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# iRhythm Holdings William Blair 46<sup>th</sup> Annual Growth Stock Conference

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June 2, 2026

iRHYTHM<sup>®</sup>

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YEARS OF  
INNOVATION

# Cautionary statement re forward-looking statements, non-GAAP measures and other matters

Certain data in this presentation was obtained from various external sources, and neither “iRhythm Holdings, Inc. (‘iRhythm’ or the ‘Company’)” nor its affiliates, advisers or representatives has verified such data with independent sources. Accordingly, neither the Company nor any of its affiliates, advisers or representatives makes any representations as to the accuracy of that third-party data or undertakes to update such data after the date of this presentation. Such data involves risks and uncertainties and is subject to change based on various factors. The trademarks included herein are the property of the owners and are used for reference purposes only. Such use should not be construed as an endorsement of the products or services of the Company.

This presentation and the accompanying oral presentation include forward-looking statements within the meaning of Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended, and the Private Securities Litigation Reform Act of 1995. Forward-looking statements give the Company’s current expectations or forecasts of future events. An investor can identify these statements by the fact that they do not relate strictly to historical or current facts. They use words such as ‘anticipate’, ‘estimate’, ‘expect’, ‘intend’, ‘will’, ‘project’, ‘plan’, ‘believe’, ‘target’ and other words and terms of similar meaning in connection with any discussion of future operating or financial performance. In particular, these include statements relating to future actions, prospective products or product approvals, future performance or results of current and anticipated products, sales efforts, expenses, strategy and plans, market size and opportunity, competitive position, industry environment, potential growth opportunities, business model, reimbursement rates and coverage, the outcome of contingencies such as legal proceedings and our expectations for future operations and results.

These forward-looking statements are subject to a number of risks, uncertainties and assumptions, including, but not limited to, those described herein and in “Risk Factors” in our most recent 10-K and 10-Q filed with the SEC. Moreover, we operate in a very competitive and rapidly changing environment. New risks emerge from time to time. It is not possible for our management to predict all risks, nor can we assess the impact of all factors on our business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statements we may make. In light of these risks, uncertainties and assumptions, the forward-looking events and circumstances discussed in this presentation may not occur and actual results could differ materially and adversely from those anticipated or implied in the forward-looking statements.

You should not rely upon forward-looking statements as predictions of future events. Although we believe that the expectations reflected in the forward-looking statements are reasonable, we cannot guarantee that the future results, levels of activity, performance or events and circumstances reflected in the forward-looking statements will be achieved or occur. Moreover, except as required by law, neither we nor any other person assumes responsibility for the accuracy and completeness of the forward-looking statements. We undertake no obligation to update publicly any forward-looking statements for any reason after the date of this presentation to conform these statements to actual results or to changes in our expectations.

This presentation regarding the Company and the accompanying oral presentation shall not constitute an offer to sell or the solicitation of an offer to buy any securities, nor shall there be any sale of these securities in any state or jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration or qualification under the securities laws of any such state or jurisdiction. Sales and offers to sell iRhythm securities will only be made in accordance with the Securities Act of 1933, as amended, and applicable SEC regulations, including prospectus requirements.

This presentation and the accompanying oral presentation contain non-GAAP financial measures. The appendix to this presentation reconciles the non-GAAP financial measures to the most directly comparable financial measure prepared in accordance with Generally Accepted Accounting Principles (GAAP). These non-GAAP financial measures include adjusted operating expenses, adjusted net income (loss), adjusted net income (loss) per share, adjusted EBITDA, adjusted EBITDA margin, and free cash flow. iRhythm reports non-GAAP financial measures in addition to, and not as a substitute for, or superior to, measures of financial performance prepared in accordance with GAAP. We believe that non-GAAP financial measures should only be used to evaluate our results of operations in conjunction with the corresponding GAAP financial measures. Other companies, including other companies in our industry, may not use this measure or may calculate this measures differently than as presented. We encourage investors to carefully consider our results under GAAP as well as our supplemental non-GAAP information and reconciliations between these presentations to more fully understand our business.

# Addressing major challenges and opportunities in healthcare



## THE PROBLEM

Heart disease is the #1 cause of death globally.

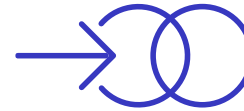
Annual cardiovascular costs in the US: \$393B, projected to reach \$1.5T by 2050.

Cost of ER visit for stroke or heart failure can exceed ~\$10K - \$20K.



## MARKET CATALYSTS

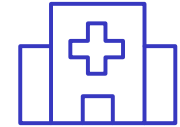
Aging population, consumer arrhythmia awareness, proliferation of therapies like pulsed field ablation, recognition of post-ablation monitoring, trends toward proactive medicine, and a growing shift to value-based care driving TAM growth.



## THE ACCESS GAP

46.3% of all U.S. counties – and 86.2% of rural counties – lack cardiologists and non-urgent cardiology wait times average 26.6 days & rising.

Holter monitoring can miss ~30–60% of arrhythmias that occur beyond the initial monitoring window.



## IRHYTHM'S IMPACT

20 years advancing cardiac diagnostics and innovating to detect, predict, & prevent disease.

Physicians agree with Zio 99% of the time.

[See appendix for sources](#)

# At the Heart of it: The Moat - iRhythm is more than a device

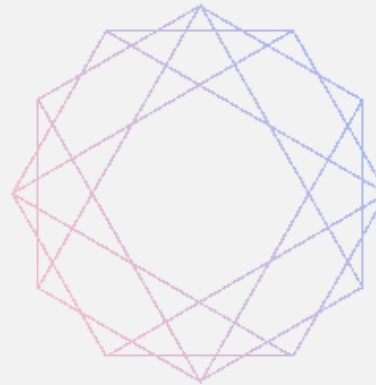


*Device*

## PATENTED WEARABLE BIOSENSORS

Single-use monitor designed for clinical accuracy and patient comfort

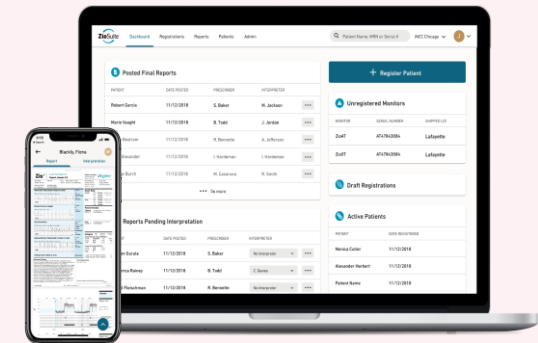
ZIO



*Data*

## ADVANCED SOFTWARE AND AI APPLIED TO A RICH DATASET

2<sup>nd</sup> Generation, FDA-cleared, deep-learned ECG detection algorithm interrogating 12M+ patient reports. Predictive care platform for population health management



