



EBR Systems, Inc.

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John McCutcheon
President and CEO

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Investment highlights

EBR is focused on executing its clear and targeted commercialisation strategy to deliver shareholder value

High value market opportunity



Unique solution

No competition as the WiSE CRT system is complementary to other leadless devices



Large markets

Targeting an initial addressable market of US\$3.6bn



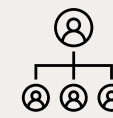
Positive results

Safety and efficacy endpoints met for SOLVE-CRT trial and Breakthrough Device Designation granted



Focused strategy

Clear pathway to achieve FDA approval and progress commercialisation activities to initial revenue

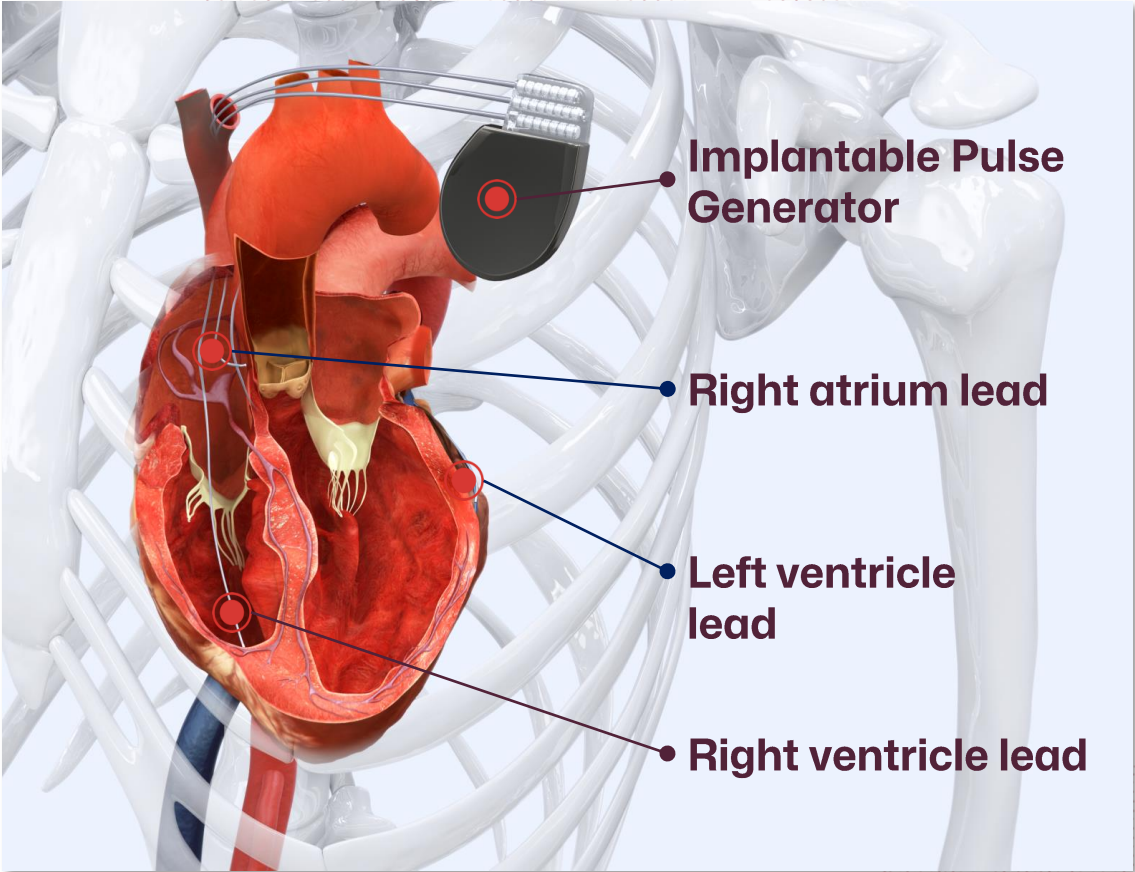
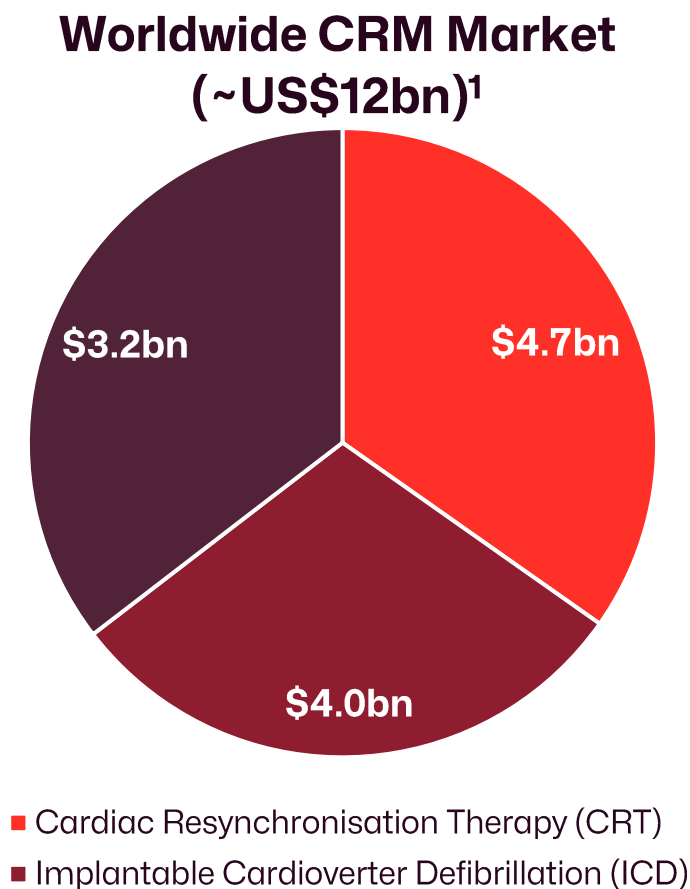


