

Assurity MRI™ Dual-Chamber Pacemaker

Merlin@home™
Transmitter
Compatible



Product Highlights - Pacemaker

The Assurity MRI pacemaker was designed to allow patients to undergo MRI scans:

- When combined with the Tendril MRI™ LPA1200M lead, the MRI-ready device:
 - Allows full-body, MRI scans*
 - Permits a maximum whole body averaged specific absorption rate (SAR) of 4 Watts per kilogram (W/kg)
- When combined with the Tendril™ 2088TC or IsoFlex™ Optim™ 1944/1948 leads, the MRI-ready device:
 - Allows MRI scans*
 - Permits a maximum whole body averaged specific absorption rate (SAR) of 2 Watts per kilogram (W/kg)
- An optional, easy-to-use hand-held device (SJM MRI Activator™ device) can be used to program the device to pre-approved MRI settings pre- and post-MRI scan, decreasing the number of workflow steps and increasing clinic efficiency
- Physician preferred size and physiologic shape minimize pocket size
- Outstanding longevity provides 9,4 years of service life,¹⁰ which is supported by an 8-year warranty¹¹
- InvisiLink™ wireless telemetry, in conjunction with the Merlin@home™ transmitter and Merlin.net™ Patient Care Network (PCN), allows for daily remote monitoring and follow-up
- CorVue™ congestion monitoring offers greater insight into pulmonary edema status by monitoring transthoracic impedance, and it provides the option for physician alerts
- The only pacemaker with programmable AT/AF alerts specifically indicated for detecting atrial tachyarrhythmias, which have been found to be associated with an increased risk of stroke in elderly, hypertensive, pacemaker patients without prior history of AF¹²
- A suite of state-of-the-art features—complete automaticity (atrial and ventricular), Ventricular Intrinsic Preference (VIP™) technology, the AF Suppression™ algorithm and SenseAbility™ technology—is designed to deliver optimal therapy for patients at implant and throughout their lives
- 6-month ERI-EOL interval

*See MRI Conditional Parameters

Ordering Information - MRI-Ready Pacing System

Model Number	Description	Dimensions (H x W x T, mm)	Weight (g)	Volume (cc)	Connector
PM2272	Assurity MRI Pacemaker	47 x 50 x 6	20	10,4 (± 0,5)	IS-1

Model Number	Description	Insulation	Fixation	Min. Introducer (F)	Connector	Length (cm)
LPA1200M	Tendril MRI Pacing Leads	Optim™	Ext/Ret helix	8	IS-1 bipolar	46, 52, 58
2088TC	Tendril STS Pacing Leads	Optim™	Ext/Ret helix	6	IS-1 bipolar	46, 52, 58
1944 (J-shaped)	IsoFlex Optim Pacing Leads	Optim™	Tines	7	IS-1 bipolar	46,52
1948 (Straight)	IsoFlex Optim Pacing Leads	Optim™	Tines	7	IS-1 bipolar	52, 58

Indications: Implantation is indicated in one or more of the following permanent conditions: syncope, presyncope, fatigue, disorientation due to arrhythmia/bradycardia or any combination of those symptoms. **Rate-Modulated Pacing** is indicated for patients with chronotropic incompetence, and for those who would benefit from increased stimulation rates concurrent with physical activity. **Dual-Chamber Pacing** is indicated for those patients exhibiting: sick sinus syndrome, chronic, symptomatic second- and third-degree AV block, recurrent Adams-Stokes syndrome, symptomatic bilateral bundle branch block when tachyarrhythmia and other causes have been ruled out. **Atrial Pacing** is indicated for patients with sinus node dysfunction and normal AV and intraventricular conduction systems. **Ventricular Pacing** is indicated for patients with significant bradycardia and normal sinus rhythm with only rare episodes of A-V block or sinus arrest, chronic atrial fibrillation, severe physical disability. AF Suppression algorithm is indicated for suppression of paroxysmal or persistent atrial fibrillation episodes in patients with one or more of the above pacing indications.

Contraindications: **Dual-chamber pulse generators** are contraindicated in patients with an implanted cardioverter-defibrillator. **Rate-Adaptive Pacing** may be inappropriate for patients who experience angina or other symptoms of myocardial dysfunction at higher sensor-driven rates. An appropriate Maximum Sensor Rate should be selected based on assessment of the highest stimulation rate tolerated by the patient. **AF Suppression™** stimulation is not recommended in patients who cannot tolerate high atrial-rate stimulation. **Dual-Chamber Pacing**, though not contraindicated for patients with chronic atrial flutter, chronic atrial fibrillation, or silent atria, may provide no benefit beyond that of single-chamber pacing in such

patients. **Single-Chamber Ventricular Demand Pacing** is relatively contraindicated in patients who have demonstrated pacemaker syndrome, have retrograde VA conduction, or suffer a drop in arterial blood pressure with the onset of ventricular pacing. **Single-Chamber Atrial Pacing** is relatively contraindicated in patients who have demonstrated compromise of AV conduction.

