# **Medical Cyber-Physical Systems**

### **Electrophysiology** basics

Lecture 10

Principles of Modeling for Cyber-Physical Systems

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Many thanks to: Zhihao Jiang, Houssam Abbas, and Rahul Mangharam, For help with preparing this module.

# Why explore cardiac modeling?

Cardiac disease is the **leading cause of death in the US** 

Around the world, **17.5 million people** die of Cardiovascular Diseases (CVD) yearly That's an estimated **31%** of all deaths

More than 75% of CVD deaths occur in low income and middle income countries Implanted devices are a leading method of treating some CVDs

# Why study cardiac devices?

These devices are life-critical  $\rightarrow$  must function correctly

Are constrained in their energy consumption  $\rightarrow$  must be low-power

Are implanted in the body  $\rightarrow$  very special design considerations (e.g., materials used, must be ex-plantable...)

Are regulated by the FDA  $\rightarrow$  must follow certain best practices, but also have some inertia

These are life-critical embedded systems

# Its shocking! Cardiac devices can have bugs











- Over 600,000 cardiac medical devices recalled from 1990-2000
   40% of which were due to software issues
- 2008-12: 15% of all the medical device recalls due to software



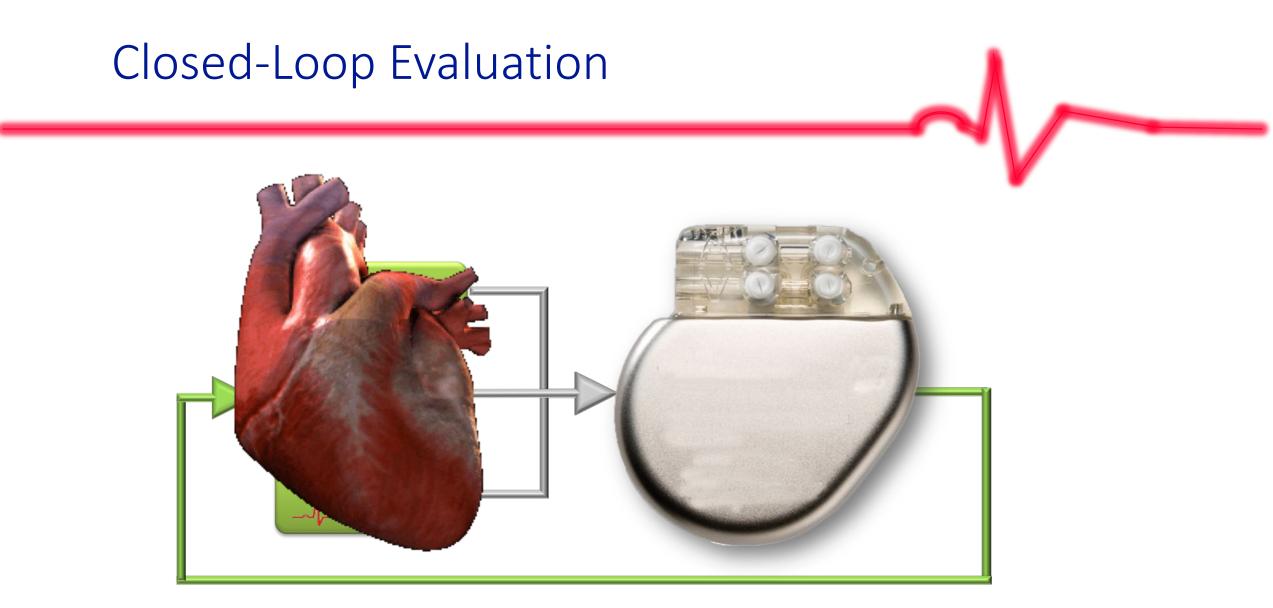


Implantable Cardioverter-Defibrillator (ICD)

#### The consequences of incorrect algorithms and implementations



Filmed and shared with patient's consent



Need a model of the heart which can capture the physiological conditions of the heart and respond to pacemaker outputs.

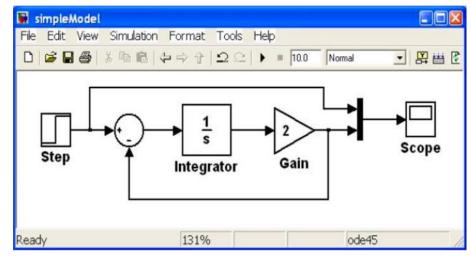
Model/implement the pacemaker

Check if pacemaker is 'safe' for different heart conditions

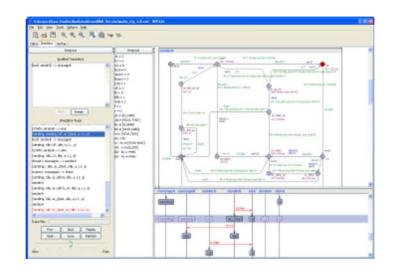
#### Let our heart (model) catch the bugs before your heart does

# This Module

- Will be challenging:
- New domain (electrophysiology)
- New tools (Simulink and UPPAAL)
- New theory (timed automata)
- New concepts (model checking)



Simulink



#### UPPAAL

# How do we go about understanding the heart?

Understand the electrical system as a circuit?

Understand the cellular activity?

Understand the molecular activity?

What does one heart tell us about other hearts?

What does a healthy heart tell us about an unhealthy heart?

What *is* a healthy heart?

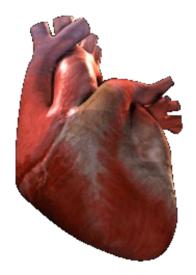
What is the purpose of our enquiry?

Need to understand the domain.

Speak the same language as the domain experts.

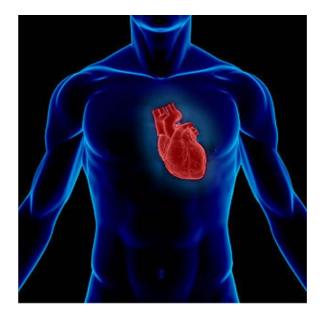
### Remember....modeling choices and 'usefulness' depend on the problem at hand.

