UNITED STATES SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported) $\underline{December~11,2024}$

QT IMAGING HOLDINGS, INC.

(Exact name of Registrant as Specified in Charter)

Delaware

86-1728920

Delaware	001-40837	00-1728720
(State or Other Jurisdiction of	(Commission	(IRS Employer
Incorporation or Organization)	File Number)	Identification Number)
	3 Hamilton Landing, Suite 160	
	Novato, CA 94949	
	(Address of principal executive offices, including Zip Code)	
	(650) 276-7040	
	(Registrant's telephone number, including area code)	
Check the appropriate box below if the Form 8-K filing is intended to simultaneously s	atisfy the filing obligation of the registrant under any of the following	provisions (see General Instruction A.2. below):
☐ Written communications pursuant to Rule 425 under the Securities Act (1	7 CFR 230.425)	
☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 C	CFR 240.14a-12)	
☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the	Exchange Act (17 CFR 240.14d-2(b))	
☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the	•	
_ ····································		
Securities registered pursuant to Section 12(b) of the Act:		
	Trading	Name of each exchange
Title of each class	Symbols	on which registered
Common stock, \$0.0001 par value	QTI	The Nasdaq Stock Market LLC
•	-	•
Indicate by check mark whether the registrant is an emerging growth company as defin-	ed in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter	r) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this
chapter).		
Emerging growth company		
If an emerging growth company, indicate by check mark if the registrant has elected no	ot to use the extended transition period for complying with any new of	or revised financial accounting standards provided pursuant to Section 13(a) of
the Exchange Act.		

Item 7.01 Regulation FD Disclosure.

On December 11, 2024, QT Imaging Holdings, Inc. (the "Company") posted to the Company's Investor Presentations section of its website www.qtimaging.com, an investor presentation containing supplemental product and operational information regarding the Company. A copy of the investor presentation is being furnished as Exhibit 99.1 to this Current Report on Form 8-K.

The information contained in, or incorporated into, this Item 7.01 of this Report, including Exhibit 99.1 attached hereto, is furnished under Item 7.01 of Form 8-K and shall not be deemed "filed" for the purposes of Section 18 of the Exchange Act or otherwise subject to the liabilities of that section, and shall not be deemed to be incorporated by reference into the filings of the Company under the Securities Act or the Exchange Act regardless of any general incorporation language in such filings.

This Report shall not be deemed an admission as to the materiality of any information in this Report that is being disclosed pursuant to Regulation FD.

Please refer to Exhibit 99.1 for a discussion of certain forward-looking statements included therein and the risks and uncertainties related thereto.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits:

Exhibit No.	Description
99.1	Investor Presentation dated December 2024
104	Cover Page Interactive Data File (embedded within the Inline XBRL document)

SIGNATURE

	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 t	the undersigned hereunto duly authorized.
--	---	---

December 11, 2024 Dated:

By: /s/ Raluca Dinu Raluca Dinu

Name:

Title: Chief Executive Officer





Quantitative Transmission Imaging

Breast Acoustic CT™ Scanner

INVESTOR PRESENTATION December 2024





Disclaimer

ABOUT THIS PRESENTATION

This investor presentation (this "Presentation") is provided for informational purposes only. The information contained herein does not purport to be all-inclusive and neither QT Imaging Holdings, inc. (the "Company", "QT Imaging Holdings", "QTI"), nor its respective directors, officers, employees, agents, advisors or affiliates, including QT Imaging, inc. ("QT Imaging"), makes any representation or warranty, express or implied, as to the accuracy, completeness or reliability of the information contained in this Presentation, which has not been verified and is subject to change at any time. Viewers of this Presentation should each make their own evaluation of QT Imaging Holdings and of the relevance and accuracy of the information and should make such other investigations as they deem necessary. To the fullest extent permitted by law, no responsibility or liability whatsoever is accepted by QT Imaging Holdings, or its directors, officers, employees, agents, advisors or affiliates for any loss howseever arising, directly or indirectly, from any use of this Presentation or such information or opinions contained herein or otherwise arising of inconnection herewith.

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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 "safe", "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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FORWARD LOOKING STATEMENTS

Certain statements included in this Presentation that are not historical facts are forward-looking statements for purposes of the safe harbor provisions under the United States Private Securities Litigation Reform Act of 1995. Forward-looking statements generally are accompanied by works such as "has the potential to", "believe", "may", "will", "stetimate", "continue", "anticipate", "intend", "expect", "should", "would", "plan", "predict", "potential", "seem", "seem", "future", "outlook", and similar expressions that indicate or predict future events or trends that are not statements of historical matters. These forward looking statements include, but are not limited to, the potential impact on existing medical technology, the company's technology, broducts, business prospects, revenue, client adoptions, commercialization, projections of market opportunity and statements regarding estimates and forecasts of other financial and performance metrics. These statements are based on various assumptions, whether or not identified in this Presentation, and on the current expectations of QT Imaging Holdings' management and are not predictions of actual performance. These forward-looking statements are provided for illustrative purposes only and are not circumstances intended to serve as, and must not be relied on by any investor as, a guarantee, an assurance, a prediction or a definitive statement of fact or probability. In addition, statements regarding the Company's products, technology, and market opportunity reflect the beliefs and opinions of QT Imaging Holdings' management on the relevant subject as of this Presentation. Actual events and circumstances are difficult or impossible to predict and will differ from assumptions. Many actual events and circumstances are beyond the control of QT Imaging Holdings' products and services; the ability of QT Imaging Holdings' business and the timing of expected business milestones; the demand for QT Imaging Holdings' products and services; the ability of QT Imaging Holdings t



Disclaimer

NON-GAAP FINANCIAL MEASURES

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 November 13, 2024, and the reconciliation of EBITDA and Adjusted EBITDA can be found on pages 38 and 39 of this presentation.

