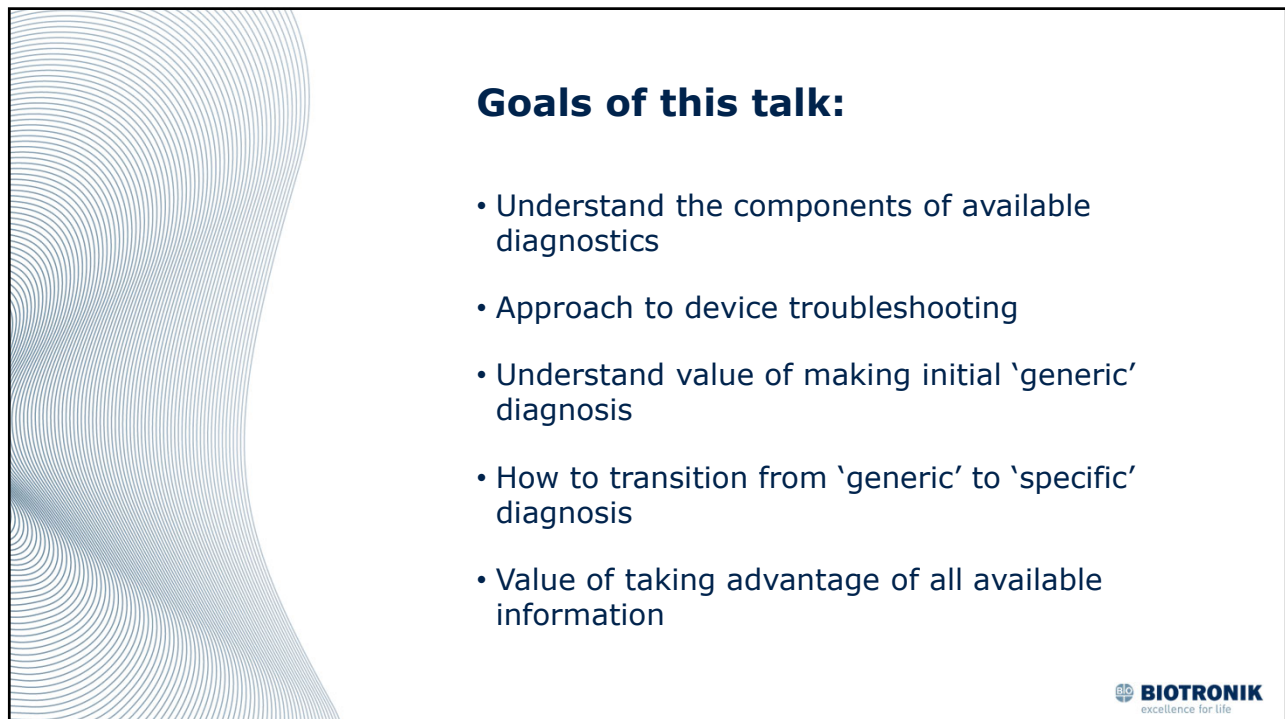


1



2

## CIED Diagnostics

- There are extensive and increasingly sophisticated diagnostics in contemporary devices
- Diagnostics are critical for determination of normal or abnormal device function
- A deep understanding of the diagnostics available from the manufacturer(s) implanted at your institution is invaluable
- In the clinical environment you will have the benefit of the patient's clinical information and programming – this talk purposefully approaches diagnostics without providing the benefit of all the usual information in the effort to develop a framework for a systematic approach

3



3

## Device Troubleshooting: *CRT builds on ICD builds on Brady*

### Approach brady devices with simple steps:

- What's the underlying rhythm?
- Single vs dual-chamber pacing?
- Which chamber(s)?
- Identifiable timing intervals?
- Make a "Generic" Diagnosis
- Target specific dx based on clinical scenario for that generic differential diagnosis

4



4

## Troubleshooting Steps to Consider: Brady

- Re-interrogate if findings don't match what you expect from programming you believe to exist
- Try other pacing modes to overcome a problem, i.e. start with simplest (VVI) and change from there
- Telemetry: EGMS, marker channel etc.
- Patient Postural Testing
- Chest x-ray
- Technical manual
- Call manufacturer 24 hour support number (1-800-547-0394)
- Intraoperative troubleshooting

5

5

## Pacing Electrocardiography

*Need to know and understand differential diagnoses of each of the following:*

- Failure to sense
- Failure to capture
- Failure to output/over-sensing
- Rate variations
- Crosstalk / safety pacing

6

6

## Loss of Capture

### Less Common

- Loose set-screw
- Exit block
- Perforation
- Battery failure
- Circuit failure
- Air in pocket (Unipolar)
- Pseudomalfunction
- Metabolic/drug

### More Common

- Lead dislodgment
- Elevated thresholds
- Inappropriate lead placement
- Lead fracture
- Lead insulation failure

7

7

## Failure to Output

### Less Common

- Loose set-screw
- Lack of anodal connector contact
- Incompatible lead/header
- Pseudomalfunction or device nuance - peculiarity

### More Common

- Over-sensing
- Crosstalk
- EMI
- Battery failure
- Circuit failure
- Lead fracture
- Internal insulation failure

8

8

## Under-Sensing

- Change in intrinsic complex, i.e. BBB, VF, VT, AF
- Myocardial infarction
- Lead dislodgment/poor positioning
- Lead insulation failure
- Magnet application
- ERI
- Functional under-sensing

9

9

## Over-Sensing

- Lead fracture
- Lead insulation defect
- EMI
- Isoelectric ventricular event
- Sensing T wave, P wave, afterpotential, etc

10

10

## **Rate Variations (i.e. variations from programmed lower rate)**

- Hysteresis
- Rate-adaptive pacing
- Oversensing
- Function of timing system, i.e. AA, VV, hybrid
- Specific Algorithms, i.e. Capture Control, Night Rate, etc.

11

11

## **ICD Troubleshooting**

- Failure to detect arrhythmia(s)
- Failure to convert arrhythmia(s)
- Overlapping Arrhythmias
- Inappropriate/unnecessary shocks