*Services*

## CLINICAL SERVICE & OPERATIONAL WORKFLOW

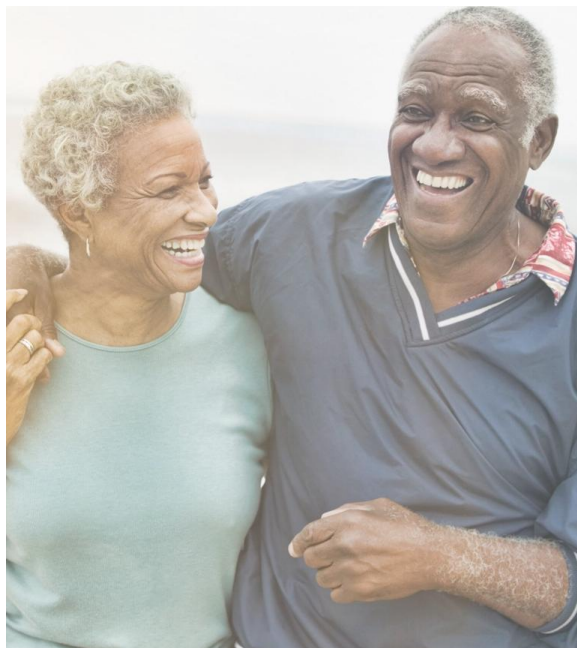
Convenient ordering and delivery  
High-quality digital reports  
Efficient-at-scale clinical oversight

**\$199.4 million**

First quarter 2026 revenue,  
a 25.7% increase year-over-year

**2.6 million**

Patient reports posted annually

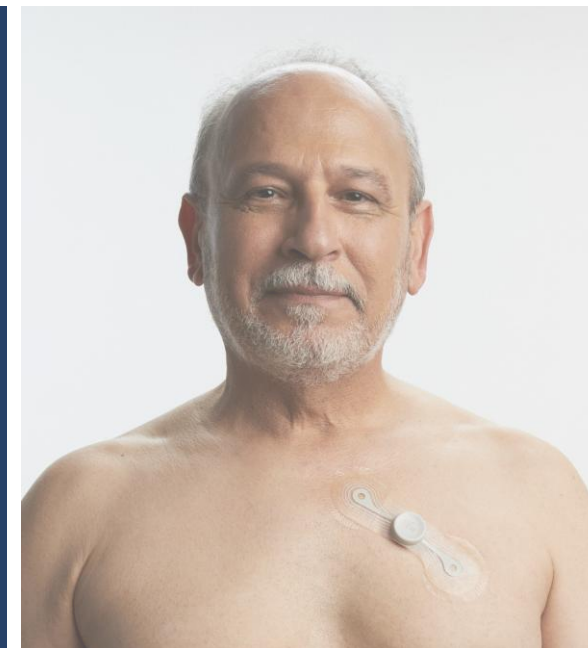


**27+ million**

Potential patients in the United  
States who could benefit from  
ambulatory cardiac monitoring\*

**3.2 million tests**

Target market opportunity  
across prioritized EU and APAC  
countries\*



**~40%**

Penetration in core  
U.S. ambulatory cardiac  
monitoring market as of  
March 31, 2026

**12+ million**

Patient reports posted since  
company inception through  
March 31, 2026



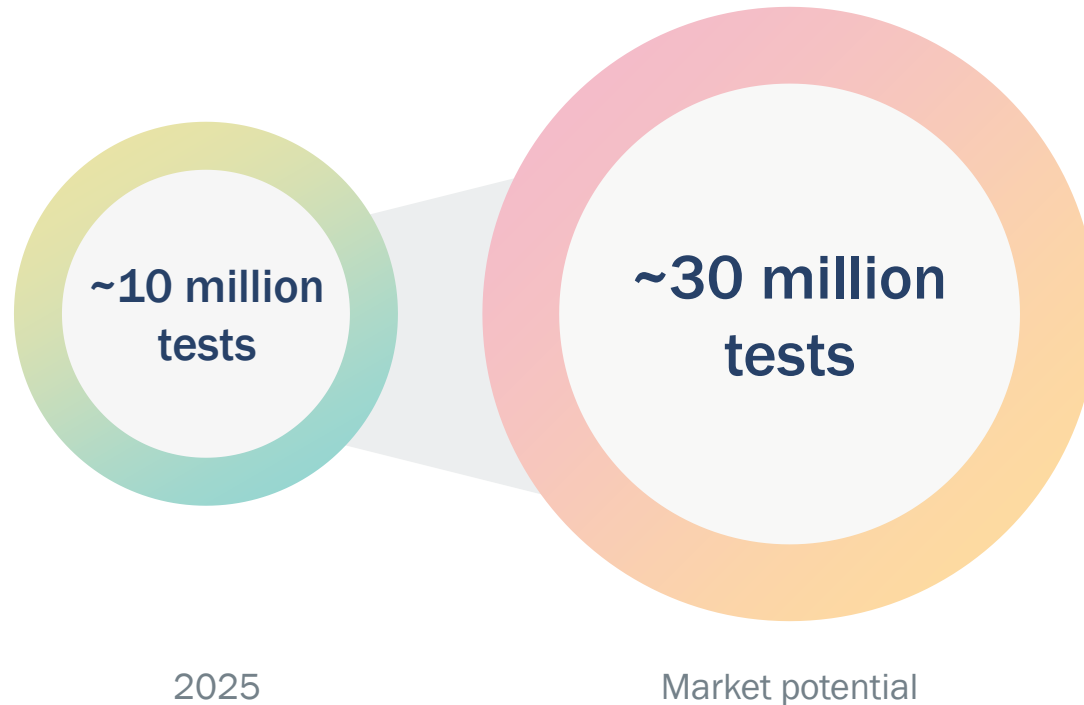
**140+**

Original scientific research  
manuscripts

**3+ billion**

Hours of curated  
ECG data since company  
inception through March 31,  
2026

# ACM market is evolving with runway for durable growth



## TREMENDOUS MARKET EXPANSION POTENTIAL AHEAD

**MARKET LEADERSHIP IN LTCM:** 72% market share in LTCM (of 3.5 million tests in US today growing high teens % YOY) with ~ 27 million undiagnosed US patients at elevated risk

