Abbott

CONFIDENCE IN EXCELLENCE

HEARTMATE 3™ LVAD

with Full MagLev™

Flow Technology

HEARTMATE 3™ LVAD

DELIVERS EXCELLENT SURVIVAL AND SAFETY OUTCOMES[1]*

MOMENTUM 3 is the largest left ventricular assist device (LVAD) trial ever conducted, demonstrating excellent survival and safety outcomes with HeartMate 3™ LVAD.[1]*

5-YEAR OUTCOMES FROM THE MOMENTUM 3 TRIAL

FIRST TIME IN LVAD HISTORY

5-year data from a prospective randomized LVAD trial

HEARTMATE 3 LVAD

is a proven **long-term, life-extending therapy** for patients with advanced heart failure

MEDIAN SURVIVAL
EXCEEDING 5 YEARS[1]

58.4%
SURVIVAL AT 5 YEARS[1]

THESE OUTCOMES REFLECT ABBOTT'S LEGACY OF DEVELOPING **LIFE-CHANGING TECHNOLOGY.**

See Important Safety Information referenced within.

Life-prolonging therapy in a patient population who otherwise would **not be expected to survive beyond 9 months**[1,2]*

Superiority of the HeartMate 3™ LVAD driven by a **reduction** in HRAEs[1]

Survival comparable to higher-risk transplant patients[1,3]

HeartMate 3 LVAD has received the highest level of recommendation (1A) from the AHA, ACC and HFSA joint societies for select Class IV heart failure patients

1 in 26

FOR EVERY 1 PATIENT RECEIVING A HEART TRANSPLANT, 25 WILL DIE**

These data provide compelling evidence that **LVAD therapy can add years to the lives** of patients who do not qualify for or are waiting for a heart transplant.

IMPROVED SAFETY PROFILE[4]

The **ELEVATE Registry** evaluated real-world experience of the HeartMate 3™ LVAD in a post-approval setting. MOMENTUM 3 trial 5-year extended follow-up showed:

STROKE[4]

10.8% IN FIRST 2 YEARS

3.6% IN YEARS 2–5

In the ELEVATE Registry, all major event types were reduced in the 2–5-year follow-up period compared to 0–2 years, especially HRAEs.[1]

UNMATCHED LEGACY IN MCS

Evolution of HeartMate™ LVAD

1998

HEARTMATE™ XVE LVAS***

Pulsatile Flow

FIRST LVAS APPROVED

FOR DT OR LONG-TERM USE

2008

HEARTMATE II™ LVAD

Continuous Flow (Axial)

OVER 26,000 PATIENTS IMPLANTED[5]

2017

HEARTMATE 3™ LVAD

Continuous Flow (Centrifugal)
With Full MagLev™ Flow Technology

OVER 36,000 PATIENTS IMPLANTED[5]

EXCELLENCE IS:

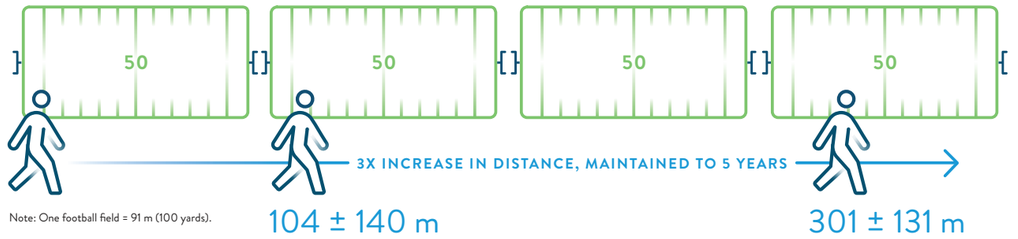
Immediate, significant and sustained improvements in functional capacity and quality of life

+30 POINTS

QUALITY-OF-LIFE SCORE IMPROVED SIGNIFICANTLY
AND WAS SUSTAINED THROUGH 5 YEARS[4]

EuroQol 5 Dimensions 5 Levels (EQ-5D-5L[†]) questionnaire

ABILITY TO WALK 3 TIMES FARTHER[4]



Note: One football field = 91 m (100 yards).

104 ± 140 m
(approx. 114 yards) at baseline

301 ± 131 m
(approx. 330 yards) at 5 years

OUTCOMES MADE POSSIBLE BY

FULL MAGLEV™ FLOW TECHNOLOGY

Full MagLev™ Flow Technology maintains gentle blood handling to minimize complications and reduce HRAEs.