Strong team

Experienced management team with significant clinical development and commercial expertise

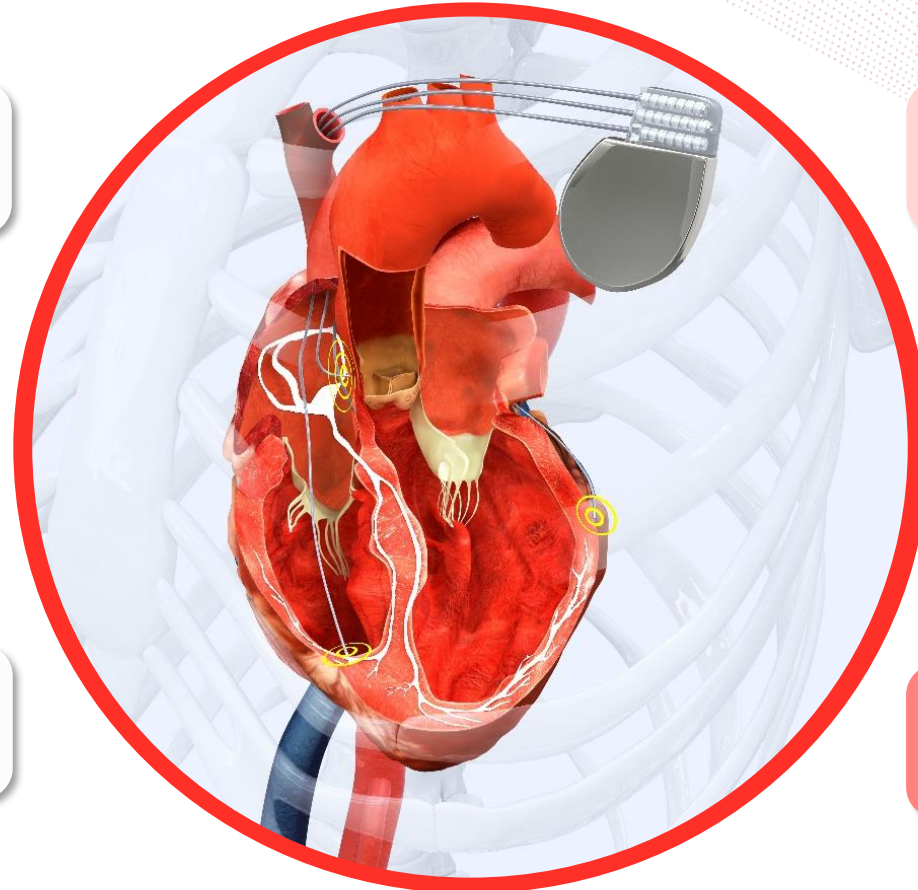
Cardiac Rhythm Management Market

Three key segments driving growth in the global cardiac rhythm management market



Traditional CRT systems are suboptimal

Traditional CRT systems use wires or leads to deliver energy to the heart, which can lead to many problems