12

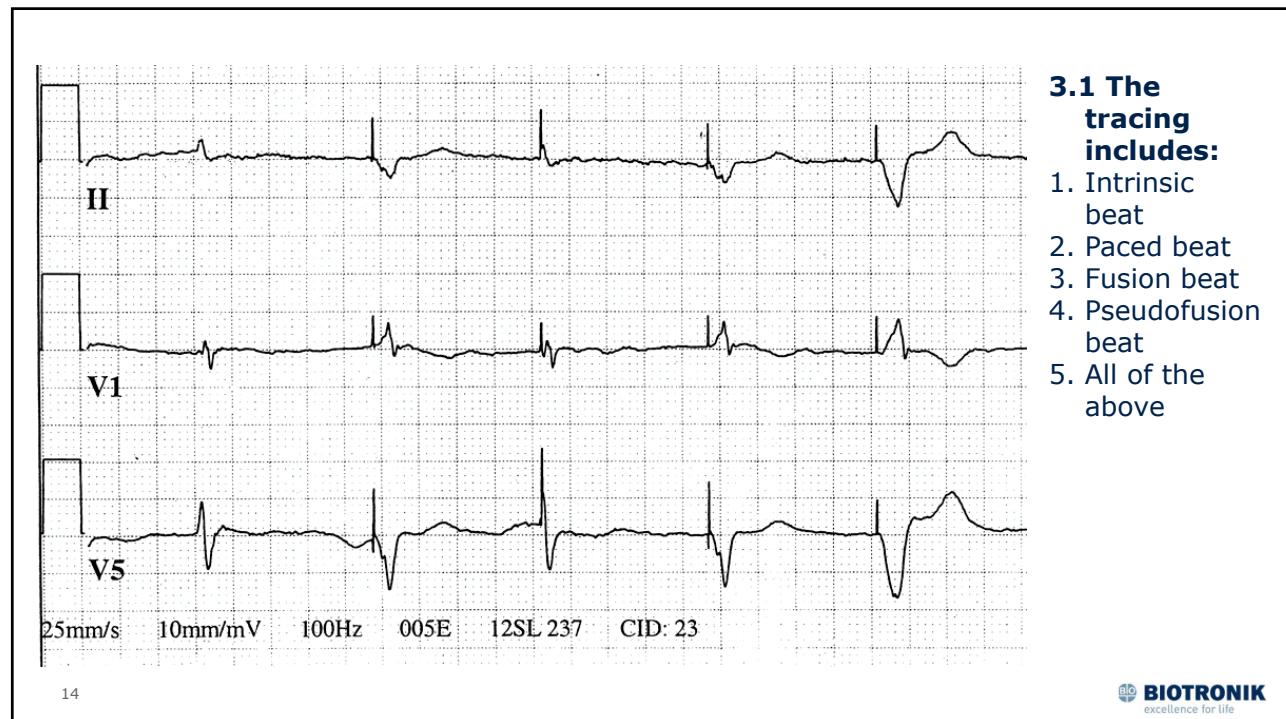
12

## CRT Troubleshooting

- Non-responder
- Suboptimal % biventricular pacing
- Atrial arrhythmias
- Ventricular arrhythmias
- Phrenic nerve stimulation

