

Lightning Learning: Long QT Syndrome





youtube.com/em3orguk



WHAT?

Congenital or acquired disorder which causes the OT interval to be prolonged (>450ms)

- **Genetic mutation** affects Na and K channels
- Drugs e.g. TCA, SSRIs, haloperidol and many more
- Electrolyte disturbance e.g. hypokalaemia and hypocalcaemia
- Heart disease

Prolonged repolarisation of the ventricles which causes arrhythmias. Diagnosed with aid of ECG and Schwartz criteria.

Romano-Ward syndrome and Jervell and Lange-Nielsen syndrome are genetic forms of long QT syndrome.

WHY?

Affects 1:2500 people worldwide. Patients may remain asymptomatic BUT Long QT syndrome is Associated with **syncope**, especially on exertion and...

SUDDEN CARDIAC ARREST

It is often misdiagnosed as epilepsy.

Management for congenital cases will be through lifestyle change, **beta-blocker** therapy and ICD.

Patients will need to avoid extreme exertion.

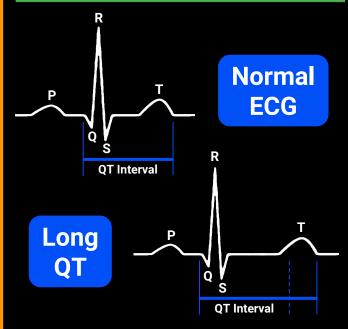
HOW?

Long QT Syndrome Workup

(Medscape) http://bit.ly/2BaBZ1s

Long QT Syndrome (Life in the Fast Lane) http://bit.ly/2AunuUX

Long QT Syndrome (BMJ Best Practice) http://bit.ly/2jYkMAX



Author: Pandora Spilman-Henham **Date:** 18.12.2017