LINQ FAMILY OF ICMs

Parameter	Reveal LINQ™ ICM[1]	LINQ II™ ICM[2]
Longevity	3 years	4.5 years*
Electrode Spacing	38 mm	40 mm
Volume	1.2 cc	1.4 cc
Mass	2.5 g	3.4 g
Episode Storage	59 min	61 min
Monitoring Option	Home monitor	Home monitor and mobile app
Patient Symptom Mark	Patient Assistant	Mobile app or Patient Assistant
Cardiac Compass™	Yes	Yes
MRI Compatibility	1.5 and 3T	1.5 and 3T
Clinician Notification	Nightly transmission/ CareAlert™ notifications	Between 5–6 a.m. clinic time daily transmission/CareAlert notifications
Telemetry	Inductive (Tel B) One-directional RF (MEDS)	Bluetooth® Low Energy Two-way communication
Detection Algorithms	P-SENSE detection TruRhythm™ ▪ Pause ▪ Brady ▪ AF	Enhanced TruRhythm ▪ Pause enhancement ▪ Tachy: Require Rapid Onset ▪ Brady: Nighttime Storage ▪ PVC burden
Remote Programming	No	Yes
CareLink™ Network	Yes	Yes

[1] Reveal LINQ LNQ11 ICM Clinician Manual M958488A001 Rev D.
[2] LINQ II LNQ22 ICM Clinician Manual. M974764A001D.

*Nominal settings.

Medtronic

Brief Statement
LINQ II™ Insertable Cardiac Monitor System (ICM) and Remote Monitoring

Indications: The LINQ II ICM is an insertable automatically activated and patient-activated monitoring system that records subcutaneous ECG and is indicated in the following cases:

- Patients with clinical syndromes or situations at increased risk of cardiac arrhythmias
- Patients who experience transient symptoms such as dizziness, palpitation, syncope, and chest pain that may suggest a cardiac arrhythmia

The device has not been tested specifically for pediatric use.

Contraindications: There are no known contraindications for the insertion of the LINQ II ICM. However, the patient's particular medical condition may dictate whether or not a subcutaneous, chronically inserted device can be tolerated.

Warnings and Precautions: Patients with the LINQ II ICM should avoid sources of diathermy, high sources of radiation, electrosurgical cautery, external defibrillation, lithotripsy, therapeutic ultrasound, and radiofrequency ablation to avoid electrical reset of the device, and/or inappropriate sensing as described in the Medical procedure and EMI Warnings, Precautions and Guidance Manual. MRI scans should be performed only in a specified MR environment under specified conditions as described in the LINQ II MRI Technical Manual.

Wireless accessories available for use with LINQ II may experience connectivity or performance issues. See product manuals for details and troubleshooting instructions.

Potential Adverse Events or Potential Complications: Potential adverse events from the LINQ II ICM include, but are not limited to, device rejection phenomena (including local tissue reaction), device migration, infection, and erosion through the skin.

There are no known adverse events associated with the use of any LINQ II ICM wireless accessory.

See the device manuals for detailed information regarding the implant procedure, indications/intended use, contraindications, warnings, precautions, and potential complications/adverse events. For further information, please call Medtronic at 1-800-328-2518 (Technical Services), 1-800-551-5544 (Patient Services), and/or consult the Medtronic website at medtronic.com.

Caution: Federal law (USA) restricts this device to sale by or on the order of a physician.

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