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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 relatively low-cost, comprehensive, no radiation, no discomfort medical imaging solution
- QTI's technology yields superior performance compared to traditional mammogram with regard to specificity and sensitivity and has similar imaging quality and diagnostic value compared to MRI but is a lower cost and more accessible solution

onwight 02024 OT Imaging Holdings by: All Dights Deserved



Our Mission

- Create disruptive innovation using technology (software, machine learning, and smart physics) to improve medical imaging and thus, healthcare quality and access
- Continue to build upon our FDA clearances to offer QTI as a breast screening imaging modality
- Expand the market opportunities beyond hospitals, imaging centers and health centers by supporting additional direct to consumer (DTC) and direct to provider (DTP) approaches
- Introduce the first comprehensive body-safe imaging technology, enabling for the first-time well-person body imaging health screening

NIH has
awarded QT
Imaging about
\$18M
for new women's
imaging solution



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Introduction to the QT Imaging Holdings Management Team



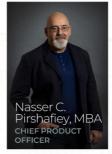
Dr. Raluca Dinu is a global business executive, with long public companies' governance experience, offering over 22 years of achievements in the high-tech industry, with an established track record of driving increased revenue and profitability, delivering strong results in turnaround or M&A situations, leading strategic growth, and consolidation in fast-paced business environments.



Mr. Budagov is serving as CFO of QTI since December 2023. He has more than 15 years of accounting and consulting experience, including consulting public and private clients. Additionally, he has 3 years of audit experience at Ernst & Young.



Mr. Steve Choate, appointed as Chief Operations Officer at QTI in April 2024, is responsible for managing the operations organization, ensuring quality, and fostering collaboration with internal, domestic, and international manufacturing partners.



Mr. Pirshafiey has been with QTI since 2017. Previously, he founded and managed a consulting firm providing sustainable practices to industries including medical device, high-tech, and consumer products for giants such as Johnson and Siemens. He has 144 inventions filed with the US patent office.



Dr. Bilal Malik has over ten years of experience in research, development, and translation of medical devices, both in academia and industry. He is an expert in leading and directing efforts in image and data science and has a track record of successfully leading innovation for medical imaging products.

QTIMAGING

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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 ultrasound principle, with quantitative measure of the intrinsic speed of sound in Breast Tissue
- Standardized scanning with operator independent images, unlike hand-held ultrasound (HHUS)
- Better resolution compared 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

CLINICAL CONSIDERATIONS

- Evidence Available: Accuracy in comparison with X-ray Mammography and DBT, Sensitivity, Specificity, and Density
- · Clinical Trials in Pipeline





Distribution Agreement with NXC Imaging A Subsidiary of Canon Medical Systems

- Under Distribution Agreement with NXC Imaging for US market
- Accessing NXC Imaging's distribution channel in the US and the US territories, this agreement provides potential to accelerate the commercial roll-out of QTI's imaging systems
- NXC Imaging provides a mature service organization to support QTI's installed base



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Successful Completion of Feasibility Study with Canon Medical Systems

- Canon evaluated the business, technical, and clinical values of QTI's ultrasound breast scanner including:
 - -product quality validation
 - -development and manufacturing studies
 - -clinical evaluation
 - -regulatory investigation
 - -market validation
- All know-how and intellectual property embodied in the QT Scanner are owned by QTI.



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Participation at RSNA 2024



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QTI's Technology Has the Opportunity to Transform Several Large Markets

2023 GLOBAL MEDICAL IMAGING MARKET SIZE: \$40B(1)

Current Market

BREAST: \$5B MARKET(2)

- 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



ORTHO: \$9B MARKET(3)

- Target replacing MRI examinations
- Primary focus on orthopedic practices



Future Markets - Body Scanner Platform Development INFANT: \$8B MARKET (4)

New market opportunity given limitations of current imaging modalities for infants

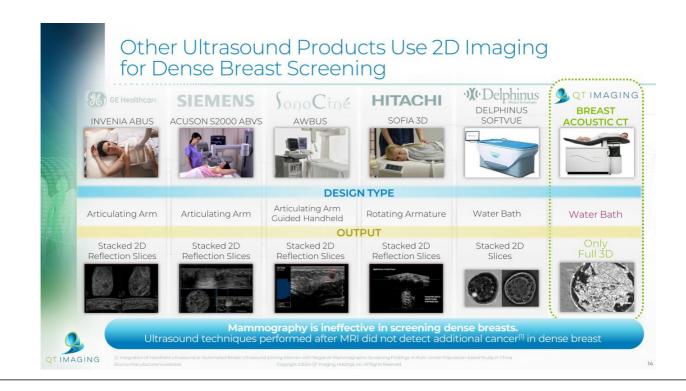


IMAGE-GUIDED PROCEDURES: \$5B MARKET⁽⁵⁾

- Commenced feasibility
- Variety of image-guided procedures including biopsies, injections and cryoablation









Current Ultrasound Technologies Have Major Deficiencies

Shortfalls of Commercial Current, Rival Systems⁽²⁾:

- Reflection images have speckle; compounding without refraction correction
- No valid true "transmission" mode use "shear wave" (low resolution) data (ABUS, AVUS, etc. are not transmission)
- Data yielded is compounded 2D not true "3D" Transmission images have artifacts.
- Low contrast-to-noise ratios (speckle)
- Specificity for masses is relatively poor
- Unable to view consistently calcifications misses 20% of cancers(1)
- No "functional" imaging features for most(doubling time, tissue identification and specific tissue volume segmentations)
- Poor reproducibility of measurement and volume data
- Operator dependence (HHUS)

