

# QTc Prolongation: Risk Stratification and Mitigation for Challenging Clinical Cases

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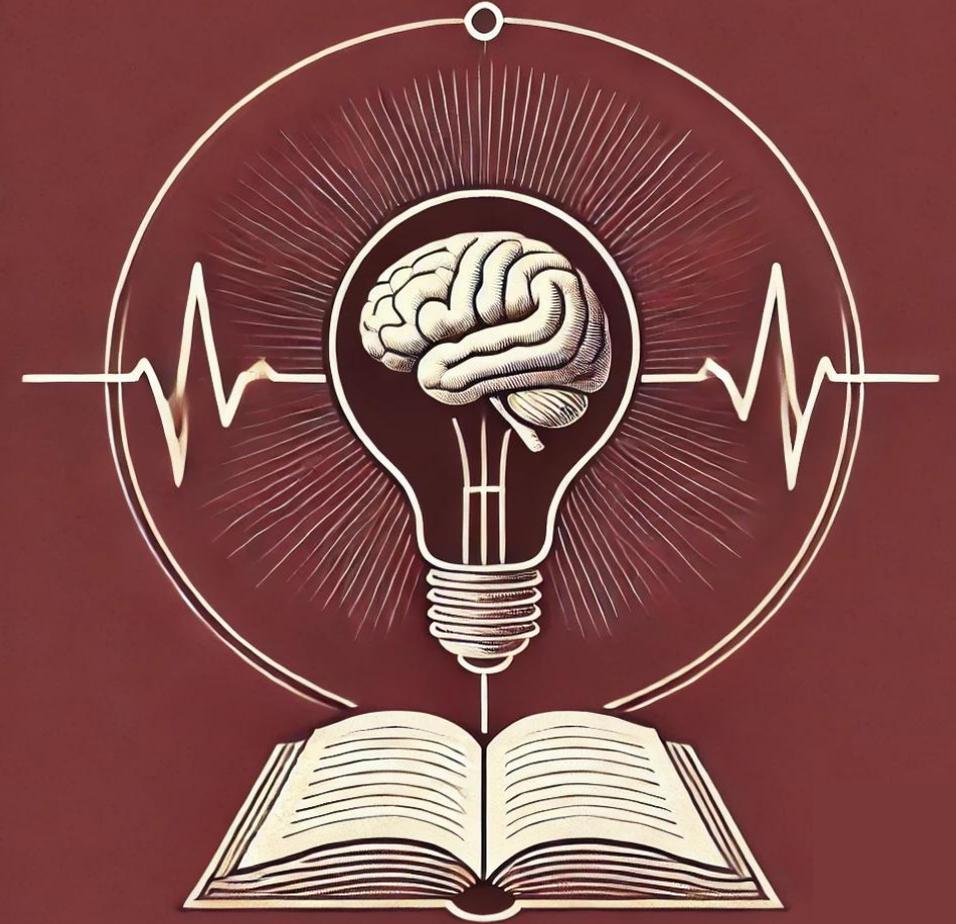
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# Types of Risk Mitigation

- Being comfortable measuring QTc
- Know when to adjust your approach (e.g. wide QRS from pacing or BBB)
- When to monitor
- Risk-Benefit Analysis
- Choice of medication
- Involve other experts
- Interventions for high-risk situations



# Clinical Case 1

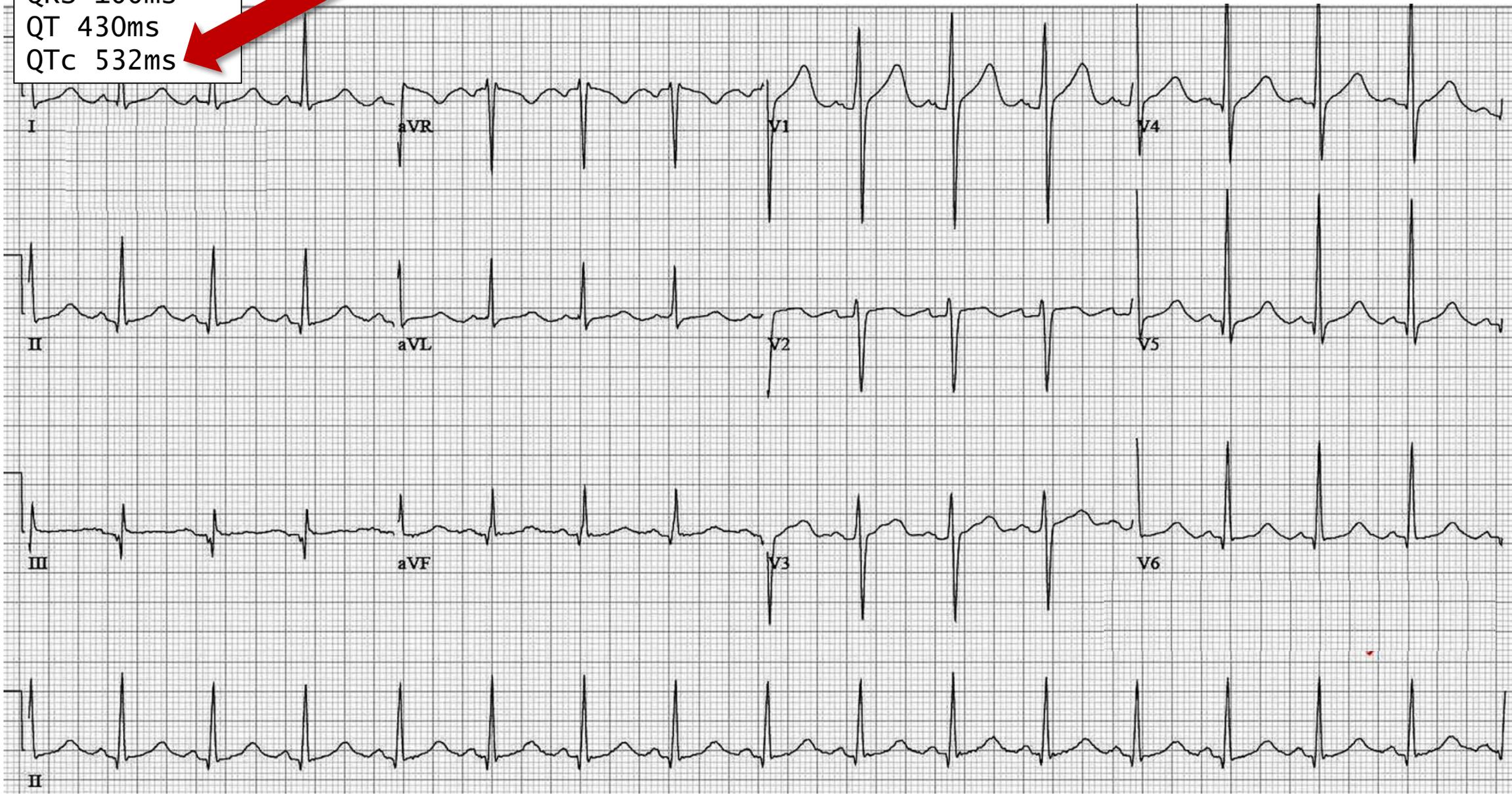
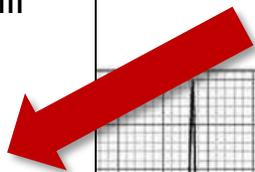
27 y/o woman with history of GAD and social phobia, well controlled on citalopram 40mg po daily, presents to your outpatient clinic to establish care after relocating for her first job out of law school.