Potential Adverse Events: The following are potential complications associated with the use of any pacing system: arrhythmia, heart block, thrombosis, threshold elevation, valve damage, pneumothorax, myopotential sensing, vessel damage, air embolism, body rejection phenomena, cardiac tamponade or perforation, formation of fibrotic tissue/local tissue reaction, inability to interrogate or program a device because of programmer malfunction, infection, interruption of desired device function due to electrical interference, loss of desired pacing and/or sensing due to lead displacement, body reaction at electrode interface or lead malfunction (fracture or damage to insulation), loss of normal device function due to battery failure or component malfunction, device migration, pocket erosion or hematoma, pectoral muscle stimulation, phrenic nerve or diaphragmatic stimulation. The following, in addition to the above, are potential complications associated with the use of rate-modulated pacing systems: inappropriate, rapid pacing rates due to sensor failure or to the detection of signals other than patient activity, loss of activity-response due to sensor failure, palpitations with high-rate pacing.

Refer to the User's Manual for detailed indications, contraindications, warnings, precautions and potential adverse events.

Assurity MRI™

Dual-Chamber Pacemaker

Product Specifications - Pacemaker

PHYSICAL SPECIFICATIONS		
Model	PM2272	
Telemetry	RF	
Dimensions (mm)	47 x 50 x 6	
Weight (g)	20	
Volume (cc)	10.1	
Connector	IS-1	
Remote Monitoring		
Compatible with Merlin@home™ Transmitter		
PARAMETER		SETTINGS
Rate/Timing		
Atrial Pace Refractory (ms)	190-400 in steps of 30; 440; 470 ²	
Atrial Sense Refractory (ms)	93; 125; 157; 190-400 in steps of 30; 440; 470 ²	
Paced AV Delay (ms)	25; 30-200 in steps of 10; 225-300 in steps of 25; 350	
Base Rate (min ⁻¹)	30-130 in steps of 5; 140-170 in steps of 10	
Far-Field Protection Interval (ms)	16 ²	
Hysteresis Rate (min ⁻¹)	Off; 30 ¹ -150 in steps of 5	
Search Interval (min)	Off; 1; 5; 10; 15; 30	
Cycle Count	1-16 in steps of 1	
Intervention Rate (min ⁻¹)	Off, Same Base Rate; 90-120 in steps of 10; Intrinsic +0; Intrinsic +10; Intrinsic +20; Intrinsic +30	
Intervention Duration (min)	1-10 in 1 minute intervals	
Recovery Time	Fast; Medium; Slow; Very Slow	
Maximum Tracking Rate (min ⁻¹)	90-130 in steps of 5; 140-210 in steps of 10	
Mode	A00(R); AA1(R); AAT(R); V00(R); VV1(R); VVTR(R); VDD(R); D00(R); DVI(R); DD(R); DDD(R); Pacing Off	
Post Ventricular Atrial Blanking (ms)	60-200 in steps of 10; 225; 250	
PVARP (ms)	125-500 in steps of 25	
Sensed AV Delay (ms)	25; 30-200 in steps of 10; 225-325 in steps of 25	
Rest Rate (min ⁻¹)	Off; 30-150 in steps of 5	
Rate Responsive AV Delay	Off; Low; Medium; High	
Rate Responsive PVARP/VREF	Off; Low; Medium; High	
Shortest AV Delay (ms)	25-50 in steps of 5; 60-120 in steps of 10	
Shortest PVARP/VREF (ms)	125-475 in steps of 25	