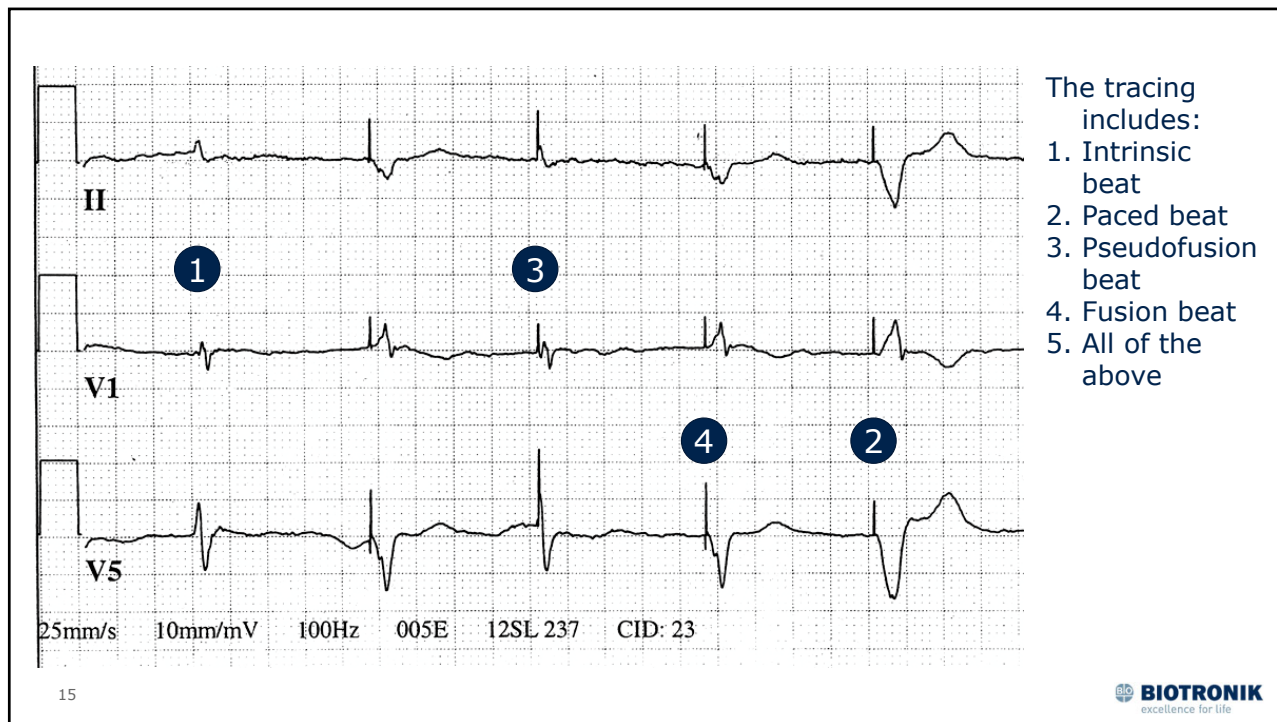
13

13

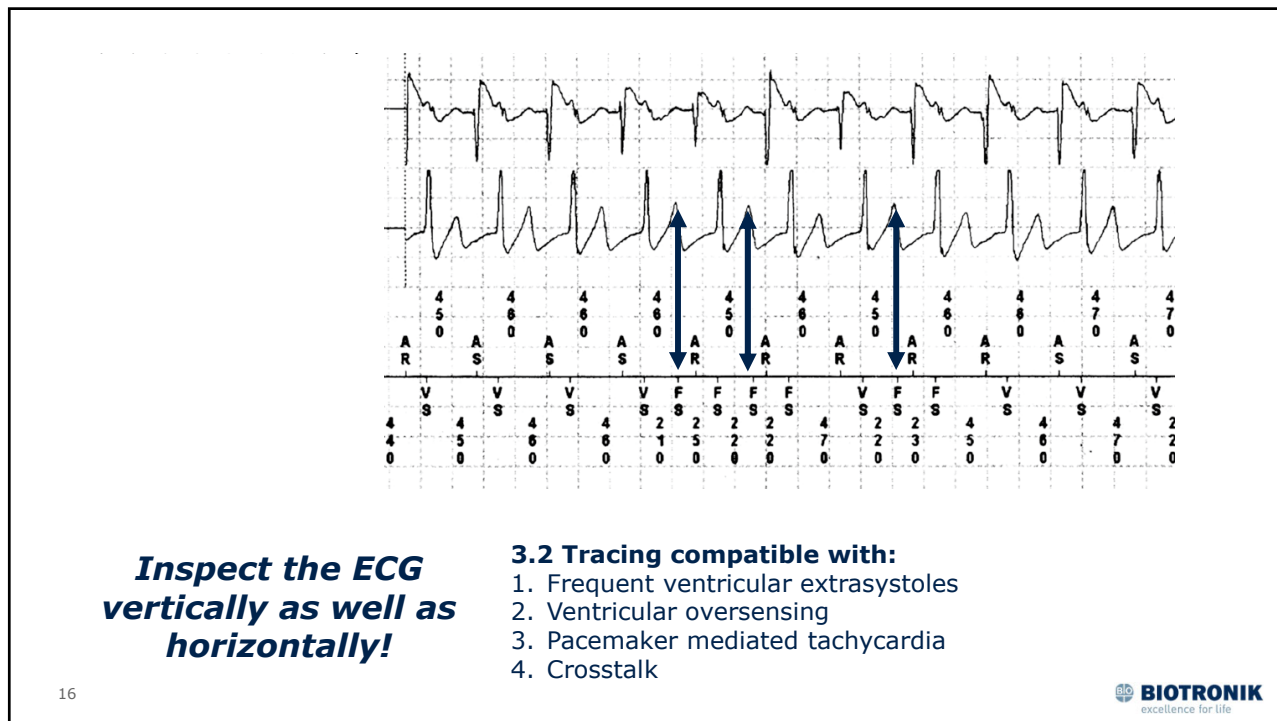


14

14

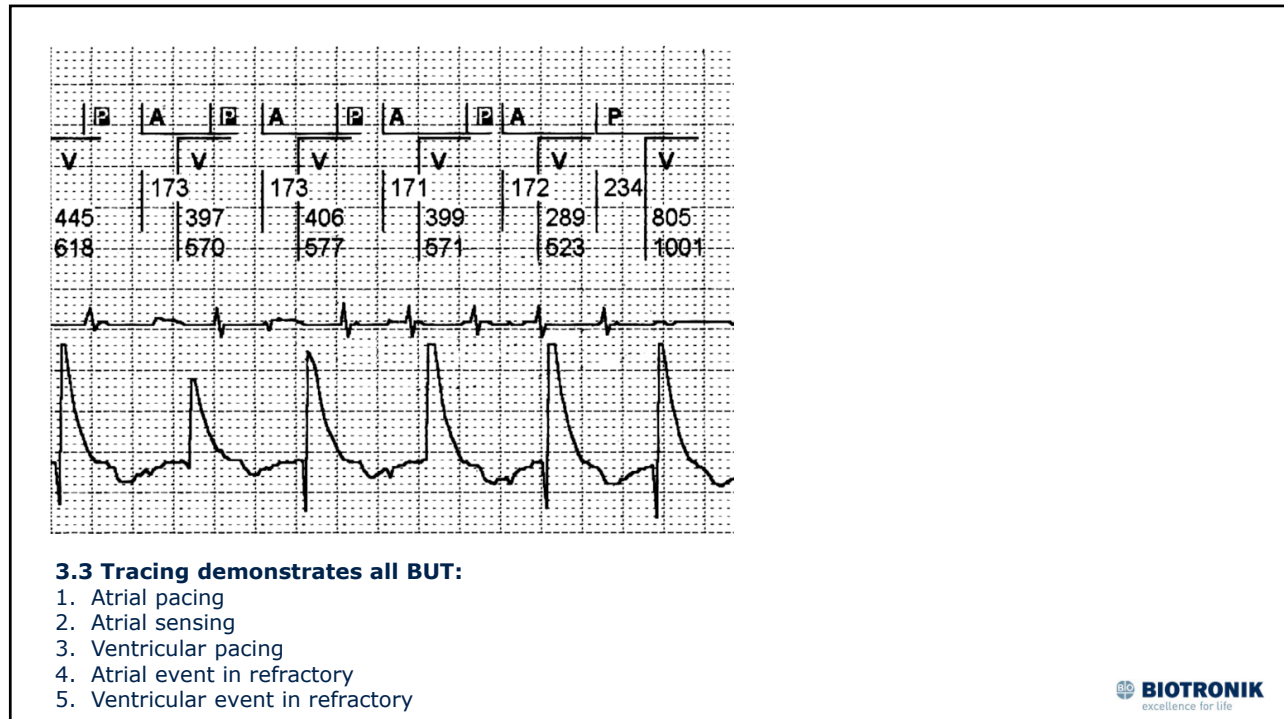


15

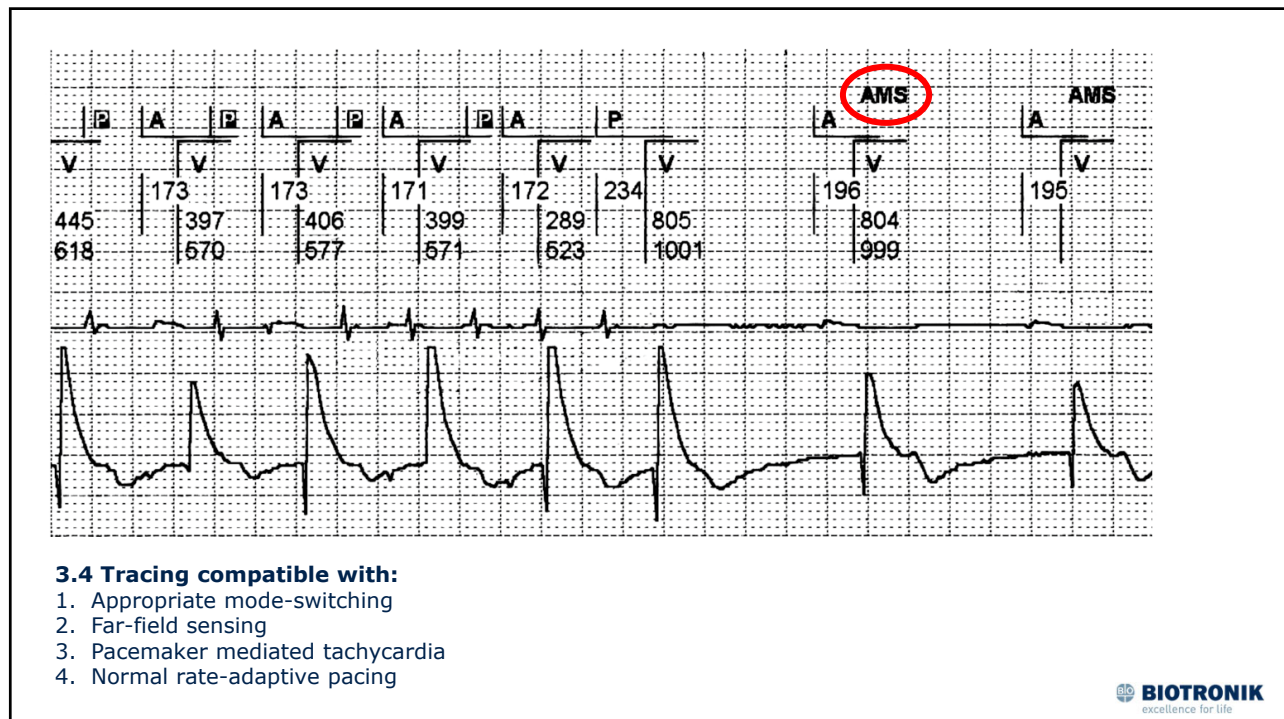


16

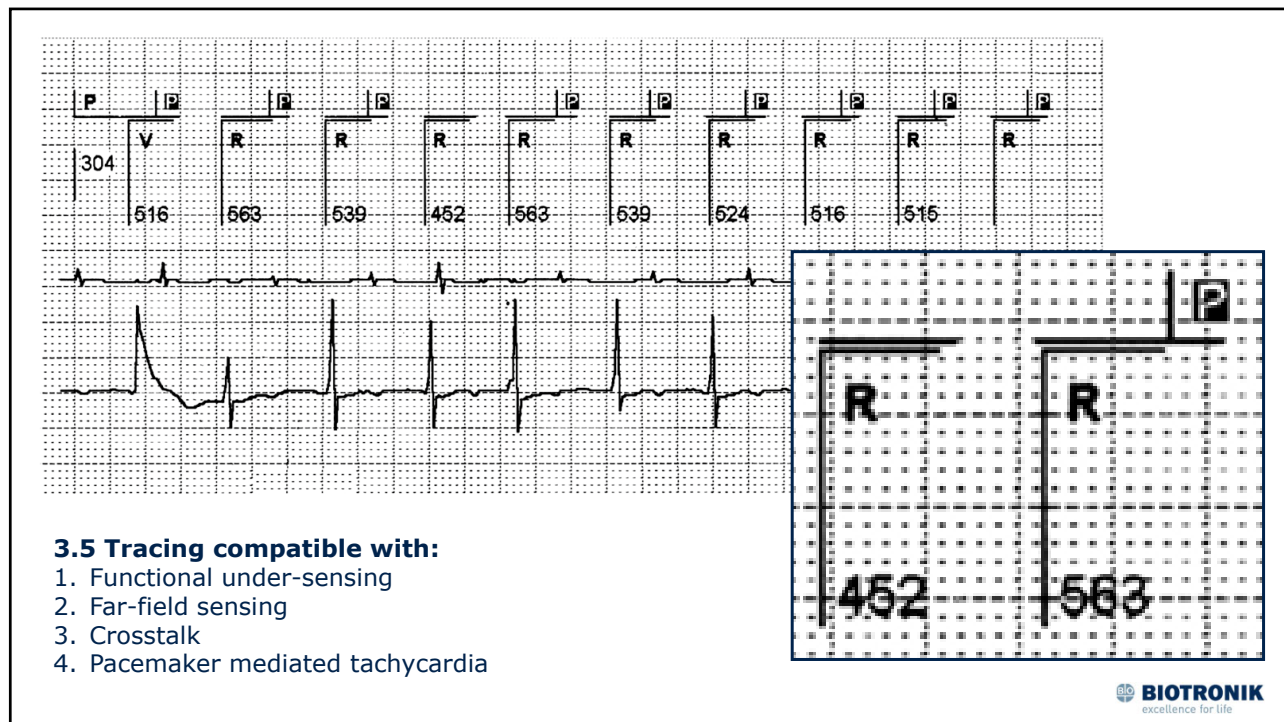




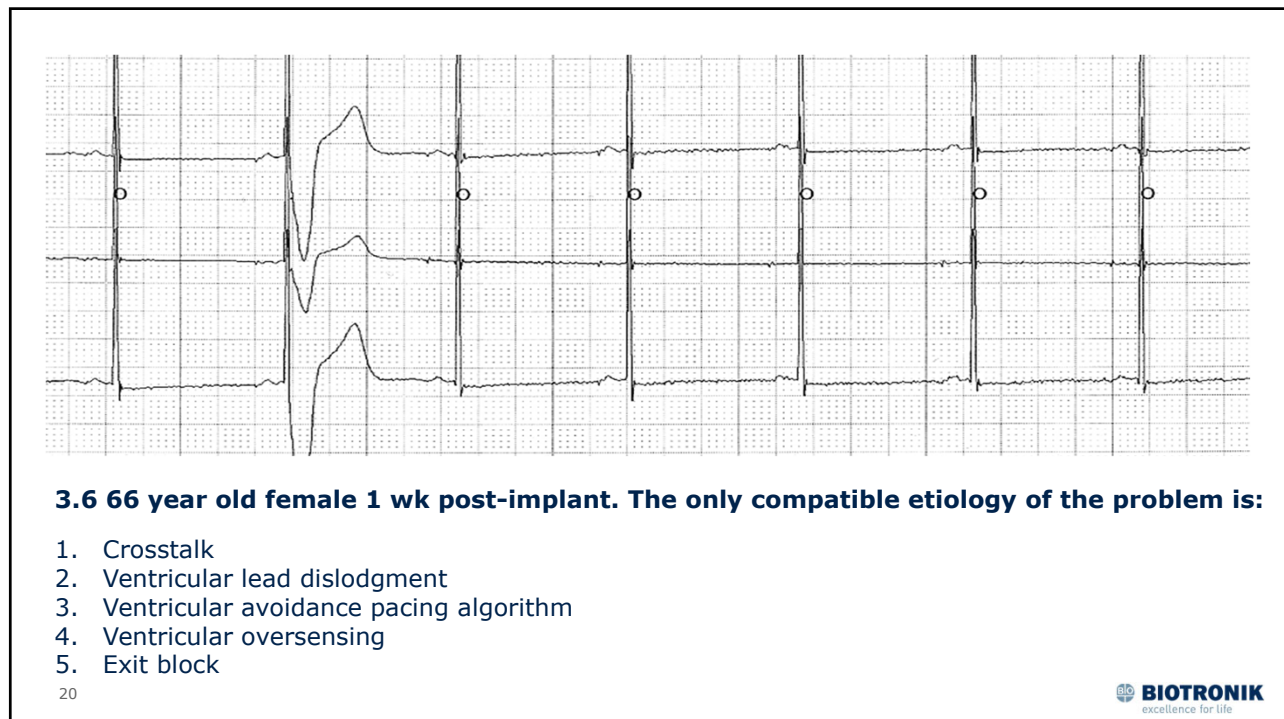
17



18