FULLY LEVITATED, SELF-CENTERING ROTOR

that does not require hydrodynamic or mechanical bearings

LARGE, CONSISTENT BLOOD FLOW PATHWAYS

to reduce shear stress[6]

INTRINSIC PULSATILITY

(30 cycles per minute) to reduce blood stasis and minimize thrombus[6,7]

FULL MAGLEV FLOW TECHNOLOGY PUMP

Large consistent blood flow pathways to reduce shear stress[6]

167 **STACKED RED BLOOD CELLS[8]**

VS.

HYDRODYNAMIC BEARING PUMP

Narrow blood flow pathways

8 **STACKED RED BLOOD CELLS[8]**

See Important Safety Information referenced within.

HEARTMATE 3™ LVAD SYSTEM

A better experience for clinicians and patients†

1. HeartMate 3™ LVAD

Connected to the left side of the heart and moves oxygenated blood from the left ventricle to the rest of the body

2. Modular driveline

Facilitates simple replacement of externalized portion

3. Batteries

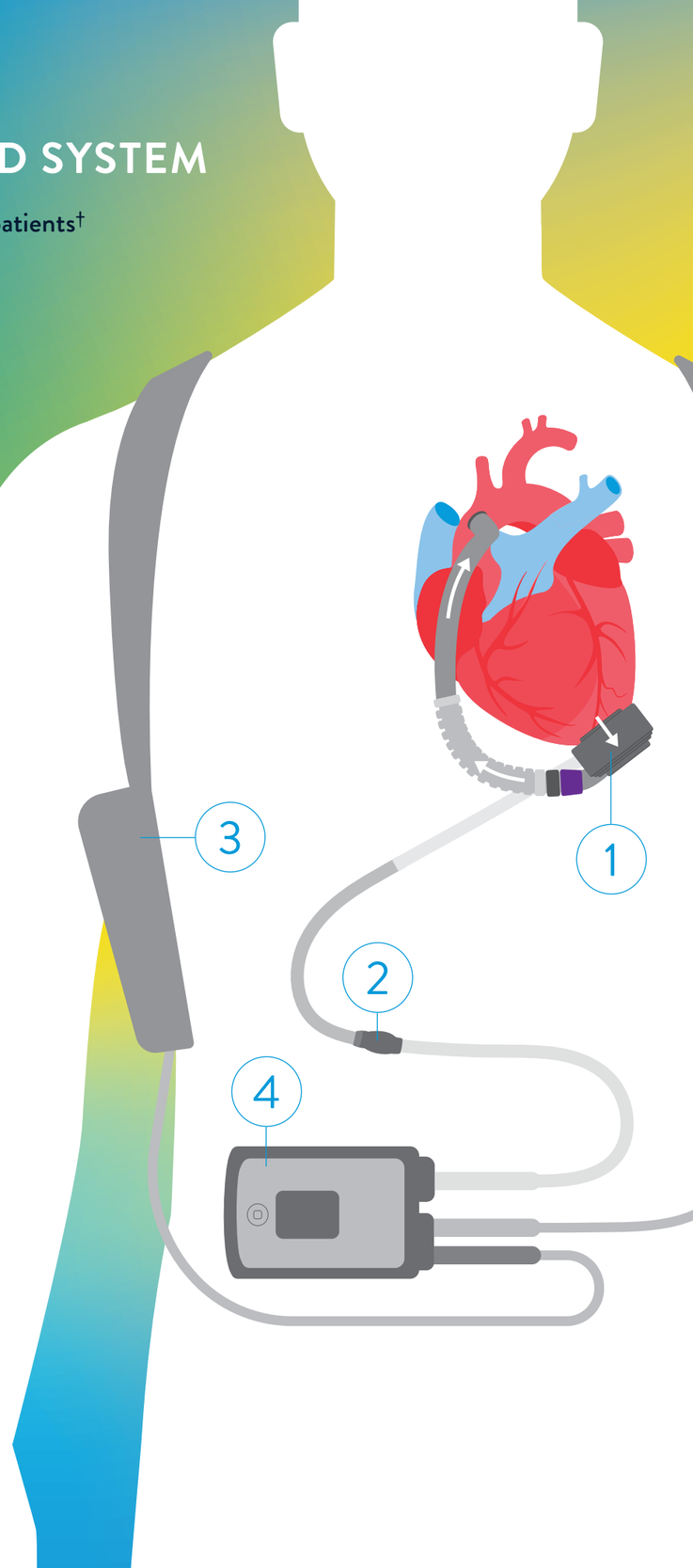
Provide up to 17 hours of uninterrupted power

4. Pocket controller

Powers and controls the LVAD and is small enough to fit in a pocket. Includes emergency backup battery

Mobile Power Unit (MPU)

Plug-in power source



EMBRACE EXCELLENCE
EVERY DAY

By choosing the HeartMate 3™ LVAD, you can help patients and their loved ones hold on to what matters most.