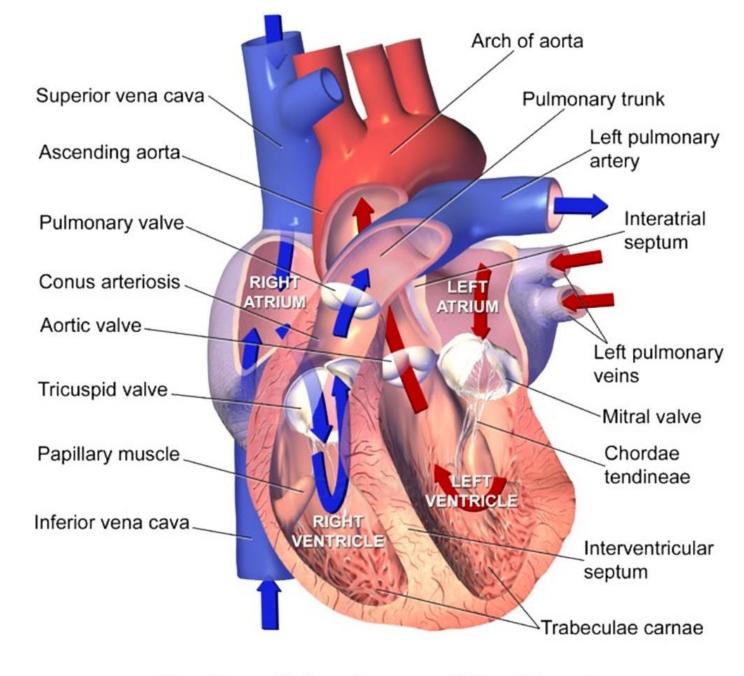
# Electrophysiology of the heart





#### First, let's examine the human heart



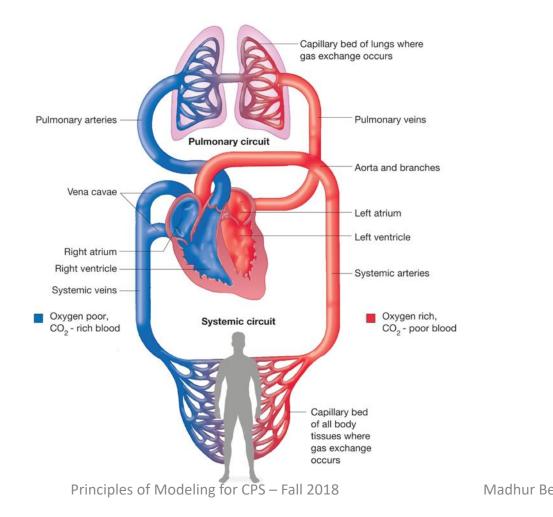


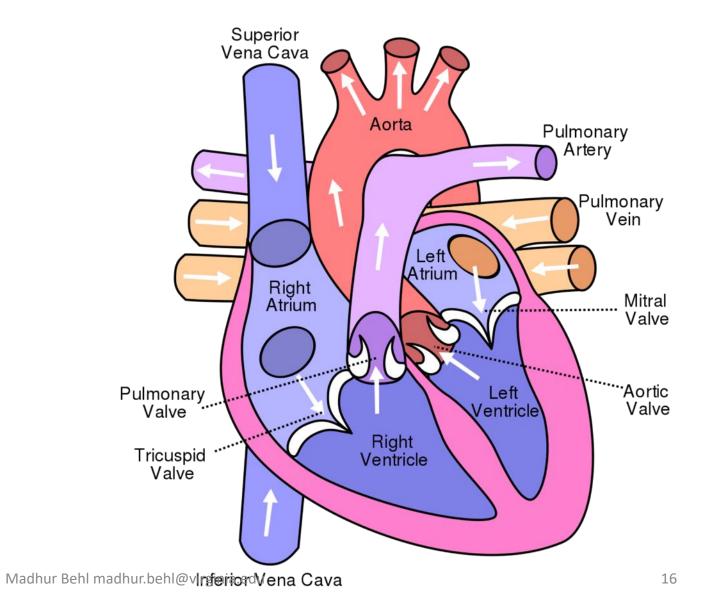
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#### **Sectional Anatomy of the Heart**

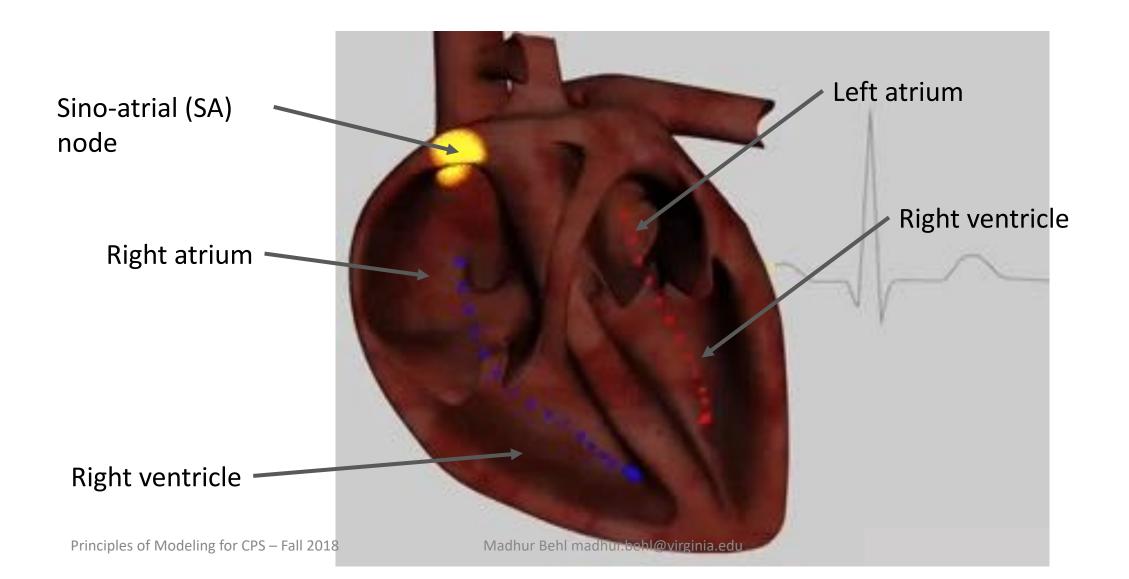
### Circulatory system and heart function