She has no significant past medical history, takes no other medications, has no family history of arrhythmia, cardiac arrest or sudden death. She has no suicide attempts, no inpatient psychiatric hospitalizations.

Your clinic just purchased an ECG machine, and the medical assistants were recently trained how to use the machine. While you believe her cardiac risk factors to be very low, since you have the resource available, you decide to obtain an ECG for the sake of completeness.

HR 92 bpm  
QRS 100ms  
QT 430ms  
QTc 532ms

!!!???



# Risk Factors for TdP

## Non-modifiable risk factors

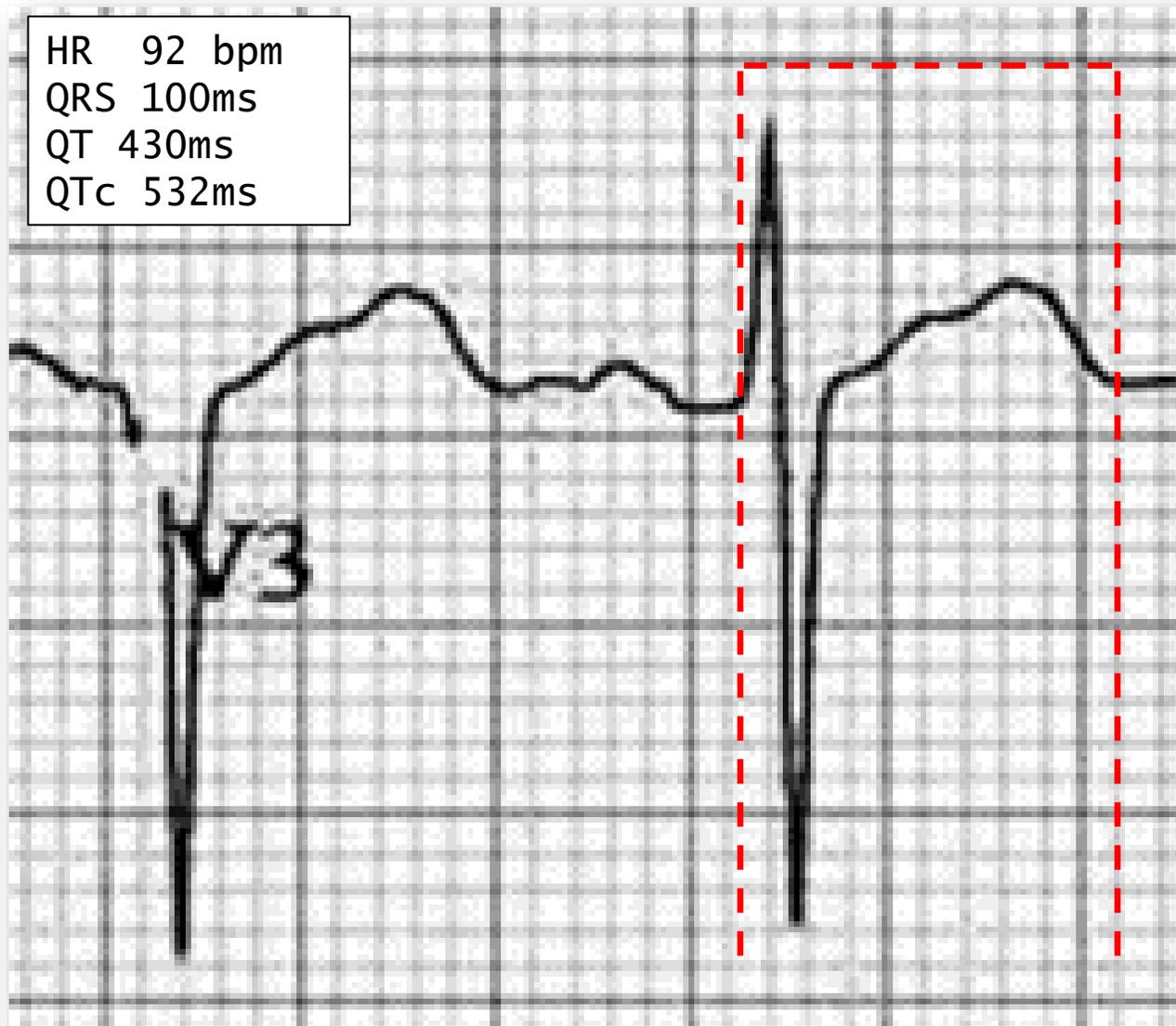
- Female sex
- Advanced age
- Metabolizer status
- Congenital long QT syndrome
- Personal history of drug-induced QTc prolongation
- Personal history of structural or functional cardiac disease
- Family history of sudden cardiac death

## Modifiable risk factors

- Use of  $\geq 1$  QTc prolonging drug(s)
- Pharmacokinetic drug-drug interactions
- Drug toxicity
- Severe acute illness
- Bradycardia
- Starvation
- Hepatic impairment
- Renal impairment, hemodialysis
- Risk or presence of hypokalemia, hypomagnesemia or hypocalcemia

**QT = 390 ms**

HR 92 bpm  
QRS 100ms  
QT 430ms  
QTc 532ms



mdcalc.com

**MD CALC** Search "QT interv" ☰

### Corrected QT Interval (QTc) ☆

Corrects the QT interval for heart rate extremes (choose from Bazett, Fridericia, Framingham, Hodges, or Rautaharju formulas).

