Medical Cyber-Physical Systems

Electrophysiology basics

Lecture 10

Principles of Modeling for Cyber-Physical Systems

Instructor: Madhur Behl

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 → 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







Implantable Cardiac Devices Recalls

- 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

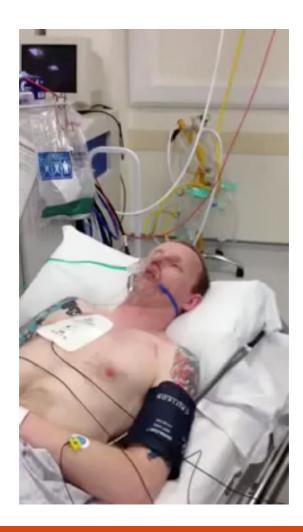






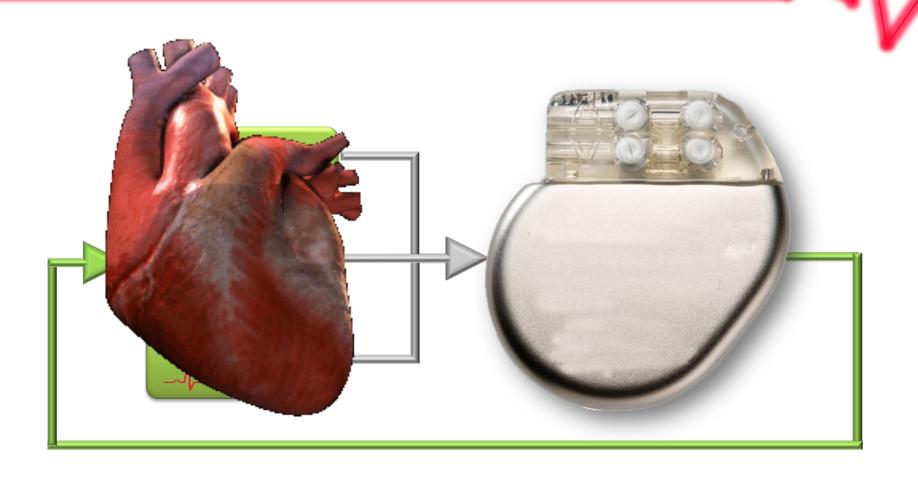
Implantable Cardioverter-Defibrillator (ICD)

The consequences of incorrect algorithms and implementations



Filmed and shared with patient's consent

Closed-Loop Evaluation



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

Model/implement the pacemaker

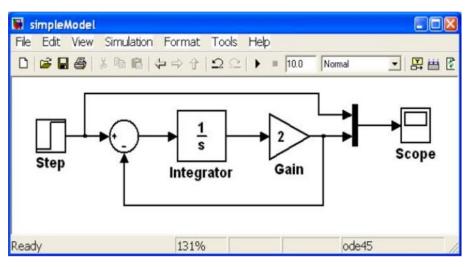
Check if pacemaker is 'safe' for different heart conditions

Let our	heart (mode	catch t	the b	ougs b	pefore '	your	heart	does
		•				,		

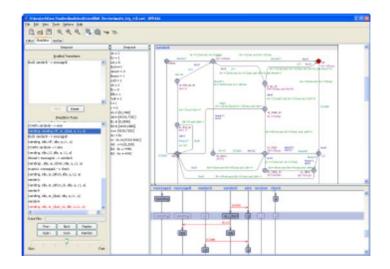
This Module

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?

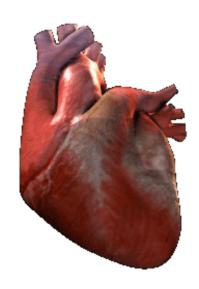
What is the purpose of our enquiry?

Need to understand the domain.

Speak the same language as the domain experts.

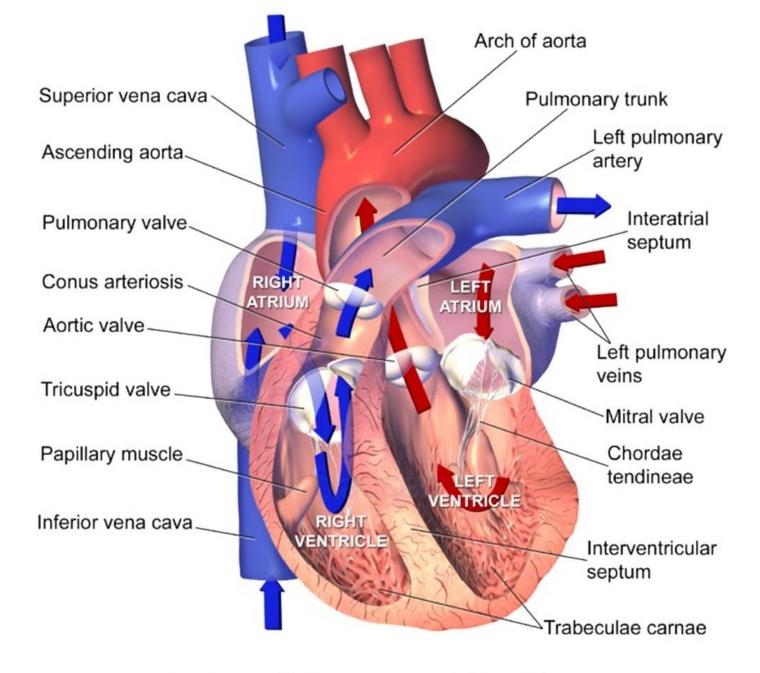
Remember....modeling choices and 'usefulness' depend on the problem at hand.

