

Quantitative Transmission Imaging

Breast Acoustic CT[™] Scanner

INVESTOR PRESENTATION August 2024



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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 "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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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

Introduction to the QT Imaging Holdings Management Team



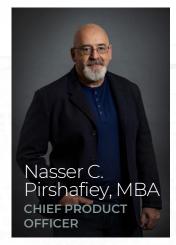
Dr. Raluca Dinu is a global business executive, with long public companies' governance experience, offering over 22 years of achievements in the high-tech industry, with an established track record of driving increased revenue and profitability, delivering strong results in turnaround or M&A situations, 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 & Johnson and Siemens. He has 14 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.



Our Mission

- Create disruptive innovation using technology (software, artificial intelligence, and smart physics) to improve medical imaging and thus, healthcare quality and access
- Continue to improve our FDA-cleared, high quality, high resolution, native 3D, reproducible image quality regardless of operator or breast size/tissue type breast imaging technology, as well as the techniques for quantifiable analysis, comparison, and training
- Introduce the first comprehensive body-safe imaging technology into the marketplace, enabling for the first-time well-person body imaging health screening, and the first health screening medical imaging for infants
- Expand the market opportunities beyond hospitals, imaging centers and health centers by supporting additional direct to consumer (DTC) and direct to provider (DTP) approaches to enable the ability to lower health care costs and increase access via personal medical imaging

• Improve medical outcomes globally by increasing access to medical imaging

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





Executive Summary

Patent-protected technology: 14 granted patents in US/Europe

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)
- Non-Inferiority over Gold Standard (X-Ray Mammography)
- Better resolution compared MRI but without any contrast agent
- Volumetric accuracy to determine mass doubling times in weeks
- More Sensitive and Specific in Dense Breast

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, Sensitivity, Specificity, and Density
- Clinical Trials in Pipeline



Agreement Signed with NXC Imaging

A Subsidiary of Canon Medical Systems

- Distribution Agreement signed with NXC Imaging marks a major milestone for QTI
- 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 will also provide a mature service organization to support QTI's installed base







Feasibility Study Agreement Signed with Canon Medical Systems

- Canon to initiate studies to evaluate 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

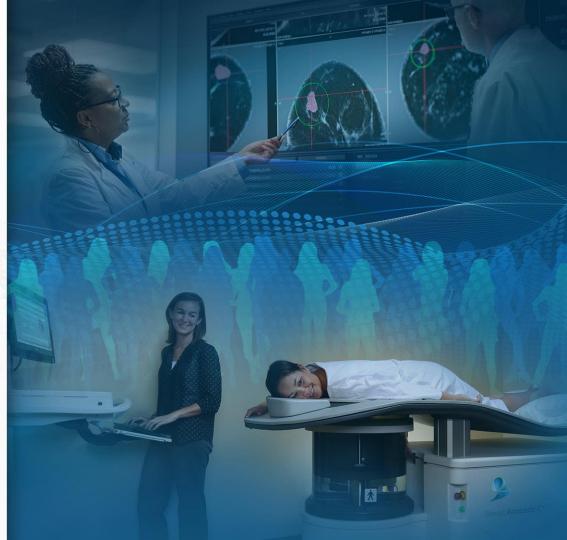


- QTI shall provide support for the feasibility study with Canon and shall use its commercially reasonable efforts to facilitate the feasibility study.
- All know-how and intellectual property embodied in the QT Scanner are owned by QTI.
- During the term of the Feasibility Study Agreement, the QTI shall give Canon first priority in any negotiations for collaborations, including joint development, contract manufacturing, and marketing, with respect to ultrasound breast scanners.





BREAST HEALTH



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

2022 GLOBAL MEDICAL IMAGING MARKET SIZE: \$29B(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



Future Markets – Body Scanner Platform Development

ORTHO: \$9B MARKET(3)

- Target replacing MRI examinations
- Primary focus on orthopedic practices



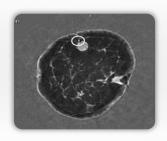
INFANT: \$8B MARKET (4)

 New market opportunity given limitations of current imaging modalities for infants



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

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





- (I) Medical Imaging Market Size, Share & Trends Analysis Report by Products (X-Ray, Ultrasound, Computed Tomography, Magnetic Resonance Imaging (MRI), Nuclear Imaging), by End Users (Hospitals, Diagnostic Imaging Centers, Other End Users), by Region (North America, Europe, Asia Pacific, Latin America, Middle East & Africa) Global Industry Assessment (2016 2021) & Forecast (2022 2028), Vantage Market Research
- (2) Contenent Market margins
- (3) Global Orthopedic Medical Imaging Systems Market Analysis Report 2022: Market to Reach \$10.6 Billion by 2026 The US Corners Orthopedic Medical Imaging Market with Adoption of Innovative Systems, Research and Markets.
- Pediatric Imaging Market Size, Share & I rends Analysis Report By Modality (X-ray, Ultrasound, WHI, C. I.), By Application (Lastroenterology, Cardiology, Uncology), By End User, By Region, And Segment Forecasts, 2020 2027, Grandwiew Research
 Images quided Theorems (Sestence Market Size) Share & Trends Analysis Departs by Product I Ultrasound Sustence Computer Services and Language Cardiology, Cardiology, Oncology), By End User, And Segment Forecasts

