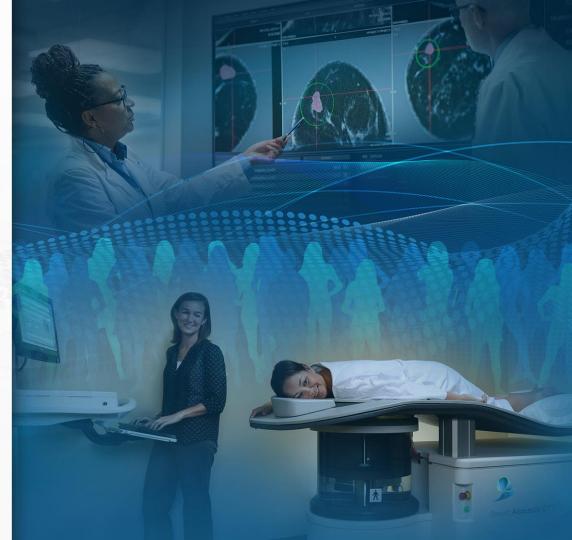


Quantitative Transmission Imaging

Breast Acoustic CT[™] Scanner

November 2025



Disclaimer

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On June 6, 2017, the U.S. Food and Drug Administration ("FDA") in response to QT Imaging's Section 510(k) Summary of Safety and Effectiveness premarket notification under the Food, Drug and Cosmetic Act, determined that the QT Breast Scanner is substantially equivalent to the predicate device. Our use of the words "safer," "safety", "effectiveness", and "efficacy" in relation to the QT Breast Scanner in this Presentation and all other QT Imaging related documents is limited to the context of the Section 510(K) Summary of Safety and Effectiveness that was reviewed and responded to by the FDA.

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This presentation includes references to EBITDA and Adjusted EBITDA, financial measures that have not been prepared in accordance with generally accepted accounting principles in the United States ("GAAP"). EBITDA is defined as loss before interest expense, income tax expense, depreciation and amortization. Adjusted EBITDA is defined as EBITDA further adjusted for equity-based compensation, net change in fair value of the derivative, earnout and warrant liabilities, and transaction expenses. Similar excluded expenses may be incurred in future periods when calculating these measures. QT Imaging believes these non-GAAP measures of financial results provide useful information to management and investors regarding certain financial and business trends relating to the Company's financial condition and results of operations QT Imaging believes that the use of these non-GAAP financial measures provides an additional tool for investors to use in evaluating projected operating results and trends and in comparing QT Imaging's financial measures with other similar companies, many of which present similar non-GAAP financial measures to investors. Investors should not rely on any single financial measure to evaluate QT Imaging's anticipated business. Certain of the financial metrics in this presentation can be found in QT Imaging's Form 8-K filed with the U.S. Securities and Exchange Commission (the "SEC") on May 13, 2025, and the reconciliation of EBITDA and Adjusted EBITDA can be found on pages 65 and 66 of this presentation.



QT Imaging Holdings (QTI) Has the Potential to Transform Medical Imaging

- QTI is a medical device company with imaging technology that has the potential to transform the industry
- QTI Scanner is the only 3D imaging device to receive FDA clearance for use as a transmission and reflection ultrasonic imaging system of a patient's breast



- QTI's patent-protected technology provides a high resolution, relatively low-cost, comprehensive, no radiation, no discomfort medical imaging solution
- QTI's technology yields improved diagnostic performance compared to traditional mammogram and has similar imaging quality compared to MRI but is a lower cost and more accessible solution.

Our Mission Transforming Breast Health For Every Woman



At QT Imaging, we are redefining what's possible in breast imaging - delivering safe, high-resolution, and cost-effective solutions where traditional technologies fall short. Our goal is to make advanced diagnostic imaging accessible to all women, including those with dense breast tissue or limited access to care. With a platform rooted in innovation, clinical validation, and Artificial intelligence (AI) integration, we are committed to building a future where early, accurate breast cancer detection is available without compromise.

NIH has awarded QT Imaging about

\$18Million

for a supplemental imaging solution for women with dense breasts



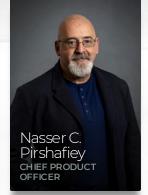
Our Management Team

















History of QT Imaging



Company founded by John Klock, MD

\$18M from NIH to develop supplemental imaging modality to resolve dense breasts

FDA Clearance QTI becomes public on NASDAQ QTI gets delisted, public on OTC

Raise \$18M NASDAQ application to relist by EOY 2025



QTI's Platform Approach



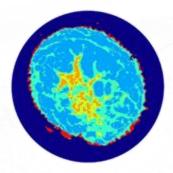
Versatile Clinical Use:

from early evaluation to diagnosis and treatment monitoring



Al-Ready Foundation:

continuous learning to improve diagnostic accuracy



Data-Rich Biomarkers:

speed-of-sound maps for tissue characterization



Upgradable & Scalable Platform:

software-defined features that adapt to evolving needs



Accessible & Affordable Design:

suitable for lowresource settings, mobile clinics, and underserved populations



Executive Summary

Patent-protected technology:

14 granted patents in US/Europe + 2 new patent applications

TECHNOLOGICAL CONSIDERATIONS

- FDA cleared for breast Imaging
 - ➤ Breakthrough Device Designation awarded by the FDA provides fast track to unique CPT codes and future clearances
- Based on safe waves, with the following biomarkers:
 - Quantitative measure of the intrinsic speed of sound in Breast Tissue
 - Quantitative measurement of fibroglandular density ratio (breast density)
- Standardized scanning with operator independent images, unlike hand-held ultrasound (HHUS)
- Resolution comparable to MRI but without any contrast agent
- Volumetric accuracy to determine mass doubling times
- Higher diagnostic accuracy in Dense Breasts

PATIENT CONSIDERATIONS

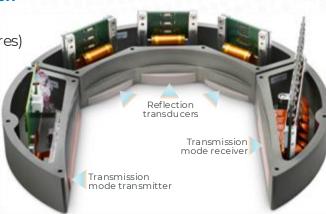
- Safe, no radiation, no contrast
- No discomfort, painless scans
- Less recalls, reduced anxiety
- Less unindicated Intervention, Biopsy
- Reduce cost of Care
- Scanning of women under 40 years not suitable for Mammography
- Useful for Cancer Therapy Monitoring



Quantitative Transmission (QT) Imaging

- What is QT Imaging?
 - Inherently 3D volumetric imaging modality due to 3D data acquisition and mathematical image reconstruction, thus its high resolution, similar to MRI
 - Two independent sources of information:
 - Uses CT-like configuration with ultrasound to acquire and reconstruct transmission images which map the speed-ofsound across the tissue volume (unique to our technology)
 - Uses reflection configuration for high-resolution depiction of tissue interfaces (ducts, Cooper's ligaments, lesion boundaries) as well as improved lesion visualization — (cancers have irregular, heterogeneous reflective signatures)
 - Overcomes operator dependence and lack of standardization associated with HHUS
 - Pain free, safe, no radiation or contrast
- Image Acquisition:
 - Prone position with breast submerged in water
 - 360-degree rotation of ultrasound arrays
 - 10-12 minutes per breast average scan time







Optimized Patient Experience

- No ionized radiation. Acoustic source only
- No breast compression and associated discomfort
- 10-12 minutes per breast exam time
- Quiet and comfortable (as compared to MRI claustrophobia, coil pressure, noise and lengthy exams)
- No contrast injection or associated risk

 (as compared to MRI Gadolinium)
- No limitations for dense breasts or implants





What's Next?







