

# Recognition of Drug-Induced Prolonged QT

### Measure QT and RR Intervals

### Correct for Heart Rate

### Check QTc versus Normal Limits

#### Risk Factors: Genetics

ECG from LQT2 patient

Current evidence suggests that 5 to 10 percent of persons in whom torsades de pointes develops on exposure to QT-interval-prolonging drugs harbor mutations associated with the long-QT syndrome. (Roden, NEJM, 2004)

#### Risk Factors: Electrolytes

Example of Hypokalemia

Imbalance in K, Ca, or Mg can lead to higher risk. In this example, lead II exhibits TP fusion while in V4-V6 these waves are clearly separate.

#### Risk Factors: Brady-Arrhythmia

Atrial fibrillation with low heart rate and prolonged QT. Long, followed by short RR, leads to Torsades de Pointes.

#### Risk Factors: Poor Liver or Heart Function

**Liver:** Poor function can lead to toxic levels of prescribed drug.

**Heart:** Cardiac ischemia can prolong QT. Low ejection fraction can increase risk of arrhythmic death due to prolonged QT.

#### Risk Factors: Drugs

- Anti-depressants
- Anti-psychotics
- Anti-biotics
- Anti-arrhythmics
- Etc.

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## Tools and Strategies for Measuring Difficult ECGs

#### Artifact

The artifact in tracing A is too excessive. The T-wave is not even visible. Advise repeat ECG. Tracing B exhibits a T-wave but the measurement of QT could be inconsistent due to baseline wobble and muscle artifact. Select the best complex for measurement.

A median complex is formed by taking the median value, at each sample time, from the superimposed raw beats. The result is a cleaner, representative complex for measurement.

#### Flat T-wave in Limb Leads - Use All Leads

Measurements based on a single lead will result in error, especially if the lead has a flat T-wave. Check all leads. Use longest interval. GE's Marquette 12SL program superimposes median complexes from each lead. The automated QT measurement is from earliest QRS onset to latest T-offset.

#### Bi-phasic T-wave - Include Terminal Portion

Do not define baseline based on the end of the T-wave. This can lead to the mistake of defining T offset within T. QRS onset should define baseline. See V2. The entire wave of repolarization includes T and T'. Use all leads to assist in defining the end of repolarization.

#### Calculating QTc in the Presence of Varying Heart Rate

QT is varying with heart rate. An overall QTc needs to be determined. Some recommend an average obtained from the shortest and longest preceding RR intervals. Others perform the average over 3 to 5 consecutive beats. Instead, GE's Marquette 12SL program uses QT from the median beat and the average RR.

#### TP Fusion - Measuring QT in Presence of High Heart Rates

At fast heart rates, the P-wave fuses with the T-wave, sometimes making it infeasible to measure the end of the T-wave. If possible, record and measure the QT at a lower heart rate. In a single lead, use tangent method to identify the end of the T-wave.

In normal sinus rhythm, leads V1 and II have the largest P waves. Other leads may reveal separation of P and T. Tick marks on 12SL medians report can be used to inspect 12SL measurements.

#### Do Not Include Normal U-waves

T and U-waves are not superimposed and can be separated. These are normal U-waves. Either use tangent method for single lead or inspect other leads without U-waves in order to determine T-offset.

#### TP Fusion - Long PR Interval

Due to a long PR interval, the P wave is superimposed on the T-wave. In lead II, the T wave appears flat. The tangent method will be difficult to apply. In V3, the tangent method will result in a longer PR interval, than inspecting all the leads.

By inspecting all leads, it is clear the T-wave shape is due to a long PR interval. In leads V1 and V5, the P and T-waves are separate.

#### Include U-wave When Abnormal or Unable to Separate from T-wave

These are abnormal U-waves. The U-waves are too large and fused with the T-wave. Do not use tangent method. Include U-wave.

### T-wave Before and After Drug Effect

**Before**

**After**

T-wave Notch