Coronary Sinus limits Left Ventricle (LV) lead placement locations



Leads can become a way for pathogens to reach the myocardium



Leads can be associated with phrenic nerve stimulation



Leads can migrate and sometimes fracture



Difficult to place



LV lead must be placed outside the heart to avoid blood clots

EBR has a wireless solution for heart failure

EBR's WiSE CRT System is the only wireless device that can synchronously pace the left ventricle

WiSE CRT System fills the gap

The only leadless solution for left ventricle (LV) pacing

Other wireless pacemakers are too big for LV pacing

Their size increases the risk of blood clots, restricting their use to right ventricle (RV) and right atrium (RA) pacing only

Complementary solution

WiSECRT System can be used in conjunction with wireless RV / RA pacemakers to deliver CRT

Strong competitive protection

WiSE CRT System is protected by over 97 issued patents globally

Left ventricle¹



EBR Systems
WiSE CRT System

Right ventricle / atrium¹



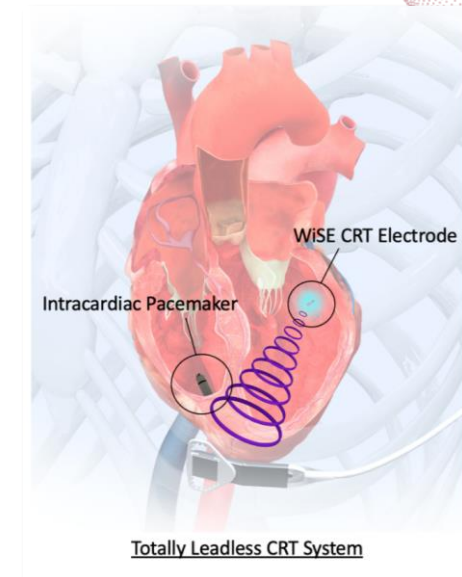
Medtronic
Micra®



Boston
Scientific
Empower®



Abbott
Aveir®



Pivotal SOLVE-CRT Study meets primary endpoints

Positive results confirm WiSE CRT System as a highly effective treatment option for patients with heart failure¹

Primary efficacy endpoint met

-16.4%

p = 0.003

Decrease in left ventricular end systolic volume (vs. -9.3% target), showing improved heart function



Success in high-risk patients

SOLVE-CRT patient pool consists of patients who have failed conventional CRT



Other key data

All data analysed to date shows consistent, positive results in reversing heart failure symptoms and physiology

Primary safety endpoint met

80.9%

p < 0.001

Patients free from type I complications (vs. 70% target)



Safety profile comparable to SoC²

Studies using SoC treatment for CRT upgrades have shown 81.3% freedom from device & procedure related complications³



Other key data

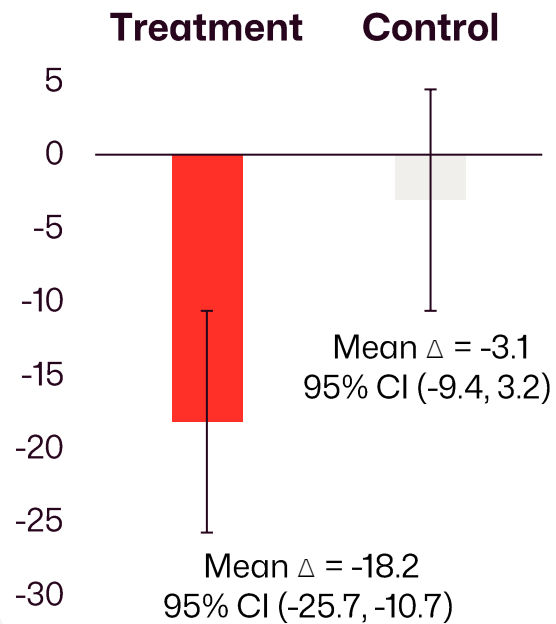
Observed complication rates decreased over time with experience

