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Content

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Operational review

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Lifecare represents high-growth MedTech opportunity

Next-generation diabetes medtech

~ USD 5 billion addressable market

Capital efficient path to commercialization

Near-term value inflection points

2nd mover advantage, initial focus on Europe

Peak revenue potential of USD 1 billion +



Diabetes – an unresolved, growing global health crisis



Diabetes is a chronic **trillion-dollar health** care challenge



~600 million people living with diabetes, of which **110 million** need glucose monitoring



Diabetes caused **3.4 million deaths** in 2024 – 1 death every 6 seconds





CGMs have become the de facto standard for diabetes management – the future is "inject and forget"

The past



Pin prick blood glucose monitoring

The present



Body-worn CGMs

The future



Lifecare seamless CGMs – inject and forget



The future of CGM: what sets Lifecare apart

Existing CGM systems

Dexcom, Medtronic, Abbott



7 – 15 days

Body-worn patch

Glucose oxidase

Annual cost: \$2.300 - 6.000

Calibration: up to 2 times/day dep. on model

MARD <10%*: Yes

First implantable

Senseonics (Eversense)



365 days

Small capsule

Flourescence

Annual cost: \$6.000

Calibration: Yes

MARD <10%*: Yes

Lifecare current product

Lifecare



180 days

Small capsule

Osmotic pressure

Annual cost: ~ \$4.000 (assumption)

Calibration:

MARD <10%*: Yes

Lifecare - future

Lifecare



550 days

Grain of rice

Osmotic pressure

Annual cost: ~\$4.000 (assumption)

Calibration:

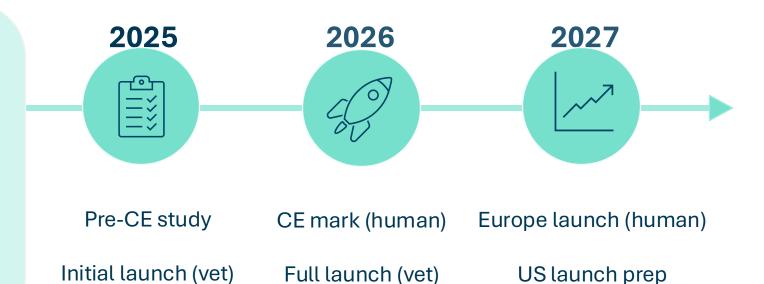
MARD <10%*: Yes



Focused, capital efficient path to commercialization

Key developments to date

- In-vitro tests confirm efficacy of miniaturized sensors
- Human study confirms accuracy with 9.6% MARD
- Biocompatibility and longevity study with CGM reference validation





Financing



Production



Partners



Highlights Q2 2025

Achieved design freeze for first-in-human trials and veterinary launch

Appointed Principal Investigator and trial sites for upcoming trial

Filed for first-in-human trials, paving the way for CE mark

Started preparations towards CE marking

Supported funding through warrant exercise



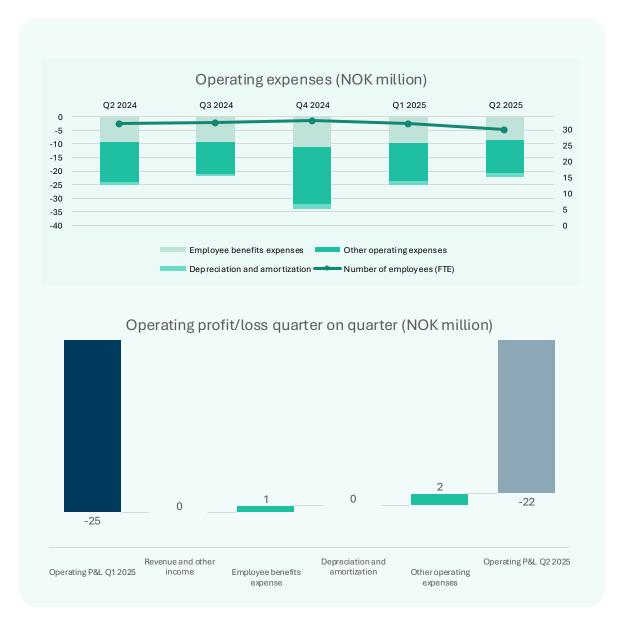




Profit & loss

Depreciation and amortization -1 Other operating expenses -12 Total operating expenses -22	260 -13 99			-19 089 -38 55 6
		2 -14 868	-26 253	-19 089
Depreciation and amortization -1	.20 .00			
	426 -1 50	2 -1 204	-2 928	-2 162
Employee benefits expense -8	540 -9 52	6 -9 056	-18 066	-17 306
Revenue and other income	7	4 6 439	12	6 850
Profit & loss (NOK 1000) Q2 2	025 Q1 202	5 Q2 2024	YTD 2025	YTD 2024

