

McGill University Health Centre

Centre universitaire de santé McGill

ECG Session Family Medicine Refresher 2022

Jacqueline Joza

McGill University Health Centre



Disclosures

Medtronic: Grant – Investigator-initiated external research program Boston Scientific – Advisory Board



- 1. Review the normal ECG
- 2. Identify the bundle branch blocks
- 3. Evaluate the underlying arrhythmia
- 4. Discover pacing
- 5. Learn how to measure the QT interval

Poll Question 1: 50 year-old asymptomatic female sent for a routine ECG.



- A. Normal ECG
- B. Left bundle branch block

- C. Right bundle branch block
- D. Long QT interval

The Normal ECG





Add ecg here with long AV delay Then one with a very long AV delay

Draw a typical RBB; typical LBBB







POLL Question 2: Is this a true left bundle branch block? (ie. Does it meet strauss criteria?)











The relationship with a His pacing lead: Left bundle pacing lead is typically 1-1.5mm further distally along the septum



Coaxial



15





What is the rhythm? Is this patient paced?



Poll Question 4: What type of device is this?

- A. Dual-chamber pacemaker
- B. Biventricular pacemaker
- C. Biventricular ICD/ defibrillator
- D. Defibrillator





Draw a typical RBB; typical LBBB





Poll Question 5: 80 year-old female with dyspnea



- A. Normal ECG
- B. Left bundle branch block
- C. RBBB + left anterior fascicular block (left axis)
- D. RBBB + left posterior fascicular block (right axis)

Baseline ECG with RBBB + left posterior fascicular block (right axis)



ECG with 'conduction system' pacing: correction of the RBBB and right axis



- A. Normal ECG
- B. Left bundle branch block

- C. RBBB + left anterior fascicular block (left axis)
- D. RBBB + left posterior fascicular block (right axis)

Baseline ECG with RBBB + left anterior fascicular block (left axis)



ECG with 'conduction system' pacing: correction of the RBBB and left axis

What else do you notice about this ECG?



Draw a typical RBB; typical LBBB







Poll 7: SVT (supraventricular tachycardia) or VT (ventricular tachycardia?



SVT or VT?





Poll Question 8: What is the underlying rhythm?



Poll 9: 70M with LVEF 60%. What type of device should you implant if any?



What is the underlying rhythm?


Poll Question 10: 83F with fatigue





83F Post-op TAVI develops complete heart block, dependent on temporary pacing wire. What should we do now?







Stable threshold, sensing, and impedance of the lead



1 week later, presents to ED with pulmonary edema.











23M, recurrent syncope 2x per month x 8 months

Short prodrome

EP study negative

so loop monitor implanted

High frequency stimulation posteromedial GP





What is the underlying Rhythm?





 $av_{1} \longrightarrow b$

Arrowhead represents initial slow descent Arrow represents later sharper rapid descent





- -During cavo-tricuspid isthmus ablation, the tachycardia terminates during the initial gradual downslope of the p wave.
- -The slow initial descent represents conduction through the isthmus
- -The sharp second descent represents penetration of the wavefront through the coronary sinus and interatrial septum, with passive depolarization of the LA

Poll Question 11: What is the rhythm?

- A. Normal rhythm
- B. SVT
- C. Atrial flutter
- D. Atrial fibrillation





25mm/s 10mm/mV 40Hz 8.0 SP2 12SL 241 HD CID: 37

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Poll Question 13:

What does the notch represent seen best on V3?



- A. Normal
- B. WPW
- C. Long QT
- D. u wave



Case: 42F

• Diagnosis: presumed long QT type 2

<u>Plan:</u>

- 1. Exercise treadmill testing
- 2. Send blood for genetic testing
- 3. Start nadolol 20mg po qd; goal 1-2mg/kg/day
 - 4. Eventual cascade screening pending genetic results
 - 5. Consider LINQ (Reveal monitor)
 - 6. Avoidance of drugs prolonging QT interval
 - 7. Lifestyle changes: reduction of exposure to sudden noises, abstention from competitive exercise, adequate hydration

Long QT Syndrome (LQTS)