Other Ultrasound Products Use 2D Imaging for Dense Breast Screening



SIEMENS

SonoCiné

HITACHI

DELPHINUS SOFTVUE















Articulating Arm

Articulating Arm

Articulating Arm

Rotating Armature

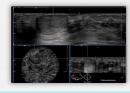
Water Bath

Water Bath

Stacked 2D Reflection Slices



Stacked 2D Reflection Slices



Stacked 2D Reflection Slices

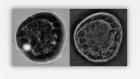
Guided Handheld



Stacked 2D Reflection Slices



Stacked 2D Slices



Only Full 3D



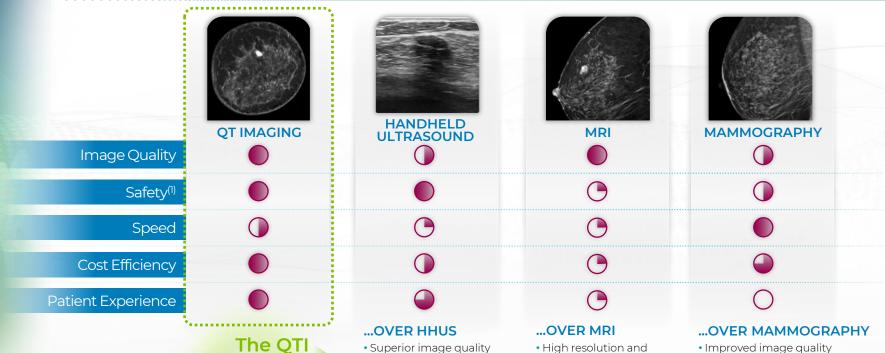


Mammography is ineffective in screening dense breasts.
Ultrasound techniques performed after MRI did not detect additional cancer⁽¹⁾ in dense breast

DESIGN TYPE

OUTPUT

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



Not operator dependent

• Quantifiable/repeatable



- contrast-to-noise ratio
- · No injection needed
- · Lower equipment cost
- No special facility or shielding requirements
- · Safer (no radiation), allowing for more frequent imaging
- Greater sensitivity and specificity
- No special facility requirements
- Quantifiable/repeatable

Imaging

Advantage

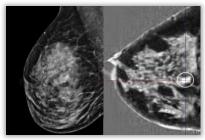
QT Imaging's FDA-cleared Solution for Dense Breasts

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 rea(\$)



"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⁾



⁽²⁾ OTI Study I Dense Breast Mass Detection

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

QTI Clinical Trials Provide Compelling Results for Adoption and Approvals

CLINICAL TRIALS

- Visual Grading Assessment of Quantitative Transmission Ultrasound Compared to Digital X-ray Mammography and Hand-held Ultrasound
- Anatomy-Correlated Breast Imaging and Visual Grading Analysis Using Ouantitative Transmission Ultrasound
- Accuracy of Cyst vs. Solid Diagnosis in the Breast Using Quantitative Transmission (QT) Ultrasound
- · Breast Cyst Fluid Analysis Correlations Using Transmission Ultrasound
- Objective Breast Tissue Image Classification Using Quantitative Transmission Ultrasound Tomography
- Quantitative Assessment of Breast Density: Transmission Ultrasound is Comparable to Mammography with Tomosynthesis
- An Exploratory Study Comparing Transmission Ultrasound to Mammography on Recall Rates and Detection Rates for Breast Cancer
- QT Ultrasound Tomography for Orthopedic Imaging
- QT Ultrasound for Whole Body Imaging

IMPLICATION OF RESULTS OR PRELIMINARY RESULTS

QT can **see more anatomy** than mammography or handheld ultrasound

QT can **distinguish specific tissues** unlike mammography or handheld ultrasound

QT can **quantify breast density** unlike mammography or handheld ultrasound

QT can identify breast and reduce recall rates better than mammography

QT can identify bone and joint structures

QT can identify internal body structures





Canada's premier Cancer centre:









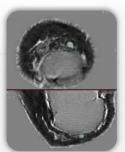
QTI Clinical Trials:

Dense Breast Imaging Studies Using DBT Show Sensitivity Close to 40%(3)

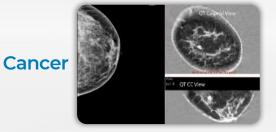
Approximately 50% of women between the ages of 40-74 in the US have dense breasts⁽¹⁾, with **traditional** mammography missing 35.6-52.2% of breast cancers in dense breast tissue⁽²⁾ making QT Scanner the only

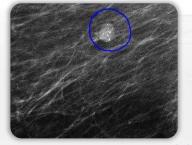
system effective at screening dense breast.