When to Use ▾ Why Use ▾

Formula

- Bazett
- Fridericia
- Framingham
- Hodges**
- Rautaharju

Heart rate/pulse 92 beats/min

Paper speed, mm/sec 25 50

QT interval 390 msec ↕

**446 msec**

**QTc (B) = 483 ms**  
**QTc (H) = 446 ms**  
**QTc (Fri) = 450 ms**  
**QTc (Fra) = 444 ms**

# Considerations for ECG Monitoring

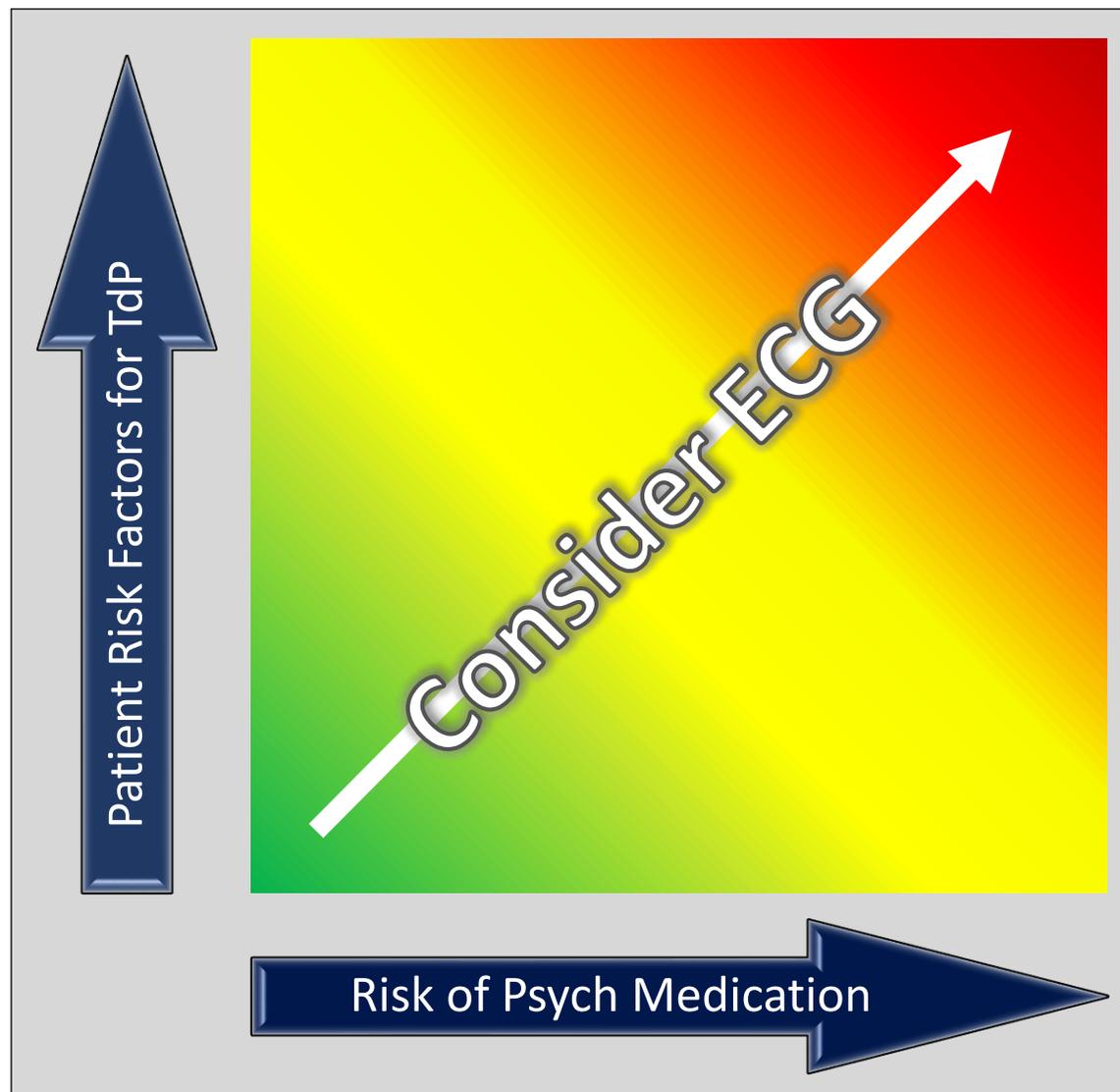
# When to Obtain An Outpatient ECG?

## Non-Modifiable Risk Factors For TdP

- Older Age
- Female Sex
- Functional structural heart disease
- Prior QTc prolongation or TdP
- Prior arrhythmia
- Prior cardiac arrest
- Family history arrhythmia, cardiac arrest, sudden cardiac death

## Modifiable Risk Factors For TdP

- Hypokalemia
- Hypocalcemia
- Hypomagnesemia
- Bradycardia
- Renal Dysfunction
- Hemodialysis
- Liver Dysfunction
- Impaired drug metabolism
- Drug-Drug interactions
- Taking one or more QTc-prolonging drugs



## Low Risk Medications

- Most SSRIs & SNRIs
- Benzodiazepines
- Valproic Acid
- Buprenorphine
- Aripiprazole
- Lurasidone