Electrophysiology of the heart



First, let's examine the human heart



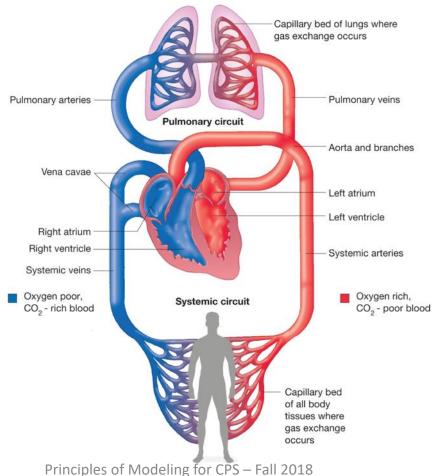


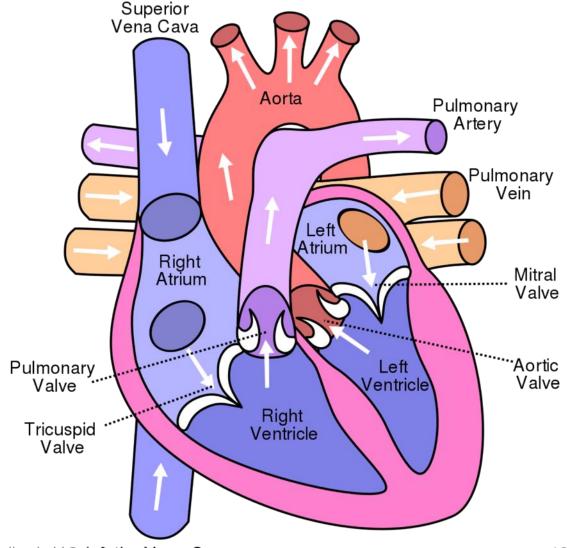
Sectional Anatomy of the Heart

Circulatory system and heart function

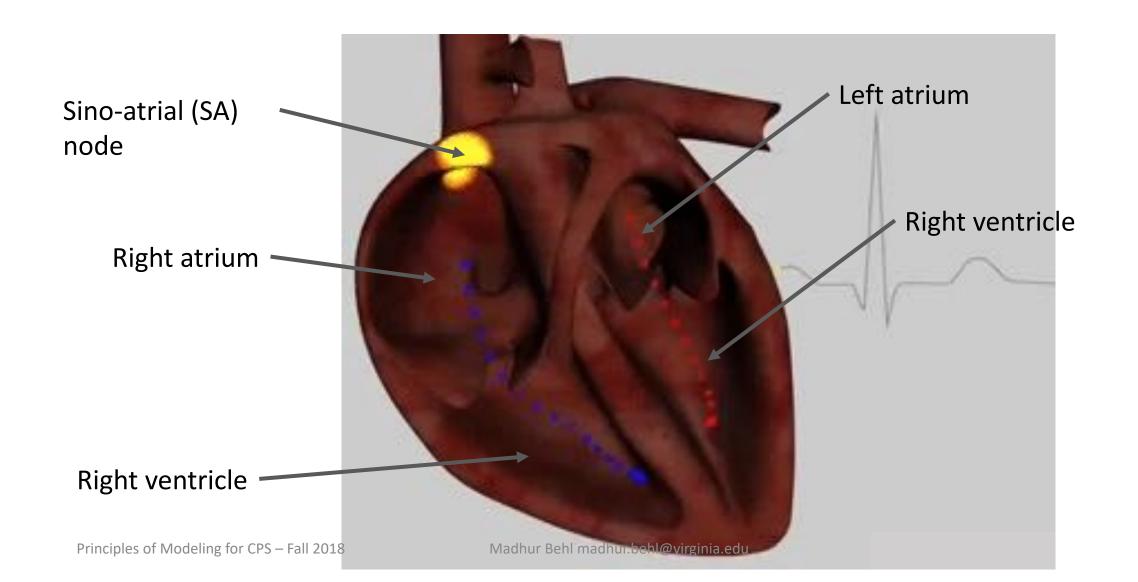
Vein: towards the heart.