Biomarker Development

- Quantitative parameters:
 Speed of Sound,
 Attenuation, Reflection,
 Nakagami parameters etc.
- AI/ML models trained on large, labeled datasets
- Validation through retrospective analysis and clinical-grade ground-truth annotations

Cloud-Based All Integration

- Deployment within QTI Cloud SaaS Integration via InteleShare's framework
- Automated pipelines for image reconstruction, feature extraction, Probability of Cancer Map generation
- Continuous learning from clinical data uploads and feedback loops

Clinical Decision Support

- Visualization in QTviewer and clinician-facing dashboards
- Automated lesion segmentation and BIRADS scoring
- FDA validation and CPT reimbursement readiness
- Cloud-delivered AI updates with no on-premise installs required



What's Next Why Biomarkers Matter for QT Imaging?

From Images to Quantitative Data

- Traditional breast imaging (mammography, MRI, ultrasound) produces qualitative visual information that relies heavily on radiologist interpretation rather than objective tissue properties
- QT Imaging's acoustic CT technology generates quantitative volumetric maps of speed of sound, attenuation coefficient, reflection intensity
- These measurable parameters serve as biophysical biomarkers, thus enabling reproducible and repeatable features of tissue composition and structure

Objective, Reproducible, and Al-Ready

- Biomarkers transform imaging from qualitative observation to objective science
- They provide numeric indicators of tissue stiffness, density, and microstructure
- Enable machine learning algorithms to detect subtle disease patterns
- Reduce inter-reader variability and allow longitudinal tracking of change over time



What's Next Why Biomarkers Matter for QT Imaging?

Clinical Utility

Biomarkers allow QT Imaging to move beyond detection toward diagnostic and prognostic precision through quantitative imaging:

- Early detection of malignancy using defined quantitative thresholds
- Therapy monitoring such as pathological complete response (pCR) prediction in neoadjuvant therapy
- Personalized medicine, by correlating imaging biomarkers with genomic and histopathologic data
- Automated lesion characterization and BI-RADS aligned standardization for consistent interpretation



Clinical Value Proposition

Clinician	Challenges	QT Solution
Radiologist	Dense breast triage, reduce recalls and false positives	Dense breast image quality powered by quantitative geometric, spectral and elastic biomarkers will allow for better triage, increased physician confidence and reduce recalls and false positives
Radiologist oncologist	Breast preservation RT, precision dose delivery and tailoring of therapy	Precision image quality and quantitative biomarkers will allow for better biology guided radiation therapy
Medical oncologist	Personalizing therapy, therapy monitoring and management of side effects	Precision phenotyping powered by quantitative biomarkers will allow for better personalizing of therapy, monitoring and management
Surgeon	Accurate surgical planning, reducing risks	Higher image quality combined biomarker driven precision phenotyping (correlated with histopathologic data) will allow for better surgical planning and risk management



From a Hardware Company...

By building a biomarker-driven platform, QT Imaging shifts from a hardware company to a precision imaging platform combining hardware, software, and cloud-based AI modules





Hardware3D acoustic scanner

Software reconstruction, denoising, analytics

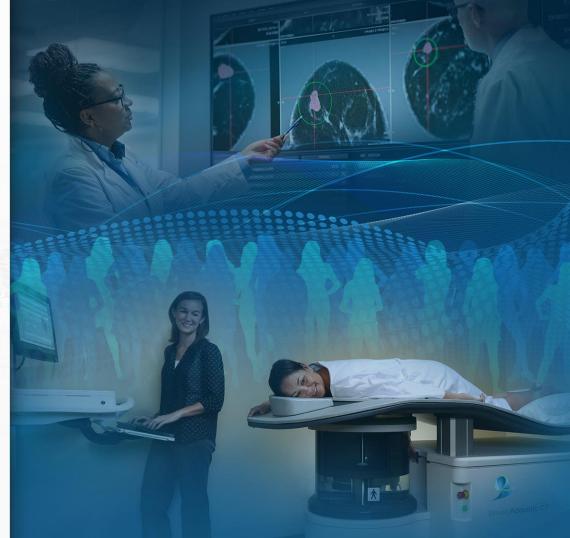
Cloud AI biomarker extraction, disease probability maps

To a Sustainable, Data-Centric Ecosystem Continuously Improving with Clinical Use





Business, Partnerships, and Footprint



Business Partnerships – Today



Under Distribution Agreement with NXC Imaging (Subsidiary of Canon Medical Systems) for U.S.A. market



Under Contract Manufacturing Agreement with Canon Medical Systems – For Canon Distribution



شَركة الخَليج الطبيَّة المحدُونة GULF MEDICAL CO. UTD.

Under Distribution Agreement with Gulf Medical for KSA (Kingdom of Saudi Arabia) market



Medical Image Management Archiving/communication systems and Cloud PACS partner



Business Partnerships



- Under Distribution Agreement with NXC Imaging (Subsidiary of Canon Medical Systems) for U.S.A. market
 - Committed quarterly minimum order quantities (MOQs) for scanners' shipments till end of 2026

 Four additional distributors signed by NXC Imaging to cover sales across all states





شَركة الخَليج الطبيَّة المحدودة GULF MEDICAL GO. TD.