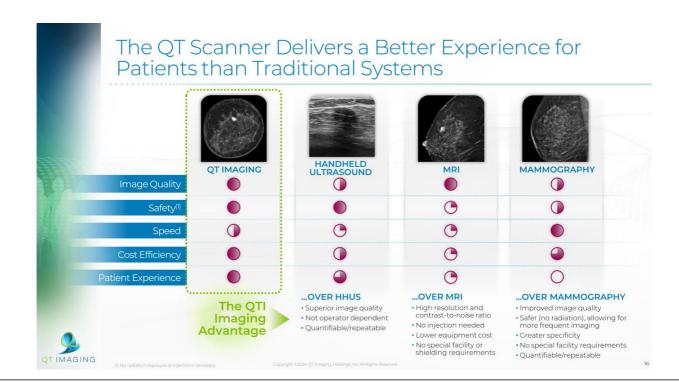
(i) A Multireader Multicase (MRMC) Receiver Operating Characteristic (ROC) Study Evaluating Noninferiority of Quantitative Transmission (QT) Ultrasound to Digital Breast Tomosynthesis (DBT) on Detection and Recall of Breast Lesions

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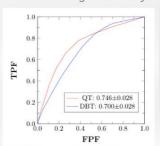
Clinical Evidence: Non-Inferiority to DBT

Non-inferiority to DBT

- Breast Abnormalities
- · Benign, non-cancer, normal without biopsy
- Cancer, abnormal with biopsy
- Different types of breast lesions (solid, cysts, complex)

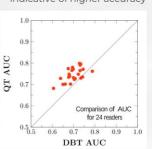
ROC curves

The line closer to 1.0 is indicative of higher accuracy



Individual reader AUC





QTI technology is a **potential alternative to mammography for breast cancer screening of women too young** to undergo DBT.

Ref.: A MultiPleader Multicase (MEMC) Receiver Operating Characteristic (ROC) Study Evaluating Nominferiority of Quantitative Transmission (27) Ultrasound to Digital Breast Tomosynthesis (DBT) o Detection and Recall of Breast Lesions: Yulei Jiang, PhD, Blain leuranow, MD, Biglial Malik, PhD, John Klock, MD, Academic Radiology, Vol 31, No. 5, June 2024.

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TABLE 3. Sensitivity and Specificity Based on Call-back vs No Call-back Decisions of 24 Readers and 177 Cases (66 Abnormal, 111 Normal)

	Modality	Average ± SD (%)	95% CI
Sensitivity	DBT	85.2 ± 6.4	[83.1, 87.1]
	QT	70.6 ± 7.2	[68.3, 72.8]
	QT-DBT	-14.5 ± 8.9	[-17.2, -11.7]
Specificity	DBT	37.2 ± 11.0	[33.6, 40.7]
	QT	60.1 ± 12.3	[56.4, 64.0]
	QT-DBT	22.9 ± 10.5	[19.8, 26.1]

CI, confidence interval; DBT, digital breast tomosynthesis; QT, quantitative transmission; SD, standard deviation.
^a Estimated from 1000 bootstrapping samples of the source data.

• Sensitivity

- Lower for QT (70.6%) compared to DBT (85.2%)

- Potentially attributable to reader unfamiliarity with QT imaging, suggesting a need for enhanced training

Specificity

- Significantly higher for QT (60.1%) compared to DBT (37.2%)
- Indicates QT's ability to better differentiate benign from malignant lesions

0.8
0.6
0.6
0.7
0.9
0.0
0.0
0.0
0.2
0.4
0.6
0.8
1.0
FPF





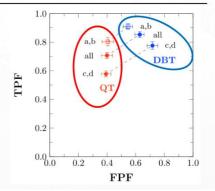


Subgroup Analysis: Dense Breasts

TARLE 4 MRMC Analysis Results by	Breast Density of 24 Readers and 177 Cases	(66 Abnormal 111 Normal)

BI-RADS Density	N	AUC ± SE ^a		95% CI	
	Abnormal/Normal	QT	DBT	QT-DBT	
c, d	28/53	0.6852 ± 0.0457	0.5987 ± 0.0447	0.0865 ± 0.0557	[-0.0227, 0.1956]
a, b	38/58	0.7912 ± 0.0335	0.7791 ± 0.0325	0.0121 ± 0.0242	[-0.0353, 0.0596]

- Both sensitivity and specificity of DBT are dependent on breast density
- Specificity of QT is independent of breast density



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Clinical Evidence:

Anatomic & Visual Grade with Comparative Modality

Normal Anatomic Comparison

- Visual Graded Analysis
- Compared QTI V/s HHUS, XRM
- Graded Equivalent or Better than XRM/ HHUS

X-Ray Mammography (XRM)

- 4 readers
- 22 breast , 20 subjects
- Lower score means better visualization



Handheld Ultrasound (HHUS)

- 5 readers
- 17 breast , 17 subjects
- Lower score means better visualization



QTI technology is **highly accurate in visualizing the ductal and glandular tissue, even in dense breasts** where such visualization can be challenging using conventional breast imaging technologies like XRM and/or HHUS.