**SHORT DURATION MONITORS YET TO CONVERT:** 1.9 million legacy technology tests still performed in US today, a ~\$500 million revenue opportunity

**MARKET SHARE OPPORTUNITY IN MCT:** 15% market share in MCT (of 1.1 million MCT tests in US today growing high single digit % YOY), with each 10 points of share = ~\$80 - \$100 million

**INTERNATIONAL MARKET EXPANSION:** 3.2 million ACM tests in active OUS markets

Increasing TAM fueled by aging population, adjacent therapies and clinical evidence driving value-based care adoption

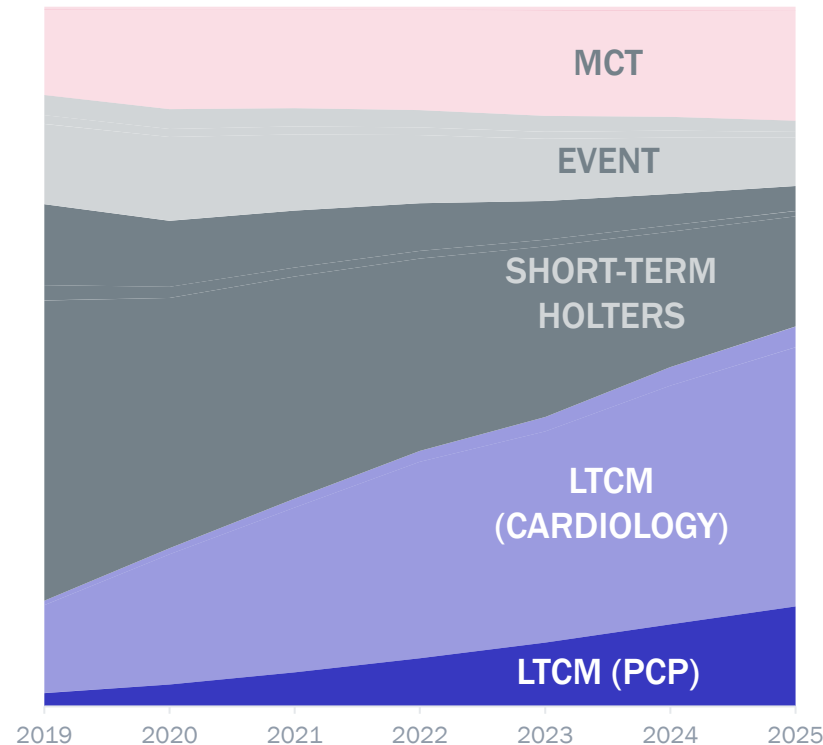
ACM = ambulatory cardiac monitoring; LTCM = long-term continuous monitoring; MCT = mobile cardiac telemetry. Estimates based off combination of Internal Data, Medicare Public-Use Files, IQVIA data, Definitive Healthcare data, Komodo Health data, and other publicly-available information.

# Primary care adoption key to accessing expansive TAM

## IRHYTHM UNIQUELY POSITIONED TO WIN IN PRIMARY CARE

- ✓ Market leader in LTCM, the preferred modality
- ✓ Rule-in/rule-out tool to streamline clinician workflows
- ✓ Home enrollment capabilities
- ✓ Scale and efficiency
- ✓ EHR integration

## TOTAL MARKET CLAIMS BY MODALITY & SPECIALTY



## APPROACHING PRIMARY CARE VIA TWO-PRONGED STRATEGY



Land-and-expand within integrated delivery networks



Integration at large national accounts

TAM = total addressable market; ACM = Ambulatory cardiac monitoring; LTCM = Long-term continuous monitoring; MCT = Mobile cardiac telemetry; EHR = Electronic health records. Estimates based off combination of Internal Data, Medicare Public-Use Files, IQVIA data, Definitive Healthcare data, Komodo Health data, and other publicly-available information.

# Precise, proactive monitoring enables population health management...



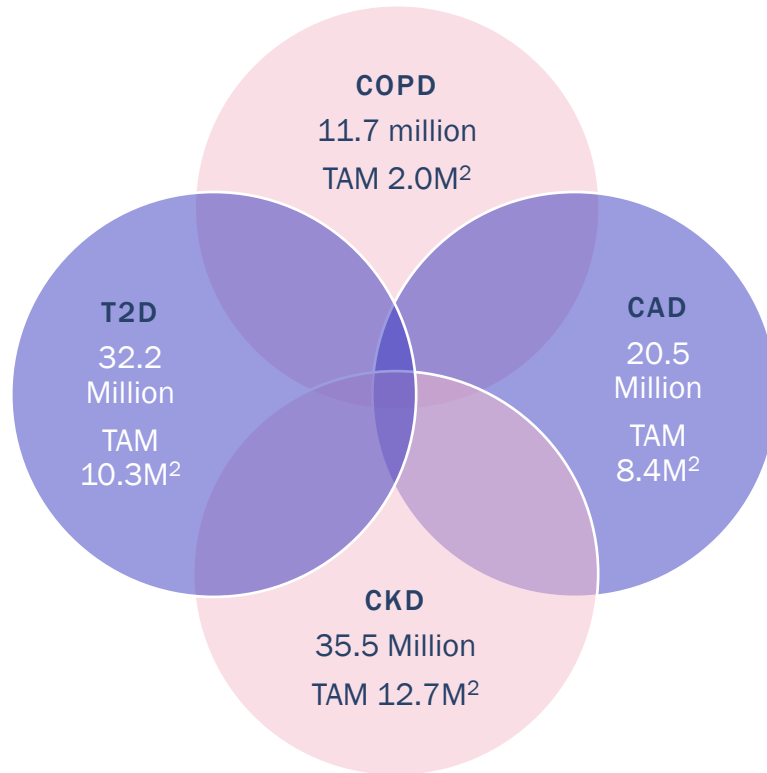
## SLEEP APNEA:

US prevalence of 40.1 million, with up to 80% of AFib patients having sleep apnea<sup>1</sup>



## HEART FAILURE:

US prevalence of 8.4 million, with 25% of heart failure patients having AFib<sup>1</sup>



## ARRHYTHMIA PATIENTS USE THE HEALTHCARE SYSTEM AT A MUCH HIGHER RATE COMPARED TO NON-ARRHYTHMIA PATIENTS<sup>1</sup>

Arrhythmia patients hospitalized more than 2x per 1,000 cohort patients per year than non-arrhythmia patients

Of those hospitalized, patient stay increased by 2-5 days for arrhythmia patients

More than 2x emergency room visits in arrhythmia cohort relative to non-arrhythmia cohorts

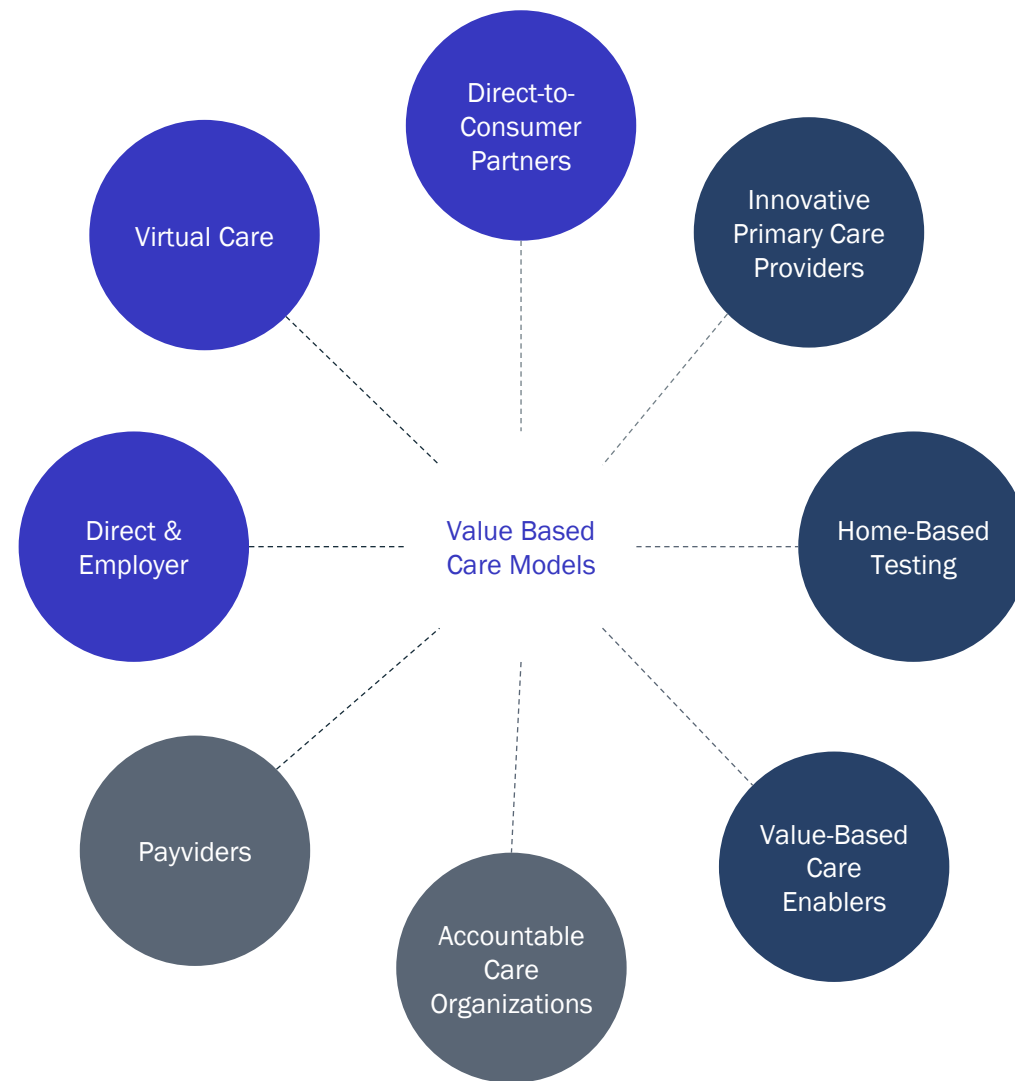
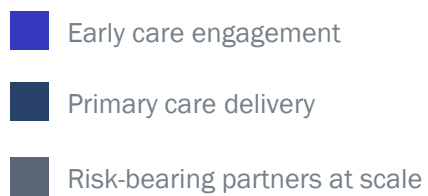
Afib = Atrial fibrillation; T2D = Type 2 diabetes; COPD = Chronic obstructive pulmonary disorder; CAD = Coronary artery disease; CKD = Chronic kidney disease. 1. [See appendix for sources.](#) 2. TAMs are not mutually exclusive as there is likely significant overlap in patient populations.

...which is being embraced by innovative channel partnerships

~85%

of patients in representative customer programs had at least one arrhythmia identified<sup>1</sup>

1. Data on file. iRhythm Technologies, Inc. 2026



# Next-gen MCT designed to extend our category leadership



**Zio AT®**



**Zio MCT\*\***

(Not yet FDA cleared, Expected H1'27)

|                          | Zio AT®                                                                           | Zio MCT**<br>(Not yet FDA cleared, Expected H1'27)                                                                                                                                                                        |
|--------------------------|-----------------------------------------------------------------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Device form factor       | Legacy patch technology                                                           | Improved form factor <ul style="list-style-type: none"> <li>• Same platform as Zio monitor</li> <li>• Better adhesion and battery</li> </ul>                                                                              |
| Wear duration            | Up to 14 days                                                                     | Up to 21 days                                                                                                                                                                                                             |
| Data transmission        | Auto-detects and transmits symptomatic and asymptomatic events during wear period | Increased maximum transmission limit and enhanced auto-detection algorithm                                                                                                                                                |
| Algorithms and reporting | Interim reports available plus final end-of-wear report                           | Enhanced arrhythmia detection and better reporting <ul style="list-style-type: none"> <li>• Advanced software for enhanced detection parameters</li> <li>• Improved final wear report with additional insights</li> </ul> |

\*Continuous, uninterrupted refers to the recording of ECG data. Zio AT Gateway transmissions may be impacted by a variety of factors. See Product Labeling for more information. †Zio AT is contraindicated for critical care patients. ‡Do not use Zio AT for patients with symptomatic episodes where variations in cardiac performance could result in immediate danger to the patient or when real-time or in-patient monitoring should be prescribed. Refer to the Zio AT labeling and Clinical Reference Manual for full contraindications. \*\*Zio MCT not yet FDA cleared.