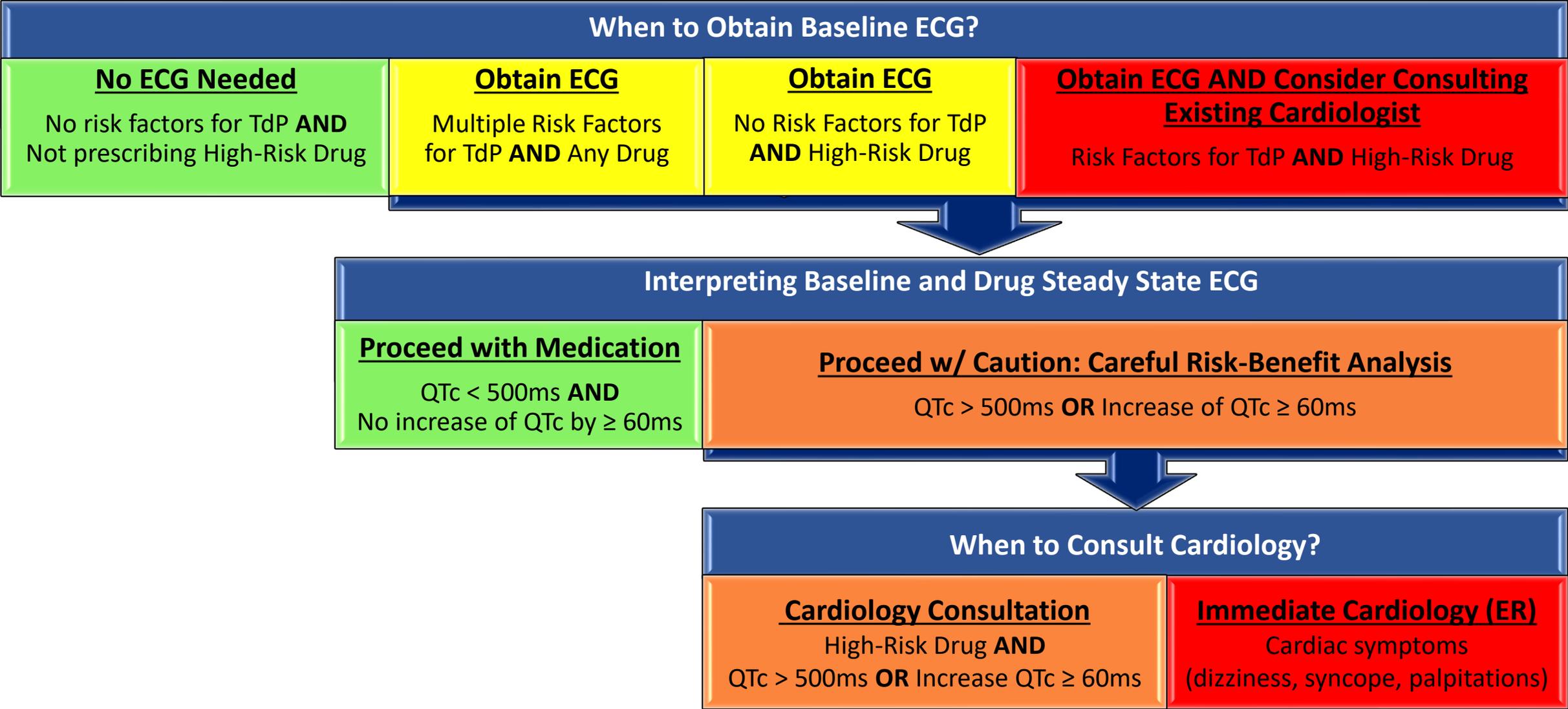
## Moderate Risk Medications

- Citalopram
- Quetiapine, Risperidone, Olanzapine, Haloperidol
- Trazodone

## High Risk Medication

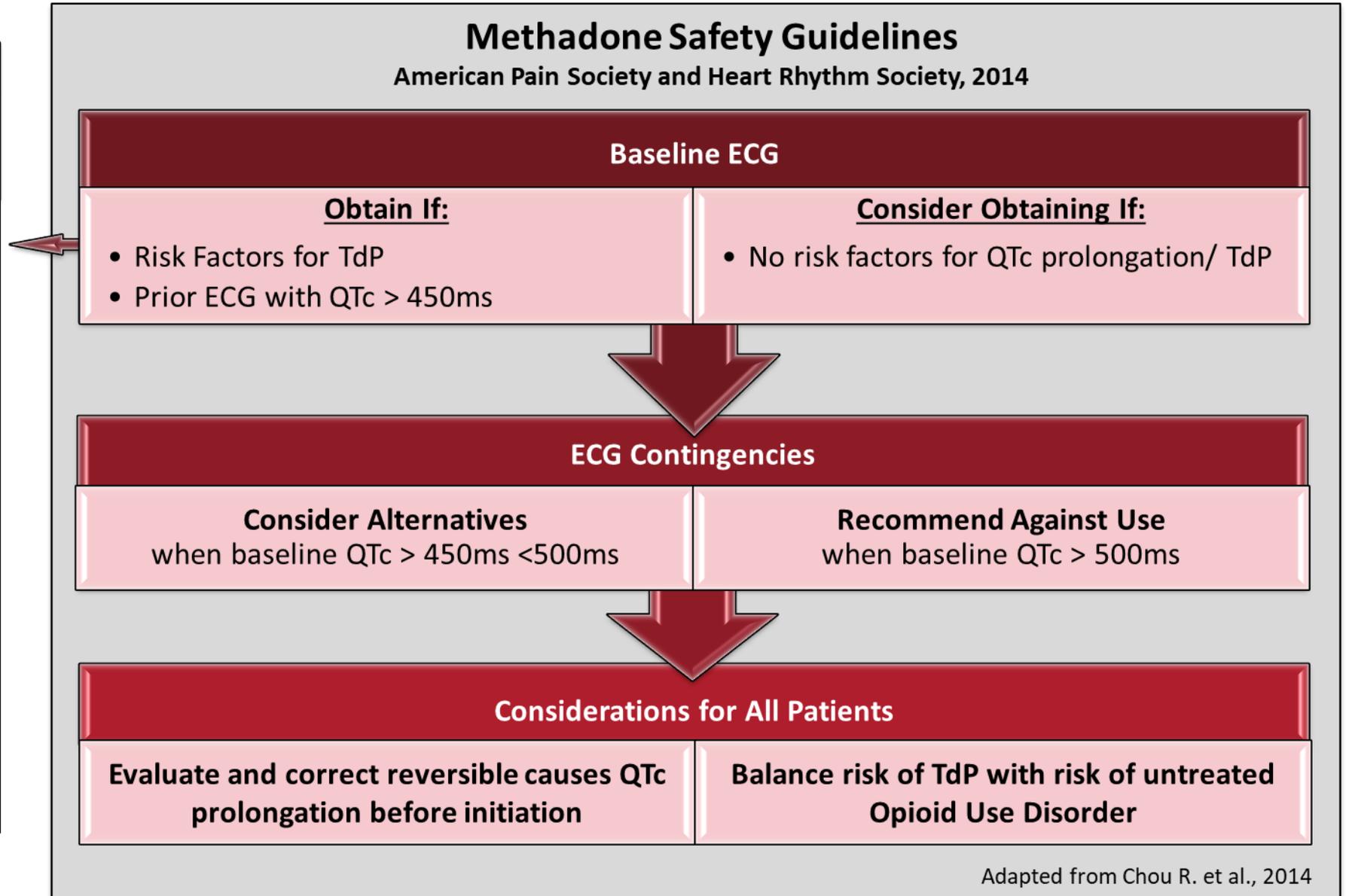
- Methadone
- Ziprasidone
- Iloperidone
- Low -Potency Phenothiazines
- Tricyclic Antidepressants

# Considerations for Outpatient ECG Monitoring



# Methadone Safety Guidelines

Risk Factors
<ul style="list-style-type: none"><li>• Older Age</li><li>• Female Sex</li><li>• Functional structural heart dz</li><li>• Prior QTc prolongation or TdP</li><li>• Prior arrhythmia</li><li>• Prior cardiac arrest</li><li>• Family history arrhythmia, cardiac arrest, sudden cardiac death</li><li>• Hypokalemia</li><li>• Hypocalcemia</li><li>• Hypomagnesemia</li><li>• Bradycardia</li><li>• Renal Dysfunction</li><li>• Hemodialysis</li><li>• Liver Dysfunction</li><li>• Impaired drug metabolism</li><li>• Drug-Drug interactions</li><li>• Taking one or more QTc-prolonging drugs</li></ul>



# Clinical Case 2

44 y/o man with a family history of cardiac arrest in his grandfather, father, and paternal aunt, tests positive for a genetic cardiomyopathy with high risk for sudden cardiac death. He has never had any arrhythmias or cardiac symptoms. He is implanted with a subcutaneous Implantable Cardioverter Defibrillator (ICD) for primary prevention.

His past medical and psychiatric history is notable for hyperlipidemia, recurrent sinus infections, and depression. He takes simvastatin, escitalopram, and is frequently prescribed a Z-pack (azithromycin) by his PCP.

He develops a viral GI illness with multiple days of vomiting and diarrhea, not able to keep anything down. He presents to the ED for dehydration where his QTc(H) is 522ms, K<sup>+</sup> is 2.7. The ED resident orders ondansetron for nausea. When another ED resident questions this decision due to Torsades risk, the first resident says “Don’t worry about it. He has an ICD.”