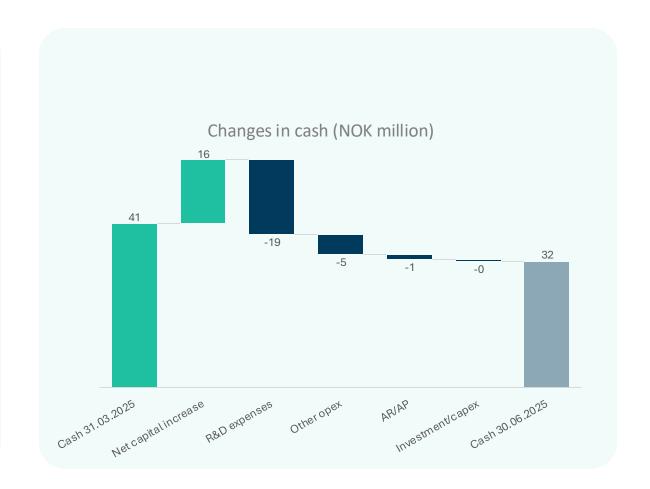
Operating expenses reflect focused investment in design freeze, trial preparations and CE-mark readiness





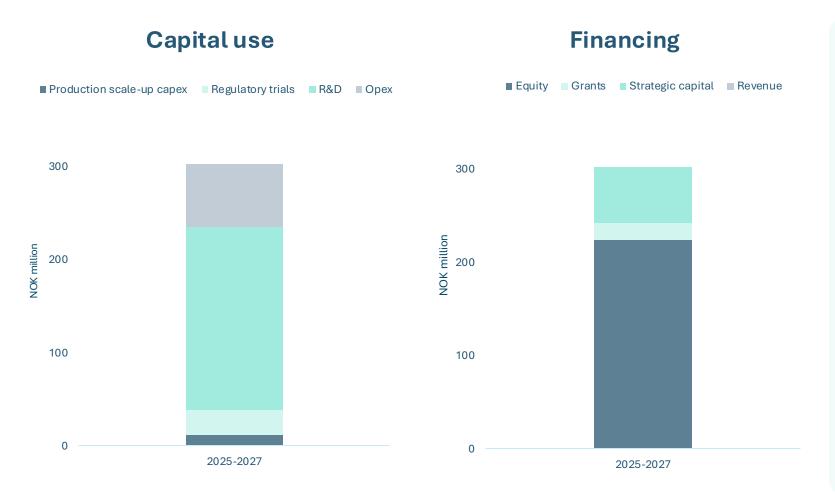
Cash flow

- Secured NOK 17 million from warrant exercise in June, with net proceeds of NOK 16 million
- R&D expenses of NOK 19 million according to plan
- Other opex of NOK 5 million stable due to lean operational structure
- Cash at quarter end of NOK 32 million
- Net cash outflow of NOK 9 million in Q2 2025





