



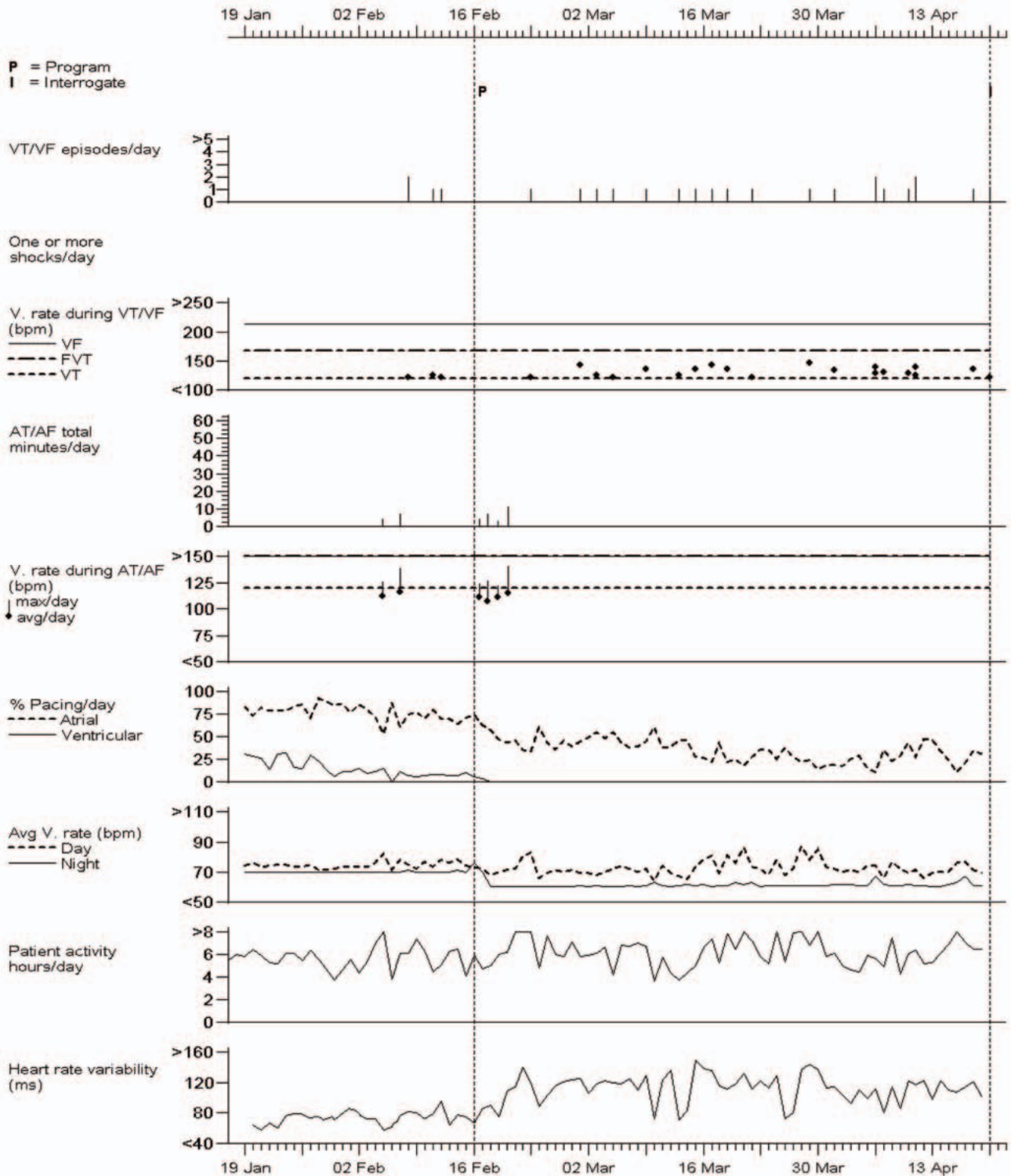
Cardiac Compass Report - Last 90 Day Zoom

ICD Model: Maximo DR

Serial Number:

Date of Visit:

Patient Name



## Brief Statements

### CardioSight Reader

#### Intended Use

The Model 2020A CardioSight™ Reader is intended for use in a clinical setting. The CardioSight Reader is indicated for use to interrogate compatible Medtronic implantable devices to collect patient and device data and send the information to the clinician. The Model 2020A CardioSight Reader cannot be used to change therapy.

#### Contraindications

There are no contraindications for the Model 2020A CardioSight Reader.

#### Warnings and Precautions

The CardioSight Reader must only be used for interrogating compatible Medtronic implantable devices. Do not use a cellular phone while the antenna is positioned over the implanted device. The CardioSight Reader is designed for use in the continental United States, Alaska, and Hawaii.

### Medtronic ICDs

#### Indications

Medtronic implantable cardioverter defibrillators (ICDs) are indicated for ventricular antitachycardia pacing and ventricular defibrillation for automated treatment of life-threatening ventricular arrhythmias.

#### Contraindications

Medtronic ICDs are contraindicated in patients whose ventricular tachyarrhythmias may have transient or reversible causes, patients with incessant VT or VF, patients who have a unipolar pacemaker, and patients whose primary disorder is bradyarrhythmia.

#### Warnings/Precautions

Changes in a patient's disease and/or medications may alter the efficacy of the device's programmed parameters. Patients should avoid sources of magnetic and electromagnetic radiation to avoid possible underdetection, inappropriate sensing and/or therapy delivery, tissue damage, induction of an arrhythmia, device electrical reset, or device damage. Do not place transthoracic defibrillation paddles directly over the device.

#### Potential Complications

Potential complications include, but are not limited to, rejection phenomena, erosion through the skin, muscle or nerve stimulation, oversensing, failure to detect and/or terminate tachyarrhythmia episodes, acceleration of ventricular tachycardia, and surgical complications such as hematoma, infection, inflammation, and thrombosis.

*See the device manuals for detailed information regarding instructions, indications, contraindications, warnings, precautions, and potential complications/adverse events.*

**Caution:** Federal law (USA) restricts these devices to sale by or on the order of a physician.

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