Randomised sub-study supports primary results

The WiSE CRT System demonstrates clinically and statistically significant evidence of reverse remodelling and electrical response within previously untreatable and high-risk patients

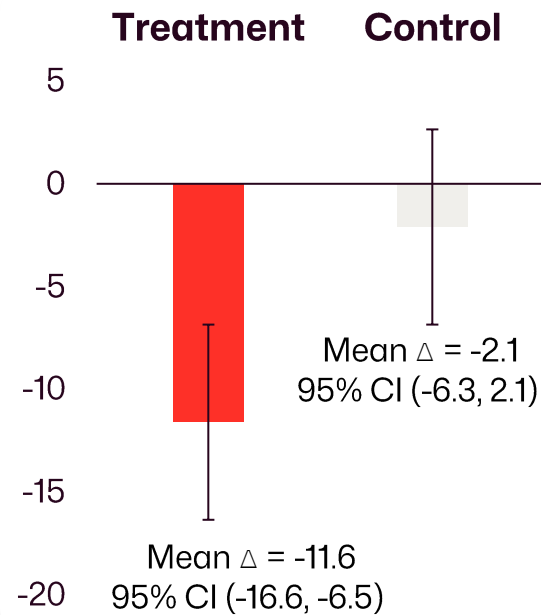
LVESV (%)

$p = 0.002^{\dagger}$



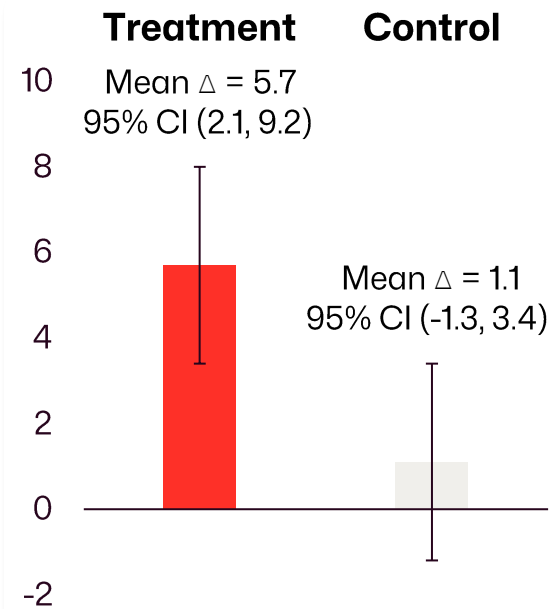
LVEDV (%)

$p = 0.004^{\ddagger}$



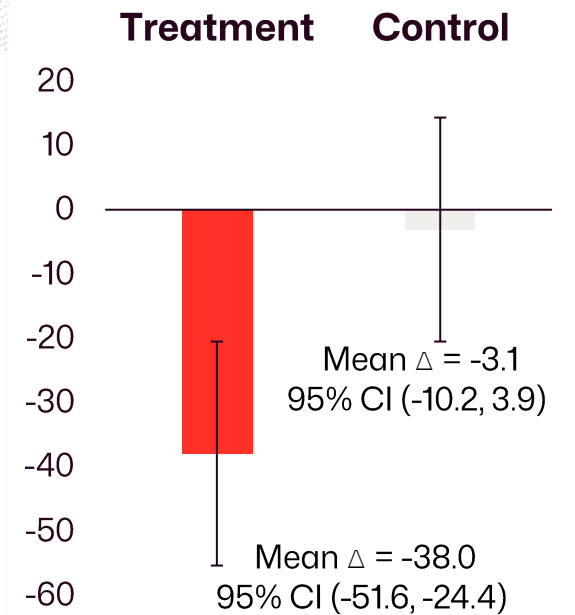
LVEF (%)

$p = 0.025^{\dagger}$



QRS (ms)

