

The Future of Cardiac Care

Second Quarter 2025
Financial Results Conference Call

NASDAQ: BEAT AUGUST 13, 2025



DISCLAIMERS

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The HeartBeam System received FDA clearance in December 2024. HeartBeam 12-lead ECG synthesis software has not yet been approved by FDA or other regulatory authorities for commercial/clinical use in the USA or other global geographies.

Agenda:

THE HEARTBEAM SYSTEM

PRODUCT VISION

COMPANY MILESTONES

COMMERCIALIZATION STRATEGY

FINANCIAL RESULTS

On the Call Today:





Groundbreaking HeartBeam Technology







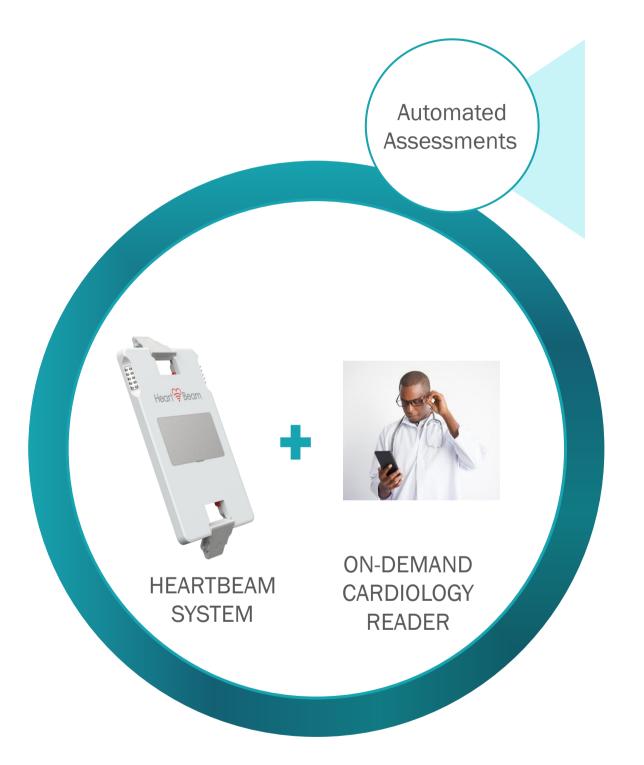
- Received December 2024
- HeartBeam System: the first ever cable-free device capable of collecting ECG signals in 3D, from 3 non-coplanar directions, for arrhythmia assessment

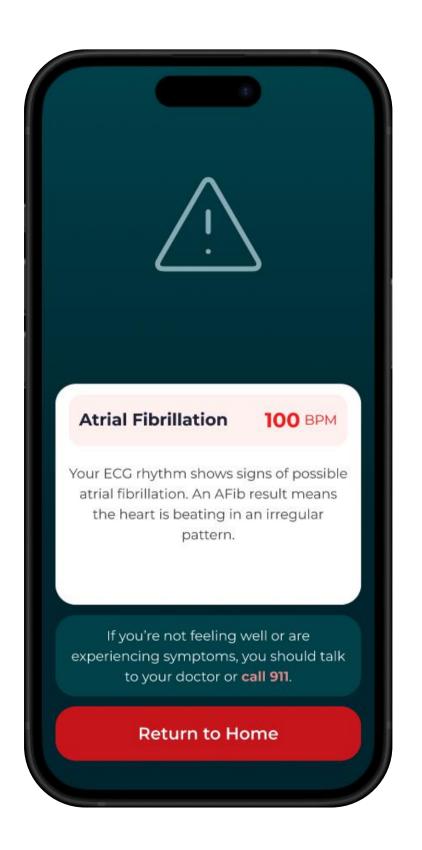
12-LEAD ECG SYNTHESIS SOFTWARE APPLICATION

- Submitted January 2025
- Software that synthesizes a 12-lead ECG from the 3D signals captured by the device for arrhythmia assessment
- Clearance anticipated by end of 2025



Leverage uniqueness of HeartBeam 12-lead ECG system to unlock unparalleled performance from anywhere and drive deeper adoption