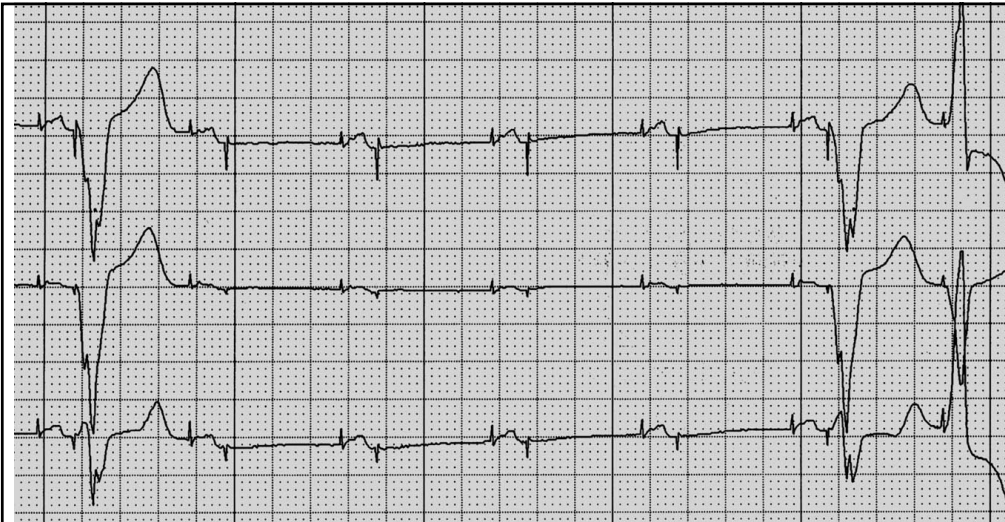


19



20

20



**3.7** 1 year post PPM; uneventful to date. Presents with recurrent syncope. All but which of the following **could** be responsible:

1. Exit block
2. Threshold  $\uparrow$  secondary to medications
3. Lead dislodgement
4. Complete break of the conductor coil