$p < 0.001^{\ddagger}$



Control n = 29, Treatment n = 22

Commercialisation pathway

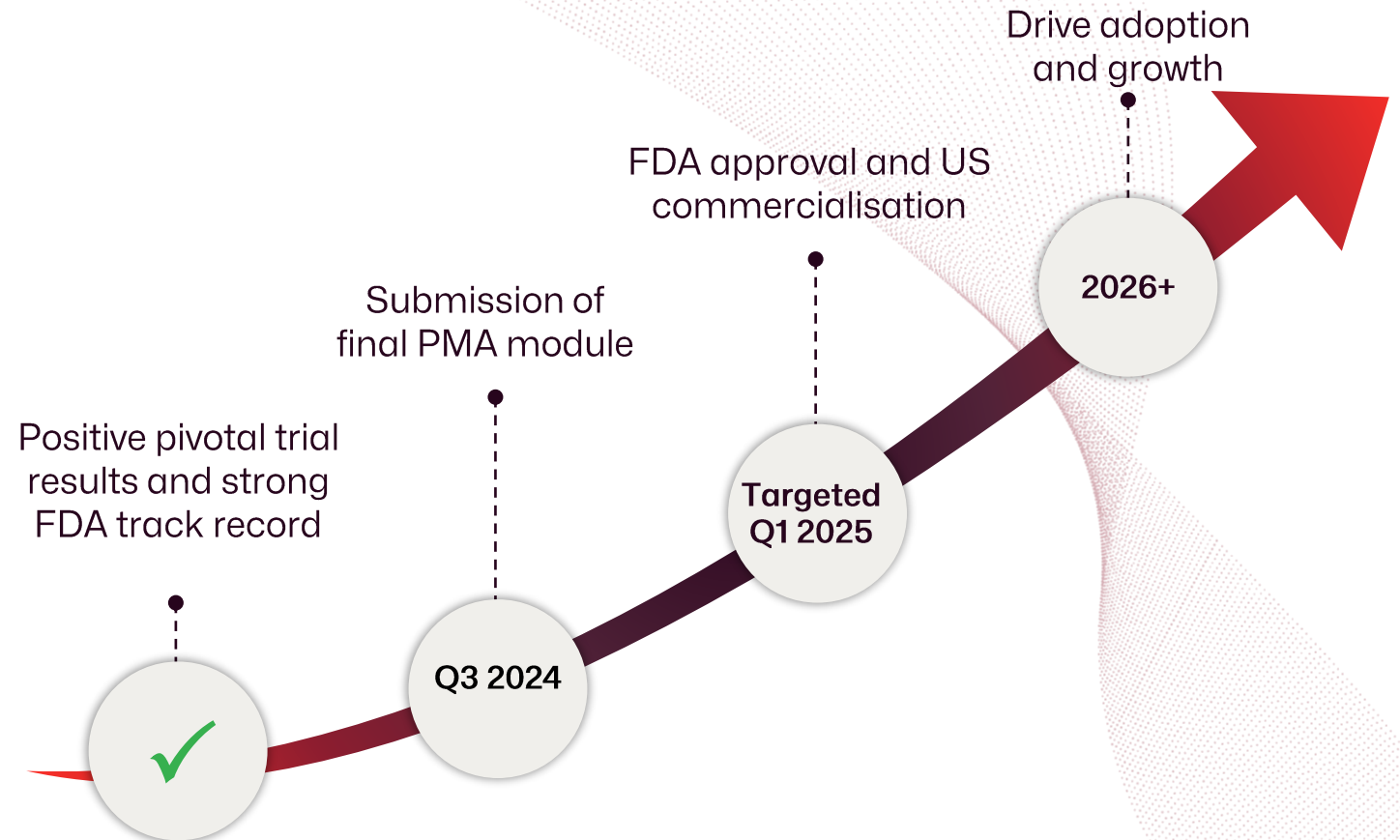
Positive pivotal trial results and strong track record with the FDA provide a clear pathway to approval and commercialisation

FDA approval process underpinned by positive pivotal trial results and a track record of successful engagement with the FDA resulting in:

- Award of Breakthrough Device Designation
- Approval of pivotal study re-design
- Approval of leadless pacemakers as a co-implant in pivotal study

EBR has submitted four out of five required modules for the PMA submission

- The final module includes biocompatibility & device verification testing
- Submission in Q3 2024



Favourable US market dynamics

Market dynamics in the US support initial adoption of the WiSE CRT System



Market validation

- Support of Key Opinion Leaders (KOLs)
- Unmet need underscored by FDA Breakthrough Device designation
- No direct competitors



Low hospital adoption barriers

- Low barrier for opening new accounts
- No capital equipment required and reimbursement available post-approval
- Proven and refined implanter training program