Vein: towards the heart. <u>A</u>rtery: <u>a</u>way from the heart

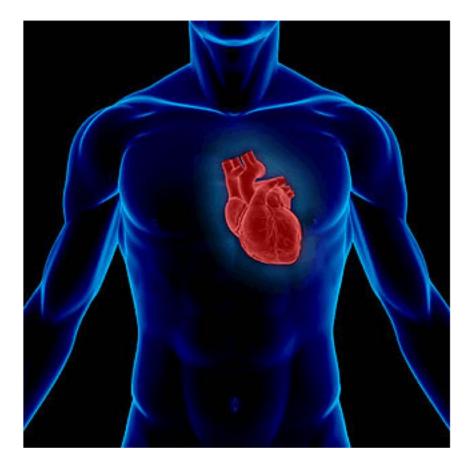


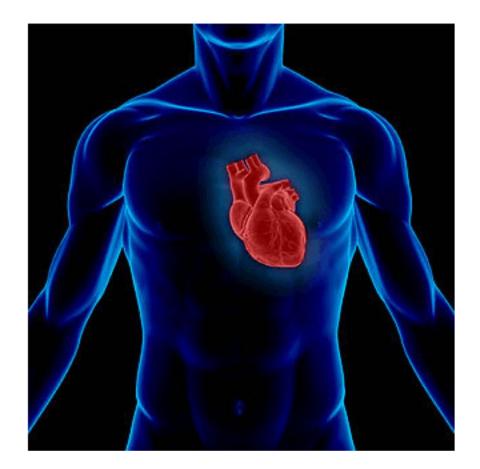


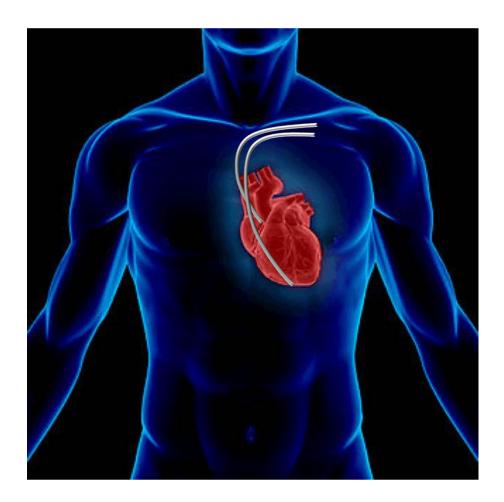
# Electrical generation and propagation



# Bradycardia: bradus (slow) + kardia (heart)





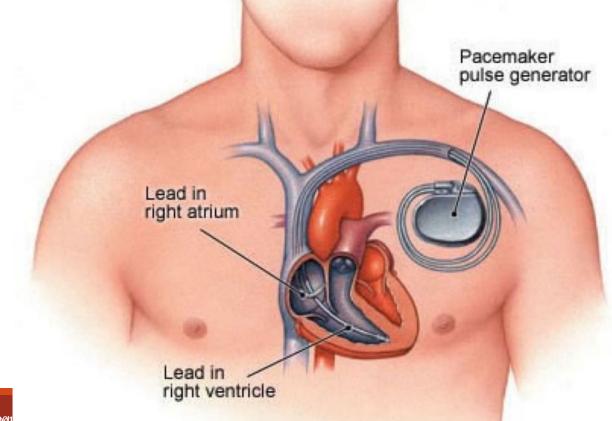