**Should they worry about it?**

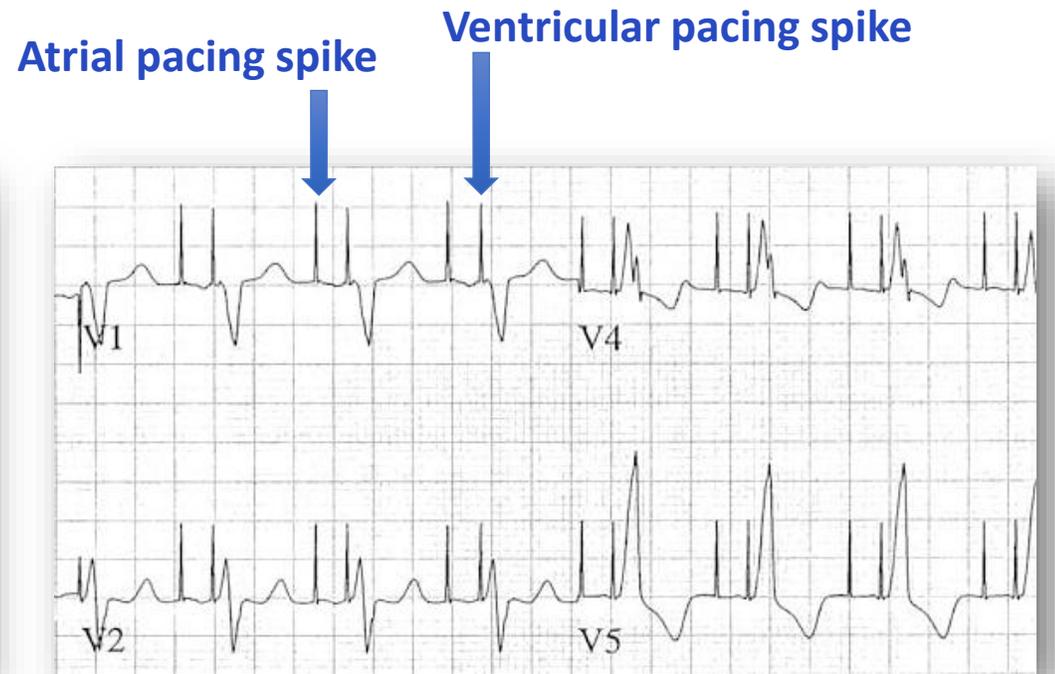
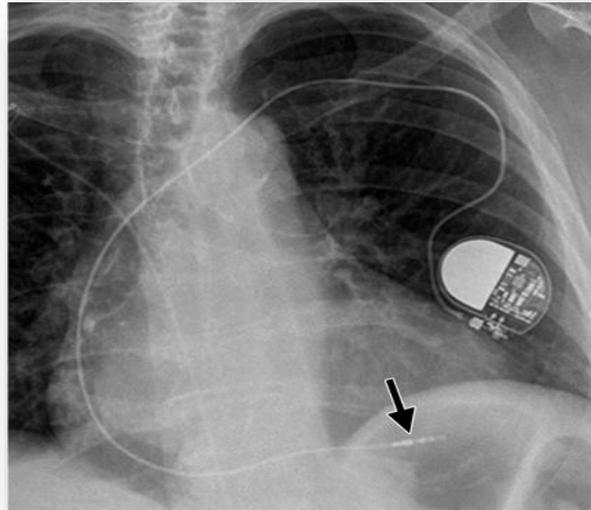
# Role of Pacemakers and Defibrillators

# Permanent Pacemaker (PPM)

**Indication:** treatment of brady-arrhythmias

**Goal:** to sustain adequate HR for daily activity

**How does it work?** Paces the heart if the HR drops below the pacing threshold (usually 30-40 bpm)



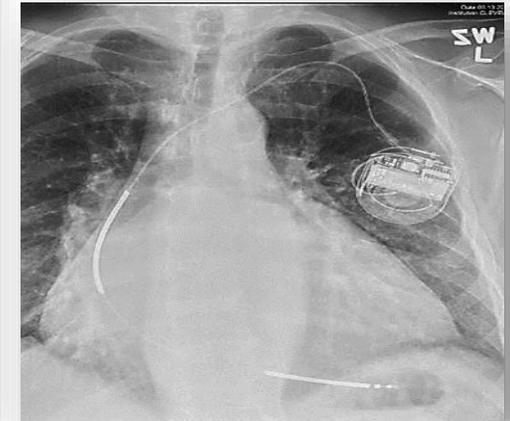
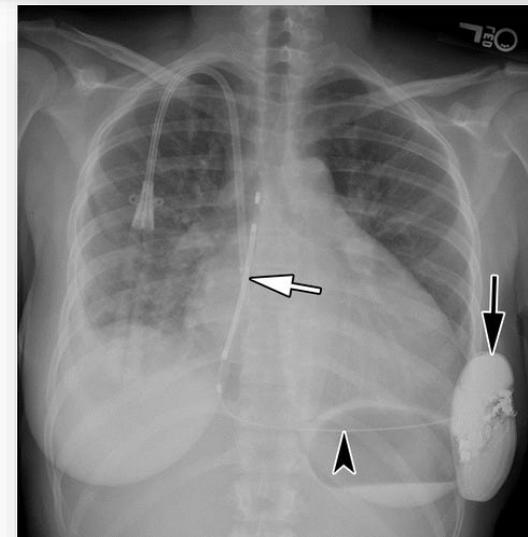
# Implantable Cardioverter Defibrillator (ICD)

## Primary Prevention:

- CAD + Heart failure LVEF  $\leq$  35%
- Dilated Cardiomyopathy
- Certain Pathogenic Gene Mutations
- Arrhythmogenic RV Cardiomyopathy
- Congenital Long QT Syndrome
- Asymptomatic QTc  $>$  500ms during  $\beta$ -Blocker
- Brugada syndrome + symptoms/ inducible AF

## Secondary Prevention:

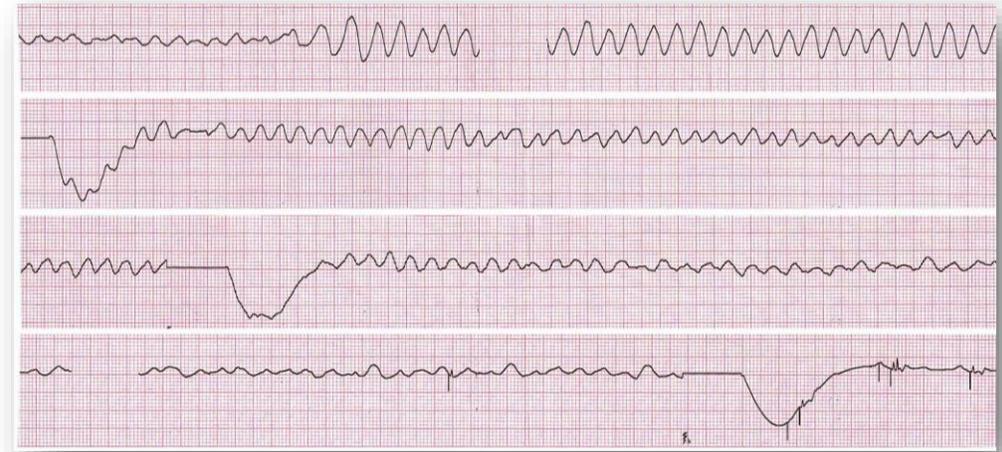
- VF sudden cardiac arrest
- sustained VT
- Hemodynamically not-tolerated VT