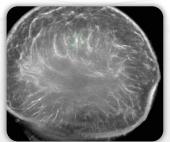




Implant
With Mass

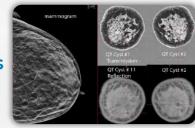






Calcification

Cysts



QTI can see calcification missed by other imaging modalities and is particularly effective in imaging dense breasts



OTIMAGING

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

⁽s) C. E. Comstock, MD, C. Gatsonis, PhD et al. "Comparison of Abbreviated Breast MRI vs Digital Breast Tomosynthesis for Breast Cancer Detection Among Women With Dense Breasts Undergoing Screening", JAMA 2020, 323 (8):746-756.

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Clinical Evidence:

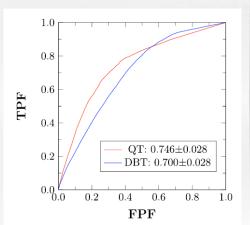
Non-Inferiority to X-Ray Mammography (XRM)

Non-inferiority to XRM

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

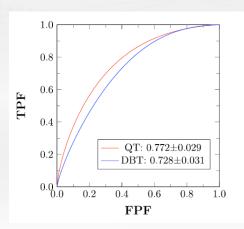
Non-parametric ROC curves

The line closer to 1.0 is indicative of higher accuracy



Proper binomial ROC curves

The line closer to 1.0 is indicative of higher accuracy



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



lef : A MultiReader Multicase (MRMC) Receiver Operating Characteristic (ROC) Study Evaluating Noninferiority of Quantitative Transmission (QT) Ultrasound to Digital Breast Tomosynthesis (DBT) or Detection and Recall of Breast Lesions : Yulei Jiang, PhD, Elaine Iuanow, MD, Bilal Malik, PhD,, John Klock, MD, Academic Radiology, Vol 31, No 6, June 2024.

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

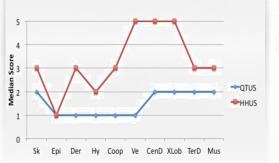
X-Ray Mammography (XRM)

- 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.



Ref : John C Klock, Elaine luanow, Kathleen Smith, Nancy A and Obuchowski (2017) Visual Grading Assessment of Quantitative Transmission Ultrasound Compared to Digital X-ray Mammography a Hand-held Ultrasound in Identifying Ten Breast Anatomical Structures. Clinical Trials 3: 015

The Current Breast Imaging Paradigm Leads to Unnecessary Concern and Costs

Screening compliance is low



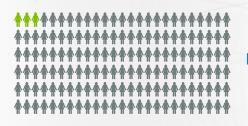
Of the **65%** of women who do get screened, many suffer through unnecessary callbacks

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 **150**



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⁾







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



Approximately 150 women out of every 1,000 screened are recalled for more tests. Out of those 150, only three or four will be diagnosed with cancer. Unnecessary recalls create stress for the patient and have other negative impacts on and the breast healthcare industry.

- Recall Rate is a metric to assess accuracy and detection rate
 - Anxiety Reducing Factor

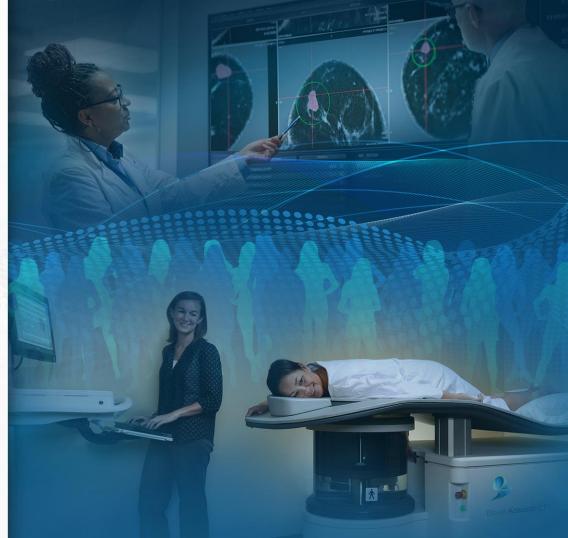
QTI technology improves non-cancer recall rates without substantially affecting cancer recall rates



*An Exploratory Multi-reader, Multi-case Study Comparing Transmission Ultrasound to Mammography on Recall Rates and Detection Rates for Breast Cancer Lesions Bilal Malik, PhD, Elain Iuanow, MD, John Klock, MD, Academic Radiology, Vol 29, No S1, January 2022



TECHNOLOGY OVERVIEW



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)













A Multireader Multicase (MRMG) Receiver Operating Characteristic (ROC) Study Evaluating Noninferiority of Quantitative Transmission (QT) Ultrasound to Digital Breast Tomosynthes bely bely on Detection and Recall of Breast Lesions and Villei at al. Academic Padiology, in press