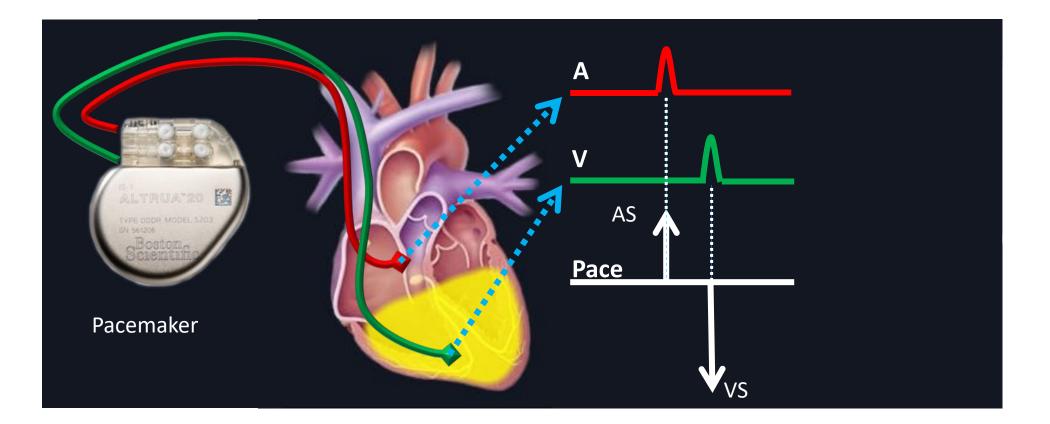


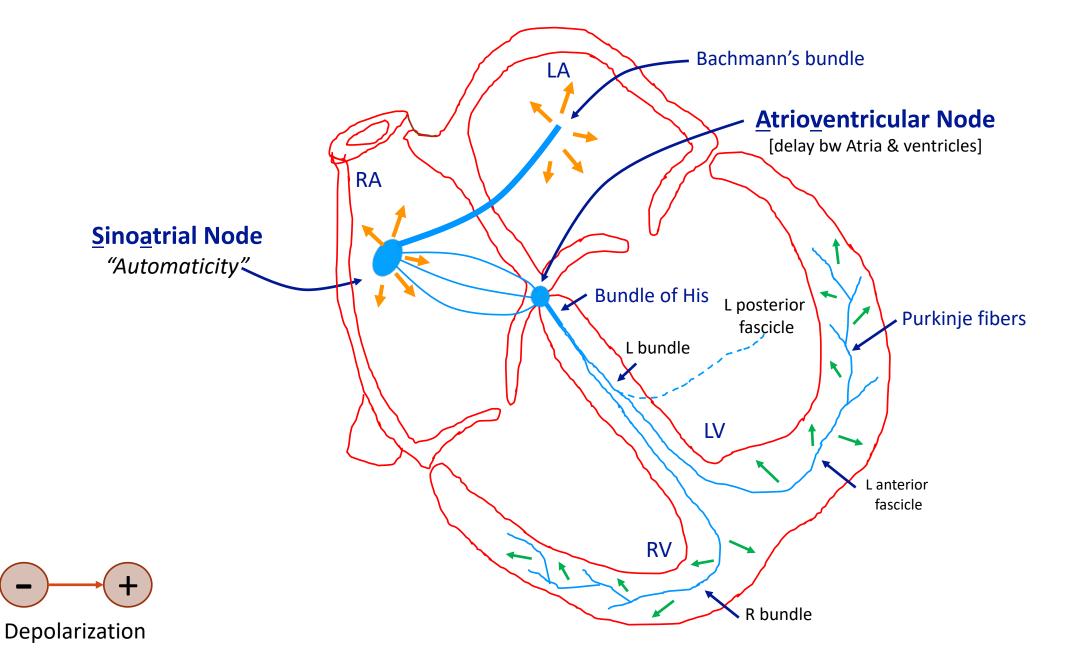
#### Two leads in heart chambers

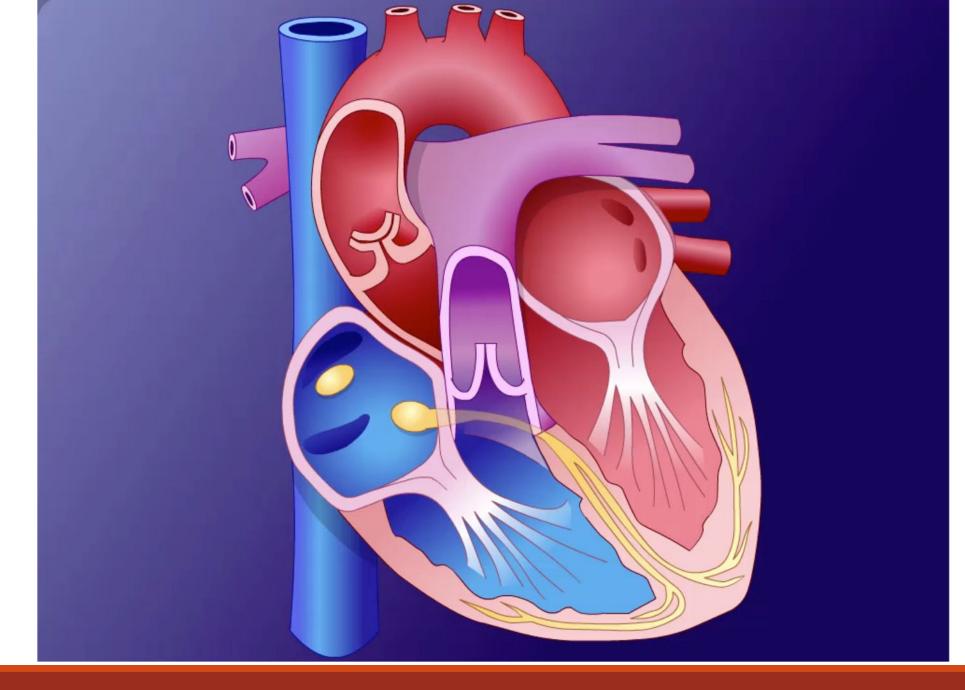
Two leads are placed in the right atrium and right ventricle

Monitors the local electrical activities of the heart and deliver therapy according to the **timing information** 





