling and platforms

New indications and platforms to be assessed

such as BCI

Populations: SCI, Parkinson's, Stroke





Note: Investigational devices, not available for commercial use; SCI = Spinal Cord Injury.

9 indications under clinical or pre-clinical evaluation, with 3 in current roadmap

Current Pipeline

Short and medium term focus Funded primarily through grants and research partners Clinical Platform Indication FDA BDD¹ Pre-clinical Human PoC **Pivotal** Feasibility² **ARC**EX **Upper Limb** Study expected to start **ARCIM Blood Pressure** early 2025 **ARCIM** Mobility / Second Indication ARCEX Mobility **ARCIM** Parkinson's - Mobility **ARCIM** Bladder Human PoC expected in ARCBCI Mobility ARCBCI Upper Limb ARCDBS Mobility ✓ BDD¹ Granted O Current Roadmap Label Expansion O Platform Expansion

Note: The company may modify the pipeline based on clinical progress and marketplace considerations

¹ BDD = FDA Breakthrough Device Designation. ONWARD has been granted four additional BDDs for ARC^{EX} Bladder, ARC^{EX} Blood Pressure, ARC^{EX} Spasticity and ARC[™] Spasticity

² Includes both early feasibility (typically before device design finalized) and feasibility (near-final or final device design) studies

³ Funded by Christopher & Dana Reeve Foundation grant, conducted by EPFL

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Achievements since July include in-licensing BCI system, raising €50M and being awarded 2024 TIME **Magazine Best Invention**

Q3 Business Update

July

Science Announced publication highlighting evidencebased programming for ARC^{EX} Therapy July

Corporate Announced publication of annual sustainability summary for full year 2023

September

Clinical Awarded Christopher & Dana Reeve Foundation grant to further study BCI system

September

Clinical Announced third implant of BCI system to restore movement after SCI September

Corporate Listed shares on Euronext Paris, in addition to existing **Euronext Brussels and** Amsterdam listings

October

Development

Secured exclusive rights to CEA's WIMAGINE BCI technology

Details to follow

October

Corporate

Welcomed former Medtronic President Rob ten Hoedt as incoming Chairman of the Board

Details to follow

October

Corporate

Successfully raised €50M in upsized capital increase including investment from Ottobock

Details to follow

October

Science

ARC^{EX} System named one of TIME's Best Inventions of 2024

Details to follow

November

Clinical

Awarded EIC grant to study ARCBCI Therapy to address upper limb movement challenges after stroke

Secured exclusive rights to CEA's WIMAGINE BCI technology

BCI License Agreement



CEA's Clinatec is a world-renowned biomedical research institute and its BCI is ideal for our applications. We can now develop a truly integrated system that is well suited for the type of study required to gain regulatory approval and bring a BCI-enabled system to market

- Dave Marver, CEO at ONWARD Medical



The early clinical feasibility research demonstrates the remarkable potential of the ARC-BCI System to restore thought-driven movement and function after paralysis... We are pleased to partner with ONWARD Medical

- Guillaume Charvet, Head of the Neurotechnology Biomedical Research Unit at CEA

BCI Selection and Potential Impact



G Courtine, PhDNeuroscientist, Co-founder and Scientific Advisor



Jocelyne Bloch, MD
Neurosurgeon, Co-founder and
Medical Advisor

Exclusive rights to CEA's WIMAGINE BCI technology provides several opportunities

In-House BCI Value Add

License agreement signed with CEA for the WIMAGINE BCI system provides ONWARD Medical with the opportunity to:

- Be first to market with a BCI-enabled system to restore thought-driven movement after paralysis
- Take full control of technology that is currently the best fit with ONWARD's indications and develop an integrated system with ARC^{IM}
- Advance technology with existing and potential new grant funding opportunities

Welcomed incoming Chairman of the Board Rob ten Hoedt and Rahma Samow

Board Changes



Rob ten Hoedt

- Former Medtronic President and Executive Committee Member
- Over 30 years of experience in medical devices
- Successful track record in technology development, commercialization, and business-model innovation
- Former Chairman of MedTech Europe, the Association representing the medical technology industry in Europe.