# Significant runway in international expansion



[See appendix for sources](#)

# Durable AI advantage creates high barriers to entry

*Clinical-grade AI isn't an algorithm, it's an end-to-end, regulated platform built over decades*

## PROPRIETARY DATASET + SIGNAL QUALITY

**>3B hours of curated ECG data across 12M+ patient reports**

- Expert-annotated multi-day recordings from real-world settings
- 98%+ analyzable signal from purpose-built biosensor
- Public datasets are small, outdated, and limited to 10-second segments

## VALIDATED AI PLATFORM

**>15 years of iterative ZEUS AI development**

- FDA-cleared AI for arrhythmia detection, AF burden, and reporting
- Clinician-validated outputs before physician delivery
- 99% physician agreement with Zio

## REGULATORY + QUALITY + CLINICAL ADVANTAGES

**Evidence at scale**

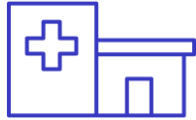
- Multiple generations of 510(k) clearances
- Higher FDA bar, but more predictable at scale
- Post-market monitoring, QMS, and surveillance built into systems
- Differentiation validated in head-to-head studies

## END-TO-END WORKFLOW + DISTRIBUTION

**Fully integrated platform**

- Device + logistics + IDTF + AI + validation + physician-ready insights
- Native EHR integration into existing clinician workflows
- Scalable onboarding, implementation, and account support

# Comprehensive evidence generation supportive of clinical use cases



## CLINICAL OUTCOME SUPERIORITY

CAMELOT and AVALON studies show Zio LTCM service demonstrate advantages over other ACM modalities in clinical performance and healthcare utilization. Data at ACC and HRS further demonstrate the benefits of Zio across multiple patient populations (cardiometabolic, post-ablation, pregnancy).



## ECONOMIC VALUE

Zio LTCM service associated with a lower probability of inpatient hospitalization, fewer ED visits, and lower all-cause healthcare costs compared to other ACM modalities.



## ADJACENT PATIENT POPULATIONS

In patients with T2D and/or COPD, patients also having arrhythmias used more healthcare resources than non-arrhythmia patients.

The average cost per arrhythmia patient was \$17.2k compared to \$1.7k for a non-arrhythmia patient.

# Next expansion opportunity: obstructive sleep apnea



## SLEEP APNEA

- Prevalence of sleep disorders<sup>1</sup>: 40.1 million
- Large prevalence of undiagnosed sleep apnea missing a convenient diagnostic pathway for many PCPs
- Natural evolution of iRhythm's platform capabilities:
  - Up to 80% of AFib patients have sleep apnea
  - ~20% of iRhythm HCPs prescribe sleep tests
  - Device/AI/IDTF services utilized for diagnosis
  - Current reimbursement established

## Home Sleep Test (HST) ordering (via third party) in ZioSuite<sup>®</sup> from iRhythm<sup>®</sup>

With HST ordering through ZioSuite, iRhythm simplifies and streamlines this important need for your patients and staff.\*



HCP = Healthcare provider. 1. Estimated 2022 prevalence in the United States. [See appendix for sources.](#)

# 1Q26 milestones reflective of execution and momentum

## ACCELERATING MOMENTUM IN COMMERCIAL BUSINESS



\$199.4 million revenue for 1Q26, reflective of continued commercial adoption and market expansion

12+ million patient reports worldwide generated to date

Opening new channel partnerships to address 27M patient opportunity

Commercialized in six OUS markets

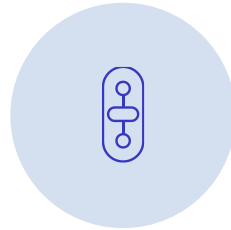
## PROVIDING A WINNING CUSTOMER EXPERIENCE



53% registration volumes from EHR-integrated accounts

Over three quarters of our top 100 customers now EHR-integrated

## BRINGING INNOVATIVE PRODUCTS TO MARKET



Strong progress supports 1H27 Zio® MCT launch timeline

Advancing next generation AI algorithm toward regulatory clearance

First health system deployment of predictive identification workflows

## GENERATING PEER-REVIEWED CLINICAL EVIDENCE



ACC and HRS data showing superiority of Zio® LTCM

Over 140 original scientific research manuscripts published to date, demonstrating leadership in ACM clinical evidence generation