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Fig. 13 hor & Dicko, Elaine Luanow, Kathleen Smith, Nancy A and Obuchowski (2017) Visual Grading Assessment of Quantitative Transmission Ultrasound Compared to Digital X-ray Mammography an

Hand-held Ultrasound in Identifying Ten Breast Anatomical Structures. Clinical Trials 3: 015

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The Current Breast Imaging Paradigm Leads to Unnecessary Concern and Costs For every 1,000 screening mammograms: Screening compliance is low 35% of women aged 40-70 do not get screened.(1) ********** CALL BACK ****** 98% of RATES ~15% call-backs 150 35% ******* Recalls are ****** 65% Avoidable rates with mammography Do not follow guidelinesFollow guidelines ****** ******* Of the 65% of women who do get Over 80% screened, many suffer through **BIOPSIES** of Callback unnecessary callbacks ~10% biopsy rate for callbacks Biopsies are Benign⁽⁴⁾ 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⁾ CANCER INCIDENCE 0.3% cancer diagnosis⁽⁵⁾



Divery Wel Health | 3 Beasons for a Mammogam Calback | Larell Scardeli | | SupubMed | False-Neigathe Rate of Combined Mammography and Ultrabound for Women with Palpable Breast Masses | Carlos H.F. Chan, Suzanne B. Coopey, Phoebe E. Freer, and Kevin S. Hugh alkatonal Breast Cancer Foundation | Breast Blogay Procedure Types, What to Expect and Results

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Clinical Evidence Recall Rate

- Recall Rates: 10% Combined Recall Rate
- · Adherence to screening compliance:
 - 16% Decrease in Non-Cancer recall
 - 2% Decrease in Cancer Recall



accuracy and detection rate

· Anxiety Reducing Factor

QTI technology improves non-cancer recall rates without substantially affecting cancer recall rates



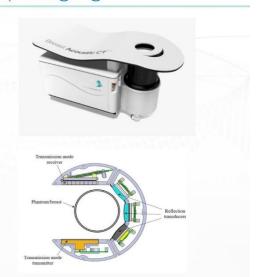




Quantitative Transmission (QT) Imaging

- What is QT Imaging?
 - Inherently 3D volumetric ultrasound modality due to 3D data acquisition and image reconstruction
 - Uses CT-like configuration with ultrasound to acquire and reconstruct **transmission images** which map the **speed-of-sound across the tissue volume**
 - High resolution, similar to MRI
 - Images tissue **without overlap**, providing more information than conventional HHUS
 - Overcomes operator dependence and lack of standardization associated with HHUS
 - Pain free, safe
- Image Acquisition:
 - Prone position with breast submerged in water
 - 360-degree rotation of ultrasound arrays
 - 10-12 minutes per breast average scan time

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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.



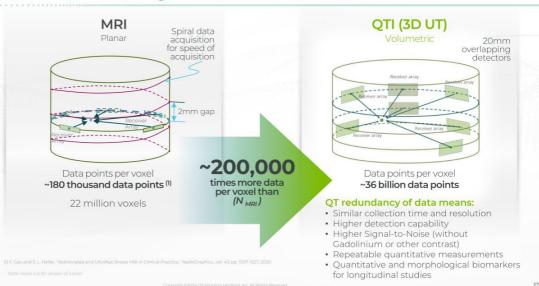




PerfeQTion Imaging Center – Haverford, PA



QTI Provides High Resolution, Similar to MRI





Technical Capabilities

- Resolution of ~600 microns in reflection compared to 800 microns⁽¹⁾ for MRI (depends on field strength, homogeneity etc)
- Contrast to noise ratio of 23:1 at 100 microns (in reflection; can detect small calcifications)
- Contrast to noise ratio of 15:1 (at resolution in transmission speed of sound)
- Speckle-free because of 360° compounding and refraction correction for reflection image
- Volumetric data acquisition (3D), not stacked 2D slices
- Volumetric reproducibility 0.2% for fibro glandular volume
- Volumetric accuracy better than 3% extrapolated from linear accuracy ~1% (vertical < 2%)



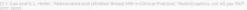
V. Gao and S. L. Heller, "Abbreviated and Ultrafast Breast MRI in Clinical Practice," RadioCraphics, vol. 40, pp. 1507-1527, 2020.

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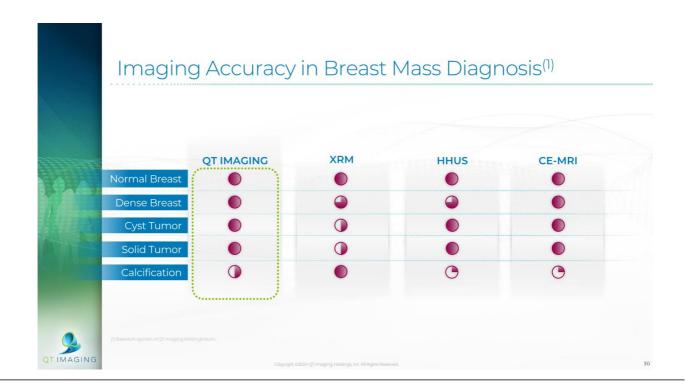
Enhanced Clinical Capabilities and Value

- High-quality and high-resolution native 3D Imaging
- Quantifiable images enables accurate analysis, comparison and trending
- Consistent and reproduceable image quality regardless of operator or breast size/tissue type
- Clinical feature detection of 50-100 microns including microcalcifications
- Functional imaging capability determine tissue type from the speed of sound
- Allows tissue doubling time assessments - similar to MRI and CT
- Highly accurate measurements, not scanner operator dependent

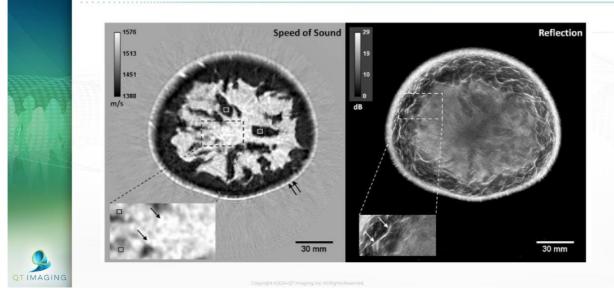


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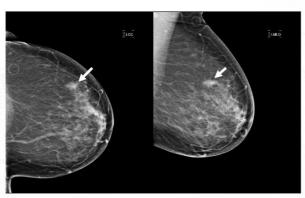


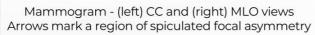














HHUS images across the lesion

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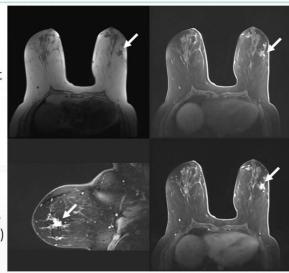
3.