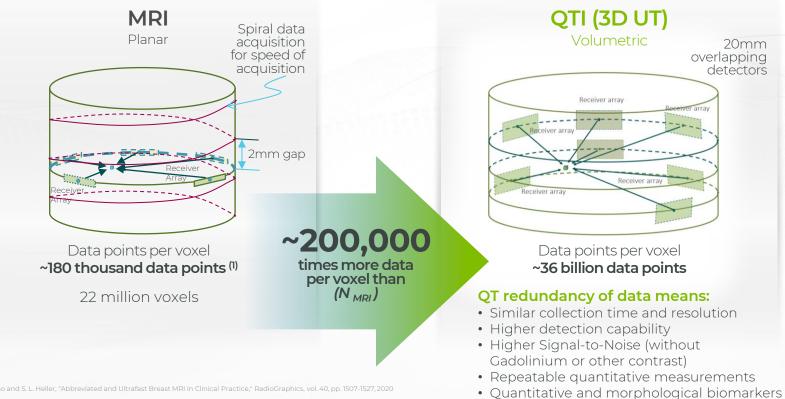
⁽²⁾ Based on opinion of QTI management. QTI believes necessary data has been obtained through 18 separate clinical trials

Critical Modality Advantages of QTI's Breast Acoustic CT⁽¹⁾

- Clinically useful sensitivity and specificity
- Presence of comparative clinical trials
- Proven success in head-to-head trials against mammography
- Ability to determine doubling times can identify slow growing cancers and help prevent cancer deaths
- Enhanced volume measurements can follow cancer treatments and provide breast density measurements
- Patented technology opens the door for potential future growth in orthopedic and pediatric imaging



Why QTI Scan Generates Better Resolution Compared to MRI: More Data!



for longitudinal studies



Technical Capabilities

• Resolution of ~600 microns in reflection compared to 800 microns⁽¹⁾ for MRI (depends on field strength, homogeneity etc)

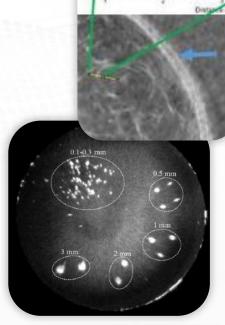
 Contrast to noise ratio of 23:1 at 100 microns (in reflection; can detect small calcifications)

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%)





(1) Y. Gao and S. L. Heller, "Abbreviated and Ultrafast Breast MRI in Clinical Practice," RadioGraphics, vol. 40, pp. 1507-1527, 2020

Enhanced Clinical Capabilities and Value

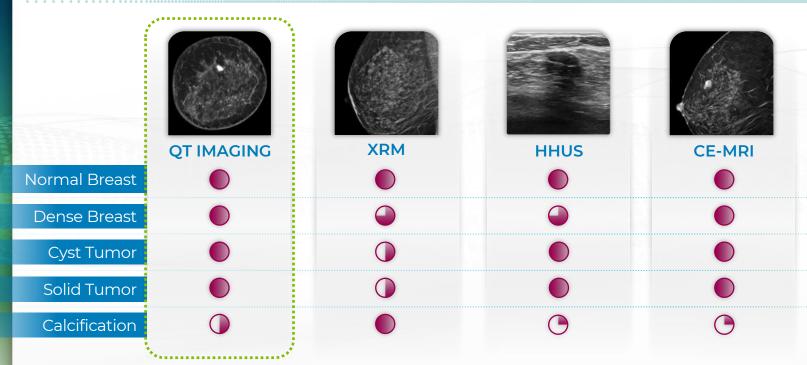
- High-quality and high-resolution native 3D Imaging
- Quantifiable images enables accurate analysis, comparison and trending
- Consistent and reproduceable image quality regardless of operator or breast size/tissue type
- Clinical feature detection of 50-100 microns including microcalcifications
- Functional 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





Y. Gao and S. L. Heller, "Abbreviated and Ultrafast Breast MRI in Clinical Practice," RadioGraphics, vol. 40, pp. 1507 27, 2020

Imaging Accuracy in Breast Mass Diagnosis⁽¹⁾



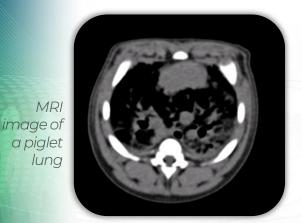




Resolution and Detectability:

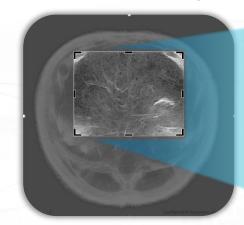
MRI vs QTI's Acoustic CT (3D UT)

MRI

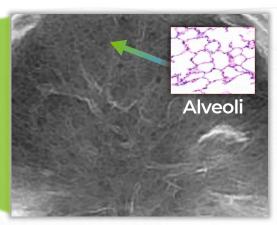


MRI resolution depends on acquisition time, B1 inhomogeneity, etc.

