Case Studies

The results presented in these case studies are specific to these individual patients. Patient results will vary, not every response is the same.

 Physicians Note: The following slides may be used as a format for your own case studies if you wish to insert clinical information for your patients who have received cardiac resynchronization therapy.

Case Study: Ischemic Cardiomyopathy

Patient:

- 57 year-old white male
- Ischemic cardiomyopathy
- NYHA Class III heart failure

Cardiovascular History/Diagnostic Evaluation:

- Anterior myocardial infarction, 1995. Cardiac catheterization single vessel coronary artery disease, S/P failed-PTCA (balloon angioplasty) with emergency coronary artery bypass graft surgery 1995.
- Aortic valve replacement 1995.
- Echo, Nov 1998 Severe LV dysfunction, EF less than 20%. LV dilatation.
- ECG, Nov 1998 asymptomatic sinus bradycardia, 55 beats per minute. RBBB. QRS duration 152 msec.
- Right hemispheric CVA, 1998.
- Hyperlipidemia.
- No indication for pacemaker or ICD; No history of atrial arrhythmias

Case Study: Ischemic Cardiomyopathy (continued)

Baseline Cardiac Medications:

- ACE inhibitor: lisinopril 20 mg once per day
- Beta blocker: carvedilol 25 mg twice daily
- Diuretic: furosemide 40 mg twice daily
- Digoxin 0.25 mg once per day
- Nitrate: isosorbide mononitrate 30 mg once per day
- Aspirin 81 mg once per day
- Anticoagulant: warfarin sodium (as directed).
- Lipid lowering: atorvastatin calcium 10 mg once per day

Cardiac Resynchronization Therapy:

- The patient received an cardiac resynchronization device Nov 1998 while enrolled in the MIRACLE trial.
- The patient was randomized to the control "OFF" arm and began receiving cardiac resynchronization therapy three months later in Feb 1999*.

^{*}Patient was enrolled in the initial phase in which patients were randomized for three months

Case Study: Ischemic Cardiomyopathy (continued)

Cardiac Resynchronization Therapy Results:

Quality of Life Score
NYHA Class
Six-Minute Walk
Peak VO₂
Peak Exercise Time
EF (core lab calc)
QRS Duration
Symptoms

Baseline (Nov 1998) 65 Ш 319 meters 12.2 ml/kg/min 619 seconds 23% 160 ms Unable to walk room-toroom, unable to carry on daily activities;

sleeping average of 22

hours a day.

12-Month Evaluation after Implant (Nov 1999) **50** Ш 391 meters 13.9 ml/kg/min 702 seconds 28% 120 ms Walks at least one mile a day, remodeling home, sleeps 6-8 hours a night, working part-time, and recently helped build a mission church in Nicaragua.

- The data reflected in the third column (12-month evaluation) corresponds to 9-months of cardiac resynchronization therapy for this patient.
- Note: This patient was enrolled in the initial 3-month randomization period of the MIRACLE trial. Crossover occurred and the patient began receiving therapy 3 months post-implant.

Case Study: Non-Ischemic Cardiomyopathy

Patient:

- 71-year-old white female
- Idiopathic dilated cardiomyopathy
- NYHA Class III heart failure

Cardiovascular History/Diagnostic Evaluation:

- Cardiac catheterization, Jan 2000 global hypokinesis with EF 25-30%, normal LV filling pressures. Luminal coronary atherosclerosis.
- Echo, Jan 2000 left atrial size 3.6 cm. Dilated left ventricle with LVEDD 58 mm. EF 30-35%. Anterior wall akinetic, no significant valvular abnormalities.
- ECG, March 2000 normal sinus rhythm, left axis deviation, LBBB, QRS 160 msec.
- Hyperlipidemia
- No indication for pacemaker or ICD; No history of atrial arrhythmias

Case Study: Non-Ischemic Cardiomyopathy (continued)

Baseline Cardiac Medications:

- ACE inhibitor: enalapril maleate 10 mg once per day
- Beta blocker: carvedilol 6.25 mg once per day
- Diuretic: furosemide 40 mg once per day
- Digoxin 0.125 mg once per day
- Aspirin 81 mg once per day
- Potassium replacement: 20 mEq once per day

Cardiac Resynchronization Therapy:

- The patient received an cardiac resynchronization device March 2000 while enrolled in the MIRACLE trial.
- The patient was randomized to the control "OFF" group and began receiving cardiac resynchronization therapy six months after the implant.

Case Study: Non-Ischemic Cardiomyopathy

Cardiac Resynchronization Therapy Results:

Quality of Life Score
NYHA Class
Six-Minute Walk
Peak VO ₂
Peak Exercise Time
EF (core lab calc)
ORS Duration

Symptoms

Baseline (March 2000)	12-Month Evaluation after Implant (March 2001)
82	4
III	1
259 meters	335 meters
15.7 ml/kg/min	16.39 ml/kg/min
215 seconds	478 seconds
34%	39%
160 ms	160 ms
Exertional dyspnea with chest discomfort, extreme fatigue with any exercise, palpitations, 2-pillow orthopnea, paroxysmal nocturnal dyspnea, inability to do housework.	Able to do daily housework, laundry, cooking, vacuuming without shortness of breath. No limitation in activity and regularly goes dancing.