Artery: away from the heart

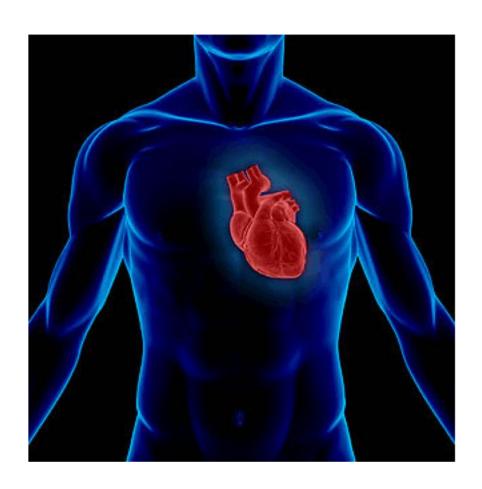


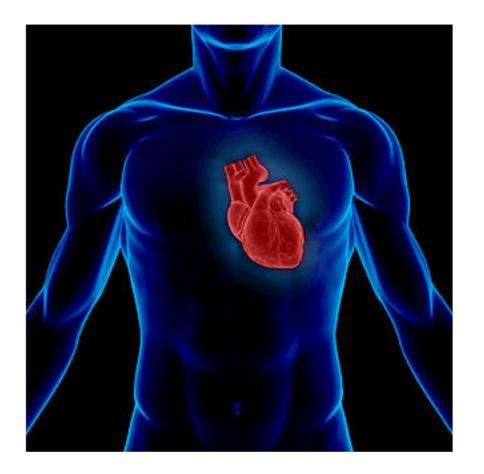


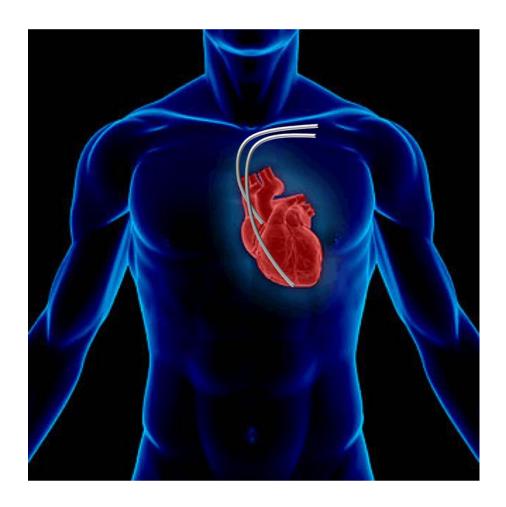
Electrical generation and propagation



Bradycardia: bradus (slow) + kardia (heart)