Canon

- Under Contract Manufacturing Agreement with Canon Medical Systems
 - In the process of bringing up large scale manufacturing with CMSC in Japan
 - QTI Novato site to continue manufacturing scanners



Business Model – HW + QTI Cloud Platform

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On Premise Intelerad. Radiology Referring MD

QTI Cloud SaaS



QTI Cloud Platform plans to offer advanced breast analysis, predicting breast cancer risk with Al precision

- Lesion-by-lesion detail
 Precision phenotyping for each lesion with biomarker based quantification and lesion risk scores
- Comprehensive breast health assessment
 At-a-glance view of characterized breast health
 by region
- Lesion scoring
 Clear and concise summary of identified lesions
 by severity to facilitate personalized treatment
- Backed by clinical evidence
 To improve prognostic value, increase physician confidence and reduce false positives (unnecessary interventions and anxiety)



QT Scanner Locations Map







COMMERCIAL CENTERS

Center For New Medicine

Dr. Leigh Erin Connealy 6 Hughes, Suite 100 Irvine, CA 92618 +1 (949) 680-1880 Website

Couri Center for Gynecology and Integrative Women's Health

Dr. Michele Couri 6708 N Knoxville Ave, Suite 1 Peoria, IL 61614 Website

Innovative Radiology

Dr. John Tentinger 7601 Office Plaza Dr, Ste 115 West Des Moines, IA 50266 +1 (515) 222-0550 Website

Longevity RX

Dr. Joel Fuhrman Rancho Santa Fe, CA +1 (858) 367-3558 Website

PerfeQTion Imaging

Dr. Jenn Simmons 346 W Lancaster Ave, Haverford PA 19041 Website

Qlarity Breast Imaging

Dr. Kristine Burke 3 Hamilton Landing #180 Novato, CA 94949 +1 (415) 842-7403 Email

Qlarity Breast Imaging

Dr. Kristine Burke True Health Center for Precision Medicine 8105 Saratoga Way, #240 El Dorado Hills, CA 95762 +1 (916) 542-1644 Website

Qlarity Breast Imaging

Dr. Yvonne Karney Vitality Renewal Functional Medicine 31 N. Virginia St. Crystal Lake, IL 60014 +1 (815) 271-7300 Website

Robinhood Integrative Health

Dr. Wiggy Saunders 3288 Robinhood Rd. Suite 202 Winston-Salem, NC 27106 +1 (336) 768-3335 Website

Vincere Cancer Center

Dr. Vershalee Shukla and Dr. Pablo Prichard Top Cancer Center in Scottsdale, AZ Vincere Cancer Center 7469 E Monte Cristo Ave. Scottsdale AZ 85260 +1 (480) 306-5390 Website

CLINICAL SITES

Mavo Clinic

Tiffany Sae-Kho, M.D. 200 First Street SW Rochester, MN 55905 +1 (855) 776-0015 Website

National Institutes of Health (NIH)

9000 Rockville Pike Bethesda, MD 20892 United States

Sunnybrook Health Sciences Center (NIH Grant)

2075 Bayview Ave North York, ON M4N 3M5 Canada

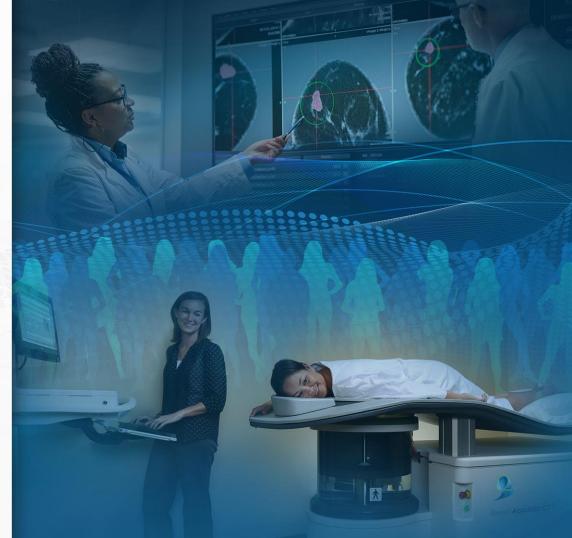
Prestigious University

in Tokyo, Japan





Regulatory Activities



Regulatory Activities

Saudi FDA (SFDA)



Submit Dossier —Jan 1, 2026 Authorized Rep via Gulf Medical; MDS-GS-004 Regulatory Review Anticipated 60–90 days Expected Marketing Authorization Target: Q2 2026

UAE MOHAP



Dossier Preparation Leverage SFDA submission MOHAP registration UAE Submission MOHAP device registration

Expected Approva Target: Q3 2026

CE Mark (EU MDR)



Notified Body Engagement BSI/Alternate; plan conformity route Submit Technical Documentation CER, Annex II/III, PMS/PMCF

Expected CE Mark Target: Q4 2026





Quantitative Transmission Imaging – Why and How It Fits In



QTI's Technology Has the Opportunity to Transform the Breast Imaging Market

CURRENT MARKET

Breast Imaging: \$6B MARKET (1)

- FDA approved as supplementary screening device for breast imaging
- Aim to revolutionize current imaging paradigm, replacing mammography, ultrasound (handheld and automated), and freeing MRI scanners time

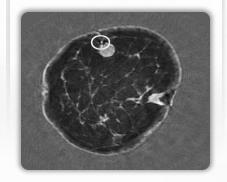


Layer	Description	2025 Value	2030 Value	CAGR
ТАМ	Total global opportunity (dense-breast supplemental screening ⁽⁴⁾ + biomarkers ⁽³⁾)	2.5	3.8	8.5%
SAM	Accessible reimbursed markets (U.S., OECD, Korea, Japan) ⁽²⁾	1.7	2.6	8.5%
QTI Penetration	~10% SAM share by 2030	0.02	0.20- 0.25	_

NEW MARKET

SaaS Biomarkers: \$1B MARKET (3)

- Move to precision diagnostics
- Move to personalized treatments with therapy guidance and monitoring





⁽¹⁾ https://www.futuremarketinsights.com/reports/automated-breast-ultrasound-system-market

⁽²⁾ https://www.grandviewresearch.com/industry-analysis/breast-imaging-equipment-market \$5.45B for 2024 with 8.9% CAGR.

⁽³⁾ https://www.archivemarketresearch.com/reports/breast-cancer-biomarkers-316622

⁴⁾ https://www.cdc.gov/breast-cancer/about/dense-breasts.htm

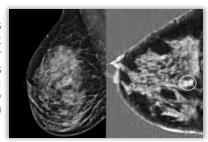
QT Imaging's FDA-cleared Solution for Dense Breasts

Many Women Have Dense Breasts, Which Mammograms are Inefficient in Screening for Cancer



50% of women between the ages of 40-74 in the US have dense breasts⁽¹⁾

In ~84% of cases observed in a recent mini-study, QT Scanner identified abnormalities in dense breasts that were not identified by x-ray mammograms⁽²⁾



X-Ray Mammogram

QT Scan

The FDA Has Recognized the Importance of Breast Density in Breast Cancer Screening

Mammograms Must Include Breast Density Information, New FDA Rule Says

About half of the women over the age of 40 in the U.S. have dense breast tissue, which can make cancer scans hard to read (3)



"the new rule advises physicians and patients to consider breast density alongside other cancer risk factors when deciding whether additional screening is necessary"

- Hilary Marston, CHIEF MEDICAL OFFICER, FDA

Mammography Misses 35.6-52.2% of Breast Cancers in Dense Breast Tissue⁽⁴⁾



OTIMAGING

(4) The Role of Ultrasound in Screening Dense Breasts. NCBI.