## EXPANDING MARKET ACCESS



Major policy shifts to provide favorable position for Zio®

Zio® covered by incremental payer policies

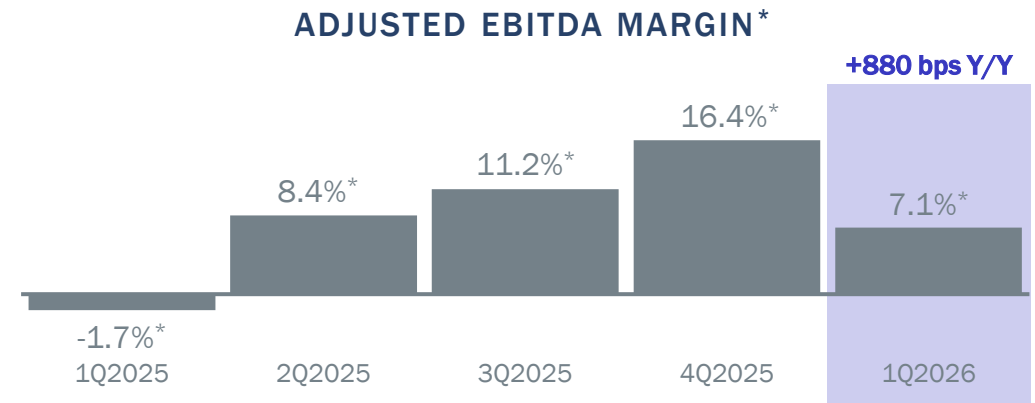
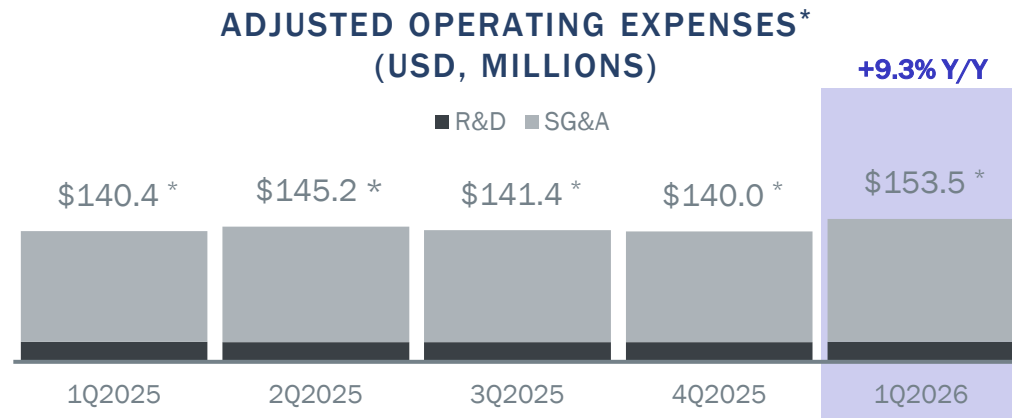
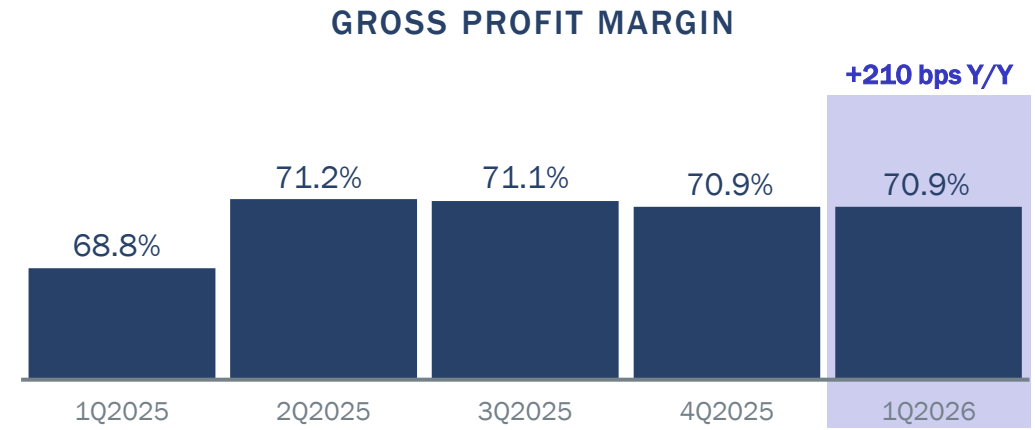
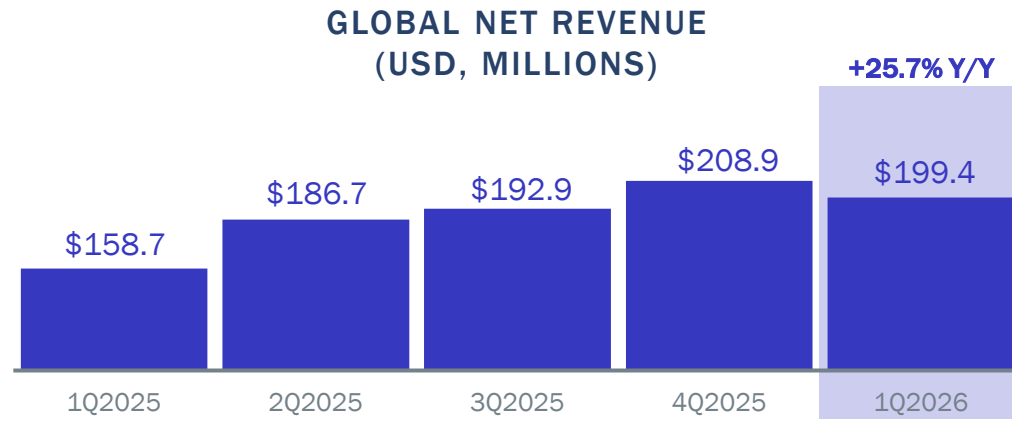
## EXECUTING WITH DISCIPLINE & EFFICIENCY



7.1% adj. EBITDA margin, demonstrating ability to deliver sustained margin expansion

Implemented additional operational efficiencies for sustained profitable growth

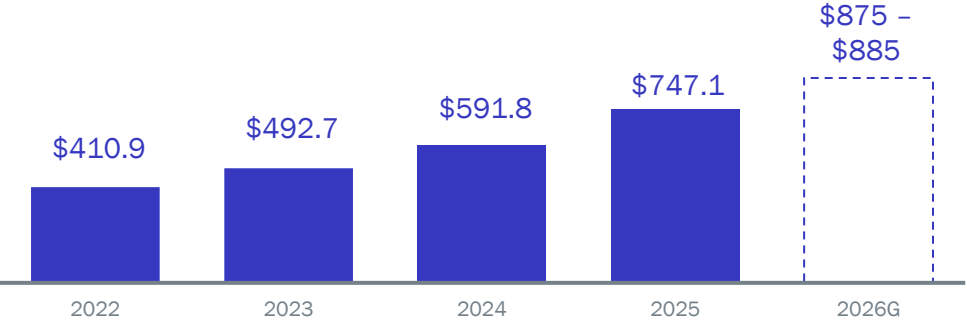
# First quarter 2026 financial performance