QTI's Acoustic CT



Higher resolution than 3T MRI in air-filled organs



QTI's Acoustic CT (3D UT) with reflection mode

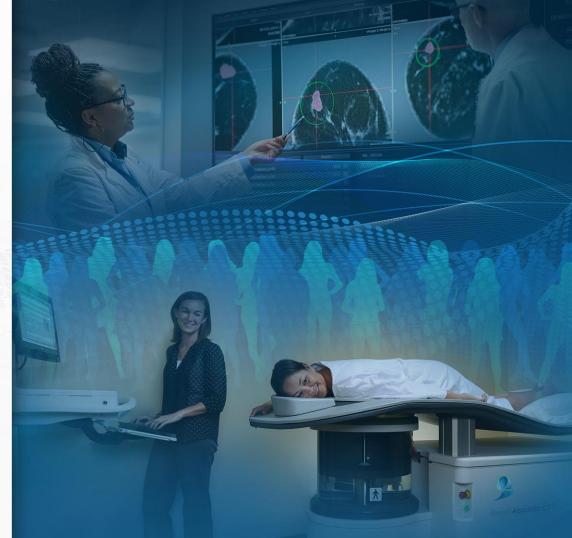
- Resolution is almost isotropic (transmission)
- Sub-mm resolution
- Detectability 0.1 mm

First time structures as small as the **Lung Alveoli** can be seen in vivo!





MARKET POSITIONING



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

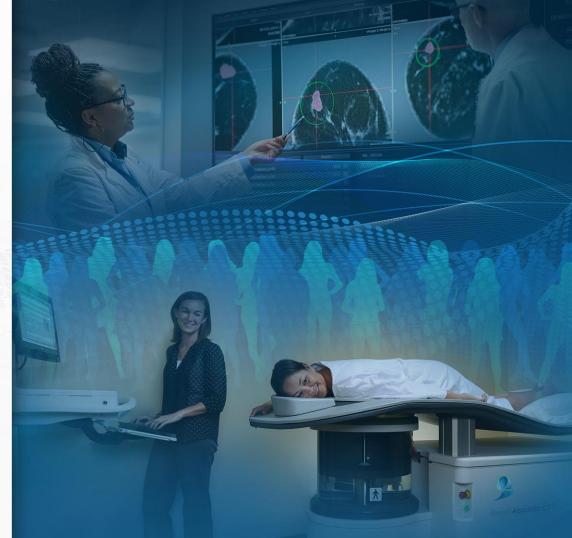
- Native 3D imaging (like MRI and CT)
- Quantifiable image analysis
- No need for specialized technologist training
- · Consistent and reproducible image quality regardless of operator







OPEN ANGLE SCANNER



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:





SPORTS TEAMS [ON THE FIELD]



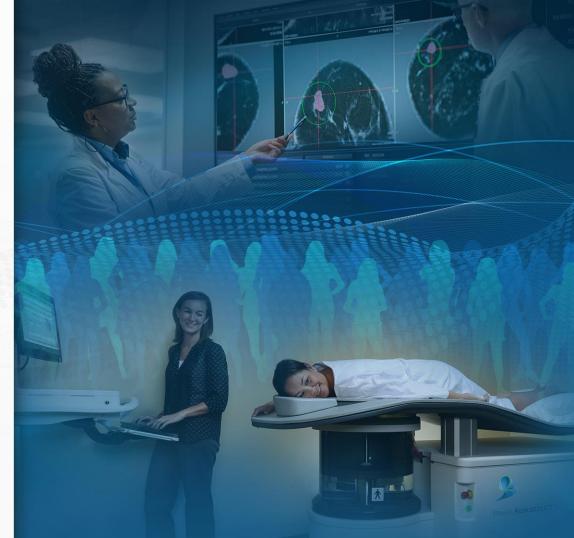
MILTARY
[SHIPS &
FIELD USE]







Q2' 24 AND YTD ENDING 06/30/24 FINACIALS



Financial Highlights for Q2'24 QTD and YTD Ending 06/30/24

- Commercial revenue was \$1.7 million during the second quarter of 2024, compared to \$1.4 million during the first quarter of 2024 and less than \$0.1 million during the second quarter of 2023.
- Commercial revenue was \$3.1 million for the six months ended June 30, 2024, compared to less than \$0.1 million for the six months ended June 30, 2023.
- Gross margin of 51% in the second quarter of 2024, compared to gross margin of 56% in the first quarter of 2024 and insignificant margin in the second quarter of 2023. The decrease in margin in the second quarter of 2024 compared to the first 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 four QT Breast Scanners during the second quarter of 2024, compared to no deliveries in the second quarter of 2023.
- Gross margin of 53% during the six months ended June 30, 2024, compared to negative margin in 2023 YTD due to the sale and delivery of seven QT Breast Scanners in 2024 YTD, compared to no deliveries in 2023 YTD.



