

# INSYNC REGISTRY

**Principal Investigator:**

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**Sponsor:**

Medtronic

**Study Purpose:**

InSync Registry: **Post market** Study Cardiac Resynchronization Therapy

**Objective:**

The purpose of the Registry is to evaluate the post market release performance of the InSync model 8040 and InSync III model 8042 and any Medtronic CRT-D systems for cardiac resynchronization therapy (CRT).

**Inclusion Criteria:** (Non-MIRACLE Patients ONLY)

- ♥ Patient implanted with InSync Model 8040, 8042 or Medtronic market released CRT-D Cardiac resynchronization therapy System
- ♥ Patient must provide written informed consent

**Status:**

Follow-up ONLY!

If you have any questions, please feel free to contact the coordinators, and they will be happy to answer any questions you have regarding this study.