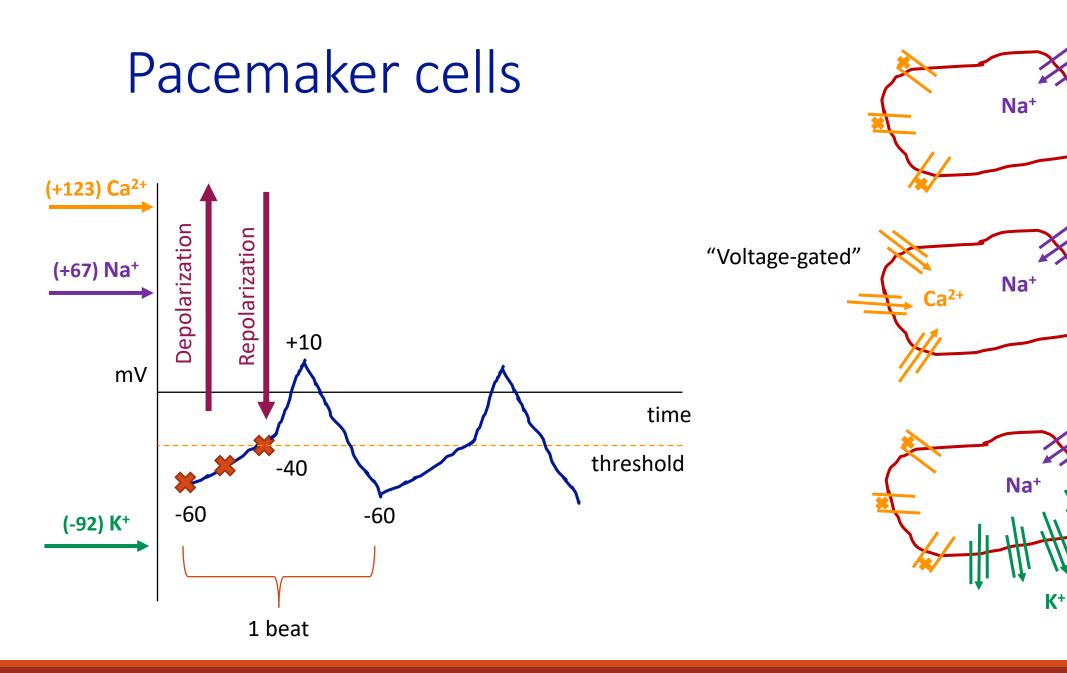


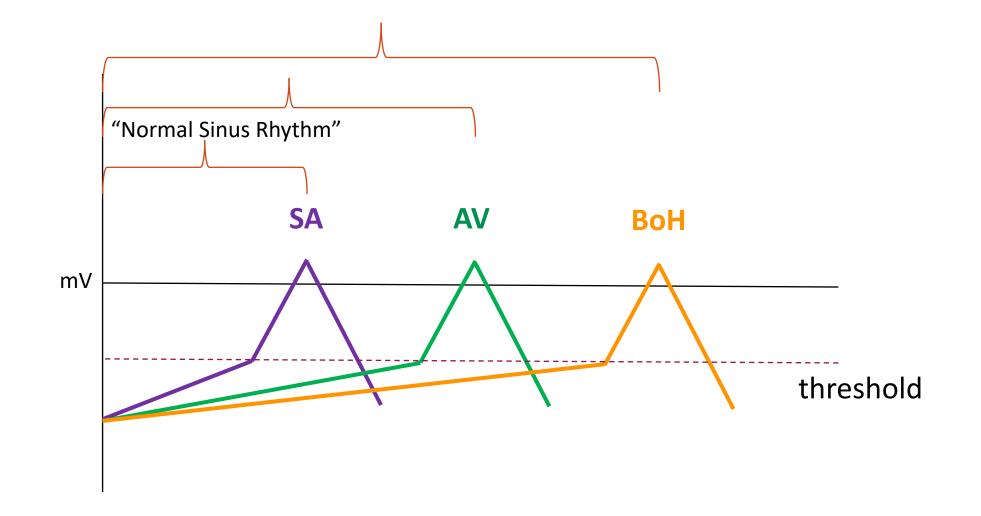


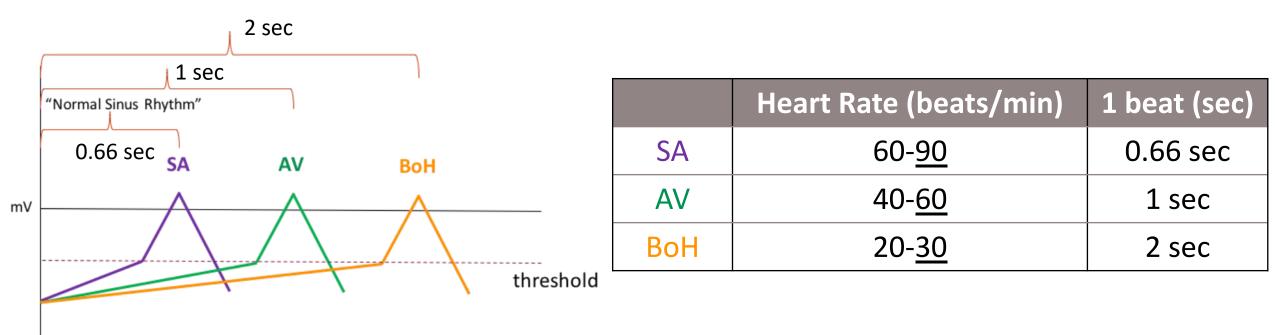
# Pacemaker cells (naturally pace the atria and ventricles) and Cardiac myocytes cells ("squeezing" the heart)

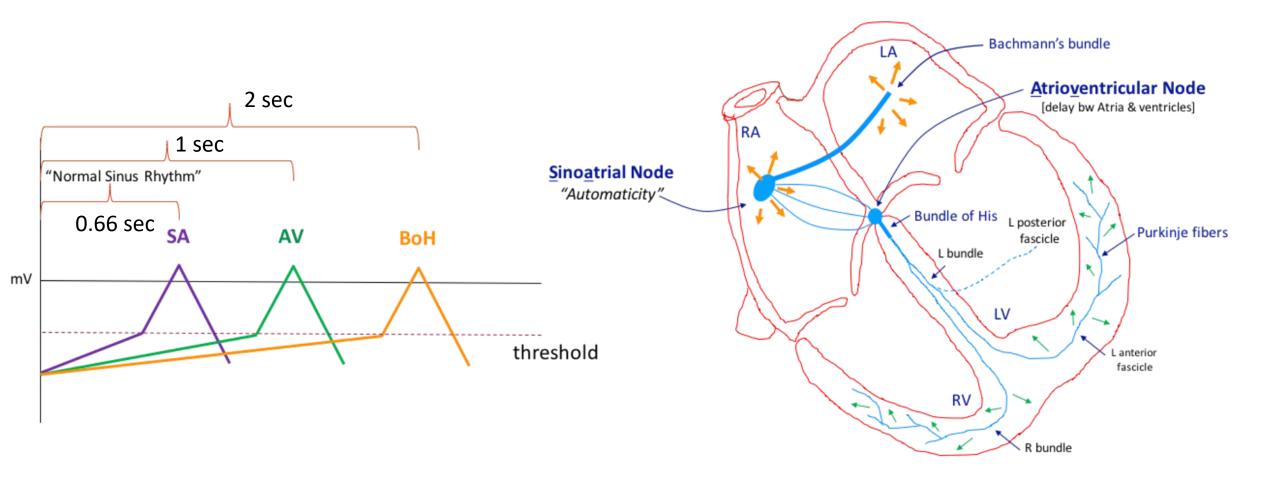
# Pacemaker cells – "Automaticity"