21

## 1 year post PPM; uneventful to date

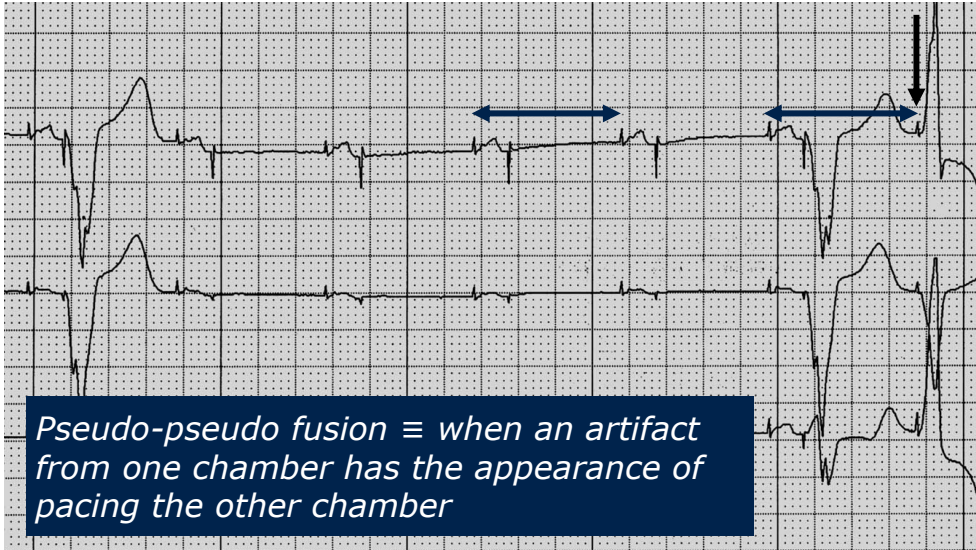
Presents with recurrent syncope. All but which of the following could be responsible:

1. Exit block - *> 1 month post-implant with failure to capture is compatible with exit block; would not likely occur at 1 year; usually earlier*
2. Threshold  $\uparrow$  secondary to medications - *failure to capture is compatible*
3. Lead dislodgement - *compatible with failure to capture*
4. Complete break of the conductor coil - *with complete transection, current would not get through and no artifact would be seen*