Scan the QR code to learn more about HeartMate 3 LVAD

ACC = American College of Cardiology‡; AHA = American Heart Association‡; DT = Destination Therapy; EuroQol = European Quality of Life questionnaire; HFSA = Heart Failure Society of America; HRAE= hemocompatibility-related clinical adverse event; LVAD = left ventricular assist device; LVAS = left ventricular assist system; MCS = mechanical circulatory support; MPU = mobile power unit

*HeartMate 3™ LVAD demonstrated superiority in event-free survival (primary endpoint) in the MOMENTUM 3 trial compared to HeartMate II™ LVAD.
**Patients on inotropes who did not receive a transplant or left ventricular assist device.
***This product is no longer available for sale or use.
†Compared to HeartMate II LVAD.

1. Mehra MR, Goldstein DJ, Cleveland JC, et al. Five-Year Outcomes in Patients With Fully Magnetically Levitated vs Axial-Flow Left Ventricular Assist Devices in the MOMENTUM 3 Randomized Trial. *JAMA*. 2022;328(12):1233-1242. doi:10.1001/jama.2022.16197
2. Hashim T, Sanam K, Revilla-Martinez M, et al. Clinical Characteristics and Outcomes of Intravenous Inotropic Therapy in Advanced Heart Failure. *Circ Heart Fail*. 2015;8(5):880-886.
3. Kilic A, Weiss ES, Yuh DD, Shah AS, Conte JV. Factors associated with 5-year survival in older heart transplant recipients. *J Thorac Cardiovasc Surg*. 2012;143(2):468-474.
4. Schmitto JD, Shaw S, Garbade J, et al. Long-Term Results in Real World Patients Treated with HeartMate 3 LVAD for Advanced Heart Failure: Data from the ELEVATE Registry. Presented at: European Association for Cardio-Thoracic Surgery (EACTS) Annual Meeting; October 8, 2022; Milan, Italy.
5. Abbott. Data on File. Based on clinical and device tracking data as of November 10, 2022.
6. Bourque K, Cotter C, Dague C, et al. Design rationale and preclinical evaluation of the HeartMate 3 Left Ventricular Assist System for hemocompatibility. *Am Soc Artificial Int Organs*. 2016;62(4):375-383.
7. Bourque K, Dague C, Farrar D, et al. *In vivo* assessment of a rotary left ventricular assist device induced artificial pulse in the proximal and distal aorta. *Artificial Organs*. 2006;30(8):638-642.
8. Based on clinical trial and device tracking data as of August 31, 2024.

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Rx Only

Brief Summary: Prior to using these devices, please review the Instructions for Use for a complete listing of indications, contraindications, warnings, precautions, potential adverse events and directions for use.

HeartMate 3™ LVAS Indications: The HeartMate 3™ Left Ventricular Assist System is indicated for providing short- and long-term mechanical circulatory support (e.g., as bridge to transplant or myocardial recovery, or destination therapy) in adult and pediatric patients with advanced refractory left ventricular heart failure and with an appropriate body surface area.

HeartMate 3™ LVAS Contraindications: The HeartMate 3 Left Ventricular Assist System is contraindicated for patients who cannot tolerate, or who are allergic to, anticoagulation therapy.

HeartMate 3™ LVAS Adverse Events: Adverse events that may be associated with the use of the HeartMate 3 Left Ventricular Assist System are: death, bleeding, cardiac arrhythmia, localized infection, right heart failure, respiratory failure, device malfunctions, driveline infection, renal dysfunction, sepsis, stroke, other neurological event (not stroke-related), hepatic dysfunction, psychiatric episode, venous thromboembolism, hypertension, arterial non-central nervous system (CNS) thromboembolism, pericardial fluid collection, pump pocket or pseudo pocket infection, myocardial infarction, wound dehiscence, hemolysis (not associated with suspected device thrombosis) or pump thrombosis.

™ Indicates a trademark of the Abbott group of companies.
‡ Indicates a third-party trademark, which is property of its respective owner.
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