
**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 March 2026

001-43033
(Commission File Number)

PULSENMORE LTD.

(Exact name of Registrant as specified in its charter)

8 Omarim St.
Omer 8496500, Israel
(Address of principal executive offices)

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

Form 20-F Form 40-F

On March 18, 2026, Pulsionmore Ltd. has posted to its website an updated corporate presentation. A copy of the presentation is furnished with this Report of Foreign Private Issuer on Form 6-K as Exhibit 99.1 and is incorporated herein by reference.

EXHIBIT INDEX

Exhibit
No.

Description

99.1	Corporate Presentation.
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SIGNATURES

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, thereunto duly authorized.

Pulsenmore Ltd.

Date: March 18, 2026

By: /s/ Eran Hirsh
Eran Hirsh
Chief Financial Officer



 Pulsenmore™
Home Ultrasound

Transforming Maternal Health Through Home Ultrasound

NASDAQ and TASE: PLSM

Legal Disclaimer- Forward Looking Statements

This presentation contains forward-looking statements. In particular, statements using words such as “may,” “seek,” “will,” “consider,” “likely,” “assume,” “estimate,” “expect,” “anticipate,” “intend,” “believe,” “contemplate,” “do not believe,” “aim,” “goal,” “due,” “predict,” “plan,” “project,” “continue,” “potential,” “positioned,” “guidance,” “objective,” “outlook,” “trends,” “future,” “could,” “would,” “should,” “target,” “on track” or their negatives or variations, and similar terminology and words of similar import, generally involve future or forward-looking statements. Such forward-looking statements include, but are not limited to, statements relating to potential commercialization and market opportunities for its products and its anticipated future milestone Company events. Forward-looking statements reflect Pulesmore’s current views, plans, or expectations with respect to future events or financial performance. They are inherently subject to significant business, economic, competitive, and other risks, uncertainties, and contingencies. Forward-looking statements are based on Pulesmore’s current expectations and are subject to inherent uncertainties, risks and assumptions that are difficult to predict, including, but not limited to, the following: the Company’s lack of operating history; the Company’s current and future capital requirements and the Company’s belief that its existing cash will be sufficient to fund its operations for more than one year from the date that the financial statements are issued; the Company’s ability to manufacture, market and sell its products and to generate revenues; the Company’s ability to maintain its relationships with key partners and grow relationships with new partners; the Company’s ability to maintain or protect the validity of its U.S. and other patents and other intellectual property; the Company’s ability to launch and penetrate markets in new locations and new market segments; the Company’s ability to retain key executive members and hire additional personnel; the Company’s ability to maintain and expand intellectual property rights; interpretations of current laws and the passages of future laws; the Company’s ability to achieve greater regulatory compliance needed in existing and new markets; the Company’s ability to achieve key performance milestones in its planned operational testing; the Company’s ability to establish adequate sales, marketing and distribution channels; security, political and economic instability in the Middle East that could harm its business; and acceptance of the Company’s business model by investors. Further, certain forward-looking statements are based on assumptions as to future events that may not prove to be accurate. For a more detailed description of the risks and uncertainties affecting the Company, reference is made to the Company’s reports filed from time to time with the SEC, including, but not limited to, the risks, uncertainties and other factors included in the Company’s Form 20-F (SEC File No. 001-43033), filed with the SEC on December 29, 2025. The inclusion of forward-looking statements in this or any other communication should not be considered as a representation by Pulesmore or any other person that current plans or expectations will be achieved. Forward-looking statements speak only as of the date on which they are made, and Pulesmore undertakes no obligation to publicly update or revise any forward-looking statement, whether as a result of new information, future developments, or otherwise, except as otherwise required by law.

This presentation shall not constitute an offer to sell or the solicitation of an offer to buy, nor shall there be any sale of these securities in any state or other jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or other jurisdiction. The 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 such products.

Some images used in this presentation are for illustrative purposes only. Individuals depicted may not be actual patients or medical staff of the service.