Rahma Samow

- President and CEO of ClearChoice Dental Implant Centers, a fast-growing
 US-based leader in dental implant therapy
- Digital health and direct-to-consumer business leader
- Former member of Executive Management Board of Swiss-based
 Straumann Group and 15 years with Siemens Healthineers

Upsized capital increase with cornerstone investment from Ottobock

October 2024 Capital Raise

- Strategic investment from Ottobock, a global leader in prosthetics, orthotics and exoskeleton technology:
 - Now the company's largest shareholder (~10% of ONWARD shares)
 - Initiates strategic relationship
- Cash runway extended to two years or more
- High-quality long-only and sector specialist investors



Strategic investment from Ottobock, with opportunities for future collaboration

ottobock.

- Over 100 years serving people with movement disabilities; global leader in prosthetics, orthotics and exoskeleton technology
- Present in ~60 countries with 9000+ employees and 400+ patient care centers
- €~1.5B in revenues and €~280M adjusted EBITDA in 2023
- ONWARD's largest shareholder with ~10% of shares since October 2024 capital raise
- Opportunity to explore future development and commercial collaboration opportunities

Strategic Investment

ONWARD Medical has the potential to become a gamechanger in the therapy of spinal cord injuries with its innovative solutions... Our investment in ONWARD is an investment in the future of medical technology."

- Professor Hans Georg Näder

Owner & Chairman of the Board, Ottobock SE & Co. KGaA



Named one of TIME's Best Inventions of 2024

ARC^{EX} Recognition



'For most people with a spinal cord injury, there's some [brain signal] conduction making it through the block, but not enough for movement,' says Dave Marver, CEO of ONWARD. 'This device serves as an amplifier.'

TIME Magazine
The Best Inventions of 2024

ARCEX FDA clearance expected in Q4 2024

Financial Update

With addition of proceeds from October 2024 capital raise, current cash position expected to fuel operations for two years or more

Q3 2024 Cash Update

Burn €8.9M used during Q3 2024 (including transaction costs¹)

Q3 Ending Balance €23.2M net cash² as of 30 Sep 2024 Current Ending Balance €65.8M net cash² as of 22 Nov 2024



¹ Transaction costs related to recent debt financing; excludes transaction costs related to October 2024 capital raise

² Net cash is defined as the sum of cash and cash equivalents and fixed term deposits included in the current assets as included in consolidated statement of financial position in the Financial Statements.



Several important catalysts expected in the next 12 months

Upcoming Milestones and News Flow

ARCEX **ARC**^{EX} **ARC**EX ARC^{EX} First commercial sale (US) FDA clearance MDR submission CE mark Upper limb Upper limb Upper limb Upper limb **ARCIM ARCIM ARC**EX **ARCIM** First participant enrollment¹ Early IDE submission First commercial sale (EU) Interim results publication feasibility study Empower BP pivotal study Upper limb Blood pressure Parkinson's mobility Blood pressure

ARCIM IDE approval

Empower BP pivotal study Blood pressure

ARCIM

First participant enrollment Empower BP pivotal study Blood pressure

ARCIM

First-in-human² Bladder

ARCBCI

Additional implants³ Upper limb and Mobility

Note: All platforms and therapies are for investigational use only

¹ Funded by Michael J. Fox Foundation for Parkinson's Research grant

² Funded by Christopher & Dana Reeve Foundation grant

³ Funded by European Innovation Council and Christopher & Dana Reeve Foundation grants









CEA's WIMAGINE BCI enables our imperative to be first-to-market with BCI-augmented movement restoration

BCI Selection Considerations



Other BCI technologies



Safety CEA's WIMAGINE system is cleared for human research with ~7 years of human safety data; already restored movement

in three humans

Invasiveness WIMAGINE implant procedure is less invasive, offering lower

risk than subdural or brain penetrating electrodes

WIMAGINE provides sufficient resolution and coverage of

motor cortex to enable thought-driven movement restoration

A limited number of other BCI platforms cleared for investigation in humans; none have yet been used to restore movement of the human body

Use subdural, brain penetrating, or intravascular electrodes, bringing higher risk than the epidural WIMAGINE platform (e.g. inflammation, perforation, thrombosis)

May provide too little or too much resolution (overkill); may face difficulties offering sufficient access to or coverage of the motor cortex

ARC^{BCI} timeline

Resolution

First-in-human Mobility 2021 First-in-human Upper Limb 2023

Additional implants with first generation technology 2024+

Pivotal study

Commercialization



Potential window for next generation technology



