

# CVRx<sup>®</sup>

**Jefferies Global Healthcare Conference in London**

**November 2025**

**NASDAQ: CVRX**



**Barostim<sup>™</sup>**  
Outsmart the heart

**CVRx**  
Outsmart the heart

# Disclaimer

## Cautionary Note Regarding Forward-Looking Statements

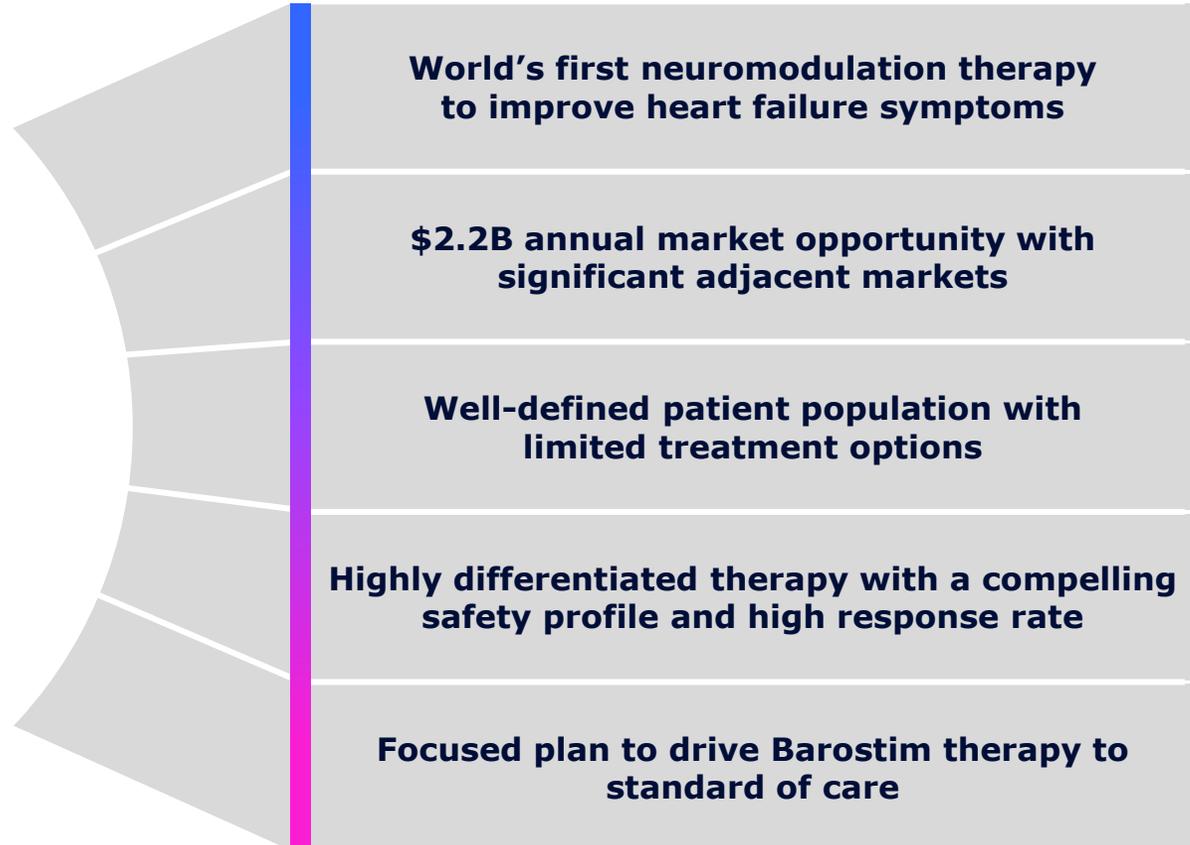
This presentation by CVRx, Inc. (the "Company") contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements other than statements of historical facts are forward-looking statements, including statements regarding our future financial performance (including, specifically, our 2025 expected operating and financial results), our anticipated growth strategies, anticipated trends in our industry, our business prospects and our opportunities. In some cases, you can identify forward-looking statements by terms such as "may," "will," "should," "expect," "plan," "anticipate," "could," "outlook," "guidance," "intend," "target," "project," "contemplate," "believe," "estimate," "predict," "potential" or "continue" or the negative of these terms or other similar expressions, although not all forward-looking statements contain these words.

The forward-looking statements in this presentation are only predictions and are based largely on our current expectations and projections about future events and financial trends that we believe may affect our business, financial condition, and results of operations. These forward-looking statements speak only as of the date of this presentation and are subject to a number of known and unknown risks, uncertainties and assumptions, including, but not limited to, our history of significant losses, which we expect to continue; our limited history operating as a commercial company and our dependence on a single product, Barostim; our limited commercial sales experience marketing and selling Barostim; our ability to continue demonstrating to physicians and patients the merits of our Barostim; any failure by third-party payors to provide adequate coverage and reimbursement for the use of Barostim; our competitors' success in developing and marketing products that are safer, more effective, less costly, easier to use or otherwise more attractive than Barostim; any failure to receive access to hospitals; our dependence upon third-party manufacturers and suppliers, and in some cases a limited number of suppliers; a pandemic, epidemic or outbreak of an infectious disease in the U.S. or worldwide; product liability claims; future lawsuits to protect or enforce our intellectual property, which could be expensive, time consuming and ultimately unsuccessful; any failure to retain our key executives or recruit and hire new employees; impacts on adoption and regulatory approvals resulting from additional long-term clinical data about our product; and other important factors that could cause actual results, performance or achievements to differ materially from those that are found in "Part I, Item 1A. Risk Factors" in our Annual Report on Form 10-K for the year ended December 31, 2024 and in "Part 2, Item 1A. Risk Factors" in our Quarterly Report on Form 10-Q for the quarter ended September 30, 2025, as such factors may be updated from time to time in our other filings with the Securities and Exchange Commission. Except as required by applicable law, we do not plan to publicly update or revise any forward-looking statements contained herein, whether as a result of any new information, future events, changed circumstances or otherwise.

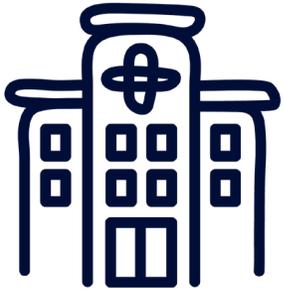
## Market & Industry Data

This presentation includes market and industry data and forecasts that the Company has developed from independent research reports, publicly available information, various industry publications, other published industry sources or the Company's internal data and estimates. Independent research reports, industry publications and other published industry sources generally indicate that the information contained therein was obtained from sources believed to be reliable, but do not guarantee the accuracy and completeness of such information. Although the Company believes that the publications and reports are reliable, the Company has not independently verified the data and makes no representation or warranty with respect to the accuracy of such information.

# Company overview



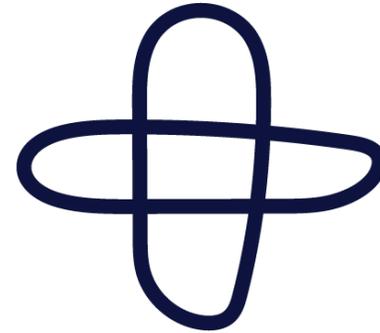
# Heart failure (HF) is a burdensome, life-limiting disease affecting over 6M people living in the U.S.<sup>1</sup>