Pulsenmore Developed the First and Only FDA-cleared Home-use Maternal Ultrasound System



The Pulsenmore Platform Technology



Patient self-scan device
Connects with smartphone to capture and securely transmit ultrasound images from home



Mobile App
Delivers personalized, step-by-step scanning instructions directly to patient



Clinician Dashboard
Physicians prescribe and review scans remotely, provide real-time feedback and guidance



Pulsenmore at a Glance



NASDAQ and TASE:
PLSM; Founded in
2014



Global leader in self-
scan home ultrasound



Strong patent portfolio;
Multi-center clinical
validation



Global Leader

220,000+ successful
patient scans
with Pulsenmore ES in
initial 4 years



Regulatory Milestones

First FDA De Novo clearance Nov
'25, Access to the U.S. market
with 3.6M annual deliveries; CE-
mark already



Product Pipeline

2nd product (IVF) in
commercialization.
3rd product (CHF) in
development



Financial Strength

NASDAQ dual listed since January
2026.
Total investments to-date \$90M
including GE HealthCare, Fujifilm



The Opportunity: Decentralized Prenatal Monitoring



Digital Health Transformation

Remote monitoring adoption accelerating across healthcare



Cost-Efficient Models

Payers shifting toward decentralized care delivery



Provider Shortages

OB/GYN shortages driving home-care solutions



Consumer Demand

Patients demanding convenient, at-home reassurance



Total Addressable Market

140M

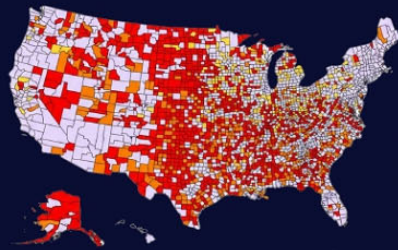
pregnancies globally

Fetal monitoring is a multi-billion global market



Access to Care in the U.S. is Unequal and Costly


3.6M
Annual U.S. Deliveries
A meaningful share of pregnancies require additional monitoring due to maternal or fetal risk factors



- Maternity Care Access
- Maternity Care Deserts (1918)
 - Low Access to Care (273)
 - Moderate Access to care (223)
 - Access to Maternity Care (2427)

The Growing Gap



35% of U.S. counties are maternity care deserts.



Rising healthcare costs



Limited ultrasound access in rural and underserved regions



CDC reports >80 severe maternal morbidity cases per 10,000 deliveries in 2019 and rising.

<https://ourworldindata.org>
<https://www.marchofdimes.org/peristats/reports/united-states/maternity-care-deserts>
<https://www.acog.org/clinical/clinical-guidance/committee-opinion/articles/2021/06/indications-for-outpatient-antenatal-fetal-surveillance>

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Healthcare Challenges Today



Growing Demand

Advanced maternal age and high-risk pregnancies increase prenatal care frequency.



Capacity Strain

Clinics face space and staffing constraints, with wait times averaging ~42 days, affecting care quality.



Limited Access

Rural and ethnic minority women face disparities in prenatal care.



Costly ED Utilization

~750,000 avoidable pregnancy-related ED visits each year, costing the U.S. health system ~\$1B annually.



Hospitals and payers face relentless pressure to reduce costs. Digital health solutions and remote imaging can decentralize care and bridge the gap in prenatal care.



U.S. Business Model

Device Sales

60%

Single pregnancy use devices (~\$1,000) for high-risk pregnancies (~20% of births) + expansion to broader prenatal segments.

Recurring SaaS Revenue 40%

Clinician dashboard subscriptions

At-scale model:

Device margin + high-margin SaaS/monitoring revenue



Device Margin
Profit from hardware sales



High-margin SaaS revenue
Subscription-based software income



Monitoring Revenue
Ongoing service and analytics



Our Competitive Advantages



FDA-Cleared Innovation with Clear Value for Patients, Providers and Payers

The first and only FDA-cleared home-use prenatal ultrasound system, with remote professional supervision.
Expands capacity with continuous clinical oversight.



Patented, validated Technology

90+ patents filed; 29 granted.
Clinical-grade imaging validated across major U.S. academic hospitals.



Road map Products

One core platform, expandable to new modalities, including IVF (already approved in Israel) and CHF (in development).



Scalable Manufacturing

World's first automated ultrasound transducer production line, enabling full manufacturing independence with scale-ready volume output.



Scalable Manufacturing

Pulsenmore developed the first & only fully automated production lines for manufacturing ultrasound transducers



Long-term competitive advantage



Full manufacturing independence



Flexibility to meet changing demand



Reduced raw material costs



Superior quality and reliability



Lowering costs.
Increasing output



Clinical Studies & Real-World Evidence

220K+

Remote Scans
Performed to date globally

1,200

Hospitalization Days Saved
at a single hospital using a hybrid care
model with Pulsenmore (including other
devices).