^{(3) &}quot;Mammograms Must Indude Breast Density Information, New FDA Rule Says". Wall Street Journal

The Current Breast Imaging Paradigm Leads to Unnecessary Concern and Costs

Screening compliance is low



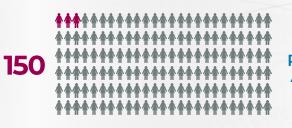
Of the **65%** of women who do get screened, many suffer through unnecessary callbacks

Aside from the discomfort of the mammogram procedure, up to 15% of women are called back for additional procedures such as ultrasound, MRI or biopsies – which can be expensive, time consuming and cause significant anxiety⁽²⁾

For every 1,000 screening mammograms:

CALL BACK RATES

~15% call-backs rates with mammography



98% of Recalls are Avoidable

BIOPSIES

~10% biopsy rate for callbacks



Over 80% of Callback Biopsies are Benign⁽⁴⁾

CANCER INCIDENCE

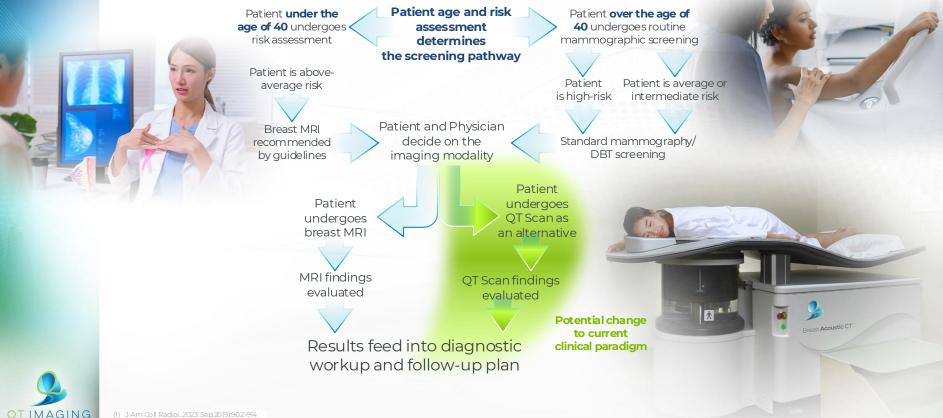
0.3% cancer diagnosis⁽⁵⁾





(1) Mammography, Center for Disease Control and Prevention
(2) Very Well Health | 13 Reasons for a Mammogram Callback | Larell Scardelli
(3) PubMed | False-Negative Rate of Combined Mammography and Ultrasound for Women with Palpable Breast Masses | Carlos H.F. Chan, Suzanne B. Coopey, Phoebe E. Freer, and Kevin S. Hughes
(4) National Breast Cancer Foundation | Breast Biopsy Procedure Types, What to Expect and Results
(5) U.S. Breast Cancer Statistics. Breast-Earner or or.

Standard of Care Today 12 How QT Scan Fits In



Current Standard of Care in Breast Imaging

Risk Category		Breast Density	Recommended Imaging Modalities	Guideline Recommendations
Average Risk	≤12–15%	Fatty Breasts	Screening Mammography (2D or 3D) annually starting at age 40	NCCN ⁽⁴⁾ : Annual mammography for women aged 40 and older. ACR/SBI ^(1,2) : Annual mammography starting at age 40. EUSOBI ⁽⁵⁾ : Biennial mammography for women aged 50–69; consider starting at 40.
Average Risk	< ≤12–15%	Dense Breasts	Screening Mammography (2D or 3D) annually starting at age 40 Supplemental Imaging: Consider Ultrasound or MRI	NCCN: Consider supplemental imaging for women with heterogeneously or extremely dense breasts. ACR/SBI: Recommend supplemental MRI for women with dense breasts and additional risk factors. EUSOBI: Recommend MRI screening every 2–4 years for women aged 50–70 with extremely dense breasts
Above Average Risk	15_19%	Any Density	Screening Mammography (2D or 3D) annually starting at age 40 Supplemental Imaging: Consider MRI or Ultrasound	NCCN: Annual mammography, consider MRI for women with a 20–25% lifetime risk. ACR/SBI: Recommend MRI for women with a 20–25% lifetime risk. EUSOBI: MRI screening for women with a 15–20% lifetime risk.
High Risk	≥20-25%	Any Density	Screening Mammography (2D or 3D) annually starting at age 30 Supplemental Imaging: Annual MRI starting at age 25–30	NCCN: Annual MRI and mammography for women with ≥20% lifetime risk. ACR/SBI: Recommend annual MRI and mammography for women with ≥20% lifetime risk. EUSOBI: Recommend annual MRI for women with BRCA mutations or equivalent risk.

⁽¹⁾ J Am Coll Radiol. 2023 Sep; 20 (9):902-914.

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⁽²⁾ J Am Coll Radiol. 2024 Jun; 21(6S): S126-S143

^{(4) 1} Natl Compr Canc Netw 2023 Sept 21/9) 900-909

QTI's Current Indications For Use

DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120 Expiration Date: 06/30/2023

See PRA Statement below.

510(k) Number (if known)

K220933

Device Name

QT Scanner 2000 Model A

Indications for Use (Describe)

The QT Scanner 2000 Model A is for use as an ultrasonic imaging system to provide reflection-mode and transmission-mode images of a patient's breast. The QT Scanner 2000 Model A software also calculates the breast fibroglandular tissue volume (FGV) value and the ratio of FGV to total breast volume (TBV) value as determined from reflection-mode and transmission-mode ultrasound images of a patient's breast. The device is not intended to be used as a replacement for screening mammography.

The QT Scanner 2000 Model A is indicated for use by trained healthcare professionals in environments where healthcare is provided to enable breast imaging in adult patients.

Broad intended use to allow breast imaging of any subject of age 18 or older

First FDA clearance for an ultrasound-based device to be able to quantify breast tissue volume



How QTI Potentially Fits Into the Current Paradigm

Risk Category

Potential Role of QTI Device

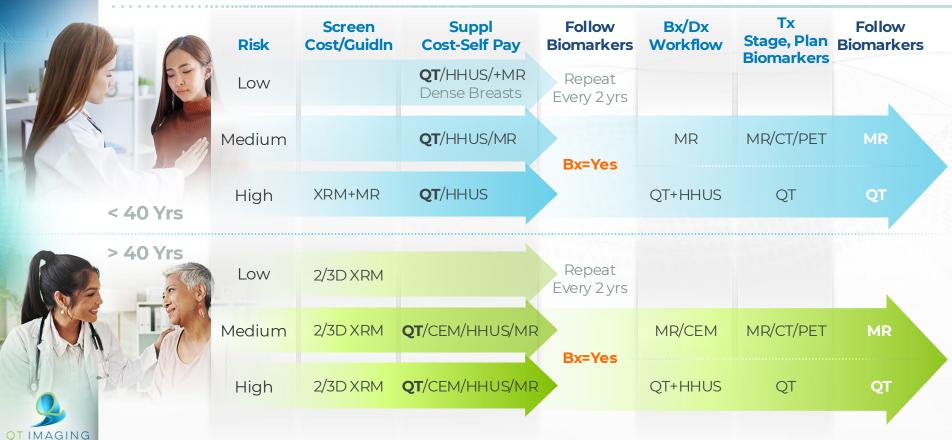
Average Risk (≤12–15%) QTI offers a non-ionizing, high-resolution alternative for supplemental imaging, especially useful in patients with dense breasts where mammography is limited. Ideal for frequent monitoring without radiation exposure.