Financial Highlights for Q2'24 QTD and YTD Ending 06/30/24

- Net loss of \$1.2 million for the second quarter of 2024, compared to net loss of \$1.3 million for the second quarter in 2023. Q2'24 net loss includes:
 - \$2.1 million of net non-cash income related to the change in fair value of the warrant, derivative, and earnout liabilities and \$0.2 million of warrant modification expense, compared to a net loss of \$1.3 million for the second quarter of 2023, which included stock-based compensation expense of \$0.2 million and one-time transaction expenses of \$0.2 million.
- Net loss of \$5.5 million for the first six months of 2024, compared to net loss of \$3.2 for the first six months of 2023. 2024 YTD net loss includes:
 - \$3.8 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 June 30, 2024, less than \$0.1 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 \$(2.1) million for the second quarter of 2024, compared to \$(0.7) million for the second quarter of 2023.
- Non-GAAP Adjusted EBITDA of \$(3.3) million for the first six months of 2024, compared to \$(1.7) million for the first six months of 2023.
- The company ended Q2'24 with \$4.6M in cash, compared to end of Q1'24 with \$5.6M in cash, thus the burn rate was about \$1.0M.



Summary of Q2 QTD and YTD Ending 06/30/24 GAAP Results

	Three Months Ended Six Months End June 30, June 30,					
\$ thousands (except share and per share amounts)		2024	2023	2024		2023
Revenue	\$	1,714	\$ 3 \$	3,076	\$	11
Cost of revenue		839	3	1,442		50
Gross profit (loss)		875	_	1,634		(39)
Operating expenses:						
Research and development		925	349	1,567		771
Selling, general and administrative		2,170	849	7,866		2,141
Loss from operations		(2,220)	(1,198)	(7,799)		(2,951)
Interest expense, net		(1,095)	(132)	(1,694)		(262)
Other expense, net		(187)	_	(208)		_
Change in fair value of warrant liability		214	_	191		_
Change in fair value of derivative liability		1,729	_	4,713		_
Change in fair value of earnout liability		310	_	(750)		_
Net loss	\$	(1,249)	\$ (1,330) \$	(5,547)	\$	(3,213)
Less: deemed dividend related to the modification of equity classified warrants		(5,186)	_	(5,186)		_
Net loss attributable to common stockholders	\$	(6,435)	\$ (1,330) \$	(10,733)	\$	(3,213)
Basic and diluted net loss per share	\$	(0.30)	\$ (0.14) \$	(0.62)	\$	(0.34)
Weighted average shares outstanding	2	1,440,447	9,540,533 1	7,333,000	9	,528,880



Summary of Q2 QTD and YTD Ending 06/30/24 Non-GAAP Results

	Three Months Ended June 30,			Six Months Ended June 30,		
\$ thousands		2024	2023	2024	2023	
Net loss	\$	(1,249) \$	(1,330) \$	(5,547) \$	(3,213)	
Interest expense, net		1,095	132	1,694	262	
Depreciation and amortization		86	116	185	233	
EBITDA		(68)	(1,082)	(3,668)	(2,718)	
Adjustments:						
Stock-based compensation		_	208	39	417	
Warrant modification expense		201	_	201	_	
Change in fair value of warrants ⁽¹⁾		(214)	_	(191)	_	
Change in fair value of derivatives(2)		(1,729)	_	(4,713)	_	
Change in fair value of earnout liability ⁽³⁾		(310)	_	750	_	
Transaction expenses(4)		_	206	4,301	562	
Adjusted EBITDA	\$	(2,120) \$	(668) \$	(3,281) \$	(1,739)	



Adjustments to EBITDA

- (1) The decrease in fair value of warrant liability during the three and six months ended June 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 during the three and six months ended June 30, 2024.
- (2) The decrease in fair value of derivative liability during the three and six months ended June 30, 2024 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 March 31, 2024 and June 30, 2024 for financial reporting purposes. The change in derivative liability was recorded as other income in the condensed consolidated statements of operations during the three and six months ended June 30, 2024.
- (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 March 31, 2024 and June 30, 2024. The net change in fair value of the earnout liability was recognized as other expense in the condensed consolidated statements of operations during the three and six months ended June 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 during the six months ended June 30, 2024. The Company recorded \$0.2 million and \$0.6 million of transaction costs during the three and six months ended June 30, 2023.