Modality Comparison – MRI Images

Non-fat sat

Fast low angle shot 3D (FL3D)

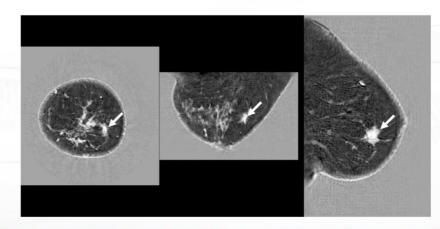


Pre-contrast

Post-contrast

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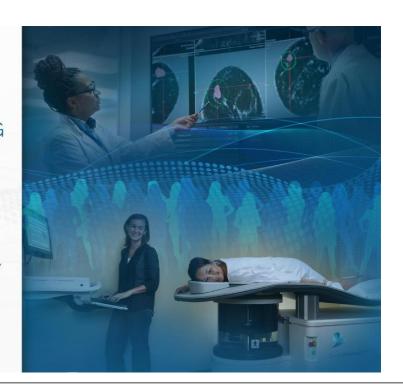


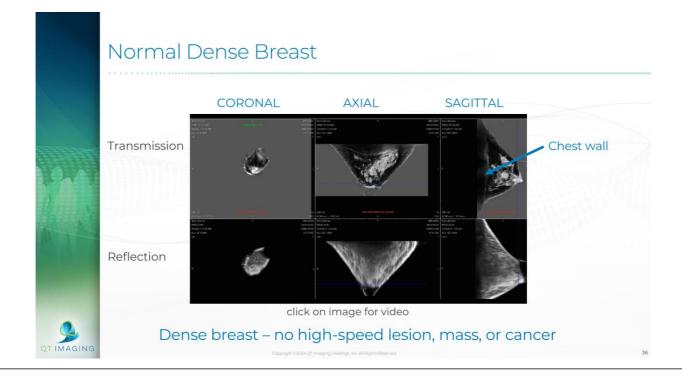


QT speed of sound image showing the mass (marked by arrows) as a region of high-speed IDC in lower outer quadrant of the left breast, 4 o'clock in the coronal view.

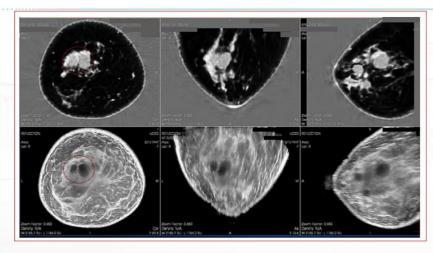
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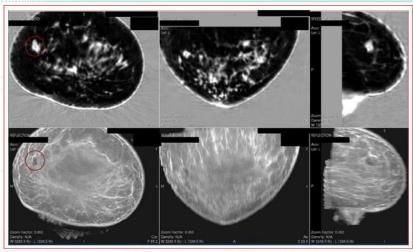


Cystic mass seen in speed of sound (top 3 panels) and anechoic in reflection (bottom 3 panels) in 3 planes – coronal, axial, and sagittal

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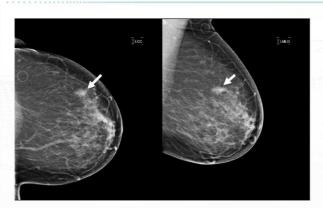


Solid Identification Using Speed of Sound



Solid mass seen in speed of sound (top 3 panels) and hypoecoic in reflection (bottom 3 panels) in 3 planes – coronal, axial and sagittal









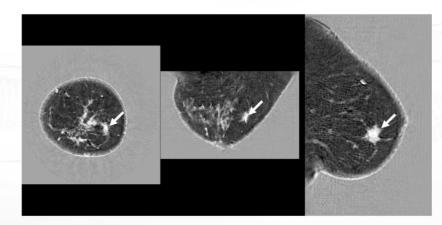
HHUS images across the lesion

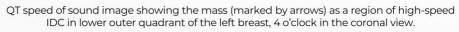
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Case 1: QTI Scan





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Case 2: MRI – Invasive Ductal Carcinoma

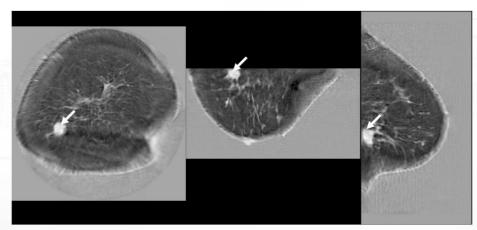
Non-fat sat

FL3D

Pre-contrast

Post-contrast

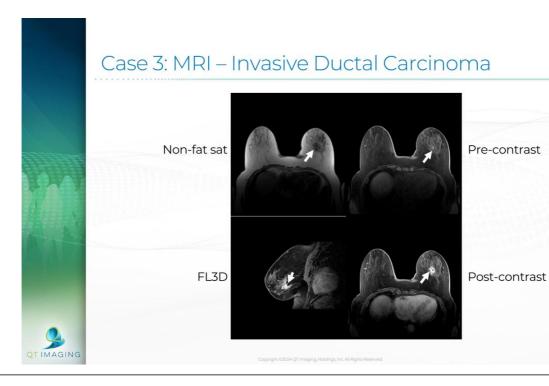
Case 2: QTI Speed of Sound



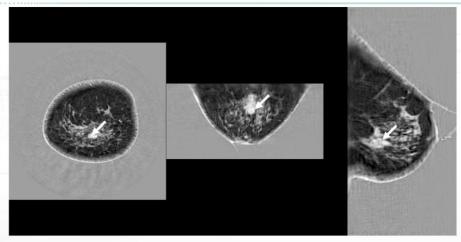
QT speed of sound image showing the mass (marked by arrows) as a region of high-speed. In coronal view at 8 o'clock. Relatively posterior mass but still all visible in the QT image.