Above-Average Risk (15–19%) QTI provides a **safer alternative to MRI for moderate-risk individuals**, including those with family history or dense tissue. It avoids gadolinium-based contrast risks, offering **functional imaging with fewer contraindications**.

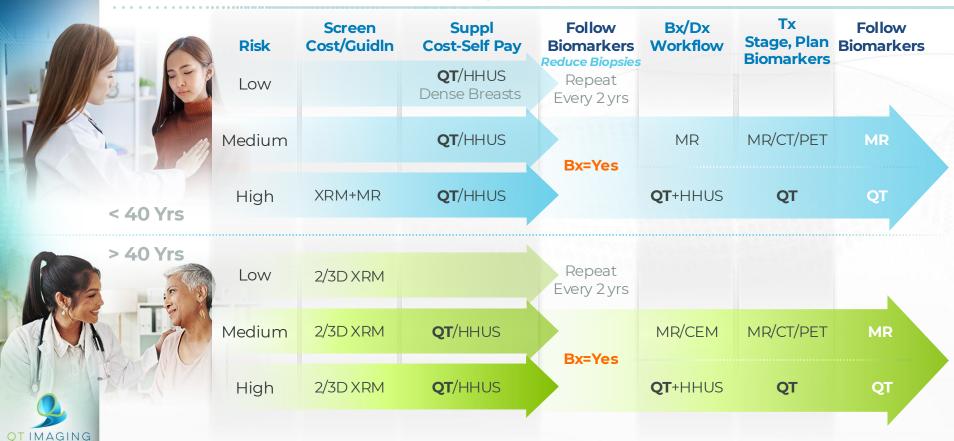
High Risk (≥20–25%) QTI may supplement or replace MRI in high-risk individuals, especially where MRI is contraindicated or poorly tolerated. Supports early, radiation-free surveillance with improved soft-tissue contrast, aligning with early screening needs.



Patient's Clinical Journey

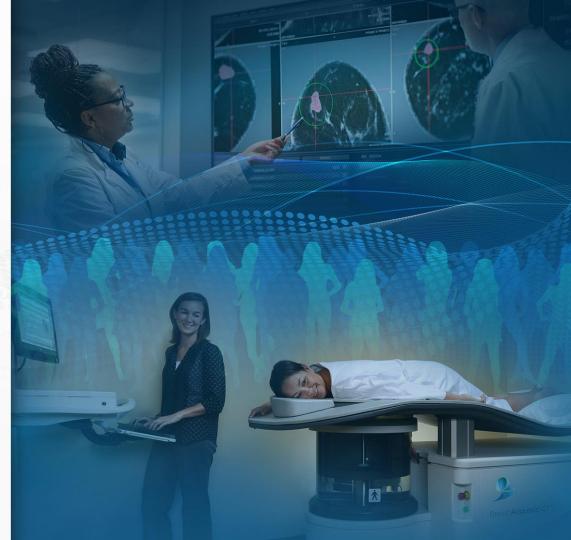


Patient's Clinical Journey



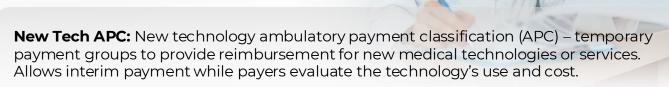


Specific Reimbursement Codes



Coding Pathways for QT Imaging

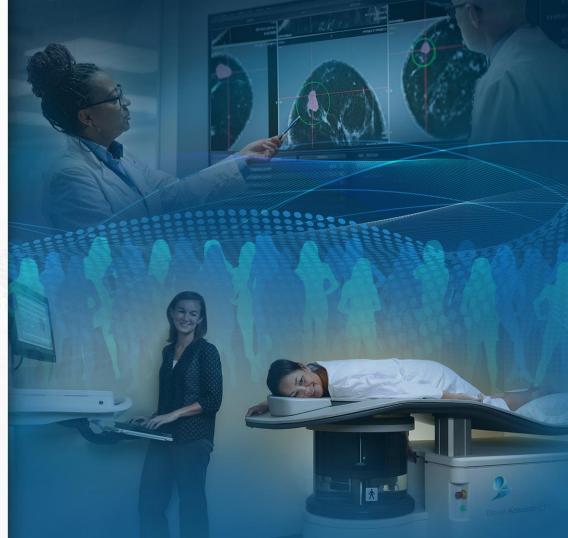
- Current State:
 - Use **CPT 76999** (unlisted ultrasound procedure)
 - QTI provided a 'guidelines' document for providers and patients (at www.qtimaging.com)
- Strategy / Next Steps:
 - New Tech APC application Submitted August 2025
 - Expected Category C-code by Q1 2026
 - Category III CPT code (early commercialization)
 - Submitted on 11/3/2025
 - Expected Cat III code by Q1 2027







Competitive Landscape



The QT Scanner Delivers a Better Experience for Patients than Traditional Systems

Underlying Technology Ultrasound

Image Quality

Safety(1)

Time Spent in the Clinic

Cost Efficiency

Patient Experience







Magnetic Resonance



X-Rav



X-Rav













30-45 min



45 min-1h





15-20 minutes











The QTI **Imaging Advantage**

...OVER HHUS

- Superior image quality
- Not operator dependent
- Quantifiable/repeatable

...OVER MRI

- · High resolution and contrast-to-noise ratio
- · No injection needed
- · Lower equipment cost
- No special facility or shielding requirements

...OVER XRM/DBT

- Improved image quality
- · Safer (no radiation), allowing for more frequent imaging
- Greater specificity
- No special facility requirements
- · Quantifiable/repeatable

...OVER BREAST CT

- No radiation breast CT radiation is significantly higher than screening mammography
- No contrast needed (compared to contrast enhanced CT)



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Imaging Accuracy in Breast Mass Diagnosis®