Summary of Q2'24 QTD and Q2'23 QTD P&L Results

Actuals		Q2 FY24	Q2 FY23
Scanners Sold:		4	-
Revenue		1,714,034	3,183
COGS		839,484	3,121
Gross profit		874,550	62
Gross margin		51.0%	1.9%
R&D		925,082	349,657
G&A		2,169,541	848,832
Operating expenses		3,094,623	1,198,489
Operating loss		(2,220,073)	(1,198,427)
Other income/expense		(187,393)	
Interest expense, net		(1,095,050)	(131,588)
Change in fair value of warrant liability		213,942	
Change in fair value of derivative liability		1,729,700	-
Change in fair value of earnout liability		310,000	-
Net loss		(1,248,874)	(1,330,015)
Weighted-average number of shares	E E E	21,440,447	9,540,533
Net loss per share - basic and diluted	\$	(0.30) \$	(0.14)
EBITDA:			
Net loss		(1,248,874)	(1,330,015)
Other expenses and fair value adjustments		(2,053,129)	-
Interest expense		1,095,050	131,588
Deprecation and Amortization		85,903	116,035
Stock-based compensation		-	208,627
Transaction expenses related to the merger recorded in G&A		-	206,361
Adjusted EBITDA		(2,121,050)	(667,404)
Adjusted EBITDA per share	\$	(0.10) \$	(0.07)

Г	
	Highlights
Α	4 scanners sold & delivered in Q2'24 vs 0 scanners in Q1'23. Revenue
	ranged \$300K-450K/unit and average cost for legacy scanners was
	\$210K/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.
L	
C	G&A increased due to new hires, bonuses, and recruiting fees as part of planned growth.
	parinod 6. Ortin
L	4004K - 150
טן	\$201K modification expense related to private warrants, offset by \$16K of income related to QTI Center.
	mount routes to Q11 conton
E	Interest expense increased due to discount amortization for the Yorkville
	and CableCar notes
F	Price of warrants decreased during Q2 from \$0.036 to \$0.021 per warrant.
۵	Derivative liability for the Yorkville note decreased due to stock price
ľ	decreasing from \$1.06/share as of 3/31/24 to \$0.739 as of 6/30/24.
Д	Earnout liability decreased due to drop in stock price during Q2.
ľ	Lamout dability decreased due to drop in stock price during Q2.
ı	Depreciation decreased due to transfer of scanners to inventory.
J	On 3/4/24, all oustanding stock options were cancelled. No new grants
1	until July 2024



Summary of YTD Ending 06/30/24 and YTD Ending 06/30/23 P&L Results

Actuals	Q	2 FY24 YTD	Q2 FY23 YTD
Scanners Sold:		7	
Revenue		3,076,197	10,747
COGS		1,441,568	49,698
Gross profit		1,634,629	(38,951)
Gross margin		53.1%	N/A
R&D		1,567,628	771,544
G&A		7,865,751	2,140,497
Operating expenses		9,433,379	2,912,041
Operating loss		(7,798,750)	(2,950,992)
Other income/expense		(208,324)	-
Interest expense, net		(1,694,019)	(261,870)
Change in fair value of warrant liability		190,819	
Change in fair value of derivative liability		4,712,800	- (
Change in fair value of earnout liability		(750,000)	- 1
Net loss		(5,547,474)	(3,212,862)
Weighted-average number of shares		17,333,000	9,528,880
Net loss per share - basic and diluted	\$	(0.62) \$	(0.34)
EBITDA:			
Net loss		(5,547,474)	(3,212,862)
Other expenses and fair value adjustments		(3,953,106)	_
Interest expense		1,694,019	261,870
Deprecation and Amortization		184,776	232,861
Stock-based compensation		38,984	417,255
Transaction expenses related to the merger recorded in G&A		4,300,703	562,140
Adjusted EBITDA		(3,282,098)	(1,738,736)
Adjusted EBITDA per share	\$	(0.19) \$	(0.18)

Highlights

- A 7 scanners sold & delivered in Q2'24 YTD vs 0 scanners in Q1'23 YTD.

 Revenue ranged \$300K-450K/unit and average cost for legacy scanners

 was \$205K/unit, which includes a mix of previously depreciated scanners
- B R&D expenses are comprised of salaries, stock-based compensation, consultants, and depreciation expense for software, science, MBHTC, and engineering cost centers.
- C 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 Q1'24 related to merger with GigCapital5.
- \$201K modification expense related to private warrants
- E Interest expense increased due to discount amortization for the Yorkville and CableCar notes
- F Price of warrants increased during Q2 YTD from \$0.01 to \$0.021 per warrant.
- G Derivative liability for the Yorkville note decreased due to stock price decreasing from \$3.53/share as of merger date on 3/4/24 to \$0.739 as of 6/30/24.
- H Q1'24 is the initial period we recorded the earnout liability. Probability and assumptions are based on revenue targets and FDA milestones.
- Depreciation decreased due to transfer of scanners to inventory.
- J On 3/4/24, all oustanding stock options were cancelled. No new grants until July 2024