Reimbursement & High ASP¹

- CMS¹ recently updated WiSE specific CPT¹ codes (e.g. 0515T, 0522T)
- CPT¹ codes already assigned to interim APC¹ codes (e.g. 5231, 5741)
- Clear pathway to NTAP¹ and TPT¹ reimbursement schemes post FDA approval
- Automatic process to reassign APC codes based on actual claims data
- WiSE CRT System target US ASP: US\$45,000²

Initial commercialisation strategy

EBR will leverage its established partnerships and presence in the US to drive initial sales growth



Clinical trial sites to drive initial sales

- CRT market is concentrated - targeting high-volume CRT procedure sites
- **2025:** Targeting US sites that have participated in the SOLVE-CRT trial and other high-volume sites with Key Opinion Leaders (KOLs)
- **2026+:** Target top 200 to 250 clinical sites, representing >50% US CRT market



Direct, specialist sales force

- Execution of commercial launch supported by specialised direct sales force to target high volume sites
- SOLVE-CRT core team in place with clinical and technical expertise of WiSE CRT System
- Grow initial sales and expand into new areas with sales force expansion over time



Manufacturing capabilities

- Manufacturing capabilities in place with capability to meet early demand
- Expand in-house manufacturing facility to meet future demand

US\$3.6bn initial addressable market

At commercial launch, EBR estimates an initial addressable market of ~US\$3.6bn⁽¹⁾

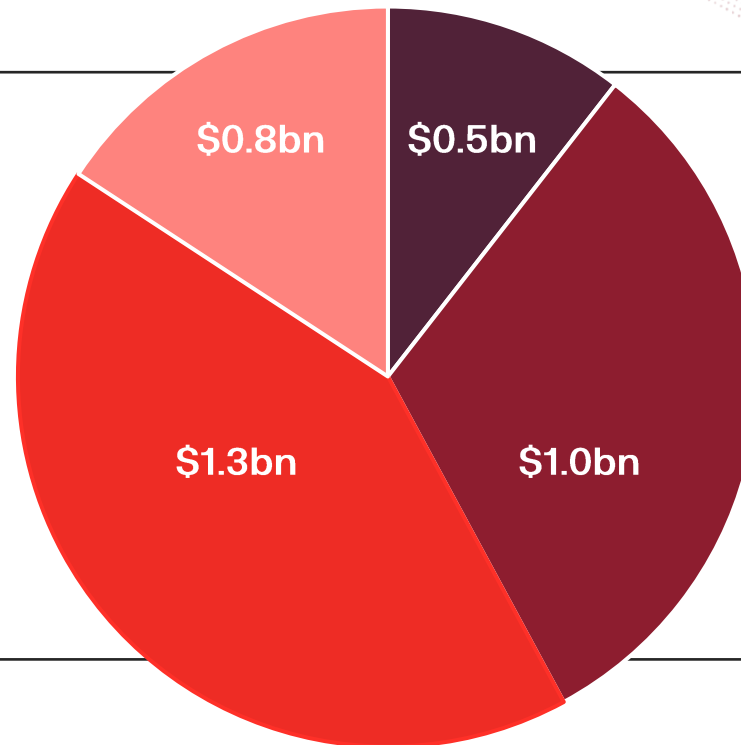
Leadless Upgrades

- Patients with a leadless right ventricle pacemaker can only upgrade with WiSE CRT System to receive effective CRT
- WiSE CRT System used in conjunction with the Medtronic Micra (single chamber) or Abbott Aveir (single chamber) device