Ventricular Blanking (ms)	Auto; 12-52 in steps of 4	
Ventricular Pace/Sense Refractory ² (Fixed) (ms)	125; 160-400 in steps of 30; 440; 470; 500 ²	
Output/Sensing		
ACap™ Confirm	On; Off; Monitor	
Primary Pulse Configuration	Bipolar	
Backup Pulse Configuration	Bipolar	
Backup Pulse Amplitude (V)	5.0	
Search Interval (hours)	8; 24	
A or V Pulse Amplitude (V)	0.25-4.0 in steps of 0.25; 4.5-7.5 in steps of 0.5	
A or V Pulse Width (ms)	0.05; 0.1-1.5 in steps of 0.1	
A or V Pulse Configuration	Unipolar (tip-case); Bipolar (tip-ring)	
A or V Sense Configuration	Unipolar Tip (tip-case); Bipolar (tip-ring); Unipolar Ring (ring-case)	
Atrial Sensitivity (mV)	0.1-0.4 ⁴ in steps of 0.1; 0.5; 0.75-2.0 in steps of 0.25; 2.5-4.0 in steps of 0.5; 5.0 ²	
Ventricular Sensitivity (mV)	0.5-5.0 in steps of 0.5; 6-10 in steps of 1.0; 12.5 ²	
Ventricular AutoCapture™ Pacing System	On; Off	
Primary Pulse Configuration	Unipolar; Bipolar	
Backup Pulse Configuration	Unipolar; Bipolar	
Backup Pulse Amplitude (V)	5.0 ²	
Search Interval (hours)	8; 24	
AutoCapture	On; Off	
Paced/Sensed AV Delay (ms)	50/25; 100/70; 120/100	
Ventricular Sensitivity (mV)	0.5-5.0 in steps of 0.5; 6-10 in steps of 1.0; 12.5 ²	
SenseAbility™ Technology	Off; On (Automatic Sensitivity Control adjustment for atrial and ventricular events)	
A Max Sensitivity (mV)	0.2-1.0 in steps of 0.1	
V Max Sensitivity (mV)	0.2-2.0 in steps of 0.1	
Threshold Start	(Atrial and Ventricular Post-Sense) 50; 62.5; 75; 100% (Atrial Post-Pace) 0.2-3.0 in steps of 0.1 mV (Ventricular Post-Pace) Auto; 0.2-3.0 in steps of 0.1 mV (Atrial and Ventricular Post-Sense) 0; 30; 60; 95; 125; 160; 190; 220 (Atrial Post-Pace) 0; 30; 60; 95; 125; 160; 190; 220 (Ventricular Post-Pace) Auto; 0; 30; 60; 95; 125; 160; 190; 220	
Decay Delay (ms)		
Rate-Modulated Parameters		
Maximum Sensor Rate (min ⁻¹)	80-150 in steps of 5; 160-180 in steps of 10	
Reaction Time	Very Fast; Fast; Medium; Slow	
Recovery Time	Fast; Medium; Slow; Very Slow	
Sensor	On; Off; Passive	
Slope	Auto (-); Auto (+0); Auto (+1); Auto (+2); Auto (+3); 1-16 in steps of 1	
Threshold	Auto (-0.5); Auto (+0.0); Auto (+0.5); Auto (+1.0); Auto (+1.5); Auto (+2.0); 1-7 in steps of 0.5	
AF Management		
AF Suppression™ Algorithm	Off; On	
Lower Rate Overdrive (min ⁻¹)	10 ³	
Upper Rate Overdrive (min ⁻¹)	5 ³	
No. of Overdrive Pacing Cycles	15-40 in steps of 5	
Rate Recovery (ms)	8; 12 ²	
Maximum AF Suppression Rate (min ⁻¹)	80-150 in steps of 5; 160-180 in steps of 10	
Atrial Tachycardia Detection Rate (min ⁻¹)	110-200 in steps of 10; 225-300 in steps of 25	
Auto Mode Switch	Off; DDD(R) to DDI(R); DDD(R) to VVI(R); VDD(R) to VVI(R)	
AMS Base Rate (min ⁻¹)	40-170 in steps of 5	

Customer Support: 46-8-474-4756

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. Devices depicted may not be available in all countries. Check with your St. Jude Medical representative for product availability in your country.