# Implantable Cardioverter Defibrillator (ICD)

## ICDs can readily terminate most VT/VF effectively, but...

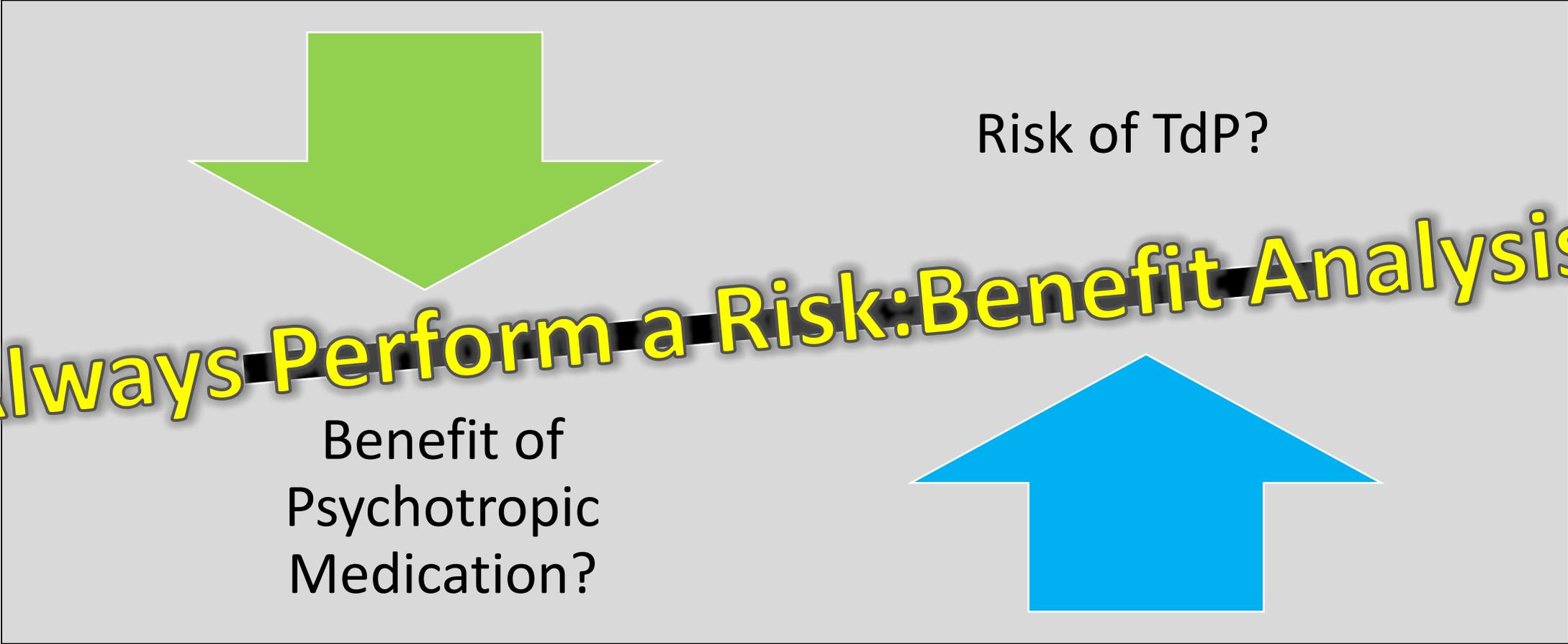
- Even appropriate shocks are independently associated with increased risks of death and worsening heart failure
- Some VT will recur if substrates are unchanged, despite “successful” shocks
- ICD “storms” are common, which lead to significant psychological morbidity (anxiety, depression, PTSD, panic)
- **A shock does not occur without morbidity!**



# Risk-Benefit Analysis

# Risk-Benefit Analysis

To Prescribe or Not to Prescribe?



# Clinical Case 2

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**Should they worry about it?**

**YES!**

# Clinical Case 2

The first ED resident keeps the ondansetron order but puts the patient on telemetry for monitoring. The patient is administered 4mg IV x 2. Forty minutes after the second dose of ondansetron, the patient suddenly yells out. Telemetry shows that he was in Torsades and shocked back to a normal sinus rhythm by his ICD.

Four minutes later, he is shocked again and in sum receives a total of 10 shocks in the span of just minutes.

He is transferred to the CCU for further monitoring and treatment. As the psychiatry consultant, you are called later in the day for management of agitation.