**>1.1M hospital discharges<sup>1</sup>**



**>1.3M emergency room visits<sup>1</sup>**



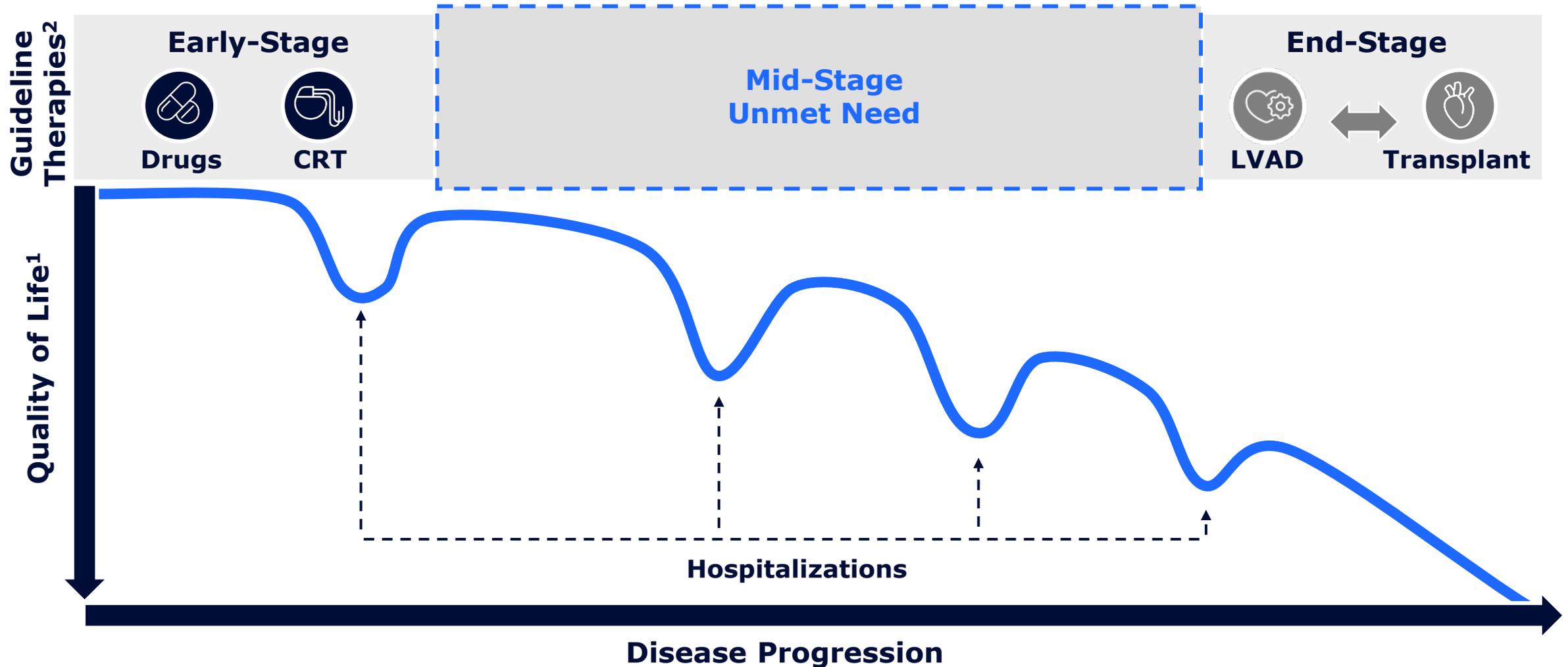
**>8M physician office visits<sup>1</sup>**



**Annual costs expected to reach \$70B by 2030<sup>2</sup>**

*All figures are annual estimates for the U.S.*

# Heart failure is a progressive disease characterized by a steady decline in quality of life (QoL) and increasingly frequent hospitalizations



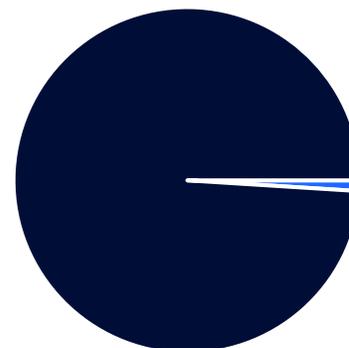
# Heart failure drug “quad therapy” has been shown to improve survival 1-6 years when taken compliantly and at optimal doses...



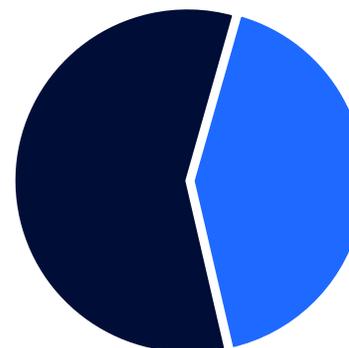
## 2022 AHA/ACC/HFSA HF Guidelines<sup>1-3</sup>

ARNI	B-Blockers
MRA	SGLT2i

**1.4-6.3** years  
Estimated aggregate mortality benefit of comprehensive quadruple therapy in HFrEF<sup>1</sup>



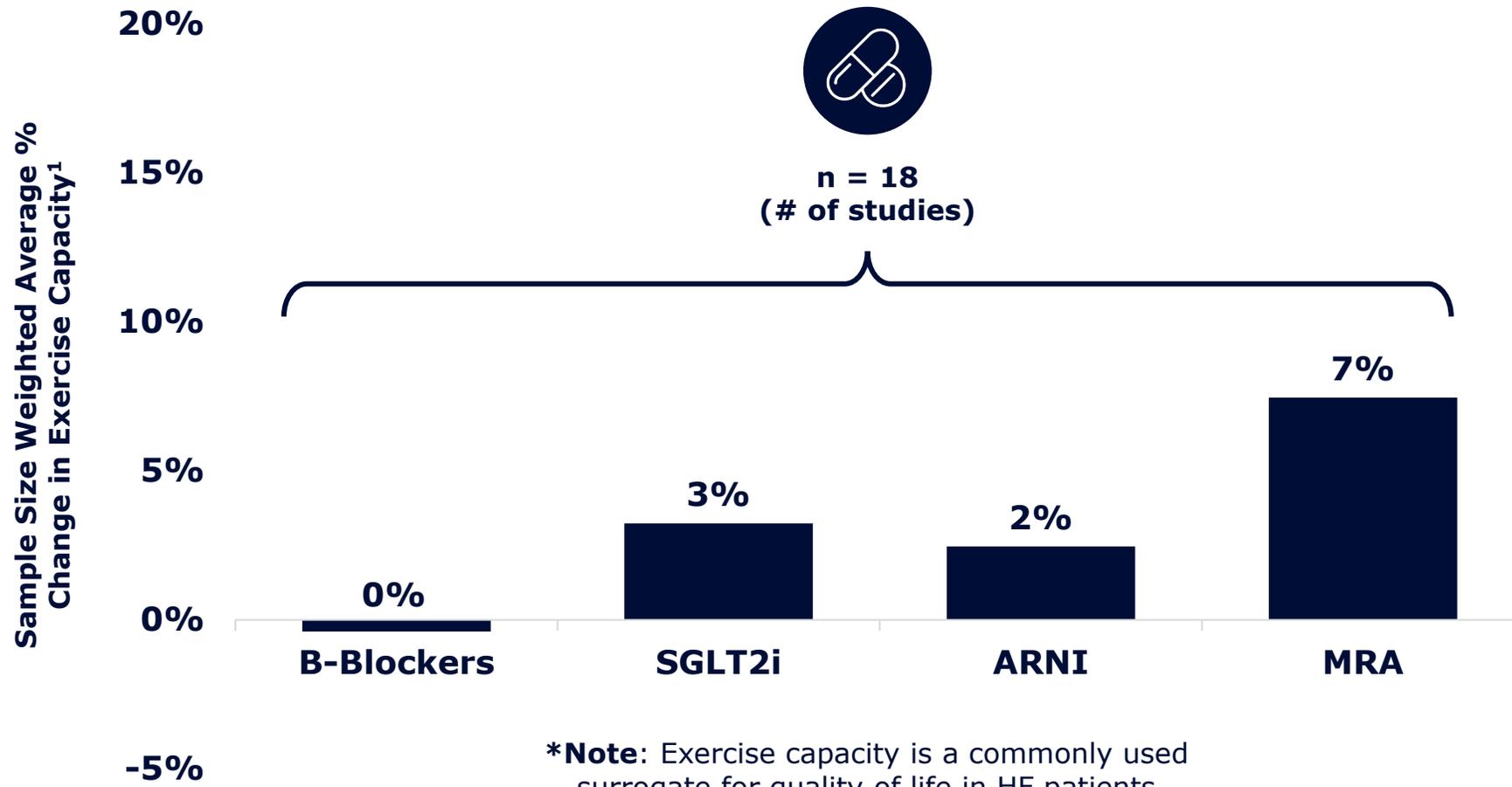
Only 1% reach optimal dose for quad therapy<sup>4</sup>