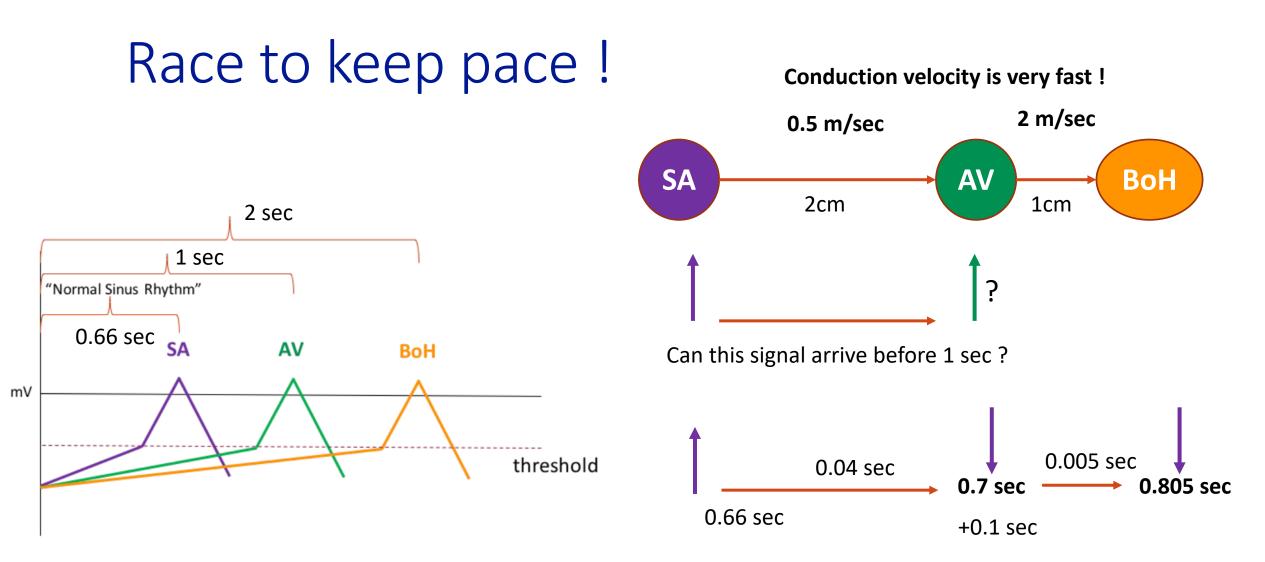
- 1) <u>Sino-Atrial</u> (SA) node
- 2) <u>Atriov</u>entricular (AV) node
- 3) Bundle of His / Purkinje fibers



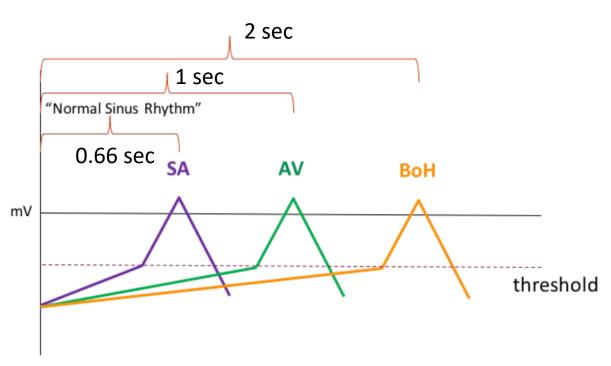




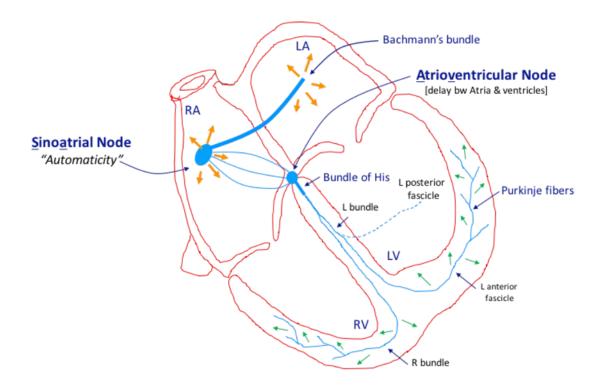




How does SA node control the heart beat ? "Normal Sinus Rhythm"



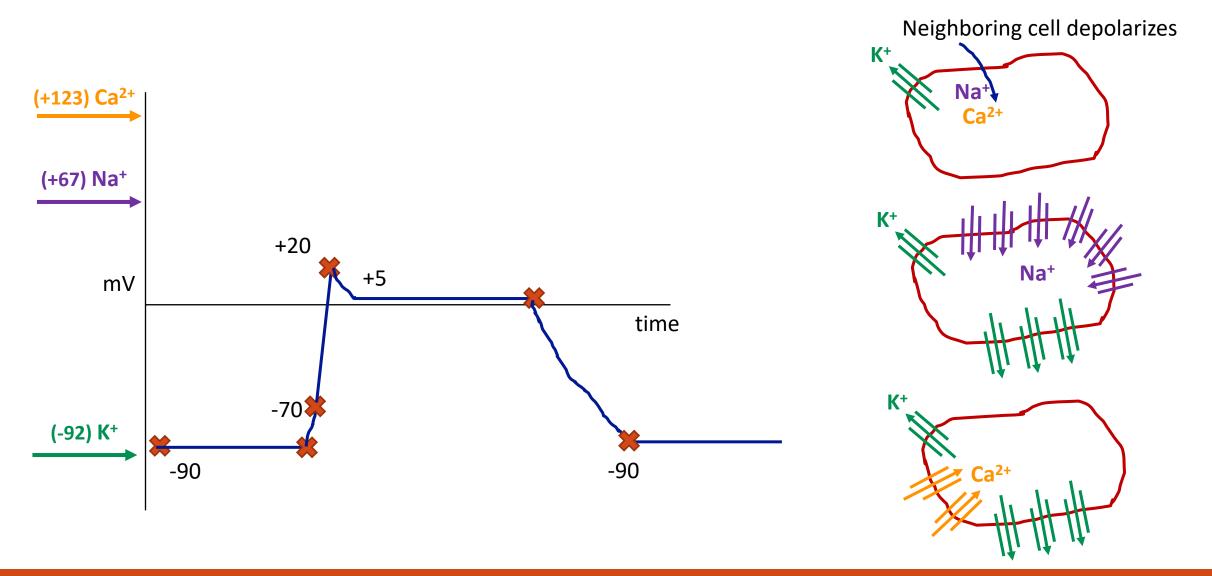
	Heart Rate (beats/min)	1 beat (sec)
SA	60- <u>90</u>	0.66 sec
AV	40- <u>60</u>	1 sec
BoH	20- <u>30</u>	2 sec