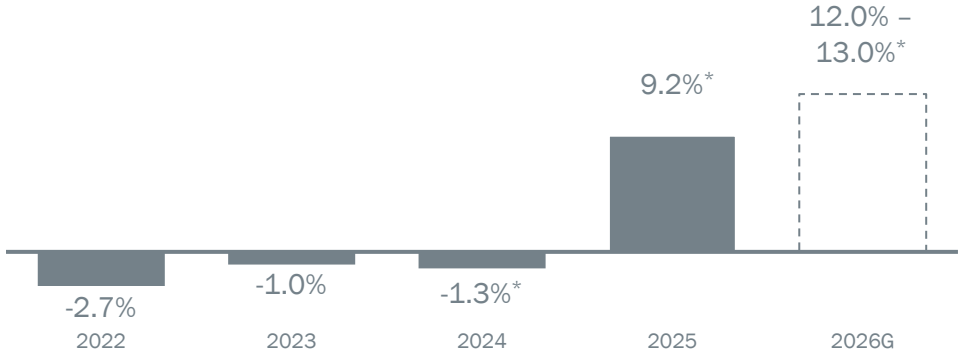
\*Adjusted operating expenses and adjusted EBITDA margin for 1Q25, 2Q25, 3Q25, 4Q25, and 1Q26 include \$0.3 million, \$1.7 million, \$0.3 million, \$0.7 million, and \$0.3 million respectively, of acquired in-process research and development expense. Adjusted operating expenses exclude impacts from business transformation, certain intellectual property litigation expenses, and impairment and restructuring charges. Adjusted EBITDA excludes non-cash operating charges for stock-based compensation expense, changes in fair value of strategic investments, impairment and restructuring charges, business transformation costs, certain intellectual property litigation expenses, and loss on extinguishment of debt. Business transformation costs include costs associated with professional services, employee termination and relocation, third-party merger and acquisition, integration, and other costs to augment and restructure the organization, inclusive of both outsourced and offshore resources.

# 2026 outlook balances growth across near-term opportunities

GLOBAL NET REVENUE (USD, MILLIONS)



ADJUSTED EBITDA MARGIN\*



## U.S. CORE COMMERCIAL BUSINESS

- Further expansion into PCP channel
- New technologies (e.g., PFA) expand monitoring
- MCT market expansion with continued innovation

## INTERNATIONAL EXPANSION

- Continued penetration in the UK and national reimbursement
- Entry into Japan, the second largest global ACM market
- Commercial ramp in select European countries

## ADJACENT MARKET OPPORTUNITIES

- Movement into proactive monitoring programs
- Initial commercial pilots into obstructive sleep apnea

\*Adjusted EBITDA margin for the years ended December 31, 2024, 2025 and 2026 include acquired in-process research and development expense. Adjusted EBITDA excludes non-cash operating charges for stock-based compensation expense, changes in fair value of strategic investments, impairment and restructuring charges, business transformation costs, certain intellectual property litigation expenses and settlements, and loss on extinguishment of debt. Business transformation costs include costs associated with professional services, employee termination and relocation, third-party merger and acquisition, integration, and other costs to augment and restructure the organization, inclusive of both outsourced and offshore resources.



Addressing  
the future focus  
of healthcare



Expanding core  
& unlocking  
adjacent markets



Growing revenue  
through global market  
expansion



Driving  
meaningful  
improvements in  
financial profile

iRHYTHM®

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# Appendix



# Reconciliation of net income (loss) to adjusted EBITDA

Adjusted EBITDA is a non-GAAP financial measure and is presented for supplemental informational purposes only and should not be considered as an alternative or substitute to financial information presented in accordance with GAAP. Adjusted EBITDA excludes non-cash operating charges for stock-based compensation expense, changes in fair value of strategic investments, impairment and restructuring charges, business transformation costs, certain intellectual property litigation expenses and settlements, and loss on extinguishment of debt. Business transformation costs include costs associated with professional services, employee termination and relocation, third-party merger and acquisition, integration, and other costs to augment and restructure the organization, inclusive of both outsourced and offshore resources.

(USD, THOUSANDS)

| ADJUSTED EBITDA RECONCILIATION*                     | THREE MONTHS ENDED MARCH 31, |                   |
|-----------------------------------------------------|------------------------------|-------------------|
|                                                     | 2026                         | 2025              |
| Net loss, as reported <sup>1</sup>                  | \$ (13,933)                  | \$ (30,700)       |
| Interest expense                                    | 3,290                        | 3,273             |
| Interest income                                     | (4,879)                      | (4,919)           |
| Changes in fair value of strategic investments      | (1,447)                      | (843)             |
| Income tax provision                                | 500                          | 665               |
| Depreciation and amortization                       | 5,042                        | 5,210             |
| Stock-based compensation                            | 21,491                       | 23,344            |
| Business transformation costs                       | 346                          | 503               |
| Intellectual property litigation costs <sup>2</sup> | 3,689                        | 832               |
| <b>Adjusted EBITDA</b>                              | <b>\$ 14,099</b>             | <b>\$ (2,635)</b> |

\*Certain numbers expressed may not sum due to rounding. 1 Net loss for the three months ended March 31, 2026 and 2025, includes \$0.3 million of acquired in-process research and development expense. 2 Excludes third-party attorneys' fees and expenses associated with patent litigation brought against the Company by Welch Allyn, Inc. and Bardy Diagnostics, Inc., subsidiaries of Baxter International, Inc.

# Reconciliation of GAAP to non-GAAP financial information

Adjusted EBITDA, adjusted net income (loss), adjusted net income (loss) per share, adjusted operating expenses, and free cash flow are non-GAAP financial measures and are presented for supplemental informational purposes only and should not be considered as an alternative or substitute to financial information presented in accordance with GAAP. Adjusted EBITDA excludes non-cash operating charges for stock-based compensation expense, changes in fair value of strategic investments, impairment and restructuring charges, business transformation costs, certain intellectual property litigation expenses and settlements, and loss on extinguishment of debt. Business transformation costs include costs associated with professional services, employee termination and relocation, third-party merger and acquisition, integration, and other costs to augment and restructure the organization, inclusive of both outsourced and offshore resources.