>40% discontinue quad therapy within the first year<sup>5</sup>

1. Heidenreich PA, et al. 2022 AHA/ACC/HFSA Guideline for the Management of Heart Failure. Circulation 2022. 2. Vaduganathan M, et al. Estimating lifetime benefits of comprehensive disease-modifying pharmacological therapies in patients with heart failure with reduced ejection fraction: a comparative analysis of three randomised controlled trials. Lancet Vol 396, Issue 10244, P121-128, July 11, 2020. 3. Rahamim E, et al. Contemporary Pillars of Heart Failure with Reduced Ejection Fraction Medical Therapy. J. Clin. Med. 2021, 10, 4409. 4. Greene S et al, Medical Therapy for Heart Failure With Reduced Ejection Fraction: The CHAMP-HF Registry, J Am Coll Cardiol. 2018; 72:351-366. 5. Savarese G et al, Heart Failure Drug Treatment—Inertia, Titration, and Discontinuation: A Multinational Observational Study (EVOLUTION HF), J Am Coll Cardiol HF. 2023; 11:1-14.

...but has been shown to have minimal impact on quality of life as measured by exercise capacity\*



# These limited treatment options leave the majority of heart failure patients suffering from significantly diminished QoL

Heart failure significantly decreases quality of life



**66%**  
have mobility problems<sup>1</sup>



**68%**  
report pain or discomfort<sup>1</sup>

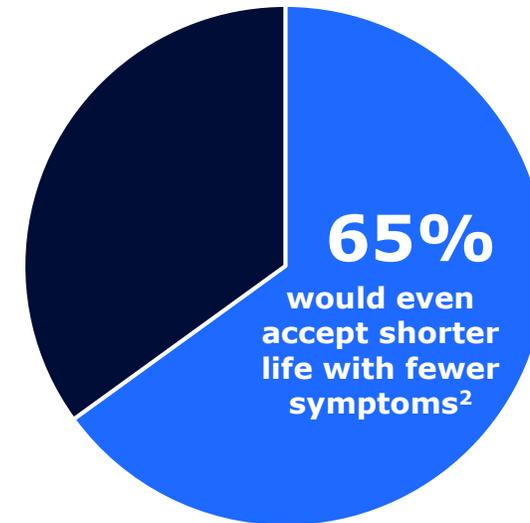


**76%**  
find activities of daily living to be difficult<sup>1</sup>

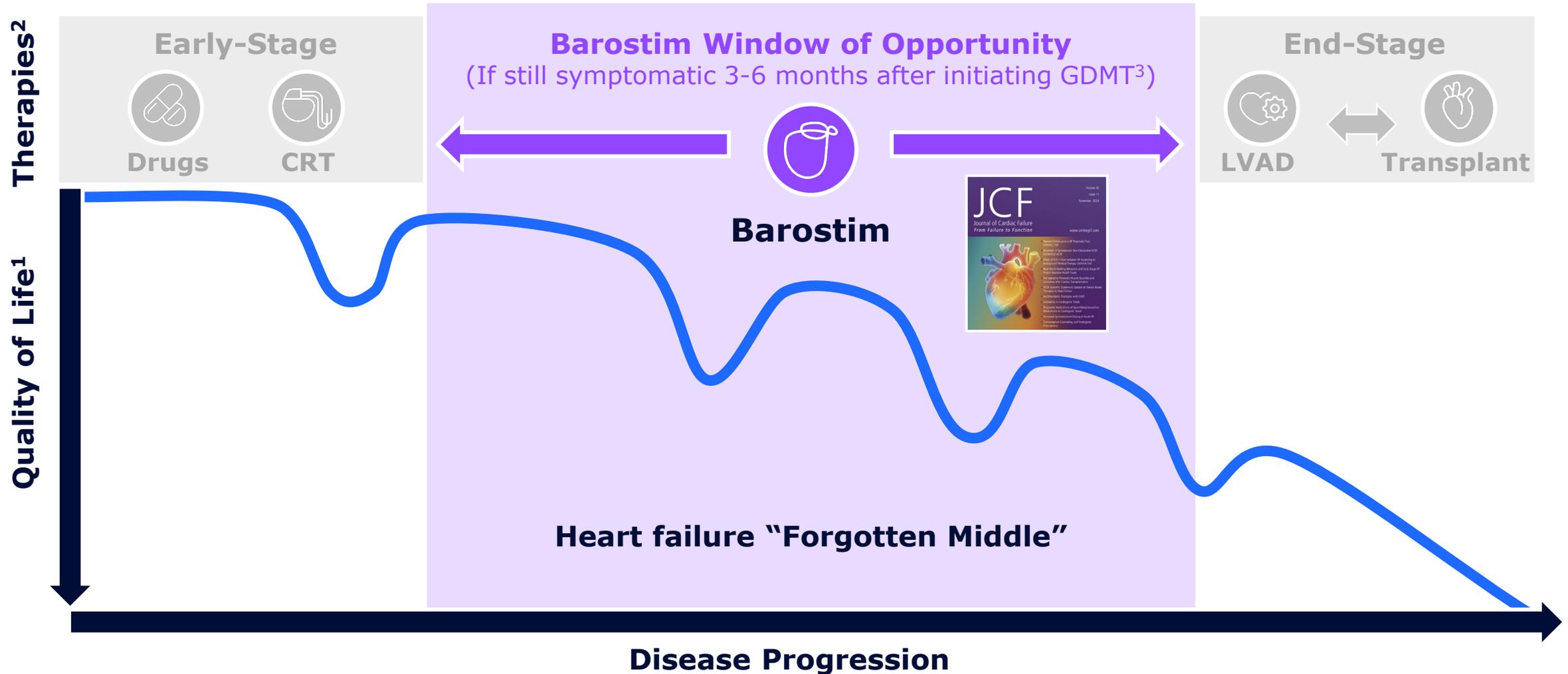


**50%**  
have anxiety or depression<sup>1</sup>

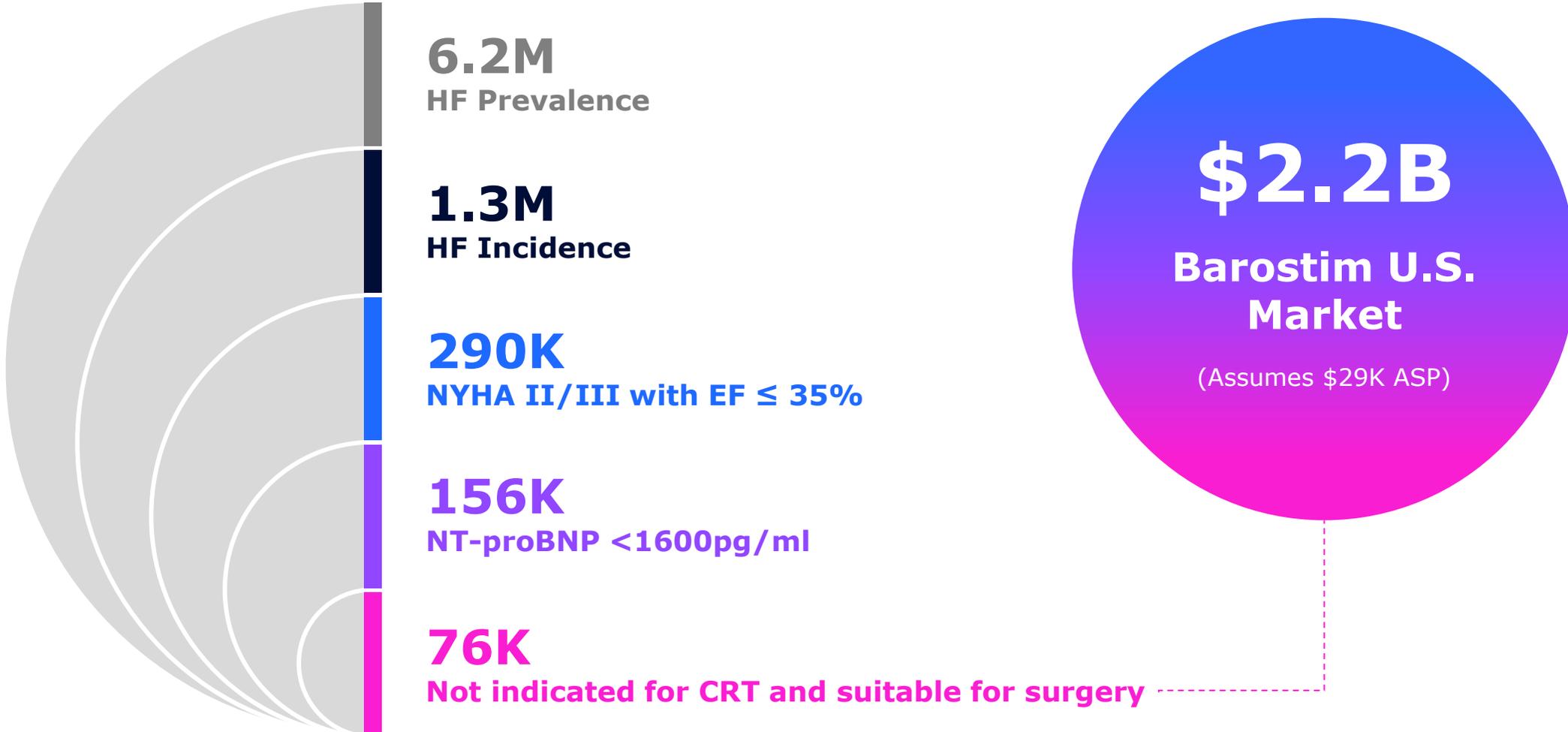
Majority of patients value symptom improvement over longevity



# Barostim addresses this significant unmet need in the heart failure treatment continuum



# We are only 2% penetrated into a \$2.2B U.S. annual net addressable market for Barostim



# Barostim Therapy



# Barostim targets the neurohormonal pathways responsible for heart failure progression



1

Weakened heart & heart failure symptoms

↓ Contractility

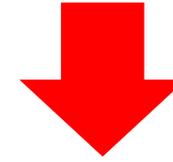
2

Reduced stretch sensed by baroreceptors

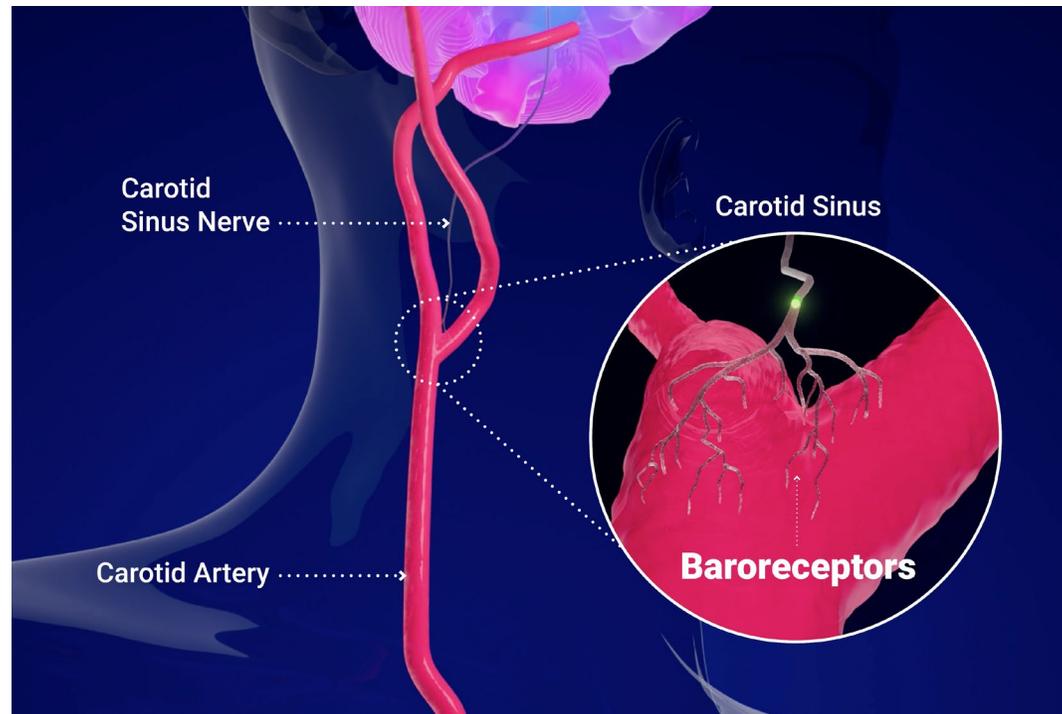
↓ Baroreceptor Signaling

2

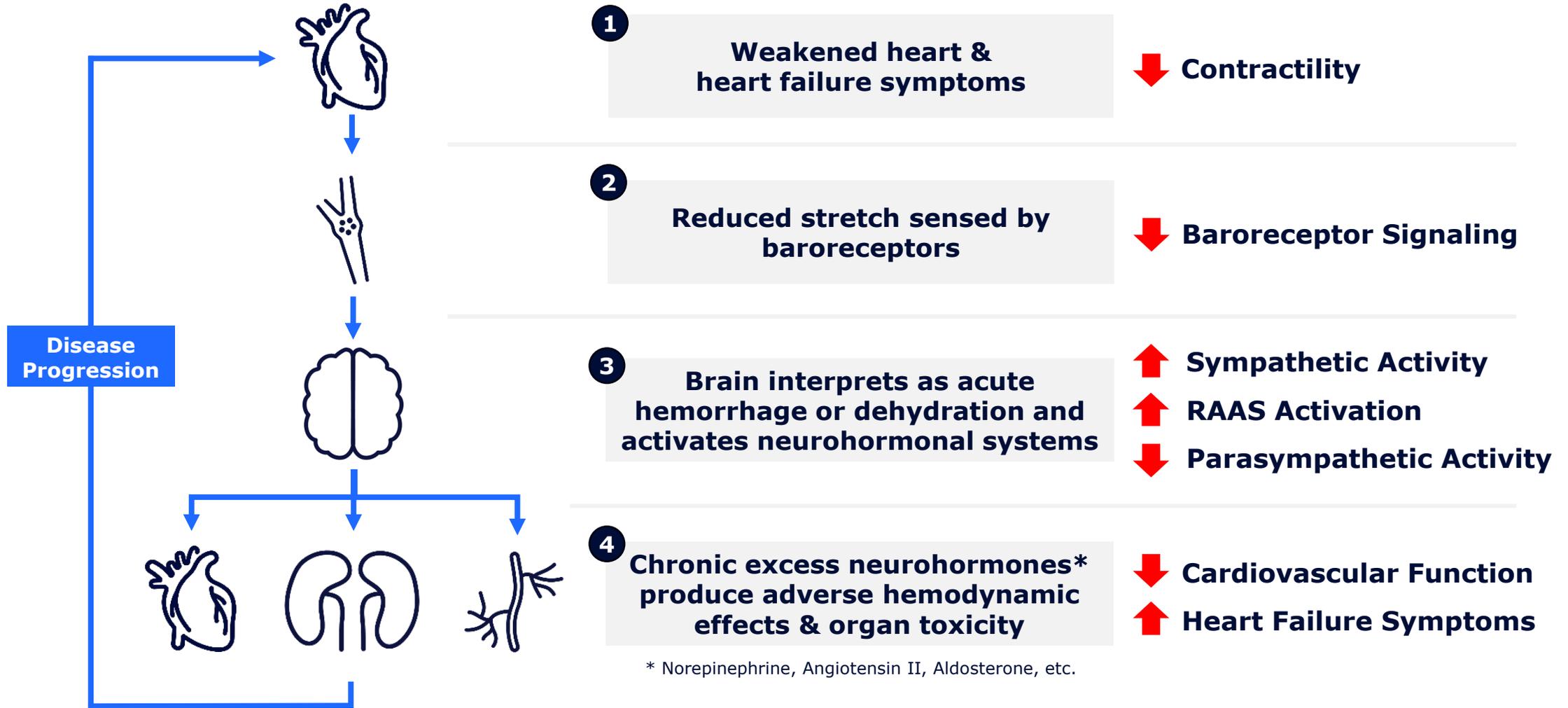
## Reduced stretch sensed by baroreceptors