Further growth potential – see next slide

Chronic Lead Failure

Traditional CRT system implanted but has ceased to provide effective CRT



Acute Lead Failure

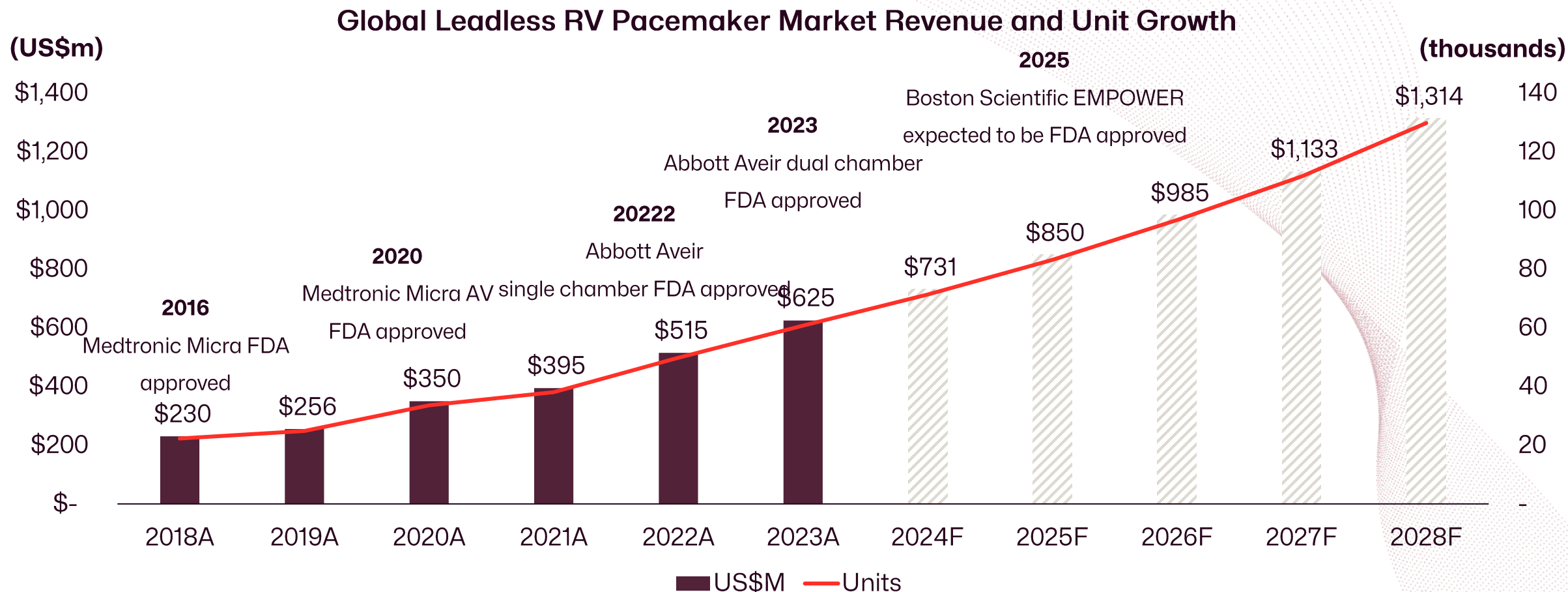
Unable to implant CRT wire in a new CRT patient

High Risk Upgrades

Patient requiring CRT, but is deemed too high risk for a conventional CS lead placement

Leadless pacemaker growth feeds EBR's addressable market

Continued global growth and adoption of leadless RV pacemakers significantly expands EBR's market opportunity



Product development: Rechargeable battery

EBR is developing a new rechargeable battery that will support the WiSE CRT System in becoming a first-line therapy option and treat a broader suite of patients