| (USD, THOUSANDS)<br>ADJUSTED EBITDA RECONCILIATION* | THREE MONTHS ENDED |                      |                       |                  |                   |
|-----------------------------------------------------|--------------------|----------------------|-----------------------|------------------|-------------------|
|                                                     | MARCH 31,<br>2026  | DECEMBER 31,<br>2025 | SEPTEMBER 30,<br>2025 | JUNE 30,<br>2025 | MARCH 31,<br>2025 |
| Net income (loss) <sup>1</sup>                      | \$ (13,933)        | \$ 5,579             | \$ (5,212)            | \$ (14,218)      | \$ (30,700)       |
| Interest expense                                    | 3,290              | 3,322                | 3,281                 | 3,278            | 3,273             |
| Interest income                                     | (4,879)            | (5,337)              | (5,944)               | (5,321)          | (4,919)           |
| Changes in fair value of strategic investments      | (1,447)            | (1,822)              | (894)                 | (2,152)          | (843)             |
| Income tax (benefit) provision                      | 500                | 447                  | 24                    | (183)            | 665               |
| Depreciation and amortization                       | 5,042              | 5,254                | 5,173                 | 5,105            | 5,210             |
| Stock-based compensation                            | 21,491             | 21,106               | 21,006                | 22,827           | 23,344            |
| Impairment charges                                  | —                  | 1,979                | —                     | 2,479            | —                 |
| Business transformation costs                       | 346                | 692                  | 913                   | 925              | 503               |
| Intellectual property litigation costs <sup>2</sup> | 3,689              | 3,070                | 3,212                 | 2,956            | 832               |
| <b>Adjusted EBITDA</b>                              | <b>\$ 14,099</b>   | <b>\$ 34,290</b>     | <b>\$ 21,559</b>      | <b>\$ 15,696</b> | <b>\$ (2,635)</b> |
| Revenue                                             | \$ 199,390         | \$ 208,890           | \$ 192,884            | \$ 186,687       | \$ 158,677        |
| <b>Adjusted EBITDA margin</b>                       | <b>7.1%</b>        | <b>16.4%</b>         | <b>11.2%</b>          | <b>8.4%</b>      | <b>-1.7%</b>      |

\*Certain numbers expressed may not sum due to rounding. 1 Net income (loss) for 1Q25, 2Q25, 3Q25, 4Q25 and 1Q26 include \$0.3 million, \$1.7 million, \$0.3 million, \$0.7 million and \$0.3 million, respectively, of acquired in-process research and development expense. 2 Excludes third-party attorneys' fees and expenses associated with patent litigation brought against the Company by Welch Allyn, Inc. and Bardy Diagnostics, Inc., subsidiaries of Baxter International, Inc.

# Reconciliation of GAAP to non-GAAP financial information

(USD, THOUSANDS)

|                                                               | THREE MONTHS ENDED MARCH 31, |                    |
|---------------------------------------------------------------|------------------------------|--------------------|
|                                                               | 2026                         | 2025               |
| <b>ADJUSTED NET LOSS RECONCILIATION*</b>                      |                              |                    |
| Net loss, as reported <sup>1</sup>                            | \$ (13,933)                  | \$ (30,700)        |
| Business transformation costs                                 | 346                          | 503                |
| Intellectual property litigation costs <sup>2</sup>           | 3,689                        | 832                |
| Changes in fair value of strategic investments                | (1,447)                      | (843)              |
| Tax effect of adjustments <sup>3</sup>                        | —                            | (91)               |
| <b>Adjusted net loss</b>                                      | <b>\$ (11,345)</b>           | <b>\$ (30,299)</b> |
|                                                               |                              |                    |
|                                                               | THREE MONTHS ENDED MARCH 31, |                    |
|                                                               | 2026                         | 2025               |
| <b>ADJUSTED NET LOSS PER SHARE RECONCILIATION*</b>            |                              |                    |
| Net loss per share, as reported <sup>1</sup>                  | \$ (0.43)                    | \$ (0.97)          |
| Business transformation costs per share                       | 0.01                         | 0.02               |
| Intellectual property litigation costs per share <sup>2</sup> | 0.11                         | 0.03               |
| Changes in fair value of strategic investments per share      | (0.04)                       | (0.03)             |
| Tax effect of adjustments per share <sup>3</sup>              | —                            | —                  |
| <b>Adjusted net loss per share</b>                            | <b>\$ (0.35)</b>             | <b>\$ (0.95)</b>   |
| Weighted-average shares, basic and diluted                    | 32,507                       | 31,590             |

\*Certain numbers expressed may not sum due to rounding. 1 Net loss for the three months ended March 31, 2026 and 2025, includes \$0.3 million of acquired in-process research and development expense. 2 Excludes third-party attorneys' fees and expenses associated with patent litigation brought against the Company by Welch Allyn, Inc. and Bardy Diagnostics, Inc., subsidiaries of Baxter International, Inc. 3. Income tax impact of Non-GAAP adjustments listed.

# Reconciliation of GAAP to non-GAAP financial information

(USD, THOUSANDS)

## ADJUSTED OPERATING EXPENSES RECONCILIATION\*

|                                                     | THREE MONTHS ENDED MARCH 31, |                   |
|-----------------------------------------------------|------------------------------|-------------------|
|                                                     | 2026                         | 2025              |
| Operating expenses, as reported                     | \$ 157,538                   | \$ 141,772        |
| Business transformation costs                       | (346)                        | (503)             |
| Intellectual property litigation costs <sup>1</sup> | (3,689)                      | (832)             |
| <b>Adjusted operating expenses</b>                  | <b>\$ 153,503</b>            | <b>\$ 140,437</b> |

## FREE CASH FLOW RECONCILIATION\*

|                                       | THREE MONTHS ENDED MARCH 31, |                    |
|---------------------------------------|------------------------------|--------------------|
|                                       | 2026                         | 2025               |
| Net cash used in operating activities | \$ (26,173)                  | \$ (7,891)         |
| Purchases of property and equipment   | (6,905)                      | (9,419)            |
| <b>Free cash flow</b>                 | <b>\$ (33,078)</b>           | <b>\$ (17,310)</b> |

\*Certain numbers expressed may not sum due to rounding.

<sup>1</sup> Excludes third-party attorneys' fees and expenses associated with patent litigation brought against the Company by Welch Allyn, Inc. and Bardy Diagnostics, Inc., subsidiaries of Baxter International, Inc.

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