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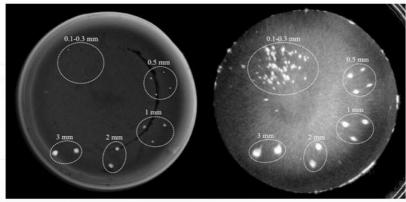




QT speed of sound image showing the mass (marked by arrows) as a region of high-speed. In coronal view, at 6 o'clock, near the center.

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Digital Mammography

QT Reflection Tomogram

Detectability of calcifications in QT Acoustic CT is superior to XRM (1)

Sensitivity of Quantitative Transmission ultrasound to detection of microcalcifications. SPIE (International Society for Optics and Photonics) Meeting Houston Texas February 20, 2018.

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Market Positioning of Breast Acoustic CT Scanner

Not intended to compete with mammography for screening, although many patients may find it preferrable for:

- Dense breasts
- Implants
- Post therapy screening where breasts can be sensitive to compression
- When concerned about radiation dose

Diagnostic alternative to MRI

- · Lower cost, faster, more accessible
- Similar image quality and diagnostic value
- More tolerable for patient (claustrophobia, noise, time, no contrast)
- Images are inherently quantitative and repeatable, and hence serve as an imaging biomarker (helps following a patient)
- Scanner is easily deployable (*2 days) and frees MRI scanners for other non-breast imaging studies

Diagnostic alternative to Hand-held Ultrasound

- Native 3D imaging (like MRI and CT)
- Quantifiable image analysis
- No need for specialized technologist training
- Consistent and reproducible image quality regardless of operator

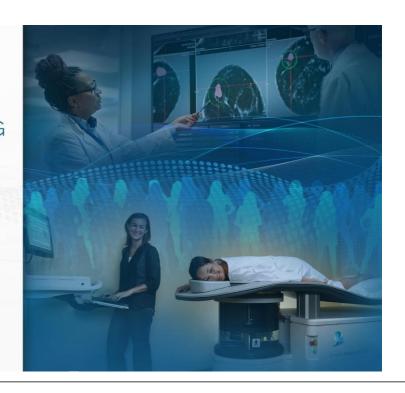




QT Scanner Locations Map









Developing an Open Angle Scanner Will Expand the Technology to New Markets

...providing significant potential to access new markets and applications

- The Open Angle Scanner uses an open, partial angle configuration which reduces the viewing field from 3600 to 3250 and provides additional capabilities for QTI technology in:
 - Orthopedic imaging
 - Prostate imaging
 - Whole body infant scanning
 - Biopsy and image-guided diagnostic and treatment procedures
- The scanner satisfies the need for better image reconstruction techniques in partial-ring tomography systems
- Potential to prevent cancers from developing into advanced stages
- · Representative point-of-care target markets include:

ORTHOPEDIC SURGEONS [IN-OFFICE]







MILTARY [SHIPS 8











Financial Highlights for Q3'24 QTD and YTD Ending 09/30/24

- Commercial revenue was \$1.0 million during the third quarter of 2024, compared to \$1.7 million during
 the second quarter of 2024 and less than \$0.1 million during the third quarter of 2023.
- Commercial revenue was \$4.0 million for the nine months ended September 30, 2024, compared to less than \$0.1 million for the nine months ended September 30, 2023.
- Gross margin of 63% in the third quarter of 2024, compared to gross margin of 51% in the second quarter of 2024 and insignificant margin in the third quarter of 2023. The increase in margin in the third quarter of 2024 compared to the second quarter of 2024 was attributable to variability in the weighted average cost related to the Company's existing inventory. The increase in margin during 2024 was due to the sale and delivery of two QT Breast Scanners during the third quarter of 2024, compared to no deliveries in the third quarter of 2023.
- Gross margin of 56% during the nine months ended September 30, 2024, compared to negative
 margin in 2023 YTD due to the sale and delivery of nine QT Breast Scanners in 2024 YTD, compared to no
 deliveries in 2023 YTD.

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5.



Financial Highlights for Q3'24 QTD and YTD Ending 09/30/24

- Net loss of \$3.6 million for the third quarter of 2024, compared to net loss of \$1.4 million for the third quarter in 2023.
 Q3'24 net loss includes:
 - \$0.1 million of net non-cash income related to the change in fair value of the warrant, derivative, and earnout liabilities and \$0.1 million of stock-based compensation expense, compared to a net loss of \$1.4 million for the third quarter of 2023, which included stock-based compensation expense of \$0.2 million and one-time transaction expenses of \$0.3 million.
- Net loss of \$9.2 million for the first nine months of 2024, compared to net loss of \$4.6 for the first nine months of 2023. 2024 YTD net loss includes:
 - \$4.3 million of net non-cash income related to the change in fair value of warrants, derivatives, and contingent consideration that were recorded as part of the closing of the business combination with GigCapital5, Inc. on March 4, 2024 and outstanding as of September 30, 2024, \$0.2 million of stock-based compensation expense, \$0.2 million warrant modification expense, and \$4.3 million of one-time transaction expenses.
- Non-GAAP Adjusted EBITDA of \$(2.2) million for the third quarter of 2024, compared to \$(0.6) million for the third quarter of 2023.
- Non-GAAP Adjusted EBITDA of \$(5.4) million for the first nine months of 2024, compared to \$(2.0) million for the
 first nine months of 2023.
- The company ended Q3'24 with \$1.6M in cash, compared to end of Q2'24 with \$4.6M in cash, which includes a \$1.5 million payment to Yorkville, of which \$1.145 million was principal and the remainder was accrued interest.