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Stored Electrograms

Options	
Priority Options	Off; Low; High
Channel	1; 2; 3
Triggers	
Advanced Hysteresis	Off; Low; High
AMS Entry/AMS Exit/AMS Entry and Exit	Off; Low; High
AT/AF Detection	Off; Low; High
Magnet Response	Off; Low; High
High Atrial Rate	Off; Low; High
Rate (min ⁻¹)	125-300 in steps of 25
No. of Consecutive Cycles	2; 3; 4; 5; 10; 15; 20
High Ventricular Rate	Off; Low; High
Rate (min ⁻¹)	125-300 in steps of 25
No. of Consecutive Cycles	2; 3; 4; 5; 10; 15; 20
PMT Termination	Off; Low; High
Consecutive PVCs	Off; Low; High
No. of Consecutive PVCs	2; 3; 4; 5
Noise Reversion	Off; Low; High
Other	
A and V Lead Monitoring	Monitor; Auto Polarity Switch
A and V Low Impedance Limit (Ω)	100-500 in steps of 25
A and V High Impedance Limit (Ω)	750-2500 in steps of 250; 3000
Lead Type	Uncoded; Unipolar; Bipolar
Magnet Response	Off; Battery Test
Negative AV Hysteresis Search (ms)	Off; -10 to -120 in steps of 10
NIPS Options	
Stimulation Chamber	Atrial; Ventricular
Coupling Interval (ms)	100-800 in steps of 10 ⁴
S1 Count	2-25 in steps of 1
SF; S2; S3 and S4 Cycle (ms)	Off; 100-800 in steps of 10 (Fixed or Adaptive)
Ventricular Support Rate (min ⁻¹)	Off; 30-95 in steps of 5
Sinus Node Recovery Delay (sec)	1; 2; 3; 4; 5
PMT Options	
PMT Detection Rate (min ⁻¹)	Off; Passive; Atrial Pace ²
PVC Response	90-180 in steps of 5
Ventricular Intrinsic Preference, VIP™ (ms)	Off; Atrial Pace ²
VIP Search Interval	Off; 50-150 in steps of 25; 160-200 in steps of 10
VIP Search Cycles	30 sec.; 1; 3; 5; 10; 30 min.
Ventricular Safety Standby	1; 2; 3
Diagnostic Trends	Off; On
Thoracic Impedance	Measured every 2 hours
CorVue™ Congestion Monitoring	Off; On
CorVue Congestion Trigger	8-18 days

MRI Settings

MRI Mode	A00; V00; D00; Pacing Off		
MRI Base Rate	30-120 bpm in steps of 5 bpm		
MRI Paced AV Delay	25 ms; 30-120 ms in steps of 10 ms		
MRI Atrial Pulse Configuration	Bipolar		
MRI Atrial Pulse Amplitude	5.0 V; 7.5 V		
MRI Atrial Pulse Width	1.0 ms		
MRI RV Pulse Configuration	Bipolar		
MRI RV Pulse Amplitude	5.0 V; 7.5 V		
MRI RV Pulse Width	1.0 ms		
MRI Conditional Parameters			
Lead	Lead Lengths	Scan Exclusion Zone	
Tendril MRI LPA1200M Lead	46, 52, 58 cm	No scan exclusion zone	
Tendril 2088TC Lead	46, 52, 58 cm	Isocenter must be inferior to L4 or 10 cm superior to C1	
IsoFlex 1944 Lead	46, 52 cm	Isocenter must be inferior to L4 or superior to C1	
IsoFlex 1948 Lead	52, 58 cm	Isocenter must be inferior to L4 or superior to C1	
Lead	Lead Lengths	Magnet	SAR
Tendril MRI LPA1200M Lead	46, 52, 58 cm	1.5T	≤ 4 W/kg
Tendril 2088TC Lead	46, 52, 58 cm	1.5T	≤ 2 W/kg
IsoFlex 1944 Lead	46, 52 cm	1.5T	≤ 2 W/kg
IsoFlex 1948 Lead	52, 58 cm	1.5T	≤ 2 W/kg

- ± 0.5 cc
- Programming options dependent on pacing mode.
- This parameter is not programmable.
- The highest available setting for hysteresis rate will be 5 min-1 below the programmed base rate.
- In dual-chamber modes, the maximum ventricular refractory period is 325 ms.
- Values 0.1-0.4 not available in a unipolar sense configuration.
- Sensitivity is with respect to a 20 ms haversine test signal.
- During atrial NIPS in dual-chamber modes, the shortest Coupling Interval will be limited by the programmed AV/PV delay.
- S1 Burst Cycle is applied at the preprogrammed S1 cycle length.
- A.V = 2.5 V @ 0.4 ms; 500 ohms; 100% DDD pacing @ 60 bpm; AutoCapture™ Pacing System OFF; SEGMS ON
- Terms and conditions apply; refer to the warranty for details.
- Healey SJ, Connolly SJ, Gold MR, et al. on behalf of the ASSERT investigators. Sub-clinical atrial fibrillation and the risk of stroke: Automatic atrial fibrillation and Stroke Evaluation in pacemaker patients and the AF Reduction atrial pacing Trial (ASSERT). N Engl J Med 2012; 366:120-129.