SYNCOPE & ECGs NOTES

Always order an ECG for syncope evaluations.

Arrhythmogenic Right Ventricular Dysplasia

KEY FACTS:

- **♦** Cardiomyopathy
- ♦ #2 Cause of Sudden Cardiac Death
- **♦** Autosomal dominant (Family Hx)
- **◆** Epsilon waves and flipped Ts V1-V3

ARVD= pt can be any age (infant to elderly). Overview of ARVD

- Autosomal dominant genetic disorder of myocardium in which there is fatty infiltration of the right ventricular free wall, predisposing to paroxysmal ventricular arrhythmias, sudden cardiac death, and biventricular failure
- Second most common cause of sudden cardiac death in young people (after HOCM), accounting for up to 10% of sudden cardiac deaths in patients < 65 yrs of age
- Prevalence ~ 1 in 5000
- Diagnosis is difficult and relies on a combination of clinical, electrocardiographic and radiological features, as defined by the (horribly complicated) 2010 Task Force Criteria

Epsilon wave in V1, due to RV conduction delay

Prolonged S-wave upstroke in V2 with localized QRS widening

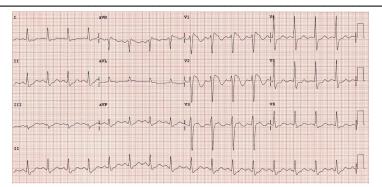
ECG features

- T wave inversion in right precordial leads V1-3, in absence of RBBB (85% of patients)
- Epsilon wave (most specific finding, seen in 50% of patients)
- Localised QRS widening in V1-3 (> 110ms)
- Prolonged S wave upstroke of 55ms in V1-3
- Ventricular ectopy of LBBB morphology, with frequent PVCs > 1000 per 24 hours
- Paroxysmal episodes of ventricular tachycardia (VT) with LBBB morphology (RVOT tachycardia)

BRUGADA

Brugada sign



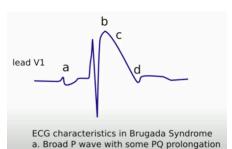


BRUGAD*A*



Mneumonic: **RBBB** with STE in V1-3 = Brugada

(when you say it out loud. it rhymes)



Coved ST segment elevation >2mm in >1 of V1-V3 followed by a negative T wave. This is the only ECG abnormality that is potentially diagnostic. It is often referred to as Brugada sign.

make the diagnosis:

This ECG abnormality must be associated with one of the following clinical criteria to

Documented ventricular fibrillation (VF) or polymorphic ventricular tachycardia (VT).

Type 1 (there are 3 types but type 1 is most fatal and most important):

- Family history of sudden cardiac death at <45 years old.
- Coved-type ECGs in family members.
- Inducibility of VT with programmed electrical stimulation .
- Syncope.
- Nocturnal agonal respiration.

b. J point elevation c. Coved type ST segment elevation d. Inverted T wave

The Strain Pattern

Left Ventricular Hypertrophy **KEY FACTS:**

- **♦** AS/HOCM have <u>exertional</u> symptoms
- ◆ Listen for aortic systolic murmur
- ◆ Both may have <u>marked voltages</u>
- **♦ HOCM** may have "needle" Q's

LVH with syncope is indicative of the beginnings of an infarct, where the vessels can't go through all that built up, hypertrophic tissue.

Therefore, LVH in the presence of a syncope event warrants a full workup with cardiology and likely a trip to the cath lab.

Pulmonary Embolus The had been a superfunction of the second The hall hall have more the transfer to the transfer t

Signs of a PE on ECG (may have all, some, or none of these signs):

- · tachycardia
- axis shift (right axis shift in the example ECG)
- deep, symmetrical T wave inversions in precordial leads
- · as it worsens, might see RBBB

Pulmonary Embolus KEY FACTS:

- **♦** ECG most commonly non-specific
- **♦** Look for signs of acute right heart strain:
 - ♦ Rightward axis shift (S1Q3T3)
 - **♦ RBBB**
 - ◆ Deep Flipped Ts V1-V4

QT Prolongation

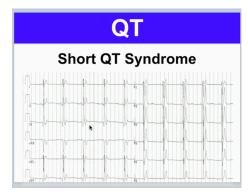
QT

QT Prolongation KEY FACTS:

- Main categories of culprit drugs:
- **♦** Antiarrhythmics
- (e.g. amiodorone)
- **♦** Antifungals ◆ Antipsychotic/antinausea (e.g. chlorpromazine)
 - (e.g. ketoconazole)
- ◆ QT Interval should be less than half of the RR interval

Quick trick on how to measure if QT interval is elongated or shortened:

The QT interval should be less than half the distance between R-R intervals.



QT

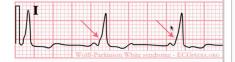
Short QT Syndrome KEY FACTS:

- ◆ Another "channelopathy" with genetic cause
- ♦ Short QT interval (< 340 ms)
- ◆ No change in QT with heart rate
- ◆ Peaked precordial T waves
- ◆ Treatment is with AICD

WPW

Wolf-Parkinson White

and other pre-excitation syndromes



WPW

Wolf-Parkinson White

and other pre-excitation syndromes

KEY FACTS:

- ♦ Not always visible on resting ECG
- ♦ Look for:
 - **♦Short PR interval**
 - **♦**Delta wave

The characteristic ECG findings in the Wolff-Parkinson-White syndrome are:

- Short PR interval (< 120ms)
- Broad QRS (> 100ms)
- A slurred upstroke to the QRS complex (the delta wave)

The ECG in SYNCOPE

6 Exotic things to look for

ARVD Arrhythmogenic Right Ventricular Dysplasia

BRUGADA Brugada syndrome

LVH Left Ventricular Hypertrophy (AS and HOCM) **Pulmonary Embolism**

QT QT Too Long / Too Short

PE

WPW Wolf-Parkinson-White