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Summary of Q3 QTD and YTD Ending 09/30/24 GAAP Results

	1	hree Monti Septemb			Nine Month Septemb	
\$ thousands (except share and per share amounts)		2024	2023		2024	2023
Revenue	\$	956 \$	25	\$	4,032 \$	\$ 35
Cost of revenue		351	24		1,792	73
Gross profit (loss)		605	1		2,240	(38
Operating expenses:						
Research and development		925	312		2,493	1,083
Selling, general and administrative		2,007	932		9,873	3,073
Loss from operations		(2,327)	(1,243)		(10,126)	(4,194
Interest expense, net		(1,455)	(133)		(3,149)	(395
Other income (expense), net		17	_		(191)	_
Change in fair value of warrant liability		9	_		200	
Change in fair value of derivative liability		87	_		4,800	-
Change in fair value of earnout liability		50	_		(700)	_
Net loss	\$	(3,619) \$	(1,376)	\$	(9,166) \$	\$ (4,589
Less: deemed dividend related to the modification of equity classified warrants		_	_		(5,186)	_
Net loss attributable to common stockholders	\$	(3,619) \$	(1,376)	\$	(14,352) \$	\$ (4,589
Basic and diluted net loss per share	\$	(0.17) \$	(0.14)	\$	(0.77)	\$ (0.48
Weighted average shares outstanding	2	1,441,416	9,541,643	1	8,712,468	9,533,185

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Summary of Q3 QTD and YTD Ending 09/30/24 Non-GAAP Results

	Three Months Ended September 30,		Nine Months Ended September 30,		
\$ thousands		2024	2023	2024	2023
Net loss	\$	(3,619) \$	(1,376) \$	(9,166) \$	(4,589)
Interest expense, net		1,455	133	3,149	395
Depreciation and amortization		20	123	204	356
EBITDA		(2,144)	(1,120)	(5,813)	(3,838)
Adjustments:					
Stock-based compensation		127	195	166	613
Warrant modification expense		_	_	201	_
Change in fair value of warrants ⁽¹⁾		(9)	-	(200)	_
Change in fair value of derivatives ⁽²⁾		(87)	_	(4,800)	_
Change in fair value of earnout liability(3)		(50)	-	700	_
Transaction expenses ⁽⁴⁾		_	315	4,301	1,186
Adjusted EBITDA	\$	(2,163) \$	(610) \$	(5,445) \$	(2,039)



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Adjustments to EBITDA

- (1) The decrease in fair value of warrant liability during the three and nine months ended September 30, 2024 relates to the liability classified private placement warrants to reflect the decrease of the publicly traded price per warrant. Additional expense related to the modification of these warrants was recorded as other expense in the condensed consolidated statements of operations and comprehensive loss during the nine months ended September 30, 2024.
- (2) The decrease in fair value of derivative liability during the three and nine months ended September 30, 2024 related to the Yorkville Prepaid 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 September 30, 2024 for financial reporting purposes. The change in derivative liability was recorded as other income (expense), net in the condensed consolidated statements of operations and comprehensive loss during the three and nine months ended September 30, 2024, respectively.
- (3) 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 as of September 30, 2024. The net change in fair value of the earnout liability was recognized as other income (expense), net in the condensed consolidated statements of operations and comprehensive loss during the three and nine months ended September 30, 2024, respectively.
- (4) 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 the 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 statements of operations and comprehensive loss during the nine months ended September 30, 2024. The Company recorded \$0.3 million and \$1.2 million of transaction costs during the three and nine months ended September 30, 2023.



Summary of Q3'24 QTD and Q3'23 QTD P&L Results

Actuals	Q3 FY24	Q3 FY23
Scanners Sold:	2	
Revenue	955,970	24,657
COGS	350,667	23,799
Gross profit	605,303	858
Gross margin	63.3%	3.5%
R&D	925,214	311,829
G&A	2,007,277	932,124
Operating expenses	2,932,491	1,243,953
Operating loss	(2,327,188)	(1,243,095)
Other income/expense	16,995	
Interest expense, net	(1,455,306)	(132,844)
Change in fair value of warrant liability	8,805	2000
Change in fair value of derivative liability	87,200	-
Change in fair value of earnout liability	50,000	
Net loss	(3,619,494)	(1,375,939)
Weighted-average number of shares	21,441,416	9,541,653
Net loss per share - basic and diluted	(0.17)	(0.14)
EBITDA:		
Net loss	(3,619,494)	(1,375,939)
Other expenses and fair value adjustments	(146,005)	
Interest expense	1,455,306	132,844
Deprecation and Amortization	19,508	122,822
Stock-based compensation	127,203	195,475
Transaction expenses related to the merger recorded in G&A		314,875
Adjusted EBITDA	(2,163,482)	(609,923)
Adjusted EBITDA per share	(0.10)	(0.06)

- 2 scanners sold & delivered in Q3'24 vs 0 scanners in Q3'23. ASP of $\sim\!\!450$ K/unit and average cost for legacy scanners was \$175 K/unit, which includes a mix of previously depreciated scanners

- Price of warrants decreased during Q3 from $0.021\,\mathrm{per}\,\mathrm{warrant}$ to $0.011\,\mathrm{per}\,\mathrm{warrant}$,
- Derivative liability for the Yorkville note decreased due to stock price decreasing from \$0.739 as of 6/30/24 to \$0.71 as of 9/30/24.
- Earnout liability decreased due to drop in stock price during Q3.