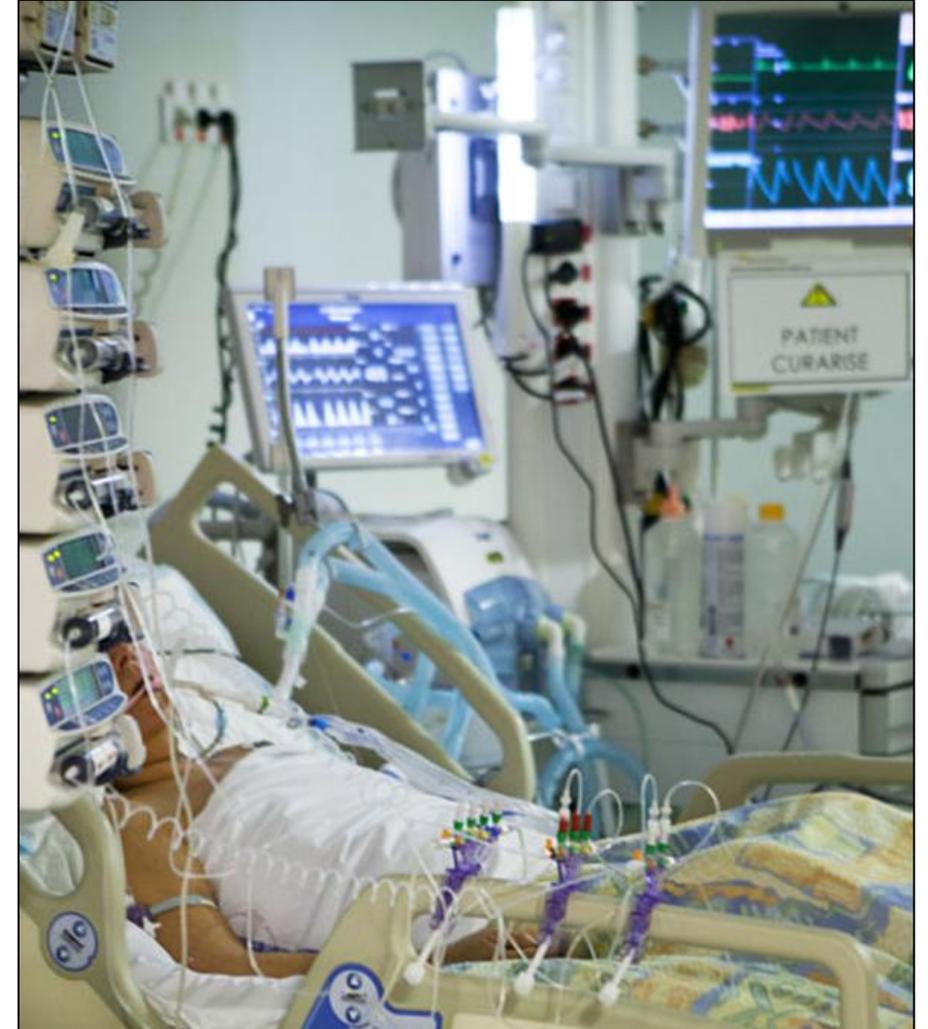
# Risk-Benefit Analysis: ICU Example



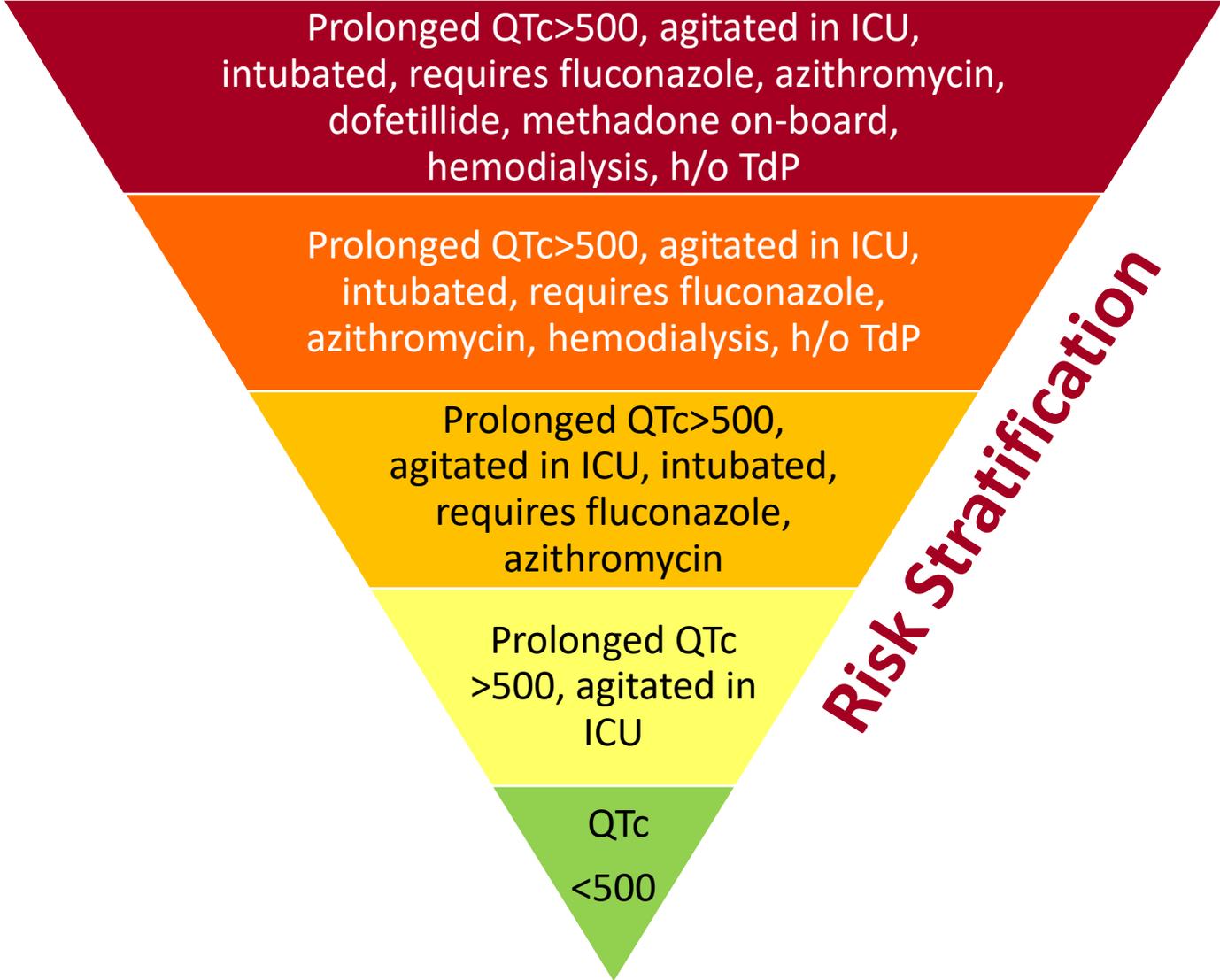
Very nature of being in ICU  
Concurrent QT-prolonging medications  
Parenteral administration  
Susceptible to fluid shifts and electrolyte imbalance  
Likely to have renal/hepatic dysfunction  
Older Age