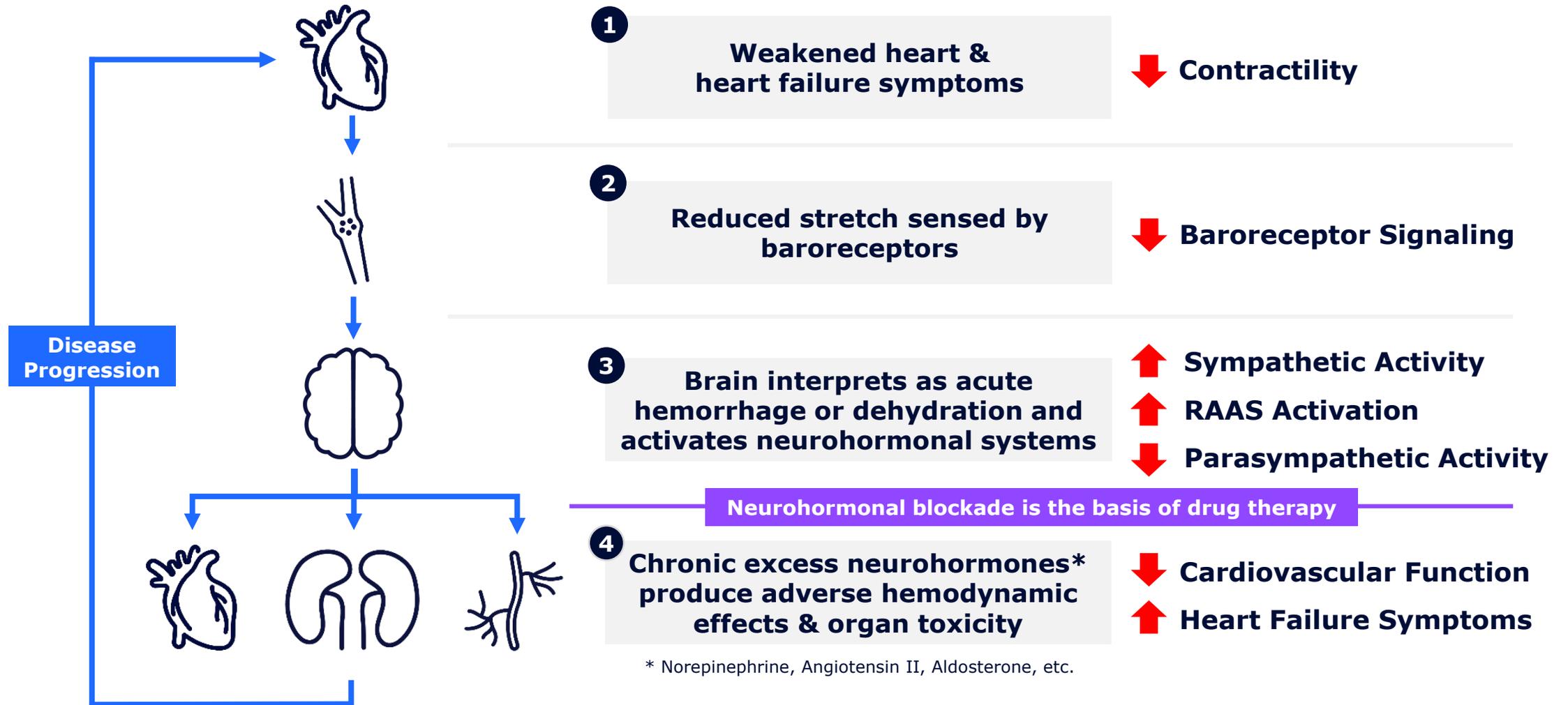
- Stretch receptors that continuously monitor **blood volume and pressure**
- Information is electrically signaled to the brain



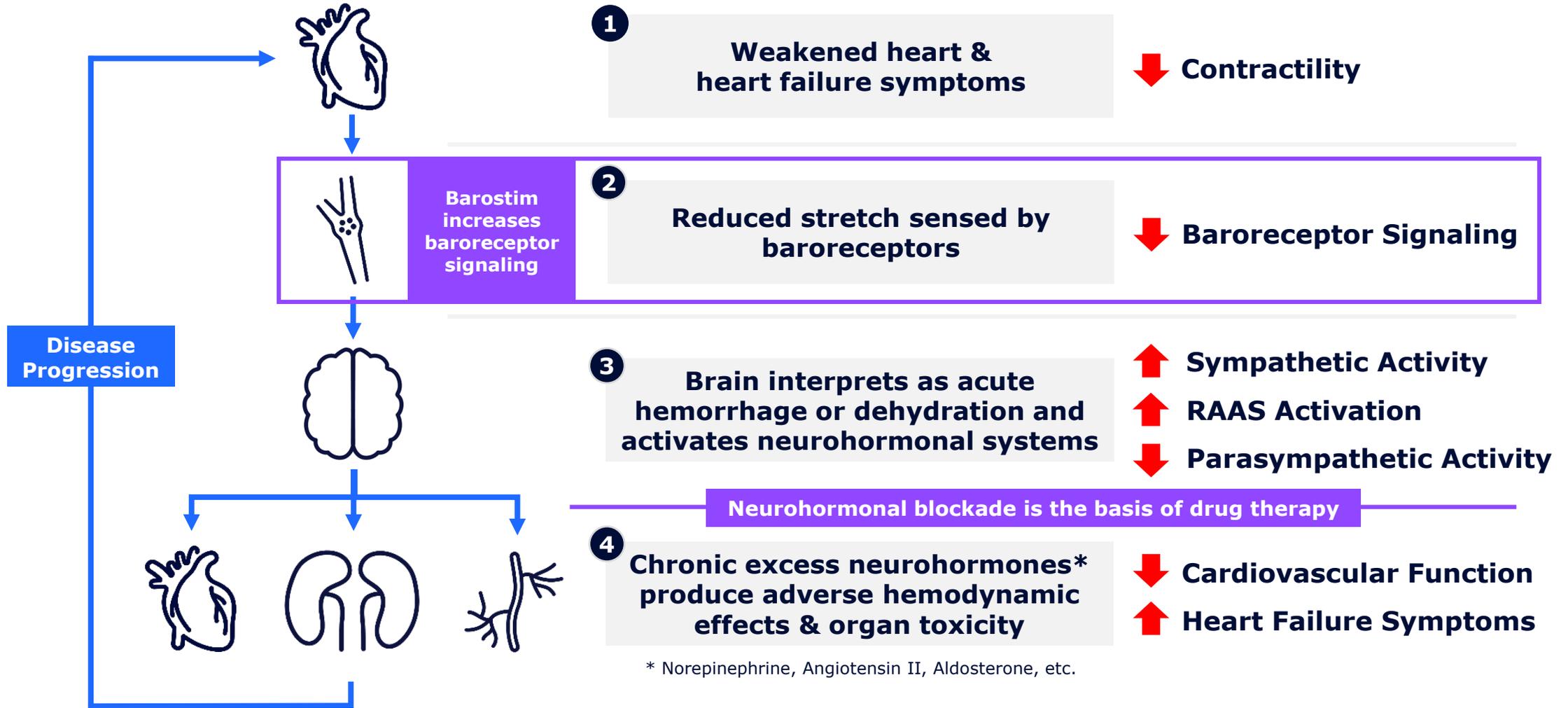
# Barostim targets the neurohormonal pathways responsible for heart failure progression