22

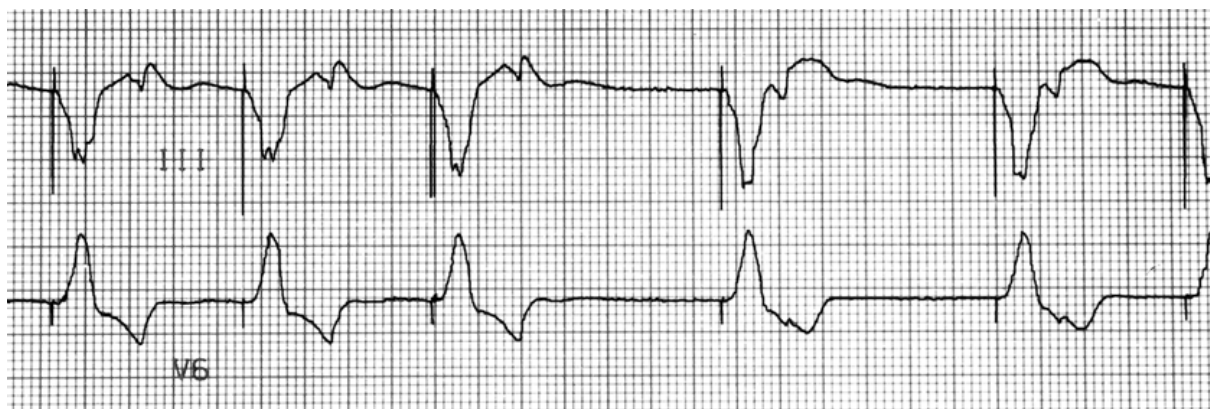
22

## What is occurring at the arrow?



**BIOTRONIK**  
excellence for life

23

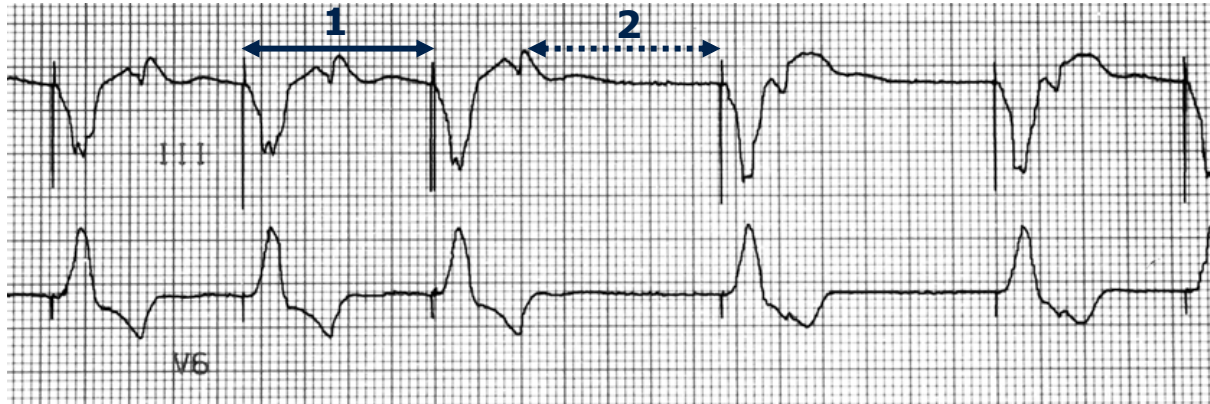


### 3.8 What is your ECG diagnosis?

1. Hysteresis
2. Over-sensing retrograde events
3. Fallback behavior
4. Normal sensor-driven pacing

**BIOTRONIK**  
excellence for life

24



### What is your ECG diagnosis?

1. Hysteresis
2. Over-sensing retrograde events
3. Fallback behavior
4. Normal sensor-driven pacing

25

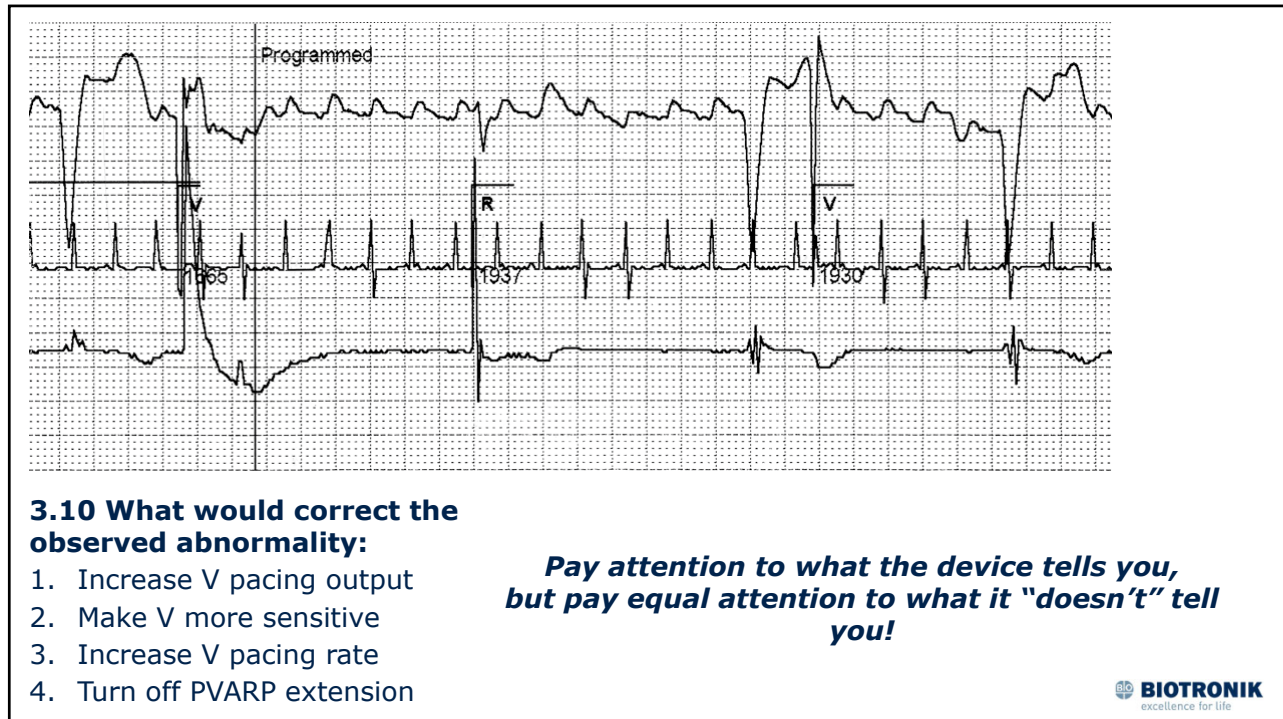


### 3.9 The ECG is obtained the morning after pacemaker implant. Which of the following is the most likely problem?

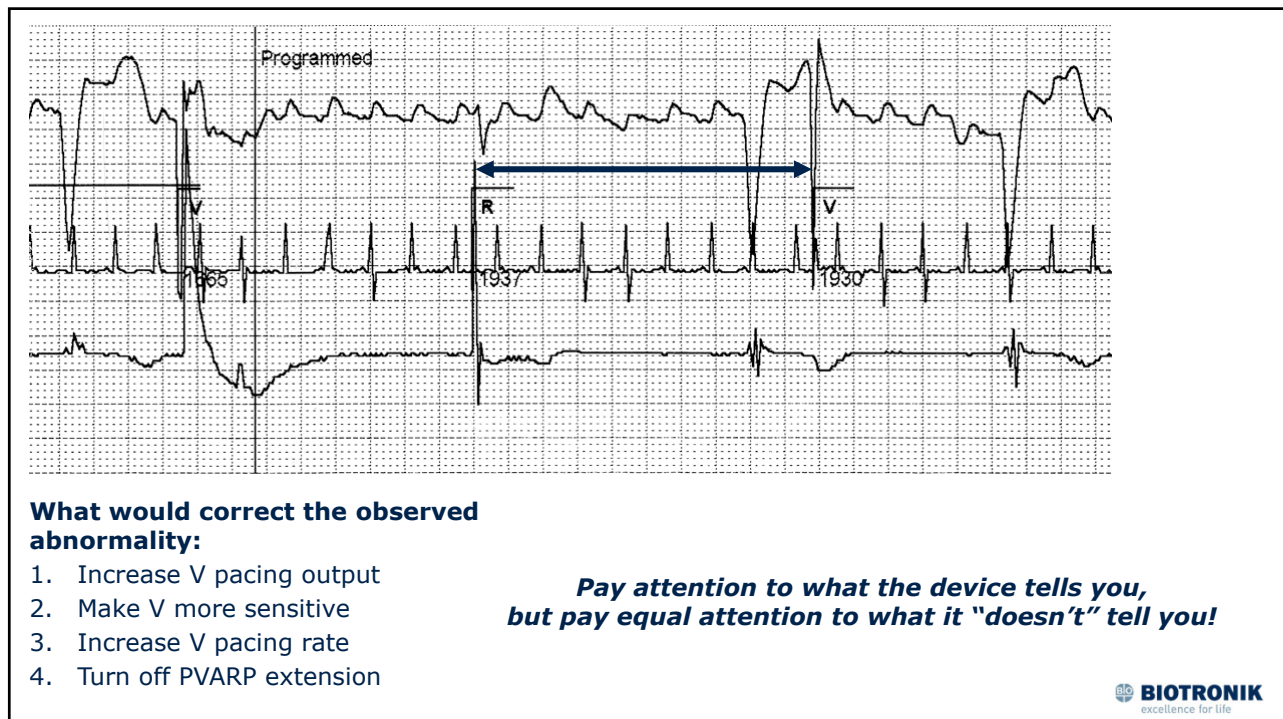
1. Crosstalk in absence of safety pacing
2. Ventricular lead dislodgment
3. Artifact
4. Ventricular lead fracture
5. Myopotential over-sensing

26

26



27



28

**3.11 The programmed P-AVI is 240 ms. Labeled QRS complex (\*) occurs in:**

1. Crosstalk sensing window
2. Post-Atrial Ventricular blanking period
3. Alert window

29

If the event occurs in the crosstalk sensing window is sensed, the V is paced after an abbreviated AVI (light orange), falling in the physiologic refractory period, ie, does not fall in the vulnerable period

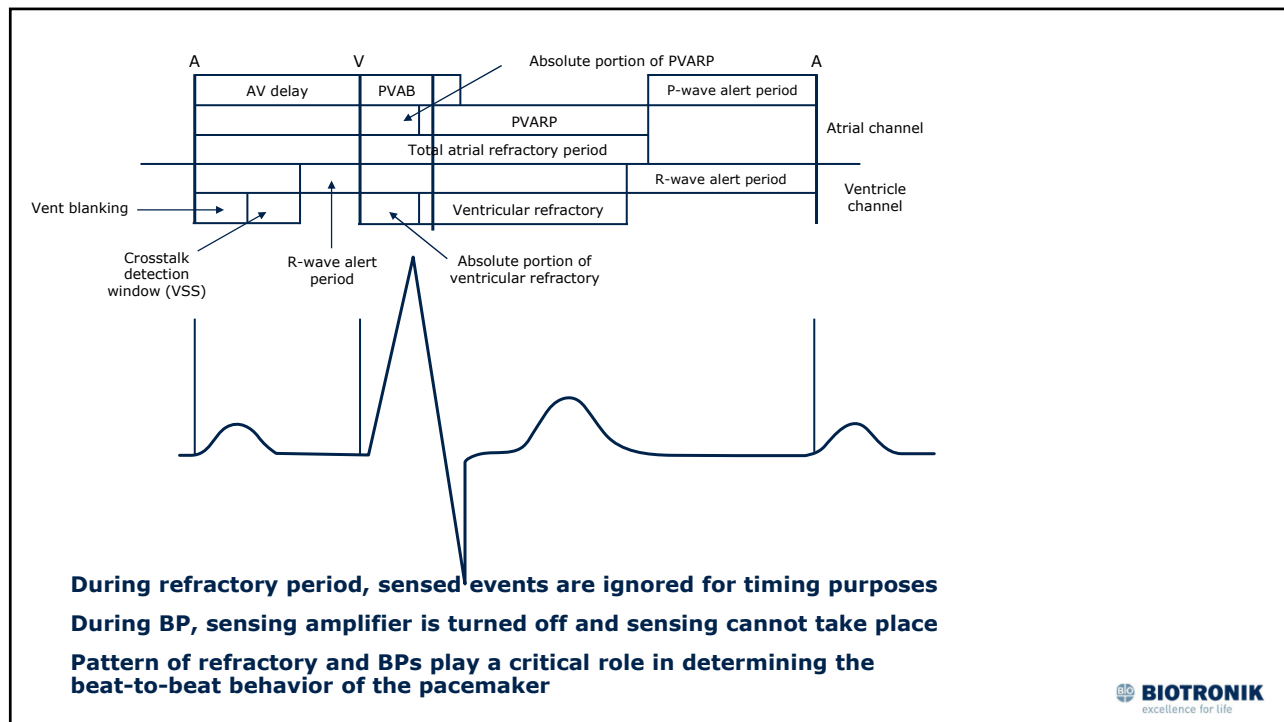
If the event occurs in the post-atrial ventricular blanking period, the sensing circuit is turned 'off' and the event is not seen (dark orange)