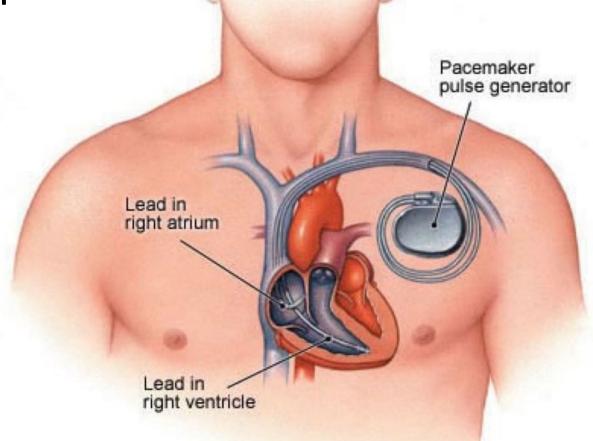


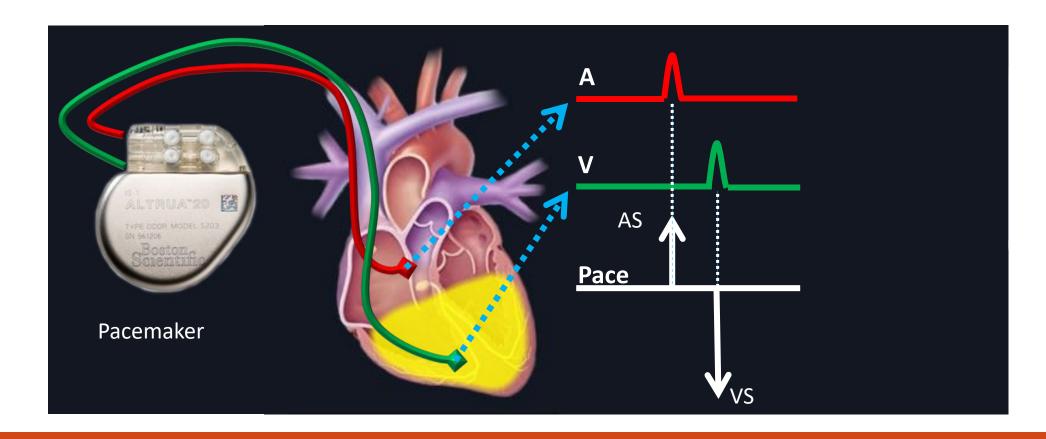
Two leads in heart chambers

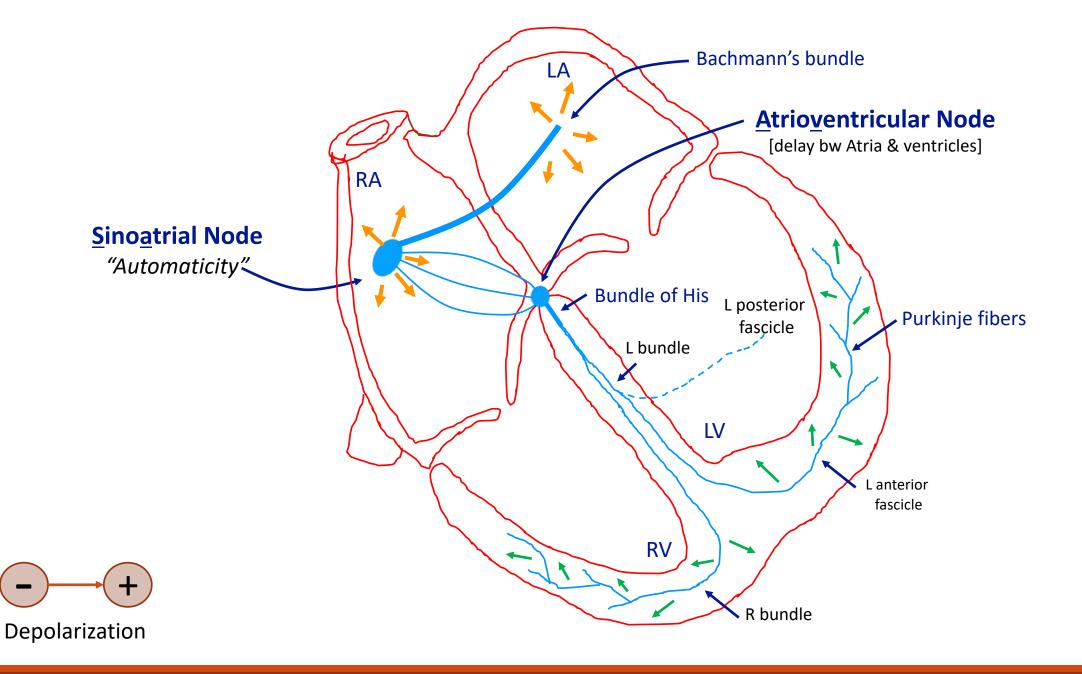
Two leads are placed in the right atrium and right ventricle

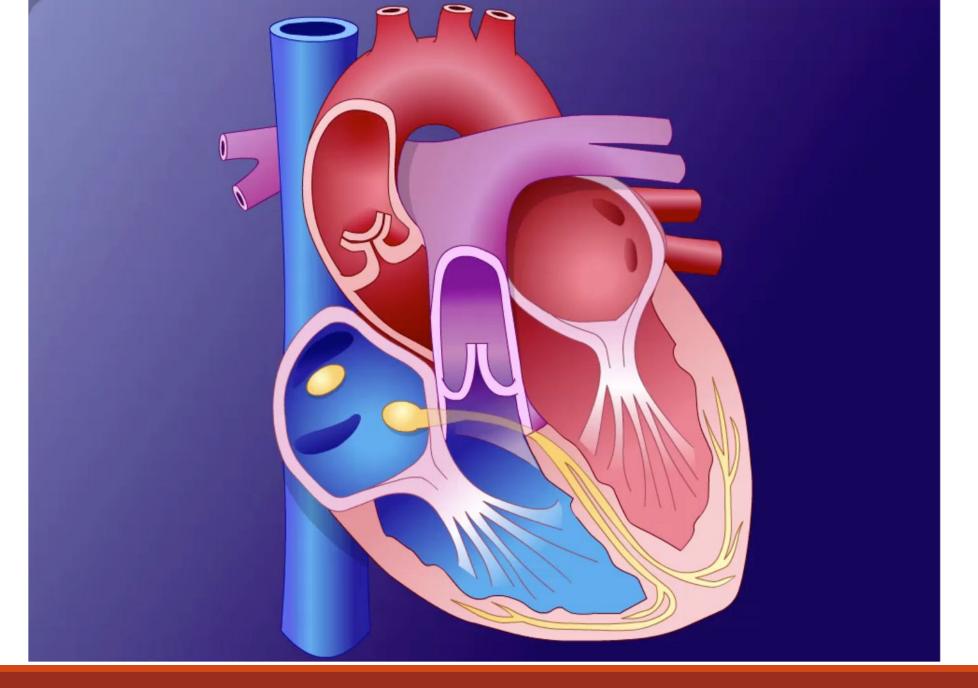
Monitors the local electrical activities of the heart and deliver therapy according to the

timing information









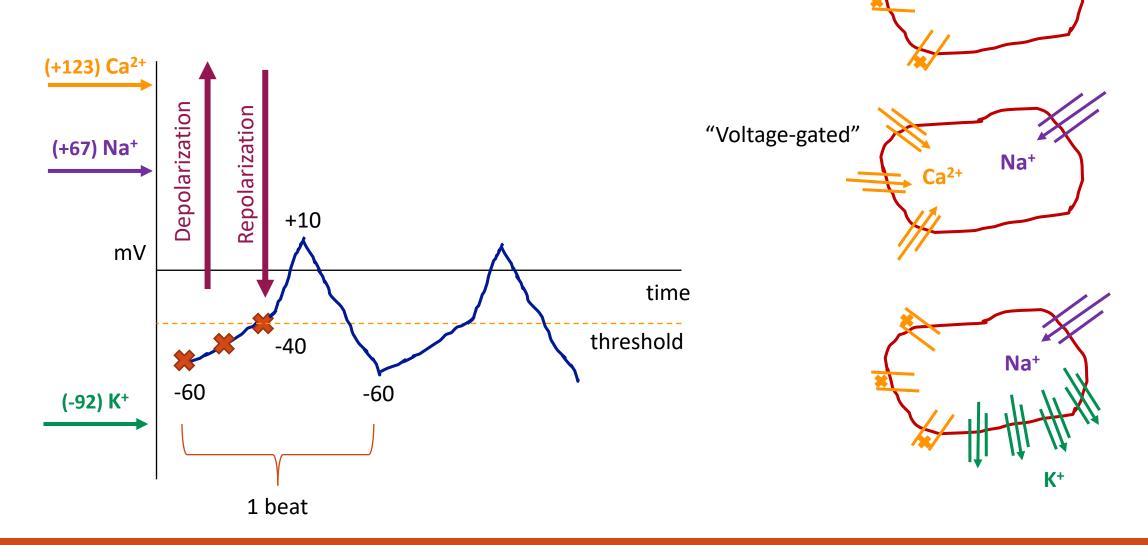
Pacemaker cells (naturally pace the atria and ventricles) and