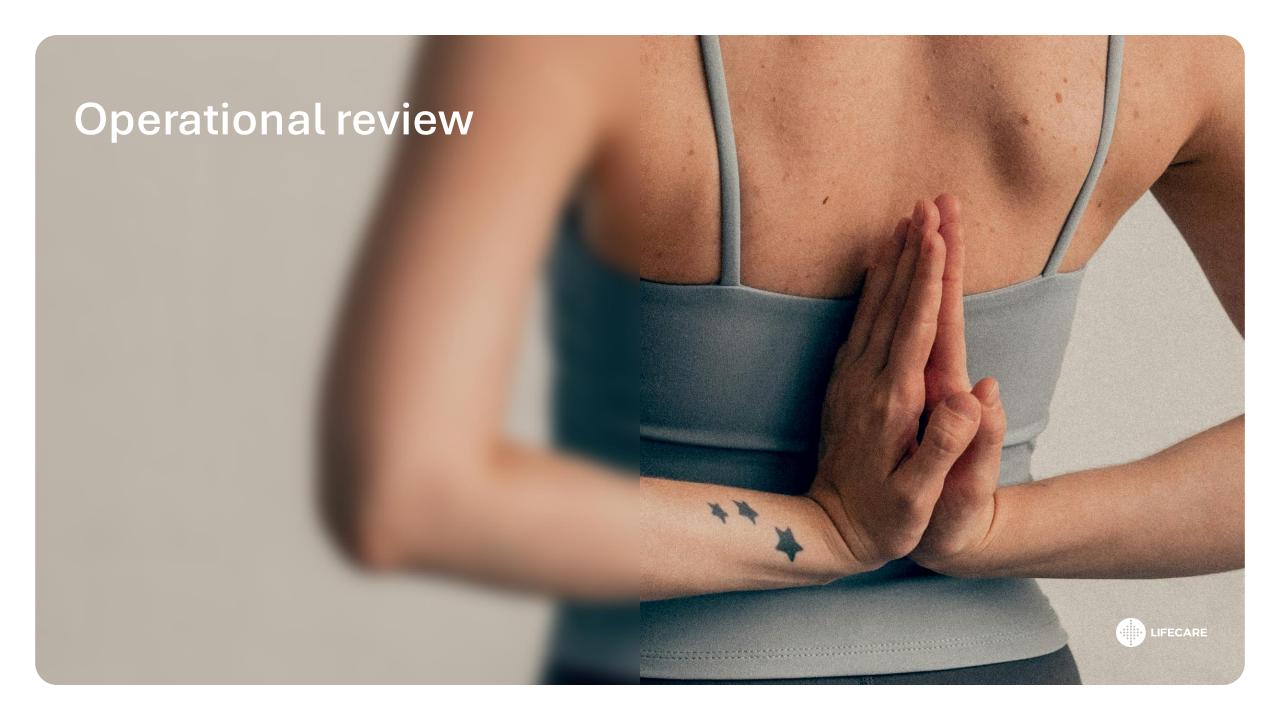
Capital plan aligned with milestones (2H 2025-2027)



- Capital use tied to clinical trials and CE-mark readiness
- Funded mainly by equity, exploring additional funding sources
- Revenue expected to contribute from 2027

Illustrative company estimates outlining primary cash flows and financing alternatives





Product development

- Finalized product design, reaching a design freeze of implantable device.
- Initial validation of the implant successfully completed.
- First-in-human expected to be concluded in 2025.
- Pivotal trial is expected to conclude in 2026.
- Commercial launch of CGM device for use is planned in 2027.



From dogs to first-in-human

- Longevity study in dogs continues, providing confidence in the safety and functional stability of the device.
- Plan to conduct trials in Norway and Germany in the second half of 2025.
- Principal Investigator appointed.
- Ethical approval to initiate first-in-human clinical trial, pending final approval to start trial.



Towards fully regulated medical device

- Started the process to achieve CE marking of our CGM system.
- Implementing a solid QMS (based on fully traceable framework).
- Regulatory compliance of electronics expected Q3 2025.