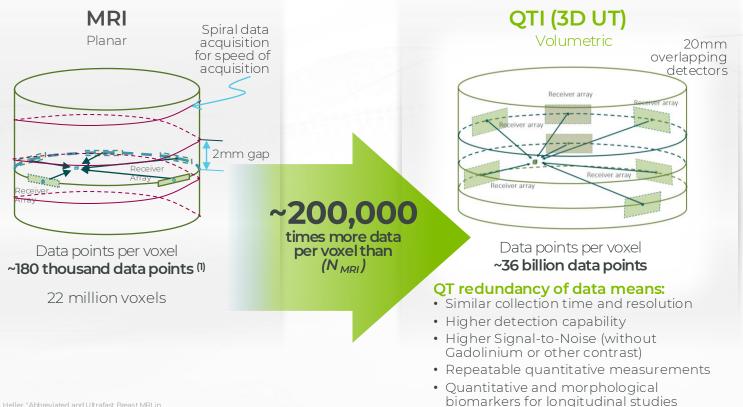
	QT Scan	XRM/DBT	HHUS	CE-MRI	СТ
Normal Breast					
Dense Breast		•			
Cyst Tumor		0		•	
Solid Tumor				•	
Calcification	0		•	0	
Quantitative Tissue / Density Characterization		•	\bigcirc	0	\bigcirc
Implant Visualization	•	0	•	•	



⁽¹⁾ Based on opinion of QT Imaging Holdings team.

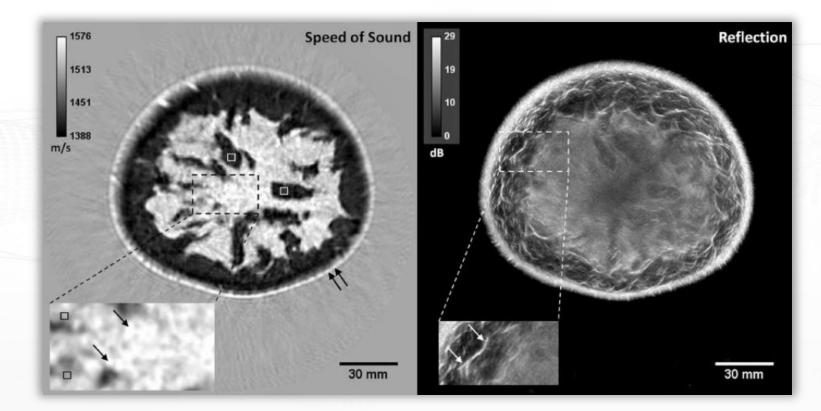
⁽²⁾ Quantitative tissue/density characterization means assessment of quantitative/volumetric breast density. Other than Mammography and QTI, there are no FDA cleared algorithms for volumetric density assessment.

QTI Provides High Resolution, Similar to MRI



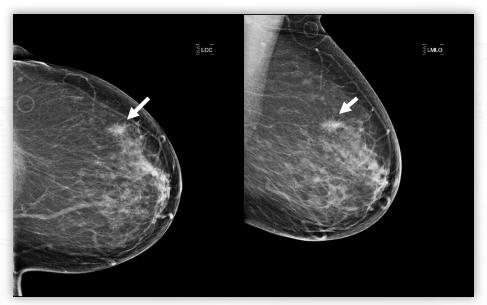


QT Speed of Sound and Reflection Images

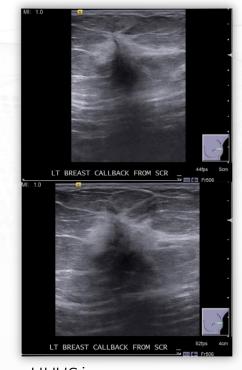




Modality Comparison – FFDM and HHUS



Mammogram - (left) CC and (right) MLO views. Arrows mark a region of spiculated focal asymmetry.

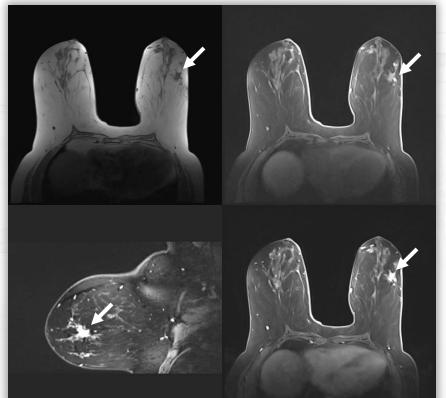


HHUS images across the lesion



Modality Comparison – MRI Images

Non-fat sat



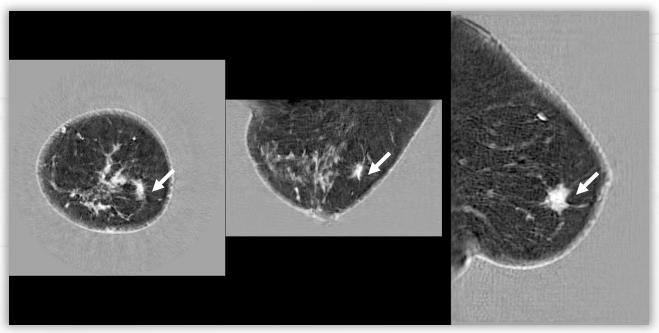
Pre-contrast

Fast low angle shot 3D (FL3D)





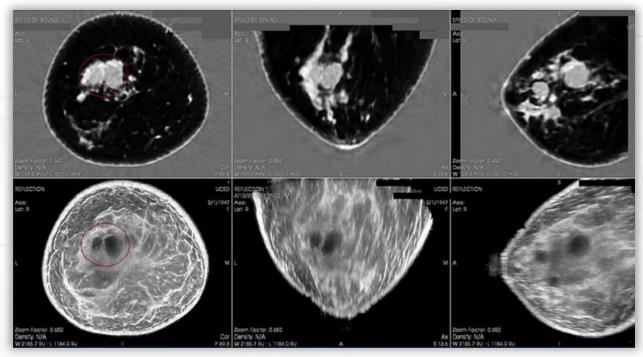
Modality Comparison – QT Image

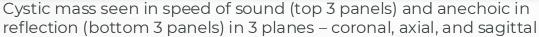


QT speed of sound image showing the mass (marked by arrows) as a region of high-speed IDC in lower outer quadrant of the left breast, 4 o'clock in the coronal view.



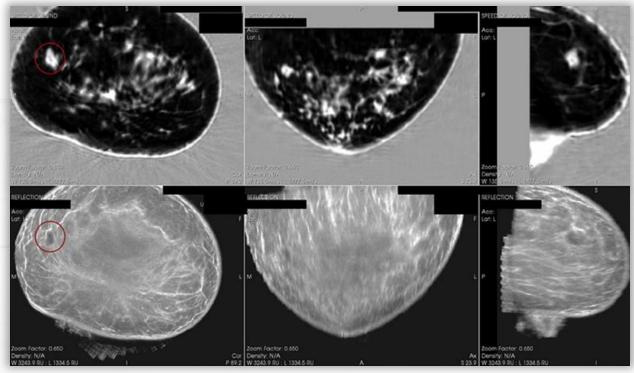
Cyst Identification Using Speed of Sound







Solid Identification Using Speed of Sound







From a Hardware Company...