# Drug therapies work by blocking specific excess neurohormones



# Barostim complements drug therapy by acting upstream to restore baroreceptor signaling



# Barostim system elements

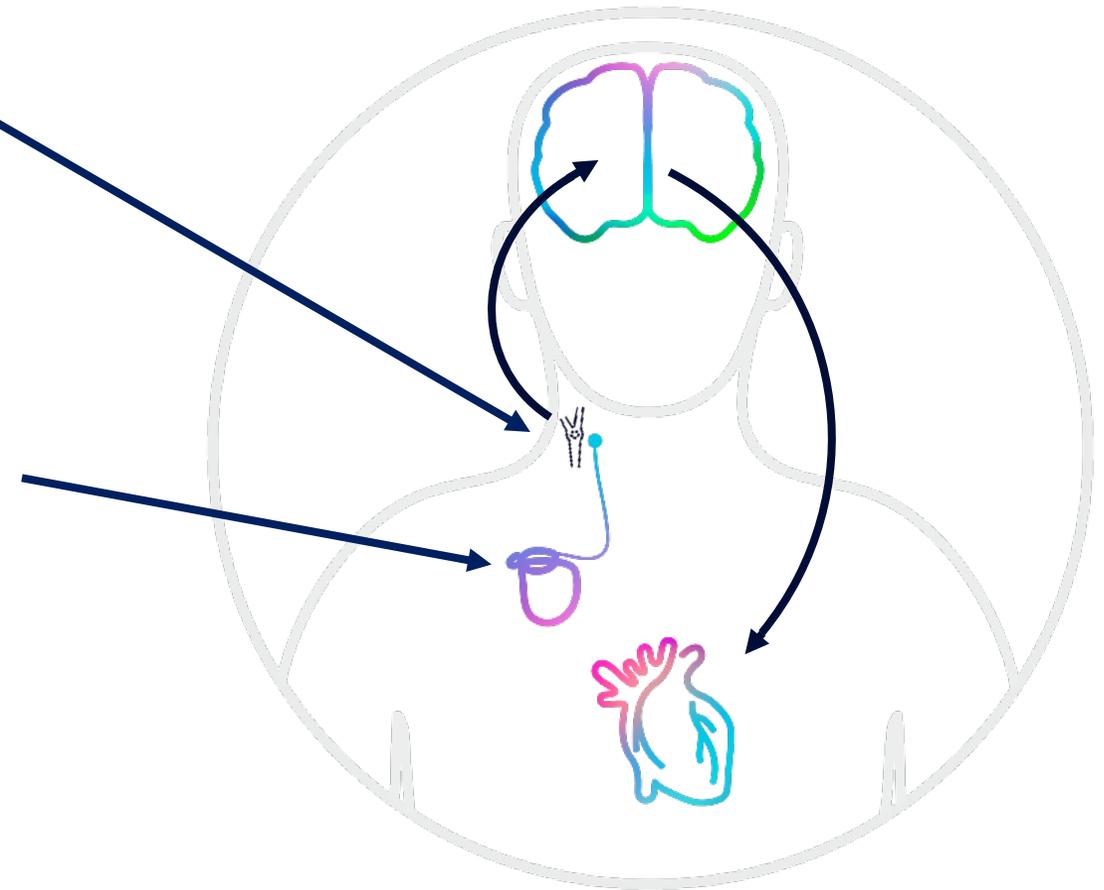
**Barostim was designed to deliver electrical stimulation to carotid baroreceptors to increase baroreceptor signaling**



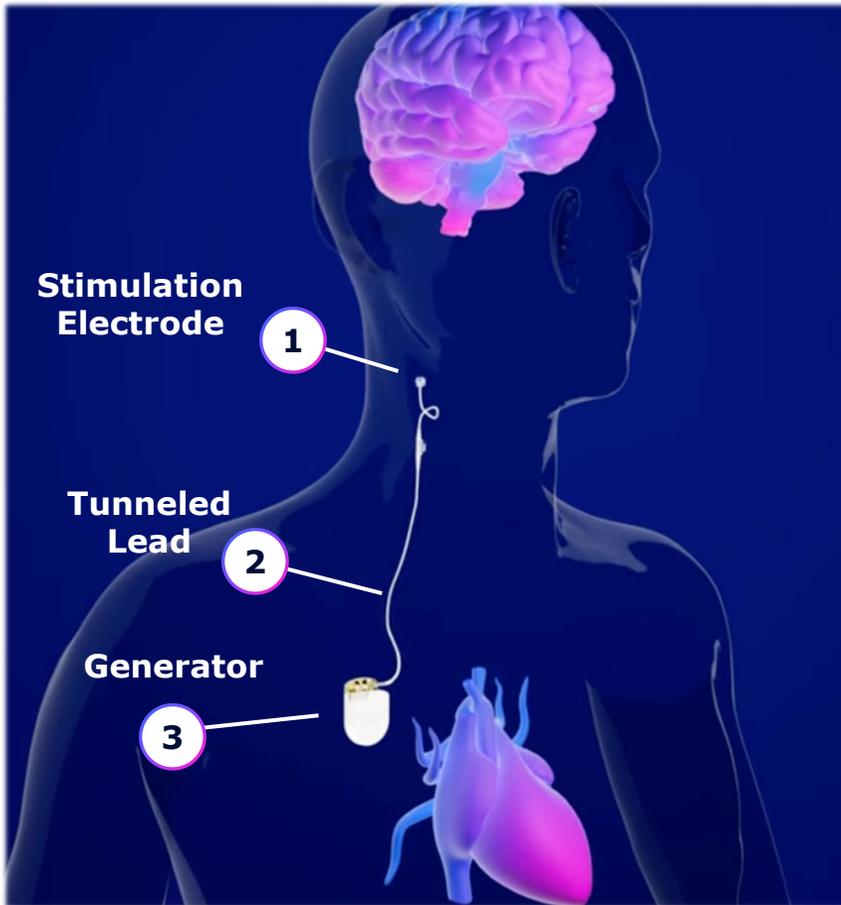
**Implantable Pulse Generator (IPG) & Carotid Sinus Lead**



**Wireless Programmer**



# Barostim is implanted in a ~60-min procedure, with 97% freedom from major complications



- **Procedure is proven safe; achieved a 97% MANCE-free rate**
- **Implanted on either an inpatient or outpatient basis**
- **Requires a small incision in both the neck and chest**
- **Entirely extravascular, with no leads in the heart or vasculature**

MANCE - Major Adverse Neurological and Cardiovascular Events includes all events that occur within 6 months of implant