Summary of Q2'24 QTD and Q2'23 QTD Cash Flow Results

Cash Flow Statement	Q2 FY24 QTD	Q2 FY23 QTD
Net loss	(1,248,874)	(1,330,014)
Adjustments:		
Depreciation and amortization	85,902	116,035
Stock-based compensation	-	208,627
Warrant modification expense	200,513	-
Fair value of common stock issued in exchange for services	3,655	-
Loss on issuance of common stock related to subscription agreer		-
Non-cash interest	902,065	10,772
Non-cash operating lease expense	(6,507)	(2,062)
Loss on disposal of assets		124
Change in fair value of warrant liability	(213,942)	-
Change in fair value of derivative liability	(1,729,700)	-
Change in fair value of earnout liability	(310,000)	-
Changes in operating assets and liabilities:		
Accounts receivable	(186,781)	5,840
Inventory	766,321	4,818
Prepaid expenses and other current assets	325,817	(5,909)
Other assets	-	5,000
Accounts payable	(162,190)	393,525
Accrued expenses and other current liabilities	1,371,144	(55,504)
Deferred revenue	(311,905)	-
Other liabilities	(465,084)	120,067
Net cash used in operating activities	(979,566)	(528,681)
Net cash used/provided by investing activities	(26,977)	(1,125)
Proceeds from sale of common stock and warrants	-	70,000
Repayment of long-term debt	(32,548)	(32,216)
Proceeds from related party payable	-	350,000
Net change in cash	(1,039,091)	(142,022)
Beginning cash balance	5,640,231	398,057
Ending cash balance	4,601,140	256,035

	Highlights
A	Scanners 18 and 22 transferred to inventory
В	Oustanding stock options cancelled on 3/4/24
С	Return of a customer deposit for scanner
D	Exercise price of private placement warrants reduced from \$11.50 to \$2.30 per warrant
E	Primarily driven by Yorkville and CableCar notes
F	Same explanation as for P&L
G	AR increased due to the open balance on two scanners shipped to NXC/Oklahoma
Н	Decrease due to 4 scanners sold
ı	D&O insurance of \$987K, less 3 month amortization
J	Reclassification of accrued interest to other current liablilities and accrued legal expenses
o	Reclass of accrued interest to other current liablities



Summary of YTD Ending 06/30/24 and YTD Ending 06/30/23 Cash Flow Results

Cash Flow Statement	Q2 FY24	Q2 FY23
Net loss	(5,547,464)	(3,212,961)
Adjustments:	(, , , , , , ,	(, , , , ,
Depreciation and amortization	184,775	232,861
Stock-based compensation	38,984	417,255
Warrant modification expense	200,513	
Provision for credit losses	1,290	
Fair value of common stock issued in exchange for services	3,718,349	<u>.</u>
Loss on issuance of common stock related to subscription agreer	206,000	
Non-cash interest	1,200,670	21,545
Non-cash operating lease expense	(11,876)	(4,124)
Loss on disposal of assets		124
Change in fair value of warrant liability	(190,819)	
Change in fair value of derivative liability	(4,712,800)	
Change in fair value of earnout liability	750,000	-
Changes in operating assets and liabilities:		
Accounts receivable	(669,138)	
Inventory	1,352,734	53,869
Prepaid expenses and other current assets	(553,691)	(40,550)
Other assets		10,000
Accounts payable	(2,280,535)	785,744
Accrued expenses and other current liabilities	51,572	(23,974)
Deferred revenue	(315,873)	
Other liabilities	(377,772)	238,814
Net cash used in operating activities	(6,955,081)	(1,521,397)
Net cash used/provided by investing activities	(26,977)	(1,125)
Proceeds from sale of common stock and warrants		1,017,850
Processed from stock subscriptions (related party)	500,000	
Proceeds from long-term debt, net of issuance costs	10,525,000	
Repayment of long-term debt	(65,018)	(64,369)
Repayment of bridge loans	(800,000)	-
Proceeds from related party payable	- '	350,000
Net change in cash	4,416,454	(219,041)
Beginning cash balance	184,686	475,076
Ending cash balance	4,601,140	256,035

	Highlights
l	
	Scanners 18 and 22 transferred to inventory
	Oustanding stock options cancelled on 3/4/24
	Extension Expense, Exit Strategy Partners, LionBay, DFIN, etc.
D	Exercise price of private placement warrants reduced from \$11.50 to \$2.30 per warrant
	Primarily driven by Yorkville and CableCar notes
	Same explanation as for P&L
	AR increased due to the open balance on two scanners shipped to NXC/Oklahoma Decrease due to 7 scanners sold
ı	D&O insurance of \$987K, less 4 month amortization
J	Repayment of aged AP and accrued expenses, most relate to transaction expenses
K	Stock subscription received from Sky D Ventures
L	Yorkville and CableCar note less issuance costs
М	4 bridge notes were repaid (Meteora, Sea Otter, Funicular, Gig4L) and 1 note was converted (USCG)
N	Net cash received from the merger after payment per non-redemption agreements and
0	Reclass of accrued interest to other current liablities
1	
1	



Investment Highlights

Cutting-edge imaging technology with multiple potential applications creates a tremendous opportunity to transform the imaging market







Thank You!