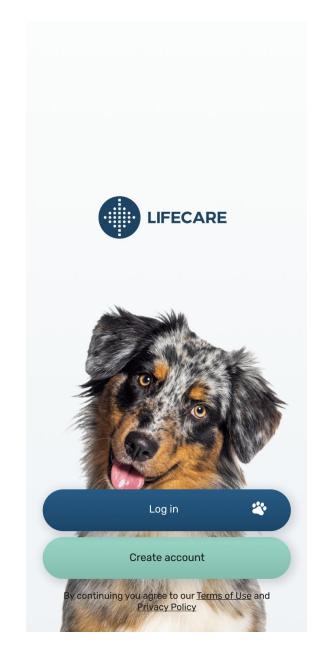






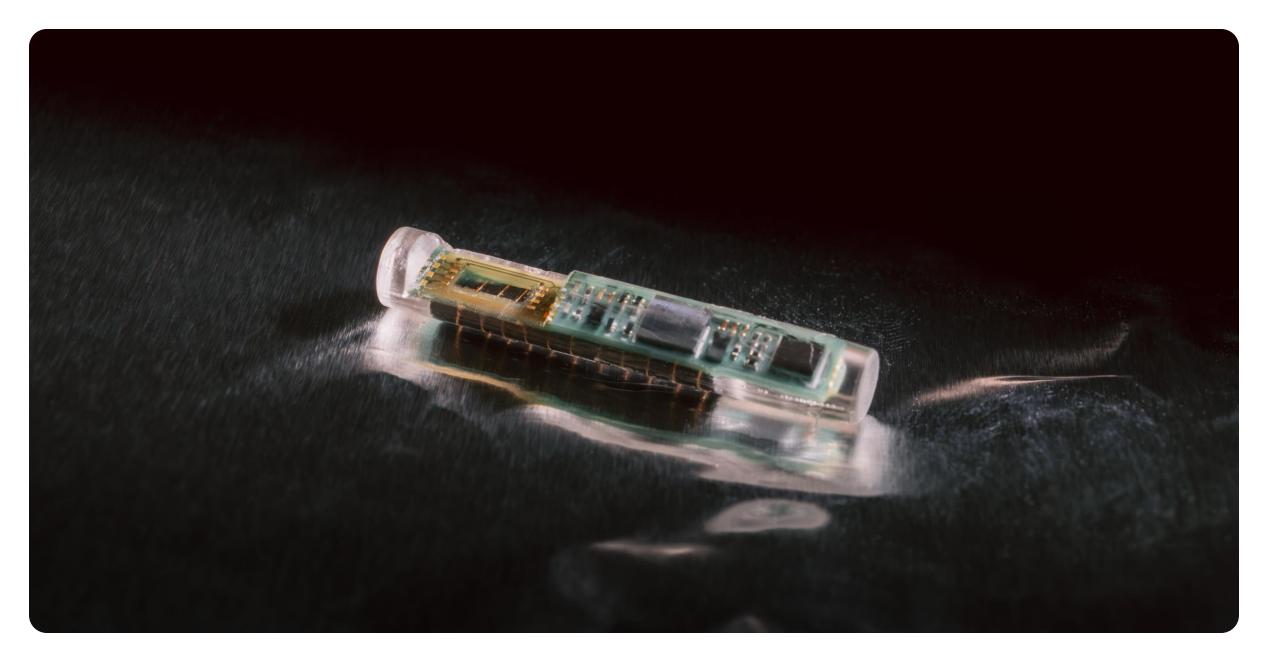










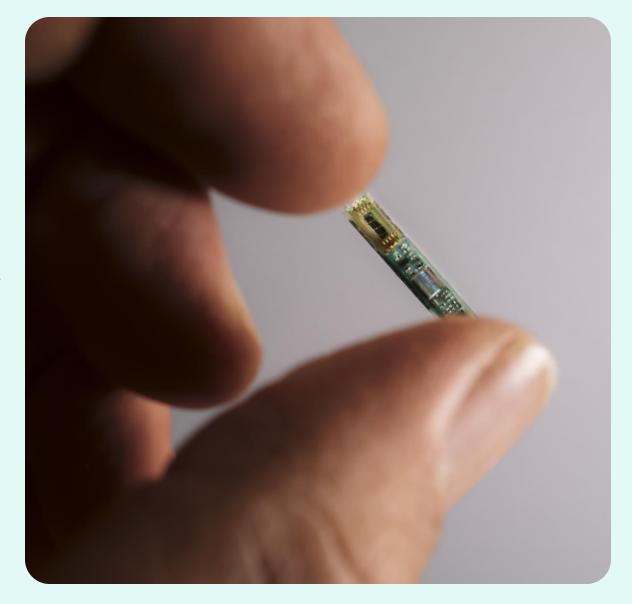






Outlook

- Strategic focus on capital-efficient development with clear milestones in production, regulatory processes, and market entry.
- Expect approval for first-in-human trials in Norway and Germany in Q3.
- Anticipate achieving compliance with all relevant standards for CE marking of electronics in Q3.
- Continue the ongoing longevity study in dogs.
- First-in-human trial planned to start in H2 2025, followed by a CE marking study in 2026, with the goal of market launch in 2027.
- Veterinary commercial launch planned before the human market to generate early revenue, gather user data, and reduce risk.
- Capital required to finance next stages of development.





Summary

- Implantables set to drive the next wave of CGM innovation
- ~ USD 5 billion adressable market
- Lifecare aims to offer the smallest, calibration free implantable
- Peak revenue potential of USD 1 billion+
- Capital-efficient model and clear path to commercialization
- Design freeze and first-in-human approval
- Entering a decisive phase with accelerating progress
- Additional capital to maintain momentum and fully capture upcoming milestones



Q2 2025 report is available for download at

lifecare.no/investor/reports-presentations/

Upcoming financial results

Q3 2025: 12 November 2025