# Barostim is an effective, predictable, and durable therapy to improve patient quality of life



**Exercise  
Capacity**



**Quality  
of Life**



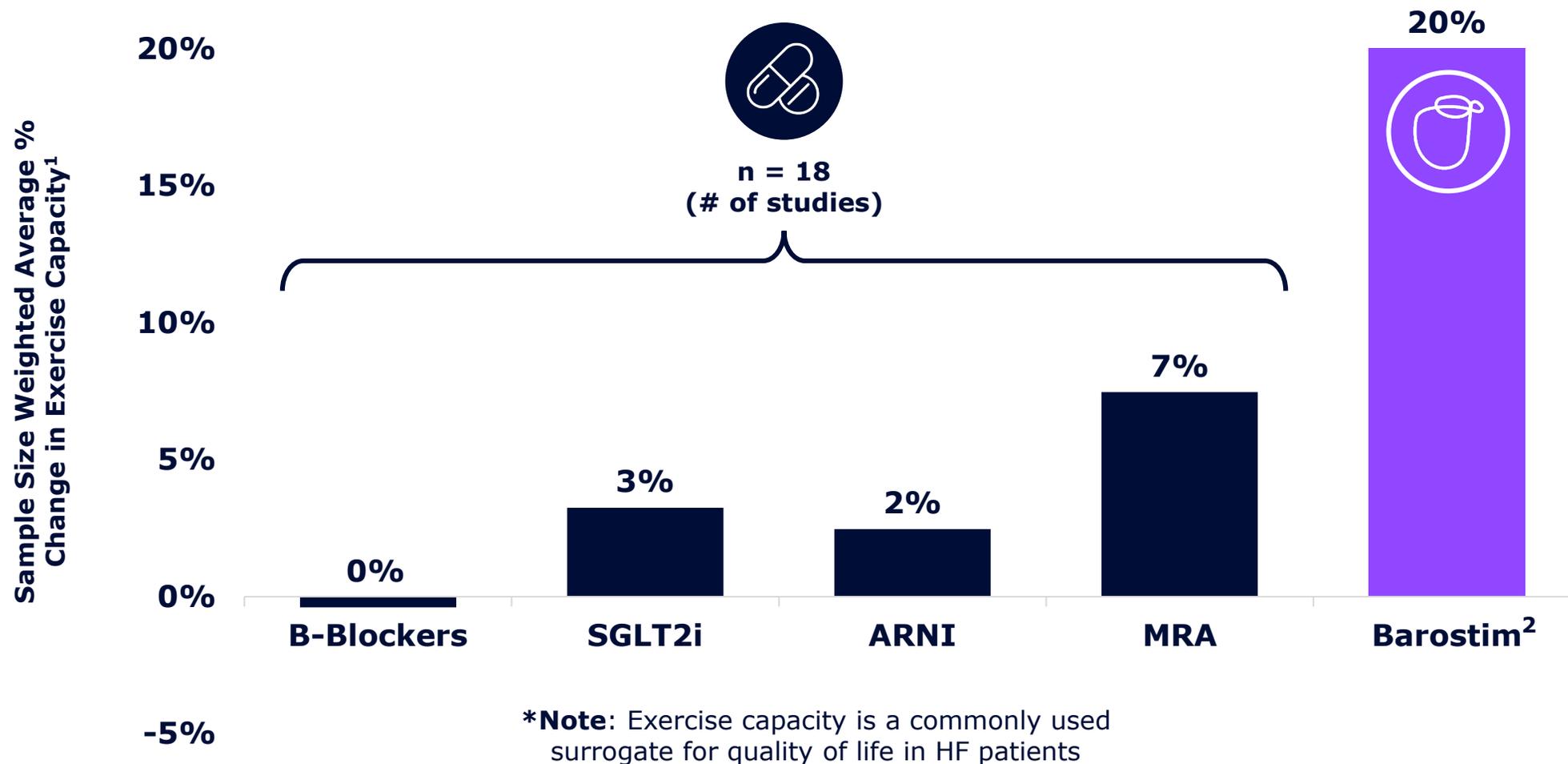
**Functional  
Status**



**High  
Response Rate**



# Improvements in exercise tolerance shown in BeAT-HF significantly exceed that of quad-therapy



# The BeAT-HF trial also showed a positive signal in all-cause death, LVAD and transplant despite confounding factors related to COVID-19

**-34%**

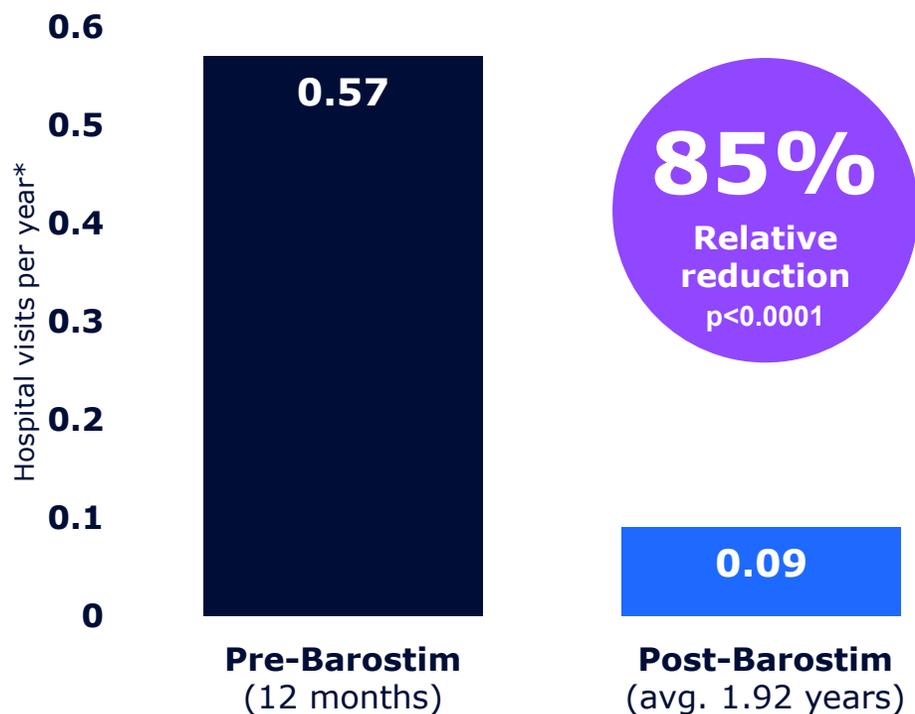
Relative Reduction in  
All-cause death, LVAD  
or transplant out to  
>4 years<sup>1</sup>

p=0.054

**Note: Not a powered endpoint**

# Recently published real-world-evidence demonstrates a significant reduction in hospitalization visits post- vs pre-implant

**Premier Healthcare Database<sup>1</sup>**  
(Post-COVID-19\*\*)  
N = 306



# Our revised go-to-market strategy is focused on driving Barostim to become Standard of Care for HFrEF

## Three Key Strategies

1

**Build a world-class sales organization**

2

**Focus on developing sustainable Barostim Programs**

3

**Address the barriers to adoption**

1

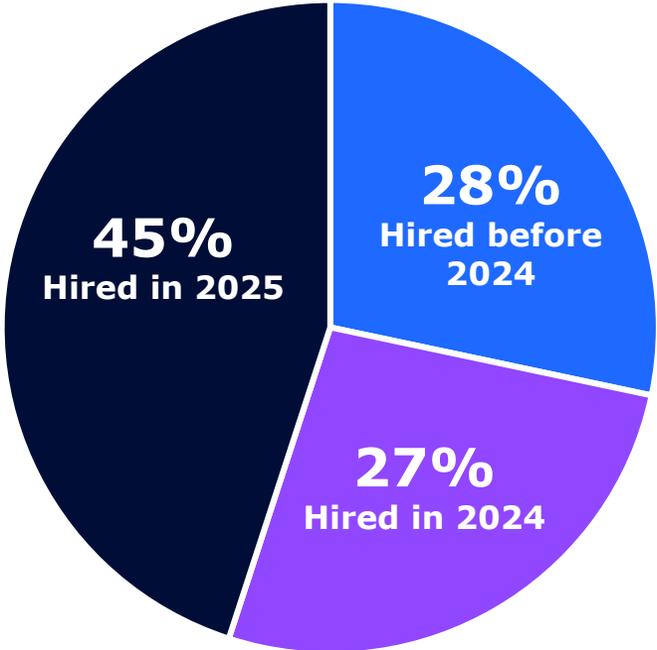
# Our efforts to transform our sales organization are largely complete and we are seeing broadening contribution from the team