By building a biomarker-driven platform, QT Imaging shifts from a hardware company to a precision imaging platform combining hardware, software and cloud-based AI modules





Hardware3D acoustic scanner

Software reconstruction, denoising, analytics

Cloud AI
biomarker extraction,
disease probability maps

...To a Sustainable, Data-Centric Ecosystem Continuously Improving with Clinical Use



Investment Highlights





respectively



Q3'25 Financials



Financial Highlights for Q3'25

- On October 3, 2025, we closed a over subscribed \$18.2 million private placement financing, which included anchoring from Sio Capital and participation from other institutional and existing company investors.
- On August 26, 2025, we received \$5.0 million in additional funding via a restated and amended senior secured term loan with Lynrock Lake, which allowed for the repurchase of the Yorkville warrant for \$5.0 million. The \$5.0 million was repaid to Lynrock Lake in October 2025.
- Commercial revenue was \$4.2 million during the third quarter of 2025, representing 339% year-over-year growth and 15% sequential quarter-over-quarter growth. The year-over-year increase in revenue was primarily attributable to the shipment of nine QT Breast Acoustic CT™ scanners during the third quarter of 2025, as per minimum order quantities ("MOQs") in the Company's Distribution Agreement with NXC Imaging, as compared to two scanners sold in the third quarter of 2024. In addition, the Company has shipped five more scanners during the month of October 2025, for a total of 28 scanners, in agreement with its distribution agreement.



Financial Highlights for Q3'25 QTD

- Gross margin of 43% in the third quarter of 2025, compared to gross margin of 63% in the second quarter of 2024
 - The decline in gross margin in the third quarter of 2025 was primarily attributable to variability in the weighted average cost related to the Company's existing inventory.
- Net loss of \$4.6 million for the third quarter of 2025, compared to net loss of \$3.6 million for the third quarter of 2024. Q3'25 net loss included:
 - \$2.3 million of net non-cash expense related to the change in fair value of earnout and warrant liabilities
 - \$0.2 million of stock-based compensation expense
- Non-GAAP Adjusted EBITDA of \$(1.4) million for the third quarter of 2025, compared to \$(2.2) million for the third quarter of 2024.



Financial Highlights for Q3'25 QTD

- Ended Q3'25 with \$1.7M in cash, compared to end of Q4'24 with \$1.7M in cash. Cash flows from operating activities was \$5.9 million year-to-date in 2025, and cash flows from financing activities was \$6.4 million, including \$15.0 million proceeds from the restated and amended senior secured term loan with Lynrock Lake, \$5.0 repurchase of the Yorkville warrant, and \$4.7 million repayment of the Yorkville and Cable Care notes.
- Reiterated plans to deliver \$18 million in revenue in 2025 (shipment of 40 scanners).
 The target is in accordance with the MOQs per our Amended Distribution Agreement
 with our strategic business and distribution partner, NXC Imaging, Inc., a wholly owned
 subsidiary of Canon Medical Systems USA.
- The Company revealed plans to deliver \$39 million in revenue in 2026 (shipment of 80 scanners). These targets are in accordance with the MOQs with the Distribution Agreement NXC Imaging in USA, as well as MOQs per our Distribution Agreement with Gulf Medical in Saudi Arabia, which was signed on August 27, 2025.



Summary of Q3'25 QTD GAAP Results

	Three Months Ended September 30,		Nine Month Septemb					
\$ thousands (except share and per share amounts)		2025		2024		2025		2024
Revenue	\$	4,192	\$	956	\$	10,650	\$	4,032
Cost of revenue		2,389		351		5,208		1,792
Gross profit		1,803		605		5,442		2,240
Operating expenses:								
Research and development		939		925		2,692		2,493
Selling, general and administrative	Т	2,516		2,007		6,487		9,873
Loss from operations		(1,652)		(2,327)		(3,737)		(10,126)
Interest expense, net		(565)		(1,455)		(1,635)		(3,149)
Other (expense) income, net		(30)		17		(8,770)		(191)
Change in fair value of warrant liability		(80)		9		(3,581)		200
Change in fair value of derivative liability		_		87		101		4,800
Change in fair value of earnout liability		(2,230)		50		(2,070)		2,970
Loss before income tax expense	\$	(4,557)	\$	(3,619)	\$	(19,692)	\$	(5,496)
Income tax expense		_		_		3		_
Net loss	\$	(4,557)	\$	(3,619)	\$	(19,695)	\$	(5,496)
Less: deemed dividend related to the modification of equity classified warrants		_		_		_		(5,186)
Net loss attributable to common stockholders	\$	(4,557)	\$	(3,619)	\$	(19,695)	\$	(10,682)
	Т							
Basic and diluted net loss per share (1)	\$	(0.47)	\$	(0.51)	\$	(2.09)	\$	(1.72)
Weighted average shares outstanding (1)		9,601,972	_	7,155,505		9,415,349	(5,245,877

⁽¹⁾ Share and per share amounts for the three and nine months ended September 30, 2024 differ from those published in prior con@nsed consolidated financial statements as they were retrospectively adjusted as a result of the Reverse Stock Split fas described below in Note1 to the condensed consolidated financial statements), Specifically, the number of shares of common stock outstanding during periods before the Reverse Stock Split are divided by the exchange ratio of 31, such that each three shares of common stockwere combined and reconstituted into one share of common stock effective October 23, 2025.



Summary of Q3'25 QTD Non-GAAP Results

		hree Mont Septem	Nine Months Ended September 30,				
\$ thousands		2025	2024		2025		2024
Net loss	\$	(4,557)	\$ (3,619)	\$	(19,695)	\$	(5,496)
Interest expense, net		565	1,455		1,635		3,149
Income tax expense		_	_		3		_
Depreciation and amortization		40	20		115		204
EBITDA		(3,952)	(2,144)		(17,942)		(2,143)
Adjustments:							
Stock-based compensation		199	127		519		166
Warrant modification		_	_		_		201
Debt modification and extinguishment expenses ⁽¹⁾		46	_		2,170		_
Change in fair value of warrant liability ⁽²⁾		80	(9)		3,581		(200)
Change in fair value of derivative liability ⁽³⁾		_	(87)		(101)		(4,800)
Change in fair value of earnout liability ⁽⁴⁾		2,230	(50)		2,070		(2,970)
Transaction expenses (5)		_	_		_		4,301
Debt issuance expense (6)		_	_		6,640		_
Adjusted EBITDA	\$	(1,397)	\$ (2,163)	\$	(3,063)	\$	(5,445)



