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3

Mode Mode Switch Detection Rate Detection Duration Blanked Flutter Search	DDDR On 150 bpm No Delay On	Stop Rate 12) ppm 10 ppm 0 ms	Rate Response Activity Acceleration Activity, Deceleration High Rate Percent ADL Rate Setpoint Upper Sensor Rate Se	30 sec Exercise 0.2% 13 tpoint 35	Ventricular Lead Amplitude Pulse Width Sensitivity Sensing Assurance Pace Polarity	3.500 0.46 5.60 On Bipol
Rates		PVARP Minimum PVARP	Auto 250 ms	- Atrial Lead	Sense Polarity Lead Monitor	Bipola Adapt	
Lower Rate Upper Tracking Rate Upper Sensor Rate ADL Rate Intrinsic/AV	90 ppm 120 ppm 120 ppm 95 ppm	PVAB Ventricular Refractory Vent. Blanking (after A. Pace) PMT Intervention PVC Response Ventricular Safety Pacing	180 ms 230 ms	Amplitude Pulse Width Sensitivity Sensing Assurance Pace Polarity Sense Polarity Lead Monitor Maximum Impedance	3.500 V 0.40 ms 0.18 mV On Bipolar Bipolar	Maximum Impedance Minimum Impedance Monitor Sensitivity Capture Management Amplitude Margin Min. Adapted Amplitude Capture Test Frequency Acute Phase	4.000 200 o 8 Adapt 2x 2.000 Day a 105 d
Paced AV Sensed AV	180 ms	Rate Response			Monitor Only		
Search AV+ Sinus Preference Sinus Preference Zone Search Interval Rate Adaptive AV	Off On 10 ppm 10 min	Optimization ADL Response Exertion Response ADLR Percent Activity Threshold	On 3 3 2.0% Medium/Low	Minimum Impedance Monitor Sensitivity Capture Management	200 ohms 8 Off	V. Sensing During Search	Adap

97 yo - 1 week s/p DC PM Implant

Battery Status			Parameter Su	ımmary			
Estimated remaining longevity: Based on Past History	7.5 years, 6 - 9 years		Mode Mode Switch	DDDR On	Lower Rate Upper Tracking Rate	90 ppm 120 ppm	Search AV+ Paced AV
Voltage/Impedance	2.79 V / 100 ohms		Detection Rate	150 bpm	Upper Sensor Rate	120 ppm	Sensed AV
Lead Summary	Atrial	Ventricular					
Measured Threshold Date Measured	Off	0.500 V at 0.40 ms 02/07/18					
Programmed Output	3.500 V / 0.40 ms	3.500 V / 0.46 ms					
Capture	Off	Adaptive					
Measured P / R Wave	<0.7 to 1.4 mV	>=80% Paced					
Programmed Sensitivity	0.18 mV	5.60 mV					
Measured impedance	457 ohms	575 ohms					
Lead Status	OK	OK					
Lead Model	407652	407658					
Implanted	1-30-18	1/30/18					

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5

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What is your generic diagnosis?

- 1. Oversensing
- 2. Undersensing
- 3. Failure to capture
- 4. Crosstalk

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97 yo - 1 week s/p DC PM Implant

What of the following would "NOT" be in your differential based on the clinical scenario and the generic diagnosis?

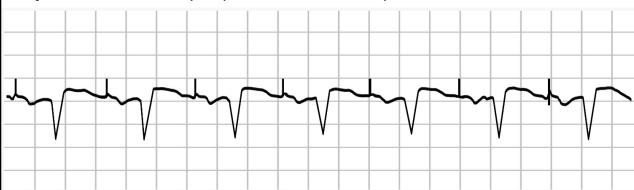
- 1. Lead dislodgement
- 2. Lead fracture
- 3. Loose set screw
- 4. Air in the header

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7

Pacemaker programmed to DDI 86 ppm; AVI = 165 ms; PVAB = 13 ms. Referred with ? of abnormal PM function

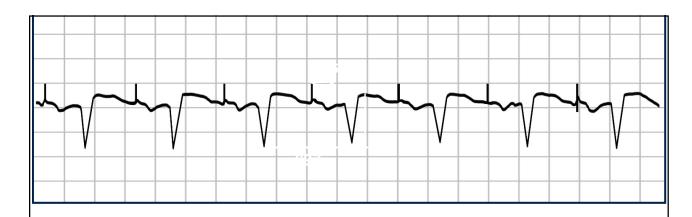
(Note - all A activity is paced & all V activity is intrinsic



The most appropriate approach would be:

- 1. Lengthen PVARP
- 2. Lengthen PVAB
- 3. Lengthen AVI
- 4. Lower the upper rate limit

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Determining the VA Interval

VV = AV + VA

VV - AV = VA

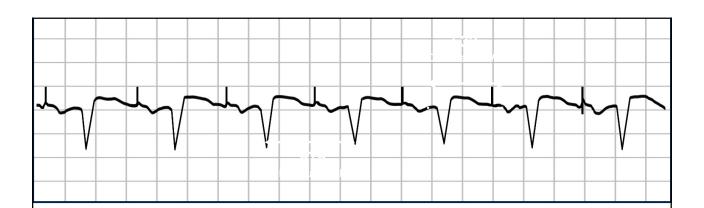
Programmed VV = 86 ppm = 697 ms

AV = 165 ms

697 - 165 = 532 ms (VA)

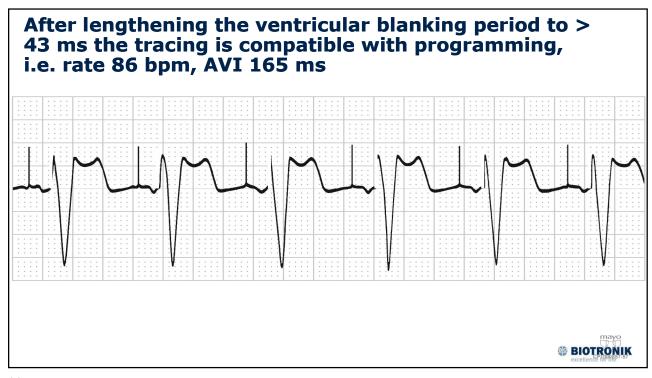
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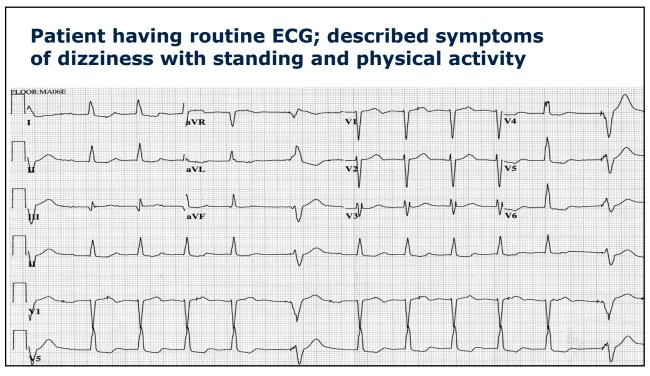
9

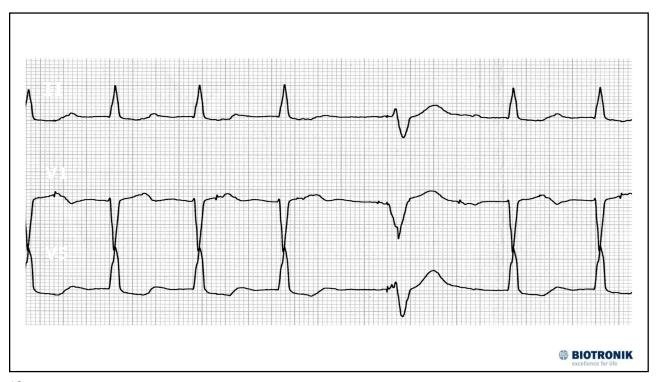


Measured RR = 575 ms = VV 575 - 532 (VA) = 43 ms (AV) Point of sensing after atrial output is 43 ms Blanking period of > 43 ms should avoid crosstalk

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Brady Programming:

Pacing Mode: DDDR ↔ AAIR
Lower Pacing Rate: 60 ppm

• Upper Rate: 110 ppm

Blanking after pace: 200 msecPaced AV Delay: 340 msec

• Sensed AV Delay: 310 msec

• PVARP: 250 msec

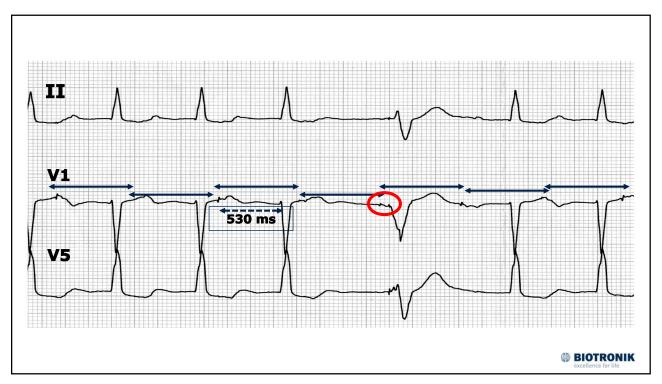
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Symptoms are related to:

- 1. Ventricular undersensing
- 2. Atrial failure to capture
- 3. Effective AV dissociation
- 4. Non-pacemaker issues

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15



Symptoms are related to:

- 1. Ventricular undersensing Any suggestion of undersensing is a function of the MVP algorithm and one would need to overlay refractory and blanking periods would be available with interrogation
- 2. Atrial failure to capture Telemetry strips should be used to definitively demonstrate that atrial capture normal; also programming to AAI or DDI and performing atrial capture thresholds should be part of troubleshooting
- 3. Effective AV dissociation As a result of MVP
- 4. Non-pacemaker issues Non-answer + there is a pacemaker explanation



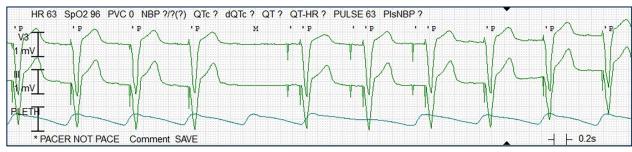
17

Pacemaker syndrome 2° to MVP

- The patient was having dizziness with standing and with physical activity. ECG demonstrates pacing spikes before and on the T wave.
- AV delay was being extended so much by the MVP mode (>400 ms), the patient would develop a 2:1 heart block at low rates and effective V rate would drop.
- Reprogrammed to DDDR and extended P-AV delay to 340 msec. Sitting the patient was Ap-Vs; standing heart rate ↑ into the 70's with AV pacing; no dizziness with standing.
- The patient will return in one month to recheck the device and see if has improvement and check AV pacing percentages.







- 1. Failure to capture
- 2. Failure to output
- 3. Undersensing
- 4. Rate variation



Mode	Rates		AV Intervals	3
Mode DDD	Lower	50 bpm	Paced AV	150 ms
Mode Switch 222 bpm	Upper Track Upper Sensor	210 bpm 145 bpm	Sensed AV	110 ms
Pacing Details	Atrial	RV		
Amplitude	2.00 V	5.00 V		
Pulse Width	0.40 ms	0.50 ms		
Capture Management	Adaptive	Adaptive		
Amplitude Margin	2.0 X	2.0 X		
Min. Adapted Amplitude	2.00 V	2.50 V		
Acute Phase Remaining	119 days	119 days		
Sensitivity	0.30 mV	0.45 mV		
Pace Polarity	Bipolar	Bipolar		
Sense Polarity	Bipolar	Bipolar		
Lead Monitor	Adaptive	Adaptive		
Min Limit	200 ohms	200 ohms		
Max Limit	3000 ohms	3000 ohms		
Device: Advisa DR MRI A	2004		Date	e of Visit: 05-Nov-2014 09:25:40
Serial Number:	ZDRUT			9995 Software Version 8.3 Copyright © Medtronic, Inc. 2014

Failure to Output

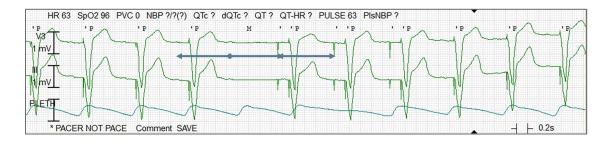
- Oversensing
 - Crosstalk
 - EMI
- Battery failure
- · Circuit failure
- Lead fracture
- Internal insulation failure

- Loose set-screw
- Incompatible lead/header
- Pseudomalfunction/ Device nuance

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21

If intervals are regular and baseline steady, less likely to be oversensing .



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Failure to Output

- Oversensing
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- Circuit failure
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- Loose set-screw
- Incompatible lead/header
- Pseudomalfunction
- Device nuance peculiarity



23

Engineering explanation:

- An occasional 'dropped VP' is specific to DDDR and DDD modes during atrial tracking (AS-VP) and may occur intermittently on an hour and 30 s schedule.
- A false ventricular sense may occur due to residual electrical disturbance on the ventricular sense amplifier and may be created by turning

Pacing spikes Misssing QRS

Misssing QRS

on/off the diagnostic EGM amplifiers during reference EGM collection. The scheduled reference EGM collection is synchronized to sensing and pacing events so that the device's sense amplifiers are blanked. However, there is no ventricular blanking following an AS event, which makes this phenomenon possible. The collection of reference EGM is non-programmable and is always active in the device.

Mehra, et al. Europace e-Pub April 16, 2015