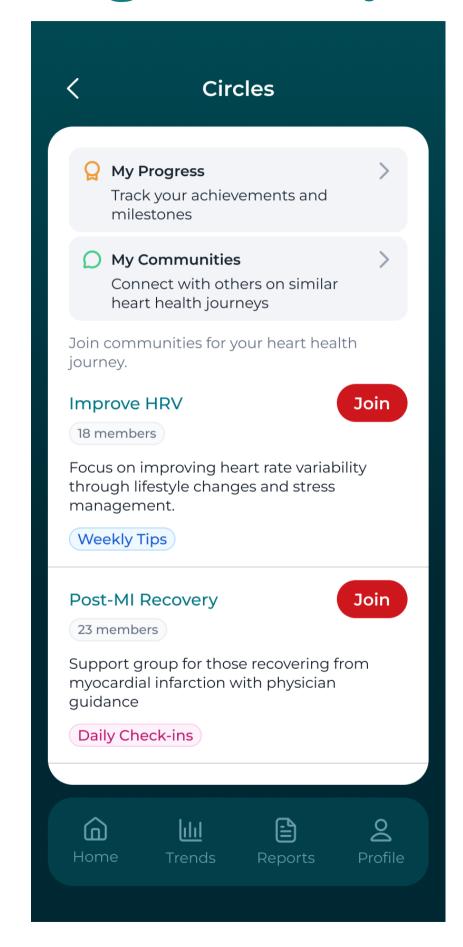




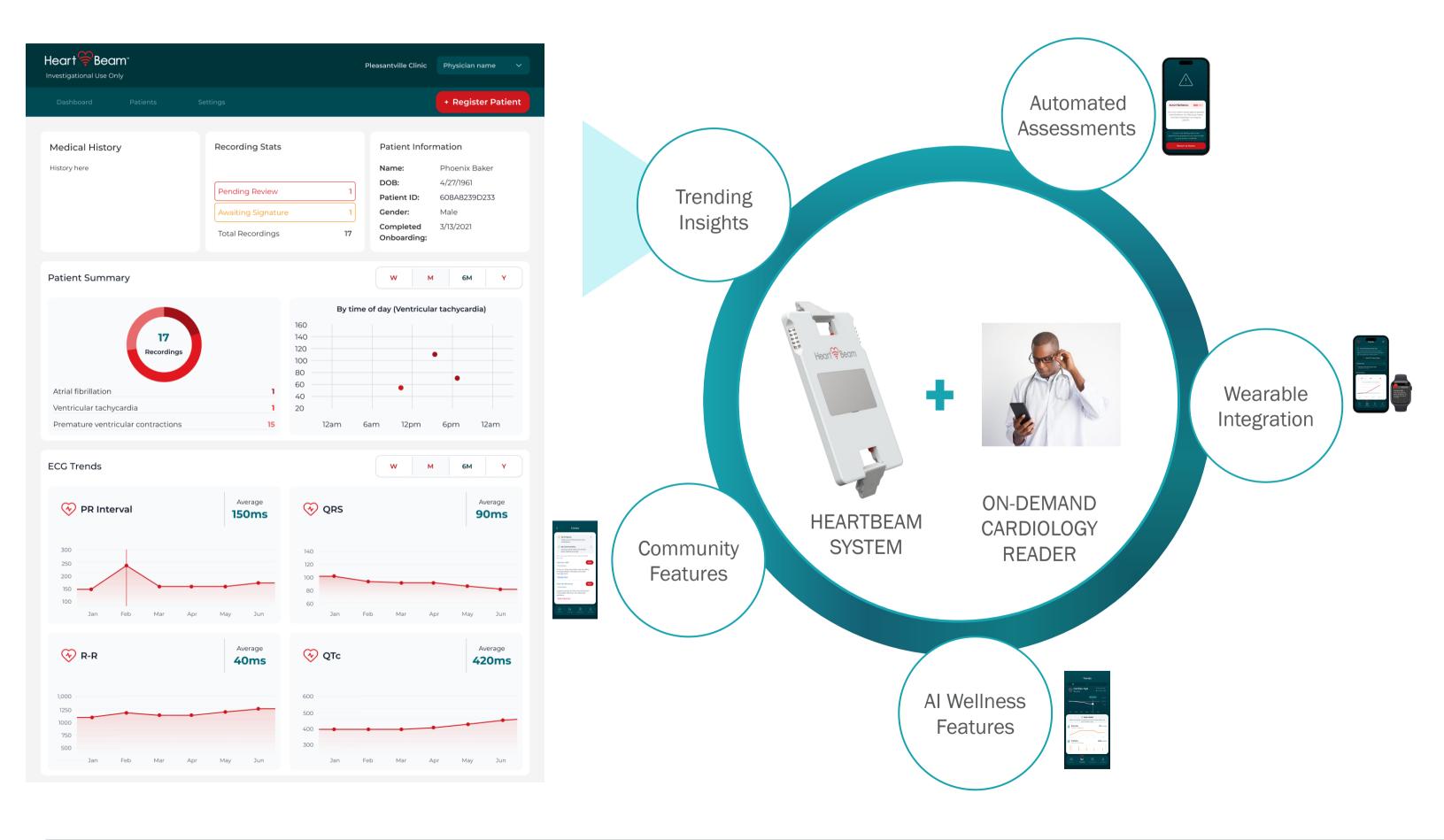




*Product concept for illustrative purposes. Final product may vary. Automated assessment not cleared by the FDA. Not available for sale in any country.









Leverage uniqueness of HeartBeam 12-lead ECG system to unlock unparalleled performance from anywhere and drive deeper adoption

HeartBeam's Long-Term Vision



Major Healthcare Trend: From Hospital Hallways to Home Hallways









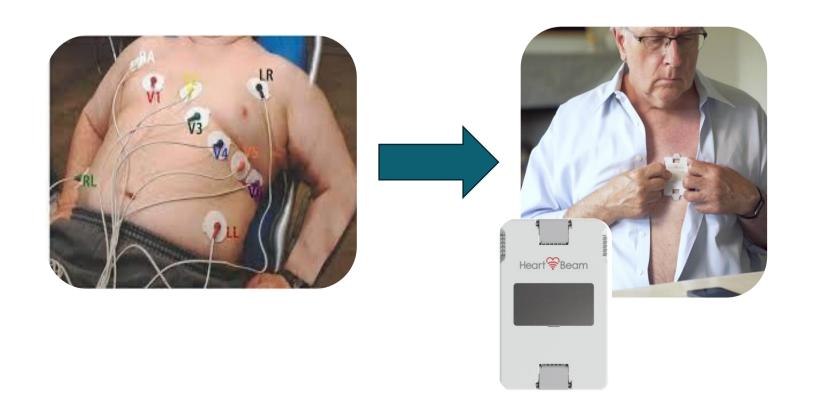
Blood Pressure Monitors



Oxygen Monitors



Portable 12-Lead ECG



- Cardiac disease is #1 global killer
- ECG is the most common cardiac test
- Current at-home gap: No easy-to-use 12-lead ECG
- HeartBeam advantage: First cable-free, pocket-size synthesized 12-lead ECG
- On-demand cardiology review: Peace of mind, immediate action
- Shifts care to home: Empowering patients, potential to improve outcomes and reduce healthcare costs

Key Developments Since Last Update

REGULATORY PROGRESS

- Continue to engage in productive discussions with FDA on 12-lead ECG Synthesis 510(k) submission for arrhythmia assessment
 - Timeline for FDA clearance remains firmly on track
- Further discussions on future clearances and indications, including the ischemia indication and the AccurKardia automated algorithm