- SA node Plan A
- AV node Plan B
- BoH Plan C

# Pacemaker cells (naturally pace the atria and ventricles) and Cardiac myocytes cells ("squeezing" the heart)

# Cardiac Myocytes — How does the heart squeeze

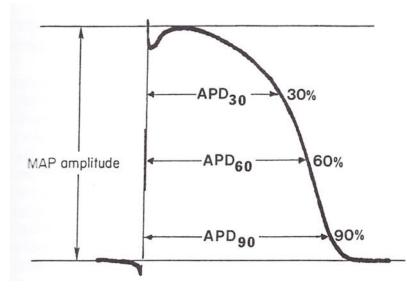


# Action Potential - Tissue

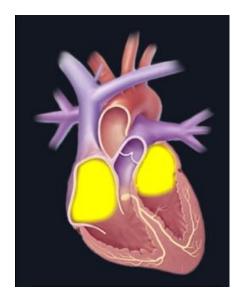
The elementary signal of the heart

It affects the *conduction velocity* and *signal passage* of the tissue.

Closely related to most cardiac diseases.



**APD:** Action Potential Duration



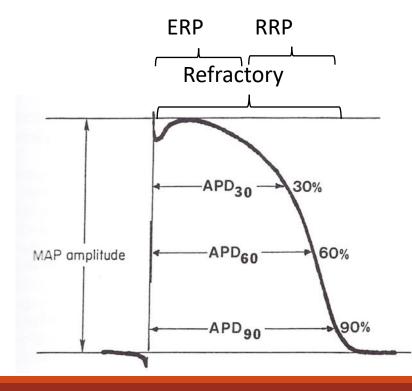
Functional syncytium

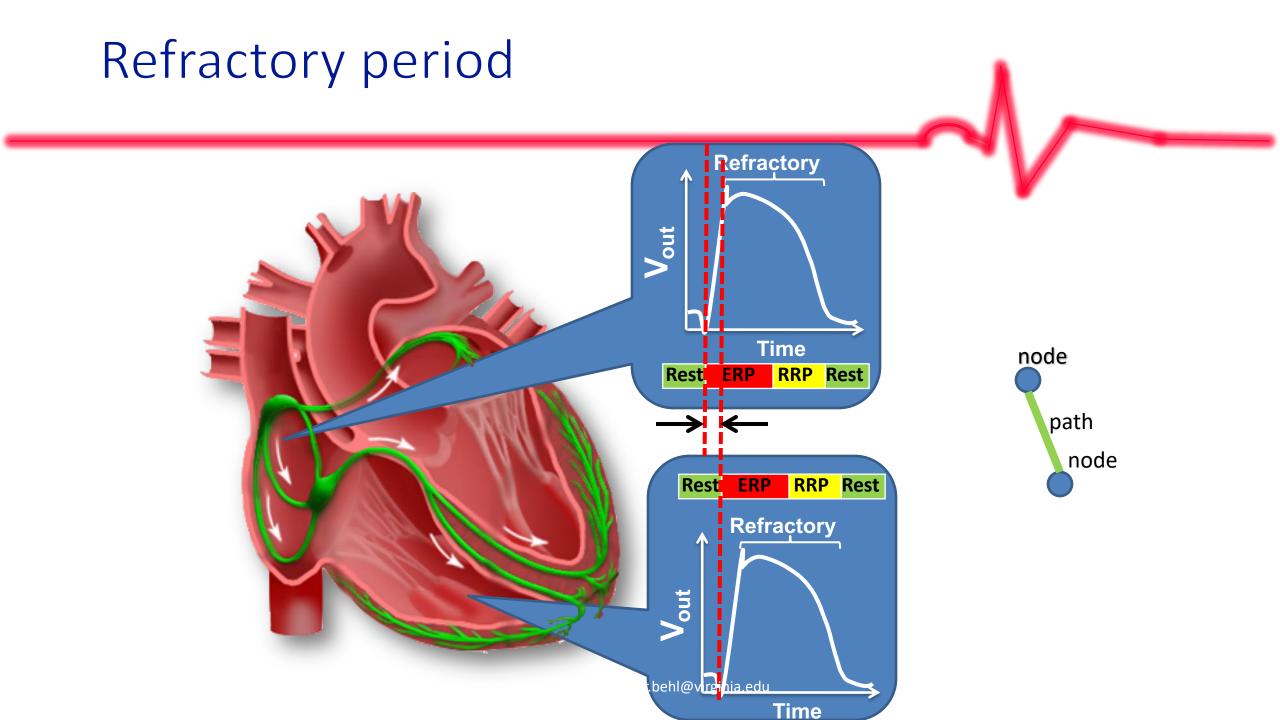
### Cellular Level - Action potential

lons refill

Divided into *Effective Refractory Period(ERP)* and *Relative Refractory Period(RRP)* for activation with certain strength

Block Interval during ERP, abnormal new action potential during RRP



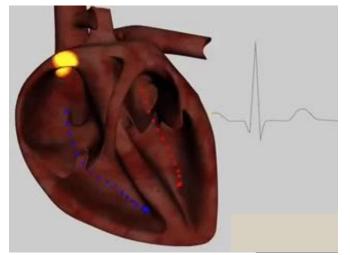


# Treating bradycardia

We want to detect when the atria or the ventricles miss a beat, and pace the chambers when that happens