98.1%



Of Scans suitable for clinical
evaluation

90%



Of Users (Callit survey)
report reduction in
pregnancy-related stress

86.3%



Of Users (Clalit survey) report
improved pregnancy
experience

Clinical Benefits

- ✔ Two RCTs (N=100) show reduced anxiety
- ✔ Reduced in-clinic visits and hospital admissions
- ✔ Early detection and intervention

Patient & Provider Value

- ✔ Time saving for patients
- ✔ Increased patient engagement
- ✔ Greater flexibility in care delivery
- ✔ More efficient resource utilization



Strong endorsement from clinicians and patients



At 39 weeks, Doria performed a home ultrasound scan. After reviewing the images, Professor Hadar advised her to come to the hospital immediately. Within 30 minutes of arrival, she gave birth - there wasn't even time for an epidural.

Doria Marley



Prof. Alfred Abuhamad

Chairman, OBGYN Department, Eastern Virginia Medical School.
Pulsenmore Medical Advisory Board

With FDA marketing authorization, Pulsenmore introduces a transformative model for prenatal care - extending ultrasound access beyond the clinic and redefining how we reach and monitor expectant mothers.



Dr. Lawrence D. Platt

Center for fetal medicine and women's ultrasound, LA. Pulsenmore Medical Advisory Board

Pulsenmore Ultrasound system marks a new era in maternal - fetal care, empowering expectant mothers with safe, guided access to ultrasound imaging from home or work, while keeping clinicians closely connected throughout the entire process.



Strategic Partnerships & Investments



Go-to-Market Strategy

U.S. Commercial Launch '26

- OB networks and hospitals
- Telehealth provider partnerships
- Employer partnerships cost savings for patients
- CPT 76815 available at launch
- Planned expansion to BPP under CPT 76819 (\$150)
- ACOG supports hybrid prenatal care models

Europe & APAC

- Expansion via leading institutions (UK, France, Australia)
- Large OB clinic partnerships
- Reimbursement pathway development



Secure early strategic provider partnerships



Demonstrate cost savings (reduced unnecessary clinic visits)



Expand payer coverage + bundled reimbursement



Rapid scale through digitally-enabled distribution



Board of Directors



Dr. Elazar Sonnenschein - CEO & Founder

Led several medical device and technology companies (NASDAQ: MDGS). PhD in Electronics & Computer Engineering, Ben-Gurion University. 70 granted patents with focus on acoustics & optics for medical applications.



Mr. Jonathan Adereth - Chairman of the board

Veteran executive (NYSE: ELT, NASDAQ: MZOR), with 27 years at Elscint as President and CEO (1994–1998). Brings deep expertise in scaling global medical imaging companies.



Ms. Linda Messalem, External Director

Financial executive with 30+ years of experience in financial reporting, taxation, M&A, and strategic advisory for public and private companies in Israel and internationally.



Ms. Racheli Guz-Lavi, Director

Managing Partner at Amit, Pollak, Matalon & Co.. Senior tax and corporate advisor to public & private companies, extensive experience in international transactions and governance.



Mr. Hagai Itkin, Director

Partner and CTO at iNEXT Capital, a venture capital firm focused on digital health. Serves on advisory boards of multiple cybersecurity and technology companies.



Prof. Anat Loewenstein, External Director

President of the Israeli Ophthalmological Society; Vice Dean, Faculty of Medicine, TLV University; Sidney Fox Chair of Ophthalmology. Leadership in academic medicine and research.

The Board combines seasoned MedTech executives, healthcare leaders, and capital markets expertise.



New Product in the pipeline-Follicles monitoring

✓ IVF and fertility preservation require 4–7 ultrasound scans per cycle, creating significant clinic load and patient burden.

✓ Home-based follicular monitoring unlocks scalable capacity and improves patient experience.

AMAR authorization in Israel secured
\$4.5M contract signed with Clalit HMO
Regulatory expansion expected in 2026



Growth & Strategy Outlook

01 

U.S.
Commercialization '26

Launch

02 

International
Expansion

EU and global market
penetration

03 

R&D Pipeline

Expanded indications to
new modalities , AI
diagnostics, enhanced
visualization

04 

Manufacturing
Automation

Industry-leading cost basis
and independent production



Long-term vision
Category leader in Digital Health and FemTech





 Pulsenmore™
Home Ultrasound

Thank you.