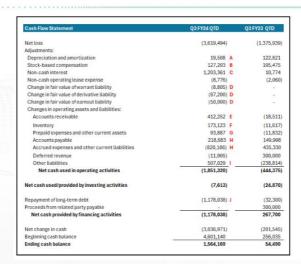
Summary of YTD Ending 09/30/24 and YTD Ending 09/30/23 P&L Results

Actuals	Q3 FY24 YTD	Q3 FY23 YTD
Scanners Sold:	9	12.
Revenue	4,032,168	35,404
COGS	1,792,234	73,497
Gross profit	2,239,934	(38,093)
Gross margin	55.6%	-510.3%
R&D	2,492,842	1,083,373
G&A	9,873,029	3,072,621
Operating expenses	12,365,871	4,155,994
Operating loss	(10,125,937)	(4,194,087)
Other income/expense	(191,330)	-
Interest expense, net	(3,149,315)	(394,714)
Change in fair value of warrant liability	199,624	
Change in fair value of derivative liability	4,800,000	-
Change in fair value of earnout liability	(700,000)	
Netloss	(9,166,958)	(4,588,801)
Weighted-average number of shares	18,712,468	9,533,185
Net loss per share - basic and diluted	(0.77)	(0.48)
EBITDA:		
Net loss	(9,166,958)	(4,588,801)
Other expenses and fair value adjustments	(4,099,111)	-
Interest expense	3,149,315	394,714
Deprecation and Amortization	204,284	355,683
Stock-based compensation	166,187	612,730
Transaction expenses related to the merger recorded in G&A	4,300,703	1,185,851
Adjusted EBITDA	(5,445,580)	(2,039,823)
Adjusted EBITDA per share	(0.29)	(0.21)

- $9 scanners sold \& delivered in Q3'24 YTD vs 0 scanners in Q3'23 YTD. ASP of $$\pi$900K.450K/unit and average cost for legacy scanners was $200K/unit, which includes a mix of previously depreciated scanners$
- R&D expenses are comprised of salaries, stock-based compensation, consultants, and depreciation expense for software, science, MBHTC, and engineering cost centers.
- G&A increased due to new hires, bonuses, and recruiting fees as part of planned growth. Includes \$4.3 million of one-time transaction expenses incurred in Q124 related to the merger with GigCapital5.
- Interest expense increased due to discount amortization for the Yorkville and CableCar notes
- Price of warrants decreased during Q3 YTD.
- Derivative liability for the Yorkville note decreased due to stock price decreasing from \$3.53/share as of merger on 3/4/24 to \$0.71/share as of 9/30/24.
- Q124 was the initial period we recorded the earnout liability. Probability and assumptions are based on revenue targets and FDA mile stones. Depreciation decreased due to transfer of scanners to inventory. New gans in July 2024 with exercise price of §0.75 price and fair value of \$0.47 per option. No new grants during the first two quarters of 2024.







New options granted in July at lower FV. Primarily driven by Yorkulle and CableCar notes Same explanation as for P&L AR decreased due to cash received in Q3 for scanners that were in AR as of Q2. Decrease due to scanners sold, partially offset by purchases Primarily driven by amortization of Insurance expense	Scanners 18 and 22 transferred to inventory
Primarily driven by Yorkville and CableCar notes Same explanation as for P&L AR decreased due to cash received in Q3 for scanners that were in AR as of Q2. Decrease due to scanners sold, partially offset by purchases Primarily driven by amortization of insurance expense Repsyment of AP and accrued expense Repsyment of AP and accrued expense Reclass of accrued interest to other current Liabilities	
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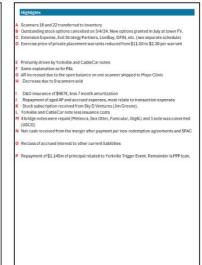
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Summary of YTD Ending 09/30/24 and YTD Ending 09/30/23 Cash Flow Results

Cash Flow Statement	Q3 FY24 YTD	Q3 FY23 YTD
Net loss	(9.166.958)	(4,588,900
Adjustments:		
Depreciation and amortization	204 283 A	355,682
Stock-based compensation	166.187 B	612,730
Warrant modification expense	200,513 D	
Provision for credit losses	1,290	
Fair value of common stock issued in exchange for services	3.718.349 C	100
Loss on issuance of common stock related to subscription agreeme	206,000 C	14
Non-cash interest	2,404,031 E	32,319
Non-cash operating lease expense	(20,652)	(6,184
Loss on disposal of assets		124
Change in fair value of warrant liability	(199,624) F	
Change in fair value of derivative liability	(4,800,000) F	
Change in fair value of earnout liability	700,000 F	
Changes in operating assets and liabilities:		
Accounts receivable	(256.886) G	(18.511
Inventory	1,525,857 H	42,252
Prepaid expenses and other current assets	(459.804) 1	(52,382
Other assets		10,000
Accounts payable	(2,061,852) J	935,742
Accrued expenses and other current liabilities	(768,614) J	411,356
Deferred revenue	(327,778)	300.000
Other liabilities	129.257 0	
Net cash used in operating activities	(8,806,401)	(1,965,772
Net cash used/provided by investing activities	(34,590)	(25,995
Proceeds from sale of common stock and warrants		1,017,850
Processed from stock subscriptions (related party)	500,000 K	0.7
Proceeds from long-term debt, net of issuance costs	10,525,000 L	
Repayment of long-term debt	(1,243,055) P	(96,669
Repayment of bridge loans	(800,000) M	15
Proceeds from related party payable		650,000
Proceeds from the Merger net of transaction costs paid	1,238,529 N	
Net cash provided by financing activities	10,220,474	1,571,181
Net change in cash	1,379,483	(420,586
Beginning cash balance	184,686	475,076
Ending cash balance	1,564,169	54,490



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