Blanking Period  
Crosstalk Sensing Window  
AVI  
Alert Period

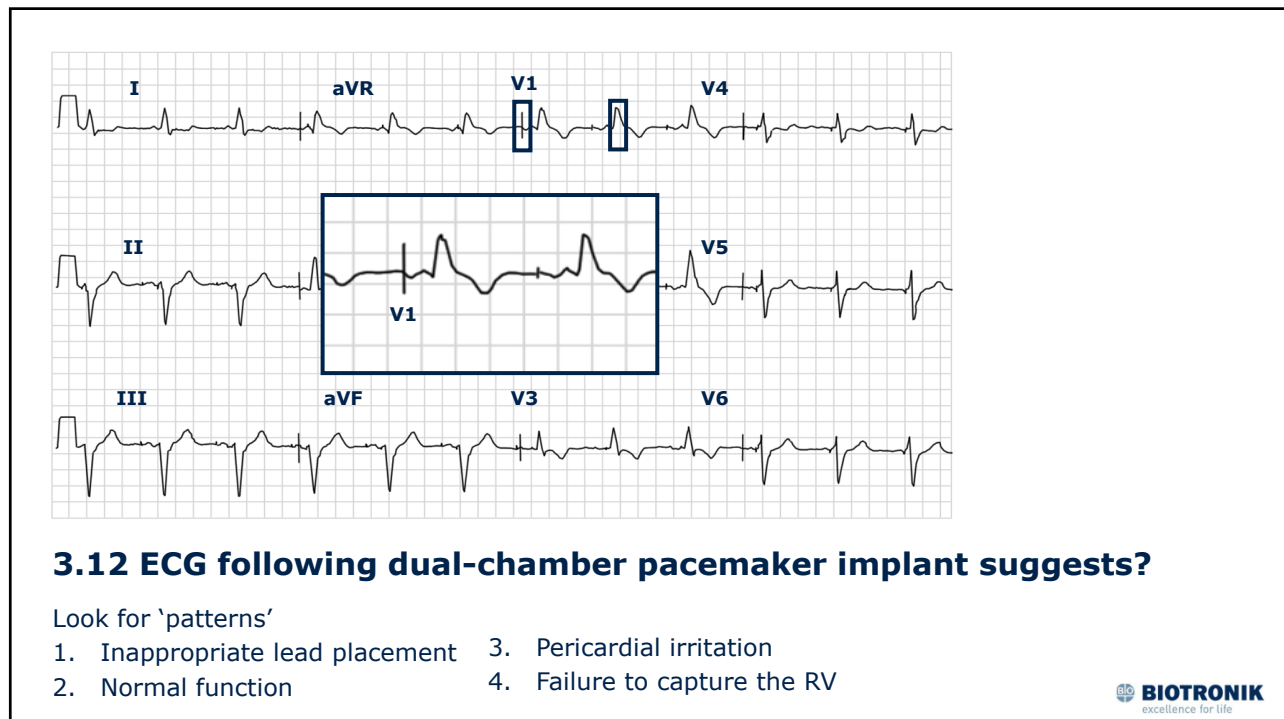
0 25 50 75 100 125 150 175 msec

If sensing occurs during the 'alert' period, ventricular output is inhibited.

30

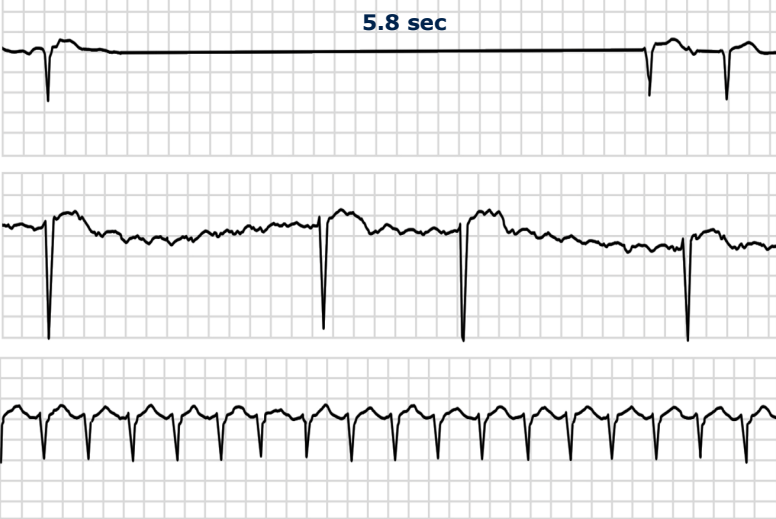


31



32





**3.13 68 year old female in hospital; 3 rhythm strips collected within a 24-hour period. You would...**

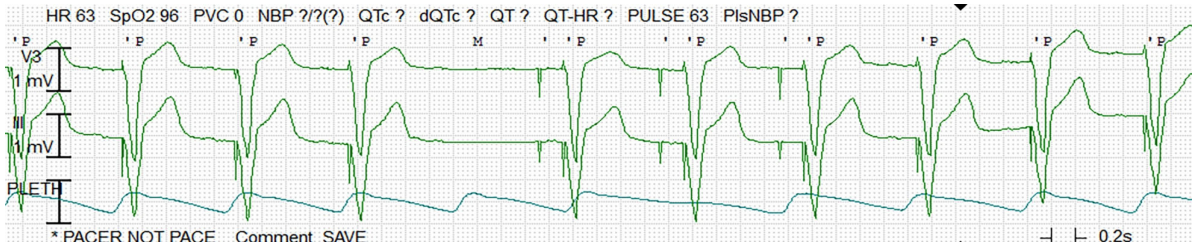
1. Start amiodarone
2. Implant ICD
3. Implant PPM
4. Start beta-blocker

33

**BIOTRONIK**  
excellence for life

33

**Rhythm strip obtained the afternoon after the pacemaker implant. Patient asymptomatic. RNs call pacemaker service.... Generic diagnosis?**



HR 63 SpO2 96 PVC 0 NBP ?/?(?) QTc ? dQTc ? QT ? QT-HR ? PULSE 63 PIsNBP ?