Commercial benefits

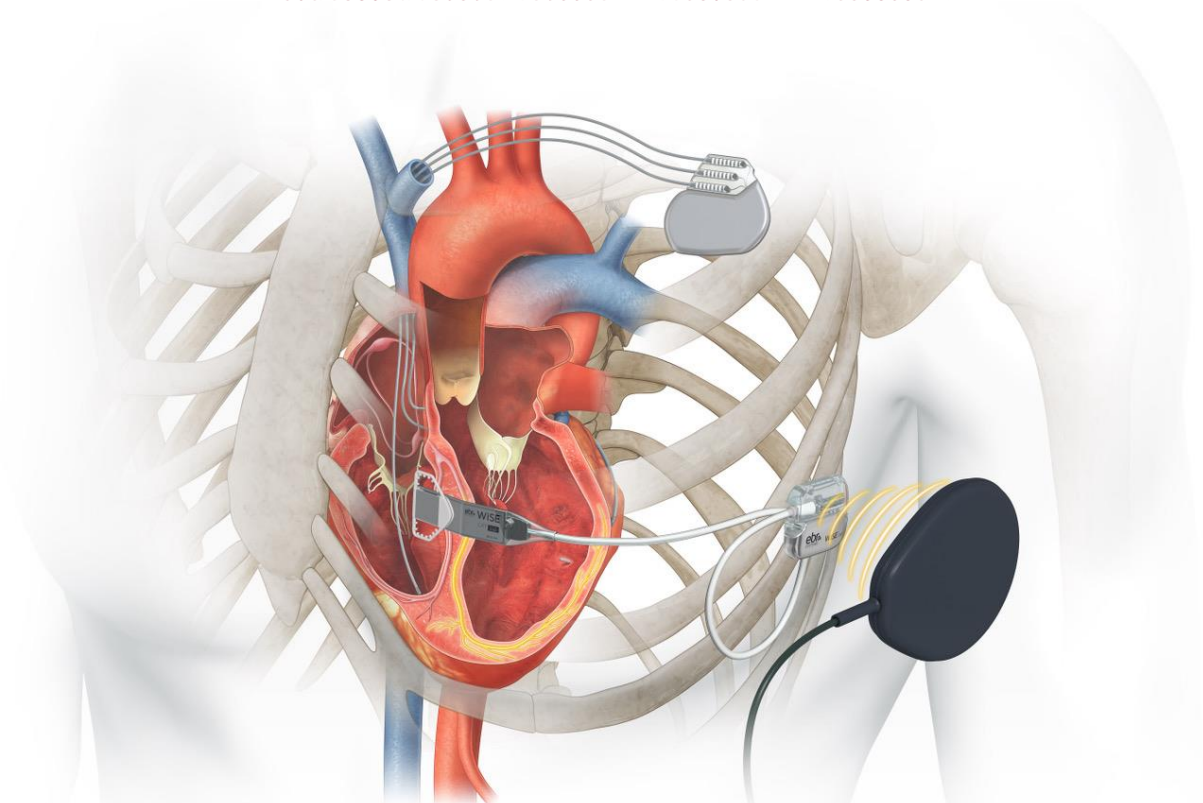
- Drives higher uptake by removing barriers to adoption
- Potential to become a first-line therapy option
- Diversifies applicability of the WiSE CRT System and grows the addressable market

Patient benefits

- Reduces need for future battery replacement surgery
- Recharge interval once per week¹
- 66% reduction in size from current battery

Development status

- Specifications and initial design completed
- Regulatory and commercial timing to be announced as project progresses



EBR's rechargeable battery will charge using a patch and external device to provide non-invasive, wireless charging

Long term growth strategy

Long term growth opportunity targeting new patient groups, indications and geographies



Product development

Grow addressable market through product development initiatives including a rechargeable battery



Pursue new indications

Progress clinical studies to expand indications and diversify product applications, opportunity to build a new market as first-line-therapy



Expand internationally

Launch in select OUS¹ markets as regulatory and reimbursement coverage is secured using US market entry as a template for success

Upcoming milestones

EBR continues to achieve significant value catalysts and pave the way to future value creation

Delivered

- ✓ SOLVE-CRT 6 month follow up completed for final patient in February 2023
- ✓ Headline data released at Heart Rhythm Society conference
- ✓ Positive trial data unlocks second tranche of growth capital facility
- ✓ Clinical module for PMA application submitted to the FDA
- ✓ Completed randomised sub study and released positive dataset
- ✓ Randomised data presented at industry conferences including Asia-Pacific Heart Rhythm Society

Upcoming

- ❑ Publication of manuscript in a peer reviewed medical journal
- ❑ Submit final PMA module including transmitter upgrades
- ❑ Additional sub-studies published using SOLVE-CRT dataset
- ❑ FDA approval in the US
- ❑ Commercial launch in the US

Next steps

- ❑ Expand manufacturing capacity
- ❑ Initiate ACCESS and TLC studies
- ❑ Drive adoption in US
- ❑ Clinical study of rechargeable battery
- ❑ Enter international markets
- ❑ Expand use of WiSE CRT System into new patient groups

Summary

EBR remains driven to deliver superior treatment for patients suffering from cardiac rhythm diseases



Developer of the world's first and only leadless pacemaker for heart failure



EBR's WiSE[®] CRT System has no direct competitors and is complementary to other pacemaker technologies



Positive pivotal trial results de-risk the regulatory pathway and validate the device as safe and highly effective



Clear commercial strategy in place focusing on high-volume procedure sites in the US, minimising execution risk



Significant market opportunity with an initial addressable market of US\$3.6bn and potential for further expansion



Funded through initial commercialisation with US\$64.1/A\$98.4m cash in bank as of 31 Mar 2024



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