Cardiac myocytes cells ("squeezing" the heart)

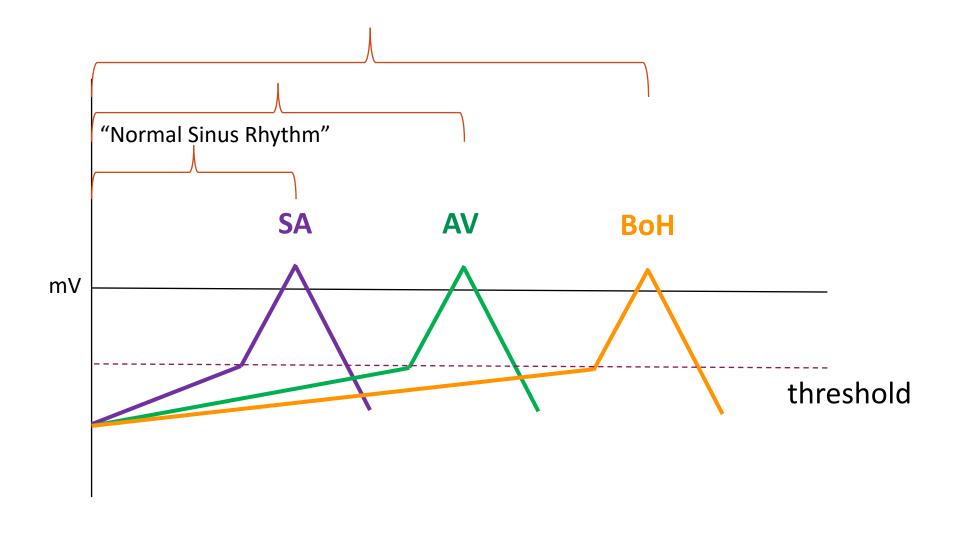
Pacemaker cells – "Automaticity"

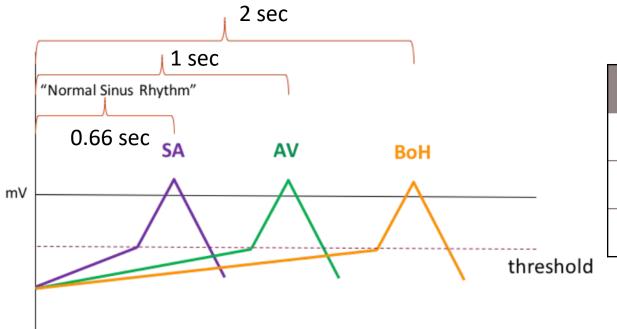
- 1) Sino-Atrial (SA) node
- 2) Atrioventricular (AV) node
- 3) Bundle of His / Purkinje fibers