Adjustments to EBITDA

- (1) The Company recorded debt modification expense of \$0.1 million primarily related to its modification of the Cable Car Note on January 9, 2025 and debt extinguishment expense of \$2.0 million related to the extinguishment of the Yorkville Note and Cable Car Note on February 26, 2025 in other (expense) income, net for the nine months ended September 30, 2025.
- (2) The increase in fair value of warrant liability during the nine months ended September 30, 2025 relates to the liability classified private placement warrants, the Lynrock Lake Warrant and Yorkville Warrant, which is primarily driven by increase in the Company's stock price from beginning of period to June 11, 2025, which is the date the Lynrock Lake Warrant and Yorkville Warrant were modified and subsequently reclassified to equity.
- (3) The decrease in fair value of derivative liability during the nine months ended September 30, 2025 related to the Yorkville Pre-paid Advance, which contained features that were bifurcated as freestanding financial instruments and initially valued on March 4, 2024 upon consummation of the Merger. The derivative liability was subsequently revalued as of February 26, 2025, prior to the extinguishment of the Yorkville Note.
- (4) The earnout liability relates to the contingent consideration for the Merger Earnout Consideration Shares pursuant to the Business Combination Agreement dated December 8, 2022, as amended in September 2023. The earnout liability was initially valued using the Monte Carlo Simulation method on March 4, 2024 and subsequently revalued using the same method.
- (5) The Company incurred transaction expenses related to the Merger with GigCapital5, Inc., which closed on March 4, 2024. These transaction expenses included a \$3.7 million of transaction costs that were settled with issuance of common stock, \$0.4 million of transaction costs settled or payable in cash and a \$0.2 million loss on issuance of common stock in connection with a subscription agreement, which were recorded as selling, general and administrative expenses in the condensed consolidated statement of operations during the nine months ended September 30, 2024. There were no transaction expenses incurred during the nine months ended September 30, 2025.
- (6) Upon the issuance of Lynrock Lake Term Loan closed on February 26, 2025, the Company recorded a loss of \$6.6 million, including debt issuance costs of \$0.2 million, in other expense, net for the nine months ended September 30, 2025.



Balance Sheets as of Q3'25 and Q4'24

\$ in thousands	Se	September 30, 2025		
Assets				
Current assets:				
Cash	\$	1,715	\$	1,172
Restricted cash and cash equivalents		20		20
Accounts receivable, net		3,244		67
Inventory		5,242		3,141
Prepaid expenses and other current assets		1,067		517
Total current assets		11,288		4,917
Non-current assets:				
Property and equipment, net		127		196
Operating lease right-of-use assets, net		667		935
Other assets		39		39
Total assets	\$	12,121	\$	6,087

\$ in thousands		September 30, 2025	December 31, 2024		
Liabilities and Stockholders' Deficit					
Current liabilities:					
Accounts payable	\$	2,019	\$ 803		
Accrued expenses and other current liabilities		5,235	3,550		
Current maturities of long-term debt		5,023	4,986		
Deferred revenue		24	49		
Operating lease liabilities, current		442	406		
Total current liabilities		12,743	9,794		
Non-current liabilities:					
Long-term debt		273	9		
Related party notes payable		3,895	3,849		
Operating lease liabilities		321	657		
Warrant liability		106	22		
Derivative liability		_	304		
Earnout liability		2,510	440		
Other liabilities		1,349	550		
Total liabilities		21,197	15,625		
Stockholders' deficit:					
Common stock (1)		1	1		
Additional paid-in capital !1)		42,558	22,402		
Accumulated deficit		(51,635)	(31,941		
Total stockholders' deficit		(9,076)	(9,538		
Total liabilities and stockholders' deficit	\$	12,121	\$ 6,087		

⁽¹⁾ Share amounts as of December 31, 2024 differ from those published in prior consolidated financial statements as they were retrospectively adjusted as a result of the Reverse Stock Split (as described below in Note 1, The Company and Summary of Significant Accounting Policies), Specifically, the number of shares of common stock outstanding during periods before the Reverse Stock Split are divided by the exchange ratio of 31, such that each three shares of common stock were combined and reconstituted into one share of common stock effective October 23, 2025.



Cash Flow Statements for Q3'25 YTD and Q3'24 YTD

		Nine Months Ended September 30,				
\$ in thousands		2025	2024			
Cash flows from operating activities:						
Net loss	\$	(19,695)	\$ (5,49			
Adjustments to reconcile net loss to net cash used in operating activities:						
Depreciation and amortization		115	20			
Stock-based compensation		519	16			
Warrant modification expense		_	20			
Loss on issuance of the Lynrock Lake Term Loan		6,640	-			
Debt extinguishment loss		2,080	-			
Debt modification expense		90	-			
Provision for credit losses		_				
Fair value of common stock issued in exchange for services and in connection with non- redemption agreements		_	3,71			
Loss on issuance of common stock in connection with a subscription agreement		_	20			
Non-cash interest		749	2,40			
Non-cash operating lease income		(31)	(2			
Change in fair value of warrant liability		3,581	(20			
Change in fair value of derivative liability		(101)	(4,80			
Change in fair value of earnout liability		2,070	(2,97			
Changes in operating assets and liabilities:						
Accounts receivable		(3,177)	(25			
Inventory		(2,102)	1,52			
Prepaid expenses and other current assets		(298)	(45			
Accounts payable		1,072	(2,06			
Accrued expenses and other current liabilities		1,856	(76			
Deferred revenue		(25)	(32			
Other liabilities		799	12			
Net cash used in operating activities		(5,858)	(8,80			

	Nine Months Ended September 30,				
\$ in thousands	2025	2024			
Cash flows from investing activities:					
Purchases of property and equipment	(47)	(35			
Net cash used in investing activities	(47)	(35			
Cash flows from financing activities:					
Proceeds from sale of common stock and warrants, net of issuance costs	679	_			
Proceeds from issuance of common stock pursuant to subscription agreement, net of issuance costs	_	500			
Proceeds from long-term debt, net of issuance costs	15,000	10,525			
Proceeds from stock option exercises	75	_			
Proceeds from warrant exercises	532	_			
Repayment of long-term debt	(4,688)	(1,243			
Repayment of bridge loans	_	(800			
Payment of deferred issuance costs	(150)	_			
Proceeds from the Merger, net of transaction costs	_	1,238			
Repurchase of warrant from Yorkville	(5,000)	-			
Net cash provided by financing activities	6,448	10,220			
Net increase in cash and restricted cash and cash equivalents	543	1,379			
Cash and restricted cash and cash equivalents at the beginning of period	1,192	185			
Cash and restricted cash and cash equivalents at the end of the period	\$ 1,735	\$ 1,564			





Thank You!