**Strengthening sales team by recruiting territory managers with strong therapy development backgrounds**

**Optimizing training & development programs to accelerate rep productivity**

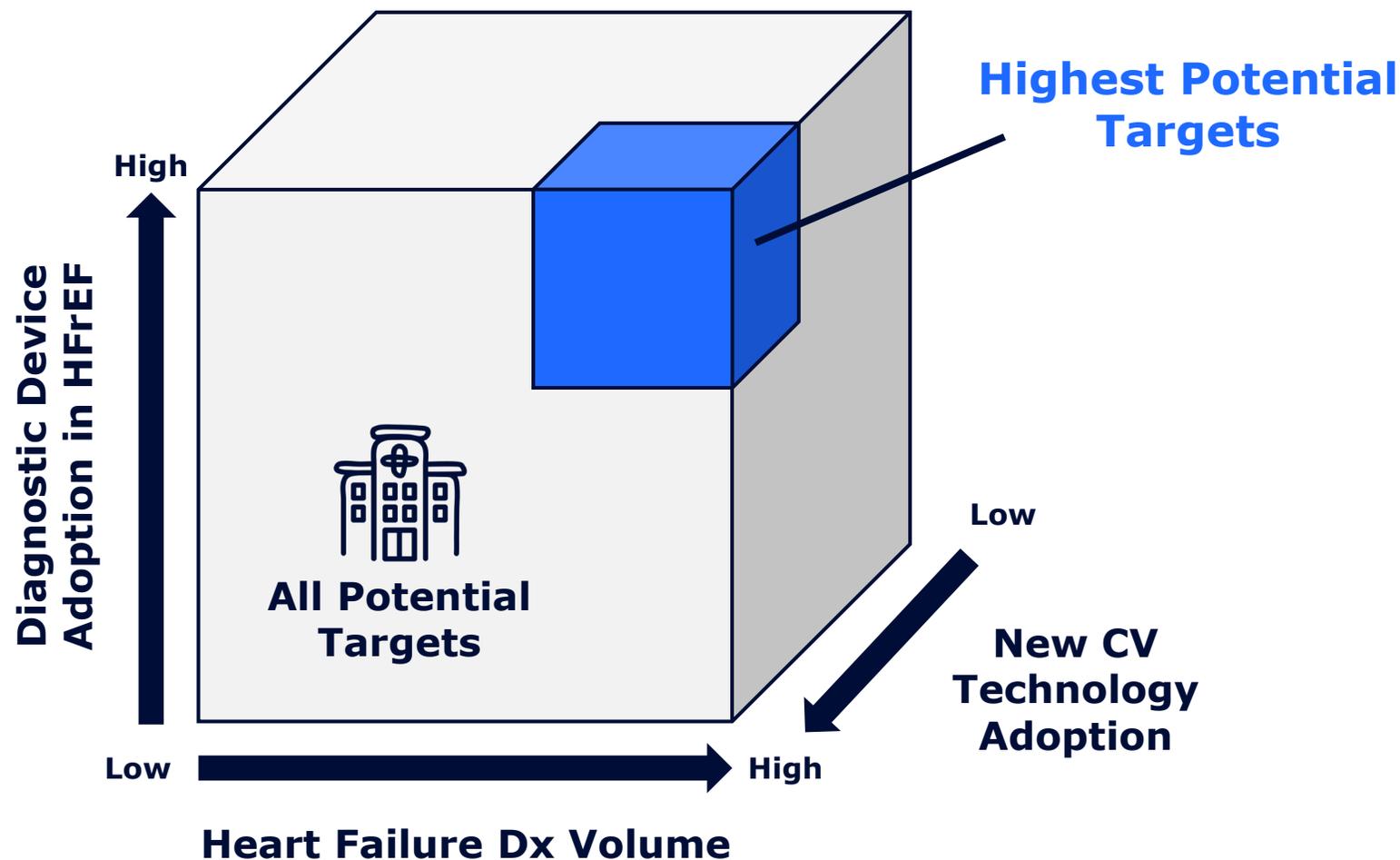
**Aligning incentives to a program-oriented sales process**

**Sales Reps by Hire Date**  
(as of October 31, 2025)



2

We are focused on developing sustainable Barostim programs by targeting centers with the highest potential for success

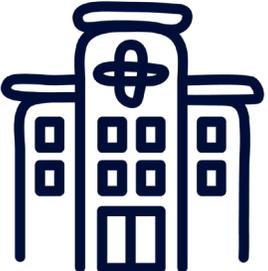


2

# ...and are seeking to replicate the elements present in Barostim centers that have achieved the deepest adoption

## Sustainable Barostim Program

  
Clinical  
Champion(s)



  
Administrative  
Champion(s)

**Referrers**  
(Identify potential patients)

**Prescribers**  
(Prescribe the therapy and send for implant)

**Implanters**  
(Technicians)

     
**HFS APP EP GC/IC**

   
**Heart Failure Specialist (HFS)\***

   
**VS / CTS**

\*May also be an EP, GC, or IC depending on account

3

**We have implemented a market development strategy focused on addressing the key barriers to adoption**

**Barrier #1: Increase therapy awareness among referrers and patients**

**Barrier #2: Develop more robust clinical evidence**

**Barrier #3: Improve patient access to Barostim**

# Barrier #1: We are focusing therapy awareness efforts on the clinicians and patients that surround targeted centers

## Referral Physician Outreach



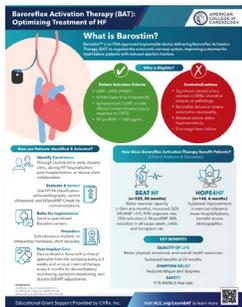
HFS



EP



GC/IC



**ACC Infographic**  
(Distributed to 50K+ ACC members)



**Medical Education Webinars**



**6 National & 100+ Local Medical Education Programs in 2025**

## Sustainable Barostim Program



**Significantly Increased Focus on APPs**

NP



PA



**Society Partnerships**



## Direct to Consumer Marketing



**Pre-Screening**



# Barrier #2: We are developing a steady cadence of clinical evidence to support Barostim therapy in two key areas

## Areas of Interest:

### Evidence of Improved Outcomes

- Improved QoL & exercise capacity
- Reduced HF and all-cause hospitalizations
- Reduced mortality
- Reduced arrhythmias
- Improved ejection fraction
- Decreased diuretic needs

### Evidence Supporting Mechanism of Action

- Fundamentally improved hemodynamic function
- Reduced sympathetic nerve activity
- Restored cardiac parasympathetic control
- Anti-inflammatory effect
- Reversed remodeling

## Evidence Generation Channels:

Real-World Evidence

Randomized Controlled Trial(s)

Investigator Initiated Research



# **Barrier #3: We have made significant progress in improving patient access to Barostim with recent positive OPPS and Physician Fee Schedule updates**

## **Milestones**



**Permanent MS-DRG assignment secured for Inpatient > \$43K (vs. \$17-\$23K)**



**Proposed to maintain assignment to APC 1580 (\$45K) in July 2025 OPPS proposed rule; CMS solicited comments regarding creation of Level 6 Neurostimulator APC**



**January 2026: CMS Category I codes effective with average physician payment of \$560+ for implant**

## **Implications**



**Aligns inpatient and outpatient reimbursement, facilitating site-of-service flexibility based on patient medical necessity**



**Current outpatient payment of ~\$45K likely maintained; potential for creation of Level 6 Neurostimulator APC (~\$43K)**



**Eliminates automatic Category III code-related denials from Medicare Advantage and commercial payers, increasing fair and consistent physician payment**

# 2025 financial status and guidance

## Q3 2025 Results

- WW Revenue: \$14.7M
- US Territories: 50
- US Active Implanting Centers: 250
- Gross Margin: 87%
- Cash Balance: \$85.1M

## Guidance as of Nov 5, 2025

- Q4 2025 Revenue: \$15.0 – \$16.0M
- Full Year 2025:
  - Revenue: \$55.6 – \$56.6M
  - Gross Margin: 85% – 86%
  - Operating Expense: \$98.0 – \$99.0M

# Barostim opportunity



**To positively impact the standard of care for a major global health condition**

**To address a significant unmet need in the heart failure treatment continuum**

**To improve the quality of life of hundreds of thousands of heart failure patients**

**To build a transformational company capable of sustaining long-term growth**