\* PACER NOT PACE Comment: SAVE

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

34

**BIOTRONIK**  
excellence for life

34

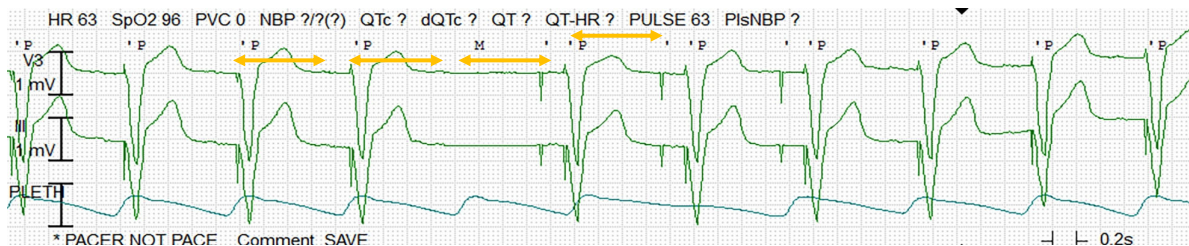
## Failure to Output

- Oversensing
  - Crosstalk
  - EMI
- Battery failure
- Circuit failure
- Lead fracture
- Internal insulation failure
- Loose set-screw
- Incompatible lead/header
- Device nuance - peculiarity

35

35

**If intervals are regular and baseline steady, less likely to be over-sensing.**



36

36

## Failure to Pace (No output)

- ~~Oversensing~~
  - ~~Crosstalk~~
  - ~~EMI~~
- ~~Battery failure~~
- ~~Circuit failure~~
- ~~Lead fracture~~
- ~~Internal insulation failure~~
- ~~Loose set screw~~
- ~~Incompatible lead/header~~
- Device nuance - peculiarity

37


**BIOTRONIK**  
 excellence for life

37

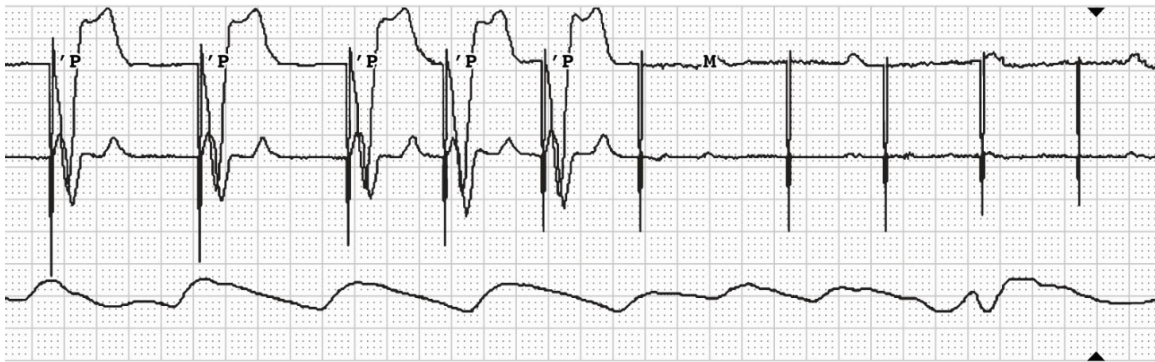
## 3.12 83-Year-Old Male with Increasing Dyspnea on Exertion

- History of coronary artery disease status post stent placement x2
- Third-degree AV block, status post pacemaker placement 8 years earlier (Medtronic dual-chamber Kappa KDR 901, atrial lead 5568, ventricular lead 4076)
- Programmed DDDR, lower rate 60 bpm, upper rate 130 bpm

38


**BIOTRONIK**  
 excellence for life

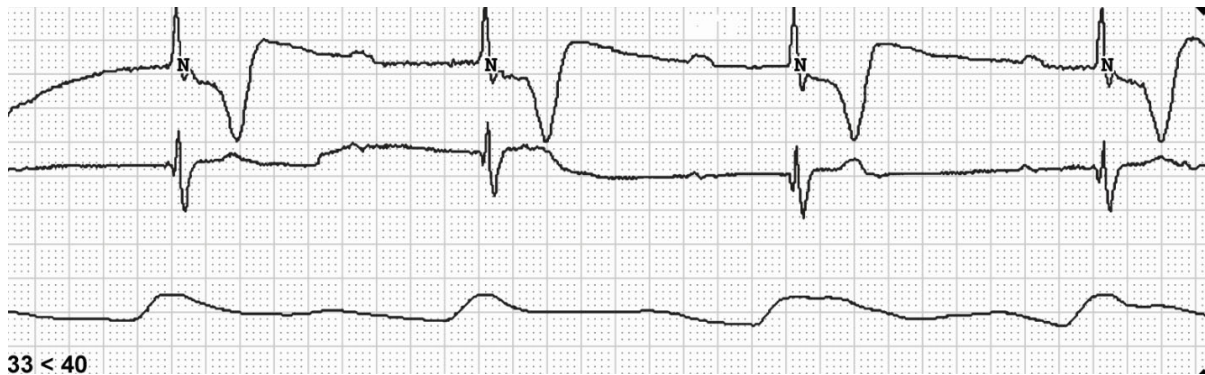
38



Prior to pacemaker interrogation, rhythm was ventricular pacing at 65 bpm. This tracing obtained when the programming wand is placed on the pacemaker. Tracing can be explained by:


1. Normal magnet function for this pacemaker
2. ERI (Elective replacement indicator)
3. EOS (End of service)
4. Ventricular lead loose in header

39



40

40

Pacemaker Model: Medtronic.Kappa KDR901	Medtronic.Kappa 900 S
Serial Number:	Copyright © Medtronic
<b>Initial Interrogation Report</b>	
<b>Patient/Device Information</b>	
Dependency:	Physician Name:
Implanted Defibrillator?:	Physician Phone:
Pacemaker Model: Kappa KDR901 PKM418309	Implanted: 07/17/15 6:49 PM
Atrial Lead:	
Ventricular Lead:	
<b>Pacemaker Status</b>	
Estimated remaining longevity: Replace Pacer	
Battery Voltage/Impedance	2.06 V / 32,125 ohms
	<b>Ventricular</b>
Amplitude/Pulse Width	2.76 V / 0.40 ms
Sensitivity	2.00 mV
Measured Impedance	680 ohms
Lead Status	Polarity Switch
<b>Clinical Status:</b>	
Diagnostic data not available.	
41	

41

## 83-Year-Old Male with Increasing Dyspnea on Exertion

- Battery voltage of 2.06 is compatible with EOS (EOL), at which point no reliable pacing occurs and diagnostic information is unreliable. It is difficult for manufacturers to give a single specific voltage at which EOS occurs because of multiple variables that may be present
- For this old Medtronic device, the company states that a measured battery voltage of 2.2 is a reasonable value to consider for EOS, but some devices may reach EOS at a higher value, and some have been identified at <2.0 V before
- Battery impedance is often ignored. In this case, the battery impedance was 32,125 ohms. As a rule, if battery impedance is  $\geq 10,000$  ohms, there should be a heightened concern that battery is approaching EOS. If battery impedance is  $\geq 20,000$  ohms, there is a high likelihood the device is at or near EOS.

| 42 |  |

42

## Basics & Beyond

David Hayes, M.D. and Beth Davenport, MSN, RN Co-Directors

**February 28<sup>th</sup> 3:00-5:00 PST (6:00-8:00 EST)**

- Pacemaker Troubleshooting
- Rate Response and CLS
- Conduction System Pacing Overview



43

## Paced ECG Interpretation: A Case-Based Approach

David Hayes, MD



© 2024 BIOTRONIK, Inc. All rights reserved. PPT745r1 1/17/2024



44