Select devices in Medtronic history. Does not include all Medtronic devices.

PCD 7216 was an investigational device in 1989 and market released in 1993.

CARDIAC

See the back side or accompanying documentation for information regarding devices supported on the CareLink Network. January 2014. Data on File.

Over 99% of Medtronic devices are compatible with the CareLink™ Network™.
The SureScan pacing systems, proper patient monitoring must be provided during the MRI scan.

For SureScan defibrillation and CRT-D systems, continuous patient monitoring is required after MRI Scan is completed. Do not leave the device in the MRI Scan environment after the scan is complete. While MRI is performed on patients, arrhythmia detection and therapies are suspended; leave the patient at risk of death from unmonitored spontaneous tachyarrhythmias. If the device is a programmer, MI may not be possible.

Potential Complications

The SureScan device is contraindicated for use after an MRI scan, except for those patients who may benefit from increased pacing rates concurrent with increases in activity.

Caution: Federal law (USA) restricts these devices to be used by or on the order of the physician.

Contraindications

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