Pacemaker cells

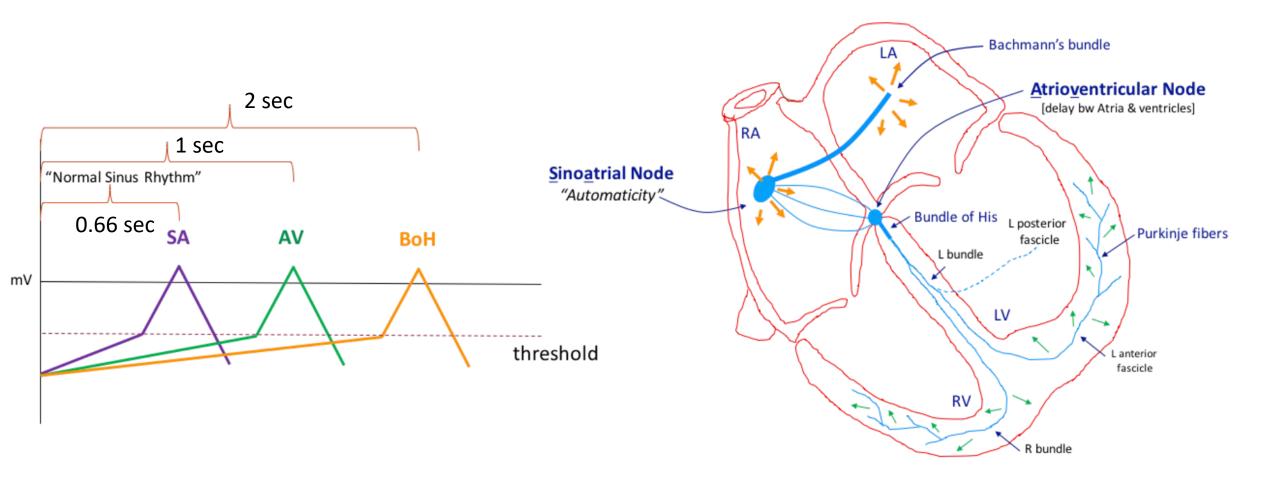


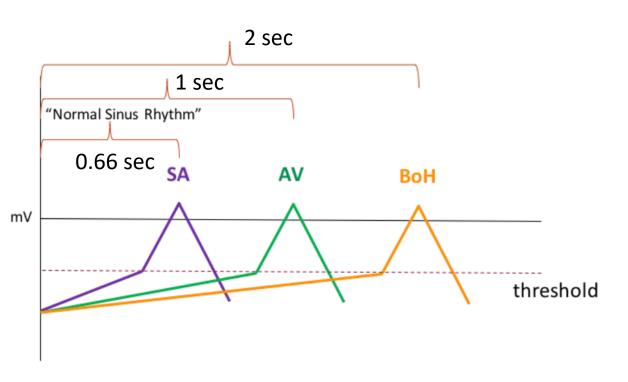
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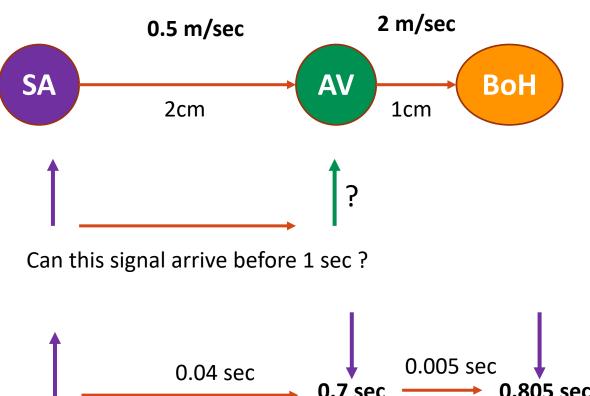
	Heart Rate (beats/min)	1 beat (sec)
SA	60- <u>90</u>	0.66 sec
AV	40- <u>60</u>	1 sec
ВоН	20- <u>30</u>	2 sec





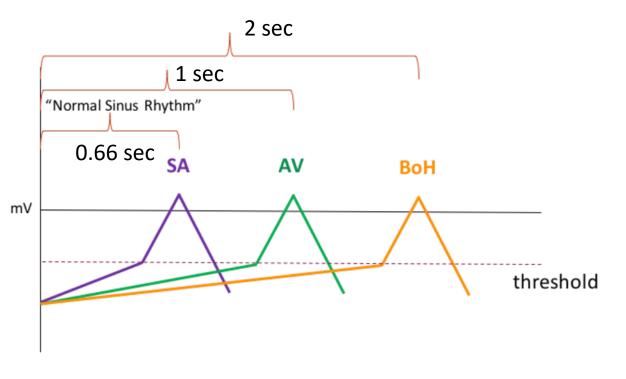
How does SA node control the heart beat ? "Normal Sinus Rhythm"

Conduction velocity is very fast!