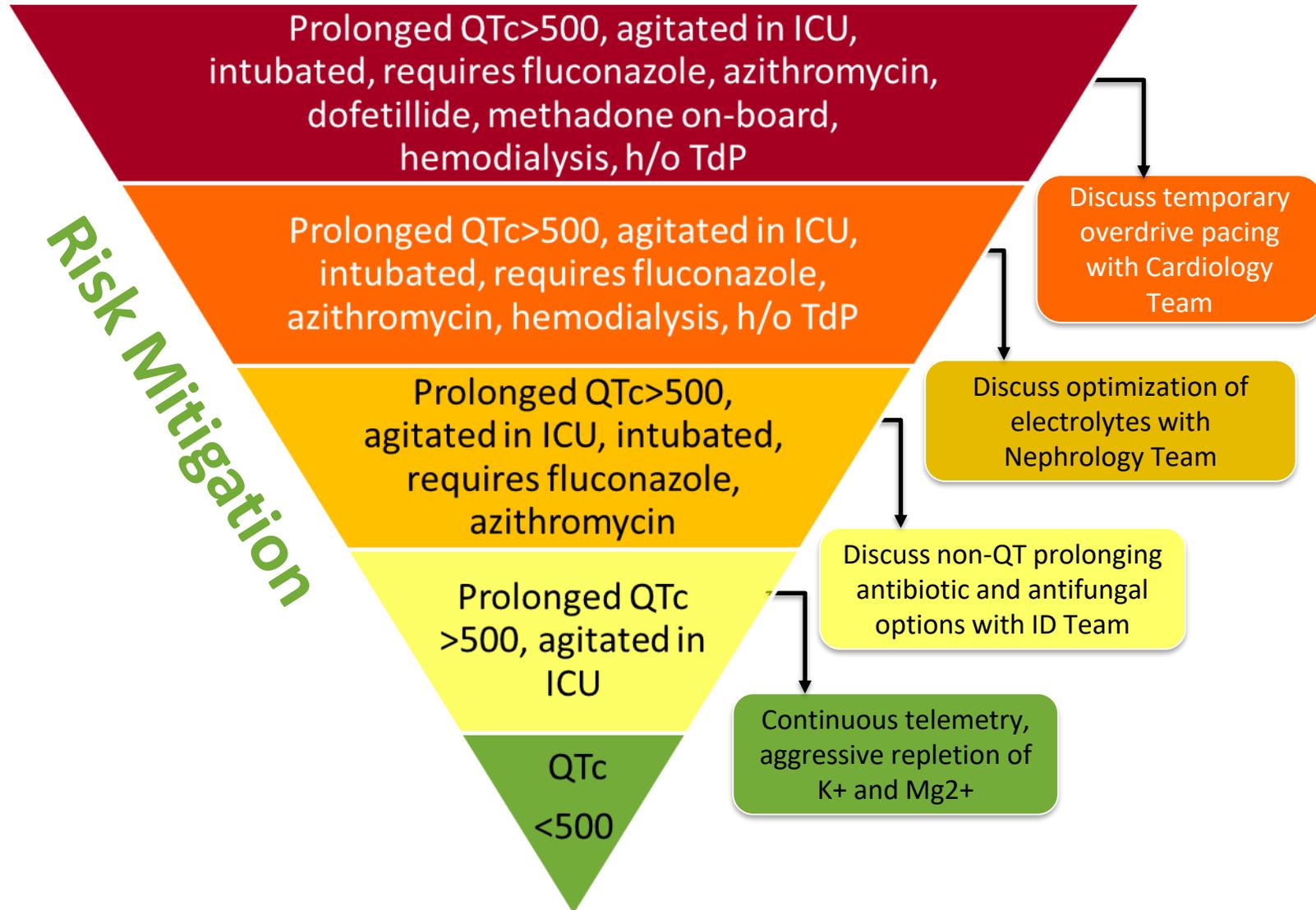
Control of Agitation  
Protection of patient from injury  
Protection of life-sustaining external lines/ devices  
Prevent re-intubation and sedation  
Prevent use of restraints



# Risk Stratification & Mitigation



# Risk Stratification & Mitigation

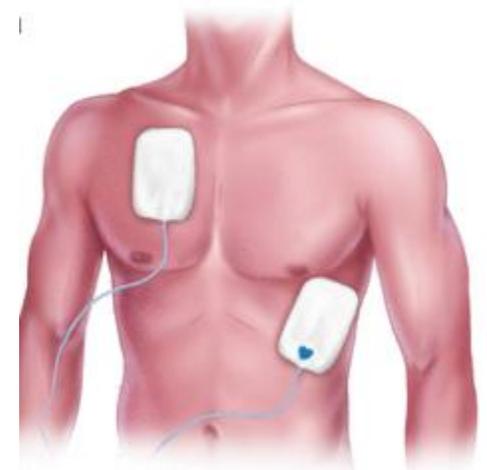


# High Risk Mitigation Strategies

# High-Risk Mitigation Strategies

## Cardiac Overdrive Pacing

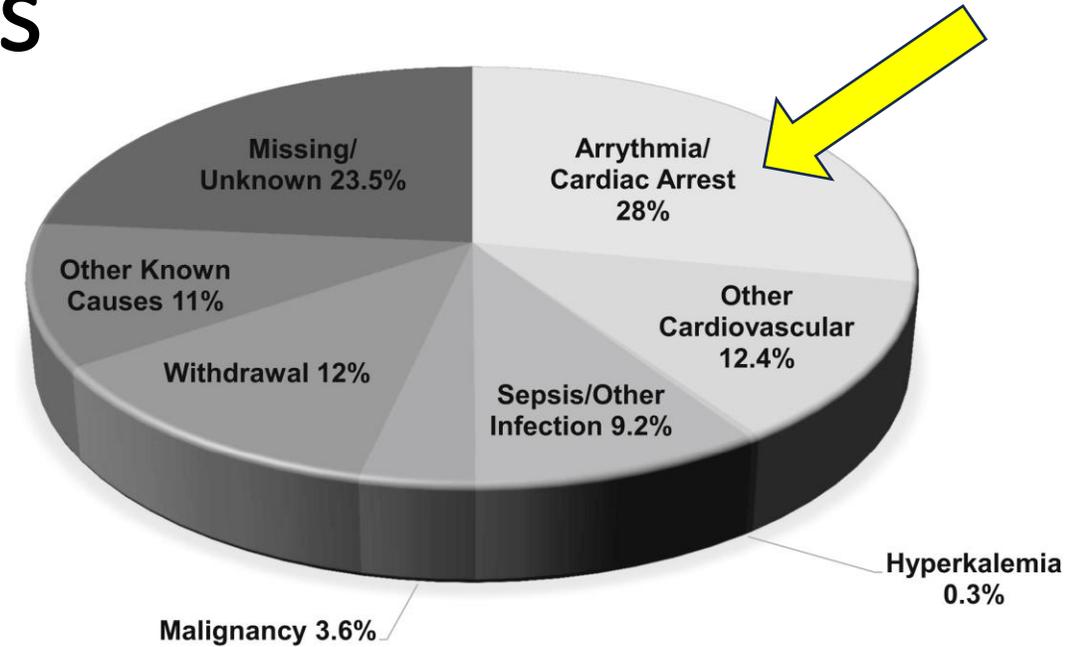
- In high-risk cases with severe bradycardia
- Overdrive pacing  $> 70$  bpm can mitigate risk of early-after-depolarizations that can lead to a reentrant rhythm
- Nearly all cardiac implanted electronic devices have capability for overdrive pacing (except subQ ICDs)
- Pacing thresholds usually set to HR 40-50 to preserve battery-life and ventricular strain, HOWEVER pacing rates can be increased in high-risk patients
- Alternatively may consider temporary transvenous pacing or “chemical” overdrive pacing with isoproterenol (less common)



# High-Risk Mitigation Strategies

## Hemodialysis

- Sudden cardiac death (SCD) is leading cause of death in hemodialysis patients
- QTc prolongation very common in hemodialysis patients
- Contradicting studies regarding impact of dialysate electrolyte concentrations and serum-to-dialysate gradient measures on QTc
- Goal K<sup>+</sup> in HD patients is 5.1
- **Work closely with nephrology in high-risk patients on HD**



# Take-Home Points

- Knowledge is power – practice ECG interpretation
- Do your own QT measurement and HR correction
- 500ms is not an absolute QTc cut-off
- Always perform a risk-benefit analysis
- Talk to your expert colleagues who can help mitigate risk



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