- Prevalence: 1 in 2000.
- QT similar between young boys and girls, where differences appear during puberty.
- In congenital LQTS, women have longer QT than men, therefore women are more often clinically diagnosed despite equal genotype sex-distribution
- The probability of a first cardiac event is higher in females by age 15, but decreases after puberty; the first cardiac event tends to be more often fatal in males than females
- The longer the QT interval, the increasing risk for malignant arrhythmias

Long QT Syndrome (LQTS)

- Although prolonged ventricular repolarization is the hallmark of LQTS, up to one third of carriers of a pathogenic LQTS mutation exhibit normal QTc values
 - These clinically 'silent' mutation carriers have a higher risk of syncope or cardiac arrest than the general population

Long QT syndrome (LQTS): measurement of the QT interval

48F several days after resuscitated cardiac arrest





≤1.0 point = low probability of LQTS

1.5-3.0 points = intermediate probability of LQTS

≥3.5 points = high probability of LQTS

Overall: QTc ≥ 460 (Bazett's)

99%: (>450 males; > 470 females)

Findings			Points		
ECG 1	QTc ²	≥480 ms			
		=460-479 ms =450-459 ms (in males)			
		≥480 ms during 4 th minute of recovery from exercise stress test	1		
	Torsade de pointes ³				
	T wave alternans				
	Notched T wave in 3 leads				
	Low heart rate for age ⁴				
Clinical history	Syncope ³	With stress	2		
		Without stress	1		
Family history	Family member(s) with definite LQTS ⁵				
	Unexplained sudden cardiac death before age 30 years among immediate family				
Total sco	ore				

Schwartz and Crotti. 2011

Gene		Affected Channel		Disease	% of LQTS		
	Almos torsac	st all dru les are	ugs causing Ikr blockers			attributed to mutation of this gene	
KCNQ1	(LOF)		Delayed rectifie	r Iks	LQTS type 1	30-35%	
KCNH2	(LOF)]	Delayed rectifier	r IKr	LQTS type 2	25-30%	
SCN5A	(gain)		Nav1.5		LQTS type 3	5-10%	

• LQTS 1,2, and 3 account for over 95% of genotype-positive cases. The yield of genetic testing in pts with a clinical diagnosis of LQTS is approx 75-80%.

Gene	Disease	% of LQTS	Gene	Disease	% of LQTS
ANK2	LQTS4	<1%	SCN4B	LQTS10	Rare (2 cases)
KCNE1	LQTS5	<1%	AKAP9	LQTS11	Rare (1 case)
KCNE2	LQTS6	<1%	SNTA1	LQTS12	Rare (3 cases)
KCNJ2	LQTS7	<1%	KCNJ5	LQTS13	Rare (2 cases)
CACNA1c	LQTS8	<1%	CALM1	LQTS14	< 1%
CAV3	LQTS9	<1%	CALM2	LQTS15	< 1%

Case: 42F judge with presyncope

RESULTS KCNH2 Pathogenic Mutation: c.307_307+1delGGinsTT SUMMARY

POSITIVE: Pathogenic Mutation Detected

INTERPRETATION

- This individual is heterozygous for the c.307_307+1delGGinsTT pathogenic mutation in the KCNH2 gene.
- This result is consistent with a diagnosis of long QT syndrome.
- The expression and severity of disease for this individual cannot be predicted.
- Genetic testing for pathogenic mutations in family members can be helpful in identifying at-risk individuals.
- Genetic counseling is a recommended option for all individuals undergoing genetic testing.





8 yr-old son Specific Site Analysis of KCNH2

RESULTS

KCNH2 SPECIFIC SITE c.307_307+1delGGinsTT Pathogenic Mutation: Detected

INTERPRETATION

This individual is heterozygous for the c.307_307+1delGGinsTT pathogenic mutation in the KCNH2 gene, which was previously identified in this individual's relative(s). This result is consistent with a diagnosis of long QT syndrome; however, the expression and severity for this individual cannot be predicted.

The only KCNH2 alteration analyzed for this individual was c.307_307+1delGGinsTT.

Genetic counseling is a recommended option for all individuals undergoing genetic testing.

Plan: Start nadolol

Implant LINQ (Reveal monitor) (long QT > 500ms)







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Plan: Start nadolol Implant LINQ (Reveal monitor) (long QT > 500ms)






General review of ECGs, pacing, QT measurements etc.