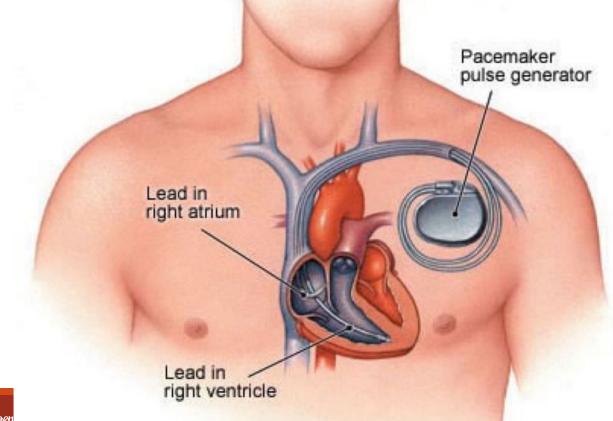
Start small and simple:

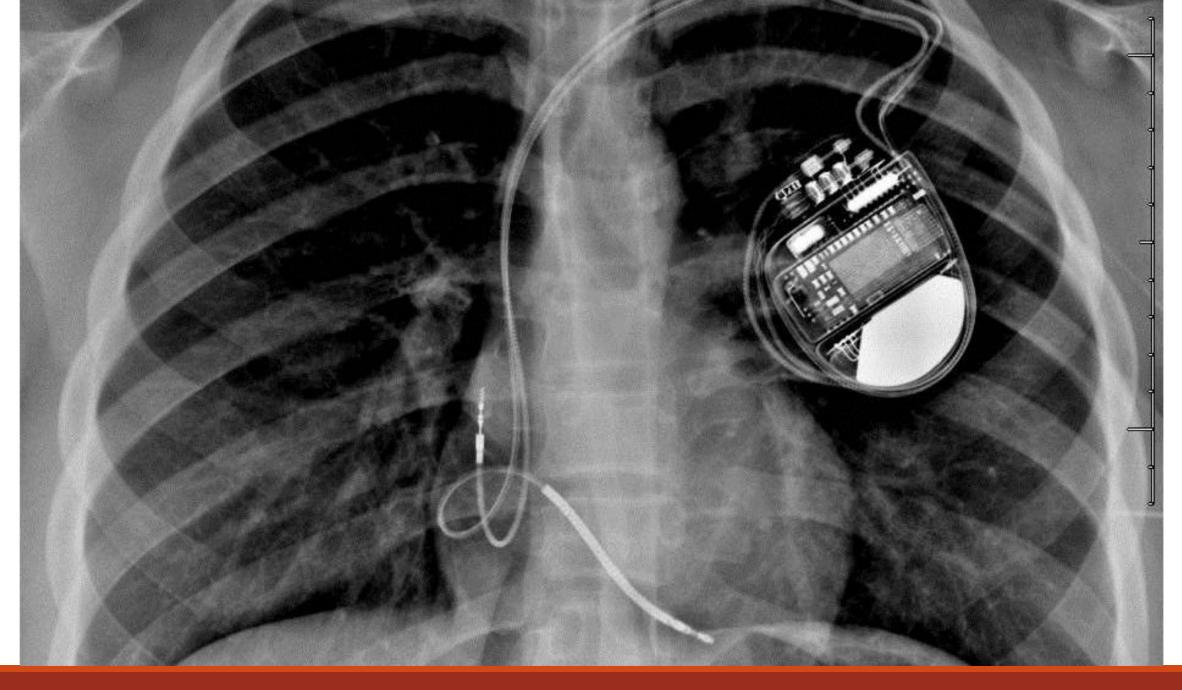
- We know that usually, the atria contract together, and the ventricles contract together → sense only in right atrium and ventricles
- Output of the standard structure
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Two leads are placed in the right atrium and right ventricle

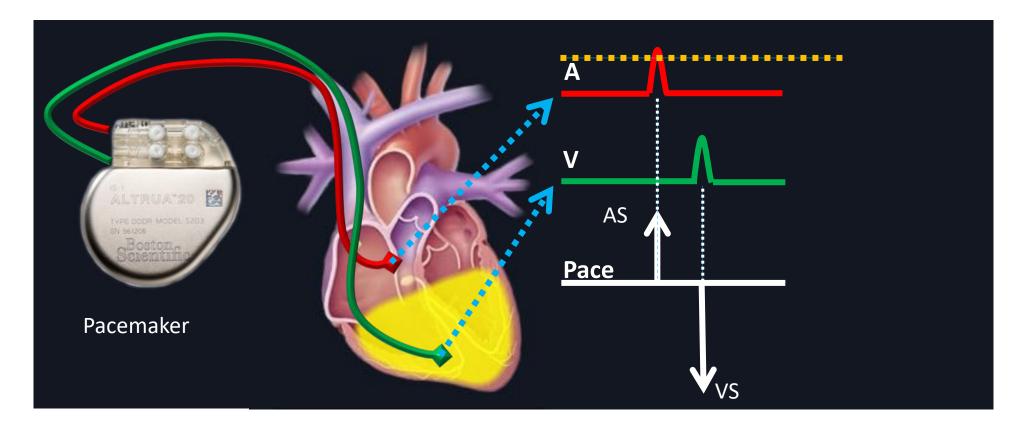
Monitors the local electrical activities of the heart and deliver therapy according to the **timing information** 





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#### Timing info for local activation



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- Simulink and State-flow tutorials have been posted on the course website.
- Not graded. No submission required.
- One week to brush up on Simulink/Stateflow.
- Go through the tutorials before the Simulink/Stateflow model walkthrough lecture next week

# Next Lecture:

- Pacemaker operation and heart conditions
- Model checking vs Model testing
- Heart modeling using timed automata