

Brady-arrhythmias

Pathophysiology of Cardiac Arrhythmias

↓ SA node automaticity (i.e. sinus bradycardia)
 (↑ parasympathetic, ↓ sympathetic stimulation)
 → If SA node rate ↓ enough, AV node & perkinje fibers initiate impulses called "escape beats".
 → A series of escape beats = "escape rhythm"

Conduction Block (i.e. AV block, BB block)
 → Delayed propagation of impulse due to electrically unexcitable tissue (from ischemia, fibrosis, inflammation, drugs)

Altered Impulse Formation

Altered Impulse Conduction

Re-entry loops

→ An impulse travels continuously around a circular (re-entrant) path in the myocardium, continuously depolarizing that cardiac region.
 → Re-entry loops occur in branched, dysfunctional/fibrotic myocardium w/:

- 1) Unidirectional block:** when impulses can't conduct forwards, but can be conducted backwards, in the piece of myocardium
- 2) Slowed retrograde conduction velocity:** backward impulse conduction speed is slow, allows normal myocardium to repolarize so that the impulse propagates in a loop

↑ automaticity

↑ SA node automaticity
 (↑ sympathetic stimulation of β1-adrenergic receptors)

↑ automaticity of latent pacemakers
 → If the AV node and purkinje fibers intrinsically depolarize faster than SA node (produce "ectopic beats"), they'll control impulse formation (produce an "ectopic rhythm") – i.e. AV Junctional Tachycardia

Abnormal automaticity (ectopic pacemakers in atrial &/or ventricular myocytes)
 → If normally non-conducting heart cells depolarize faster than SA node, they'll produce an abnormal ectopic rhythm
 → Usually due to myocyte injury

Triggered activity ("R on T phenomenon")
 → "after-depolarizations" cause extra ventricular contractions during their repolarization.
 → Early after-depol's in Long QT pts → torsades de pointes
 → Delayed after-depol's in high-Ca²⁺-pts → idiopathic V-tach

Tachy-arrhythmias

More on Re-entry loops:

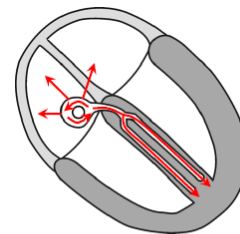
→ Rate of re-entrant circuits is only limited by the refractory period of the tissues involved. Thus, re-entry can ↑ contractions >300bpm!

→ Re-entry loops around distinct anatomical pathways → monomorphic tachycardia on ECG (each QRS looks the same)

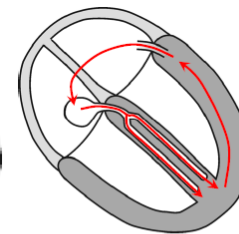
→ Re-entry loops that are disorganized and constantly changing → Polymorphic tachycardia on ECG (no distinct QRS complexes visible)

Ex. VT due to ventricular scar, A-flutter, AVNRT, AVRT (WPW)

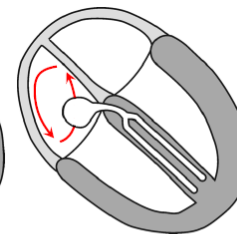
Ex. Polymorphic VT, V-fib, A-fib



AV nodal reentry tachycardia (AVNRT)



AV reentry ("reciprocating") tachycardia (AVRT)



Atrial flutter & some atrial tachycardias