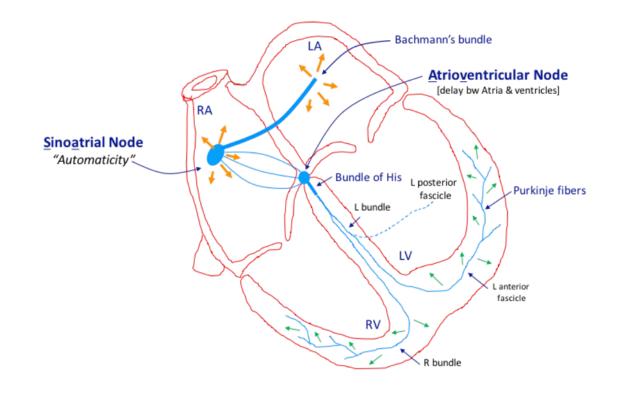


+0.1 sec

0.66 sec



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

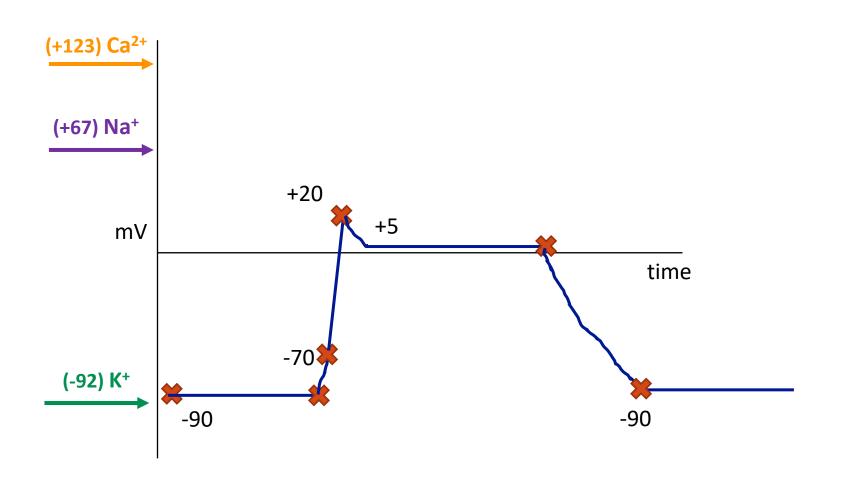


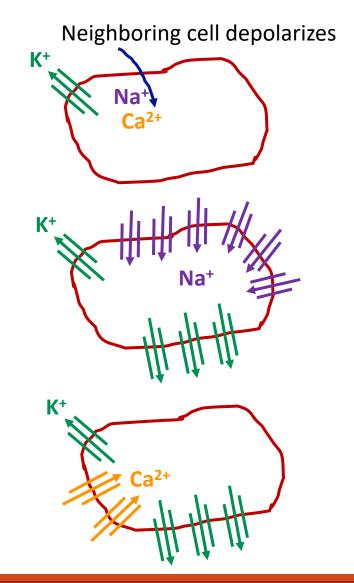
- SA node Plan A
- AV node Plan B
- BoH Plan C

Pacemaker cells (naturally pace the atria and ventricles) and

Cardiac myocytes cells ("squeezing" the heart)

Cardiac Myocytes — How does the heart squeeze



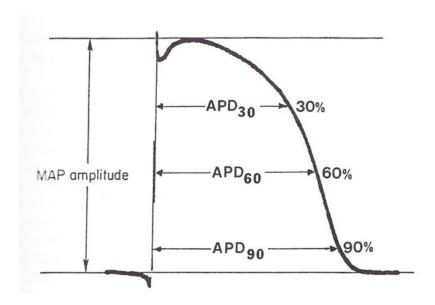


Action Potential - Tissue

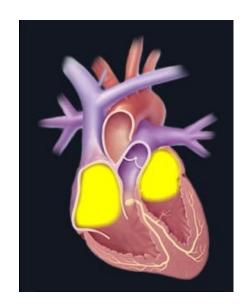
The elementary signal of the heart

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

Closely related to most cardiac diseases.



APD: Action Potential Duration



Functional syncytium

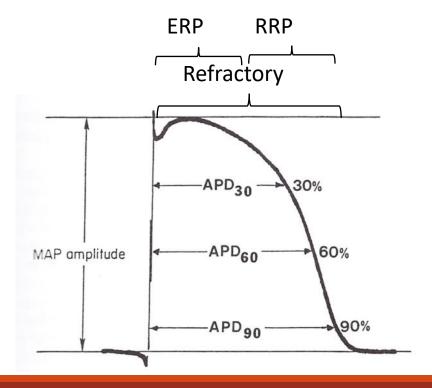
Pot in cut hipplies Bethi | Not read that it in glot feeling (GPS) in g Fixibit . 2012.8

Cellular Level - Action potential

Ions refill

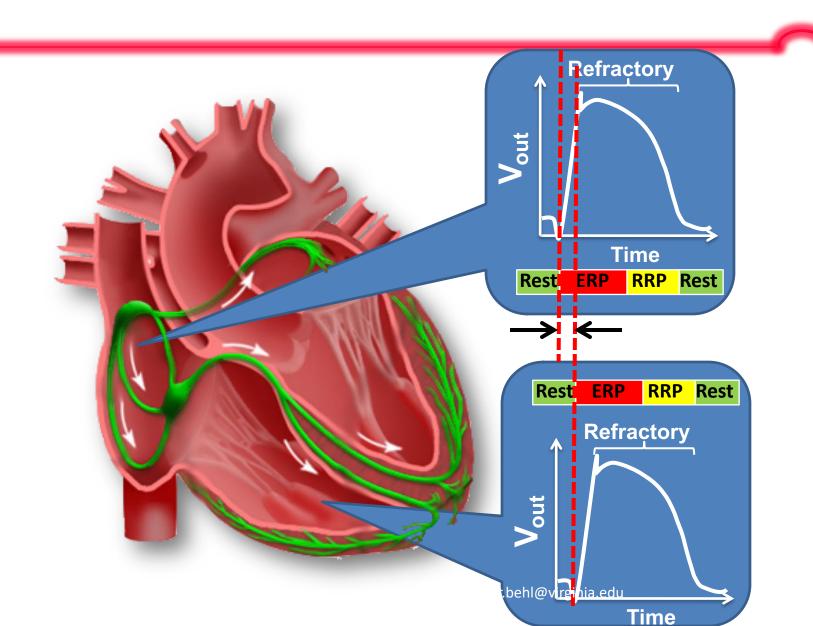
Divided into *Effective Refractory Period(ERP)* and *Relative Refractory Period(RRP)* for activation with certain strength

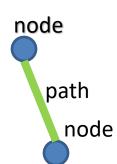
Block Interval during ERP, abnormal new action potential during RRP



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Refractory period



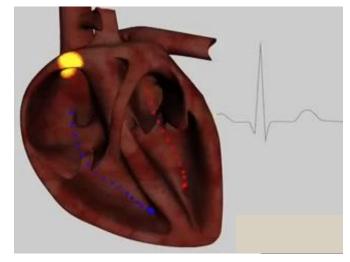


Treating bradycardia

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

Start small and simple:

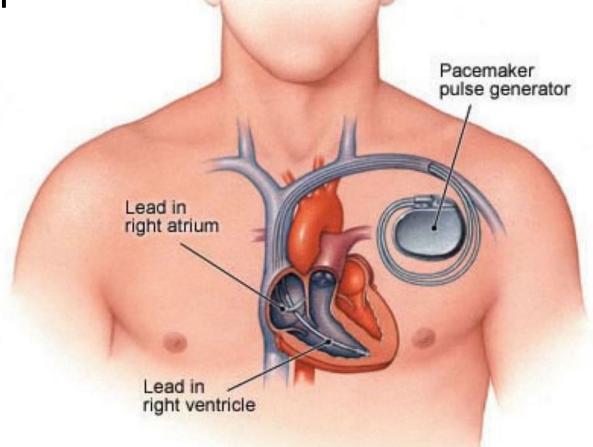
- We know that usually, the atria contract together, and the ventricles contract together → sense only in right atrium and ventricles
- Usually, depolarization is synchronous with contraction → measure the electrical activity as a proxy for the mechanical activity
- Usually, depolarization in part of the atrium (or ventricle) is propagated to the rest of that chamber → measure only in one location of the chamber

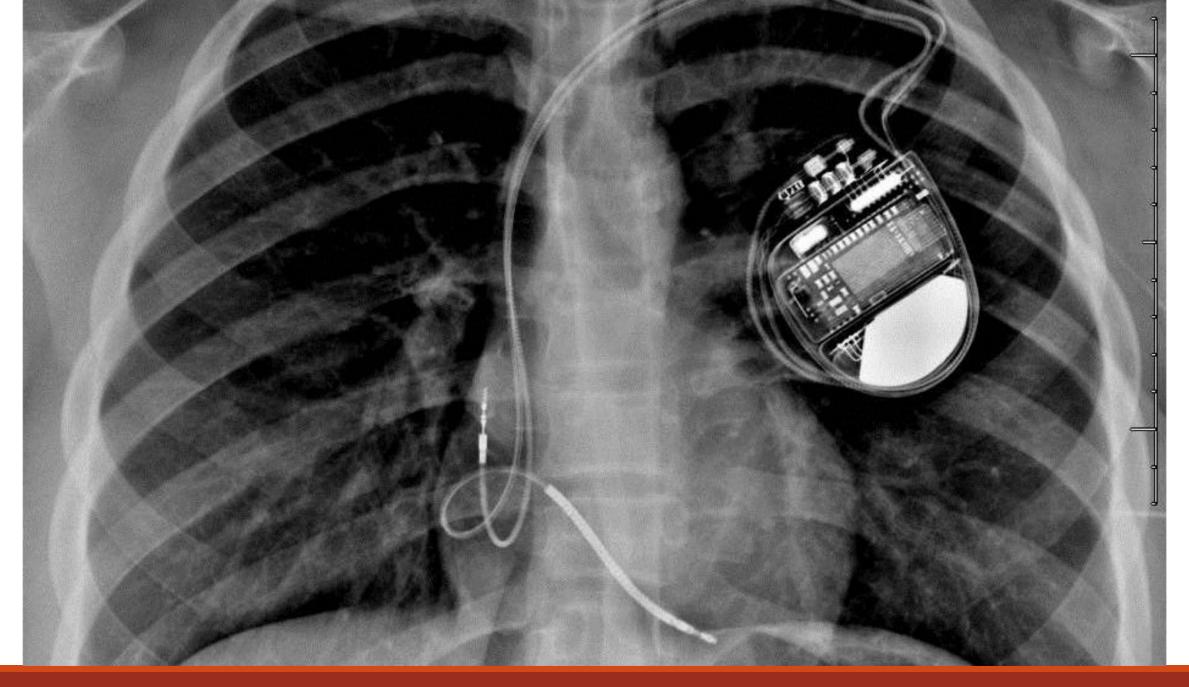


Two leads are placed in the right atrium and right ventricle

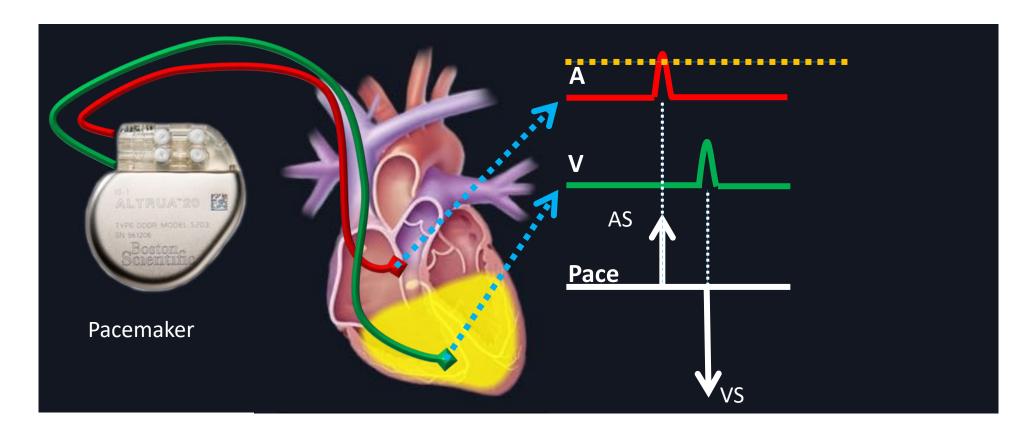
Monitors the local electrical activities of the heart and deliver therapy according to the

timing information





Timing info for local activation



- 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