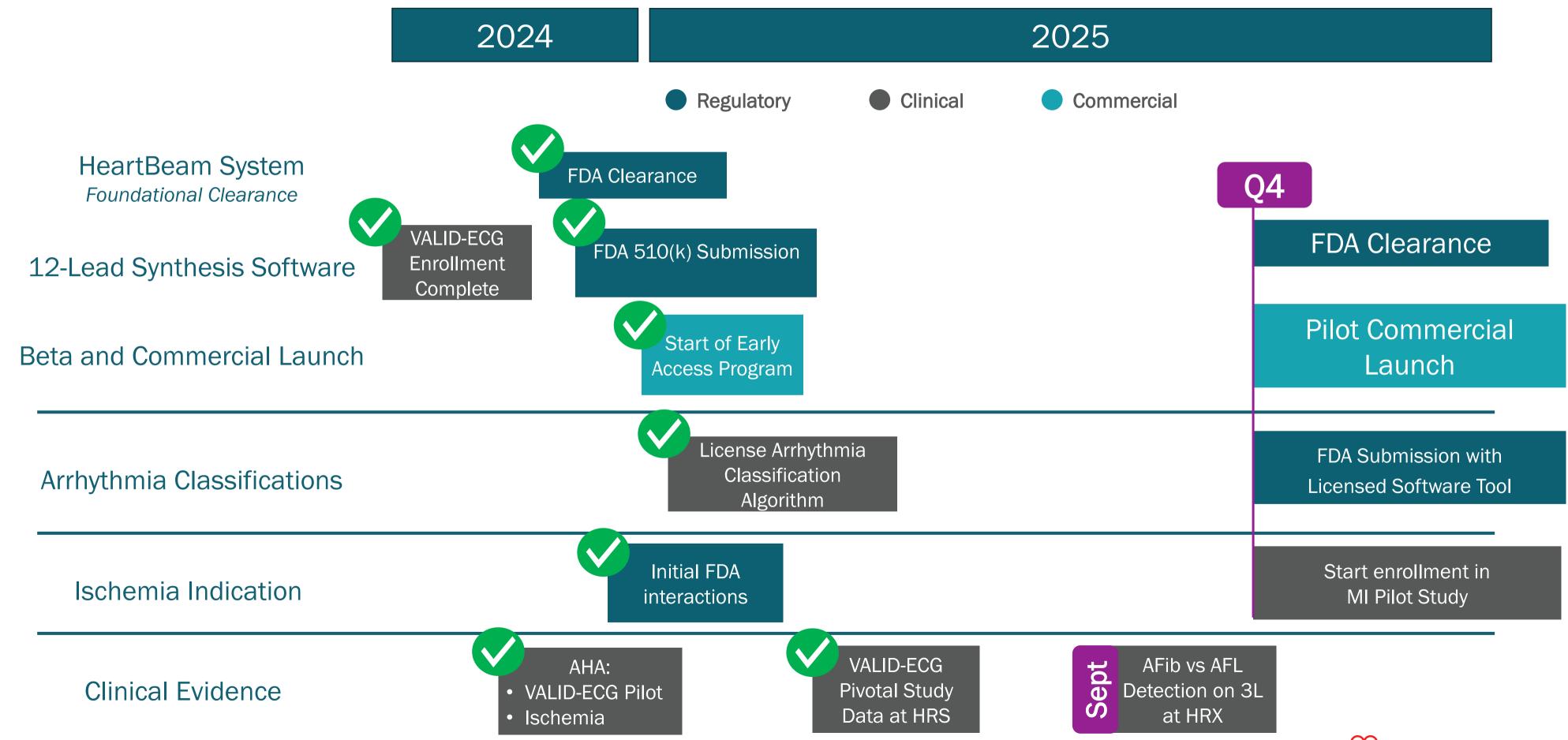
GO TO MARKET/ BUSINESS DEVELOPMENT

- Executing on commercial readiness plans
- Business development activity intensifying as FDA Clearance and commercialization are approaching

OTHER PROGRESS

- Abstract on AI algorithm accepted for presentation at HRX Live 2025 in September
- One new international patent issued, total of 21 issued patents worldwide
- Continued industry validation with two additional industry award recognitions

Continued Achievement of Stated Milestones



Continued Recognition for HeartBeam's Innovative Technology

- Awarded Innovation in Remote Cardiac Diagnostics as part of 2025 Medical Device Network Excellence program
 - Recognizes forward-thinking companies that drive positive change and set new standards for excellence across the global medical device sector
- Named finalist in 2025 Octane High Tech Awards
 - Recognizes top innovators, entrepreneurs and technology leaders with presence in Orange County
- Growing list of recent industry recognitions:











Executing on our Go-To-Market Strategy

INFRASTRUCTURE

- On-demand cardiology reader service
- Customer service
- Contract manufacturing
- Logistics and fulfillment

PRODUCT

- Commercial software updates
- Commercial hardware
- Quality enhancements

PILOT MARKETS

- 2 markets: Florida and Southern California
- Meeting with target concierge practices

PRICING MODEL

- Establish premium pricing and subscription model
- Validating with concierge practices

CONCIERGE MARKET

- 1.5 million US concierge patients
- 1/3 cardiac: 500,000 people

\$250 - \$500 million annual revenue*

Overall direct patient market: \$1.3-\$2.6 bn in annual revenue**

* At \$500-\$1,000 / year price

** 1/3 with elevated cardiac risk, top 5% income: 2.6 million people

Significant Increase in Business Development Activity

Strategic Partners

- Significant inbound interest from industry partners given proximity to FDA clearance
- HeartBeam is an ideal platform technology
- Multiple applications and ability to use at home

Areas of Interest

- Data and Al
- Companion products and services
- Complementary diagnostics and treatments
- Distribution

Expanded the business development team with recognized industry expert

Capital Efficiency to Create Value for Shareholders

- Net cash used in operating activities of \$3.4M for Q2 FY'25.
 - 23% decrease quarter-over-quarter.
 - Recurring baseline expenditure of ~\$3.1M, below historic baseline of \$3.5M.
 - Investments in commercial readiness activities and manufacturing capabilities of ~\$0.3M.
- Continue to strategically fund Company.
 - Strong financial discipline in place as we continue to de-risk business.
 - Made substantial cuts to ensure runway through key milestones.
 - Continuing to time commercial team spend in line with FDA Clearance.
 - Focused on minimizing further dilution to shareholders.

\$ in Thousands	Quarter Ended		Quarter Ended	
	June 30, 2025 ⁱⁱⁱ		March 31, 2025	
Operating Expenses:				
General & Administrative	\$	1,711	\$	2,012
Research & Development		3,326		3,492
Total Operating Expenses		5,037		5,504
Loss from Operations		(5,037)		(5,504)
Interest and Other (expense)		63		20
Net loss	\$	(4,974)	\$	(5,484)
Less non-cash items and timing differences		1,529		1,007
Net cash used in operating activities	\$	(3,445)	\$	(4,477)
\$ in Thousands	June 30, 2025		March 31, 2025	
Cash & Cash Equivalents	\$	3,256	\$	4,390
Short-term Investments	\$	1,797	\$	3,760
Total Cash and cash equivalents and short-term investments	\$	5,053	\$	8,150

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Quarter Ended June 30,						
2025		2024				
\$	1,711	\$	2,246			
	3,326		2,844			
	5,037		5,090			
	(5,037)		(5,090)			
	63		134			
\$	(4,974)	\$	(4,956)			
	1,529		1,406			
\$	(3,445)	\$	(3,550)			
June 30, 2025		June 30, 2024				
\$	3,256	\$	9,212			
\$	1,797	\$	-			
\$	5,053	\$	9,212			

- i. Net cash used in operating activities is calculated as Net loss, less non-cash items such as stock-based compensation and timing differences.
- i. Based on maturity dates, a portion of cash was considered held as short-term investments at June 30, 2025.
- iii. Refer to the Form 10-Q unaudited financial statements for the quarter ended June 30, 2025.

HeartBeam: Q2 2025 Summary

- FDA clearance for 12-lead ECG synthesis software submission for arrhythmia assessment remains on track, expected by end of year
- Completing commercial readiness activities: product, infrastructure, target practices
- Marked increase in interest from industry partners, as we get closer to our 12-lead ECG synthesis software clearance and commercialization
- Heightened capital efficiency to create value for shareholders and minimize dilution
- HeartBeam poised to be a fundamental advance in cardiac care

Q&A

Company

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