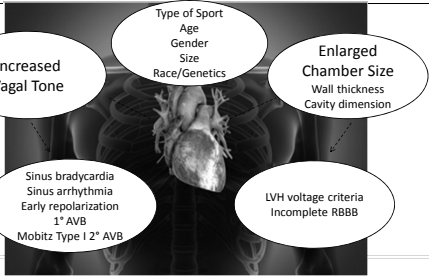




Applying the International Criteria for ECG Interpretation in Athletes to a pre-participation screening program

DAVE SIEBERT, MD, CAQSM 2018 COXHEALTH SPORTS MEDICINE CONFERENCE
 ASSISTANT PROFESSOR
 DEPARTMENT OF FAMILY MEDICINE UNIVERSITY OF WASHINGTON SPRINGFIELD, MISSOURI, USA
 UW HUSKY TEAM PHYSICIAN JUNE 23, 2018

Background – “Athlete’s heart”



Increased Vagal Tone

Type of Sport
Age
Gender
Size
Race/Genetics

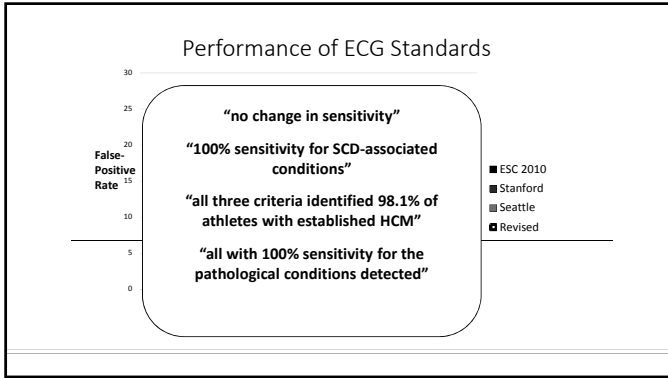
Enlarged Chamber Size
Wall thickness
Cavity dimension

Sinus bradycardia
Sinus arrhythmia
Early repolarization
1° AVB
Mobitz Type I 2° AVB

LVH voltage criteria
Incomplete RBBB

Ultimate question

- In the context of a **highly trained athlete**, which screening ECG changes can be considered **normal manifestations** of the “athlete’s heart,” and which should be considered **pathologic**?

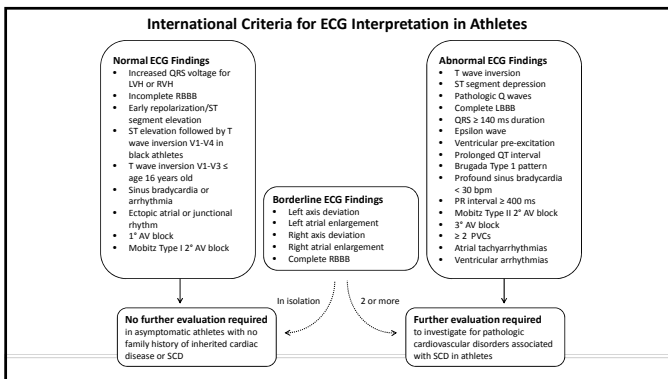


Performance of ECG criteria

Accuracy of the ECG for differential diagnosis between hypertrophic cardiomyopathy and athlete's heart: comparison between the European Society of Cardiology (2010) and International (2017) criteria
 Alessandro Zorzi,¹ Chiara Calore,¹ Riccardo Vio,¹ Antonio Pelliccia,² B/SM; Domenico Corrado² 2017

	ESC 2010	International Criteria 2017	
Specificity	86.9%	95.9%	p<0.001
Sensitivity	95.5%	93%	p=0.47

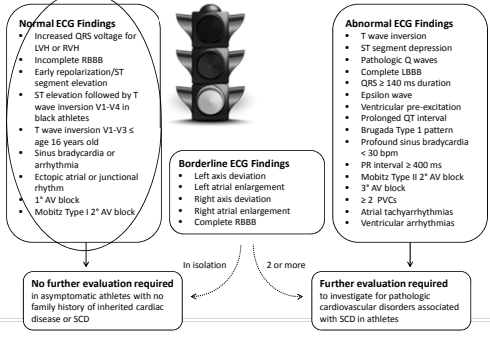
2017 International Criteria improved the specificity and reduced the number of unnecessary investigations by 69% (from 1:8 athletes to 1:24 athletes)



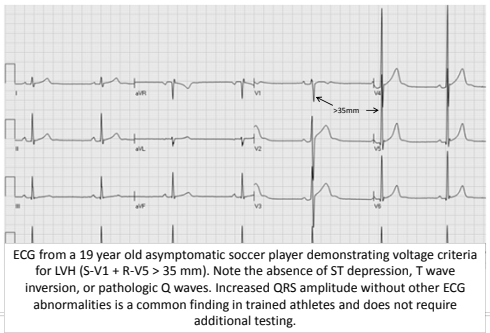
Clinical questions when interpreting ECGs

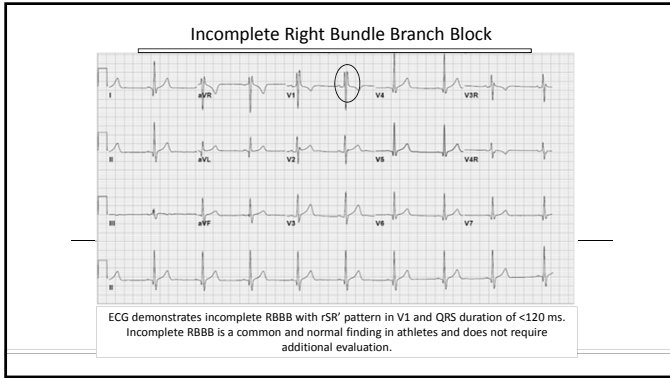
- 1) Is the ECG classified as:
 - A. Normal – no further evaluation needed
 - B. Abnormal – further evaluation needed
- 2) If the ECG is “abnormal”:
 - A. What is the specific ECG abnormality?
 - B. What is the appropriate next step in evaluation?
- 3) Relevant clinical information
 - A. Age, race, and sex of athlete
 - B. Asymptomatic and no family history of inherited cardiac disease or SCD?

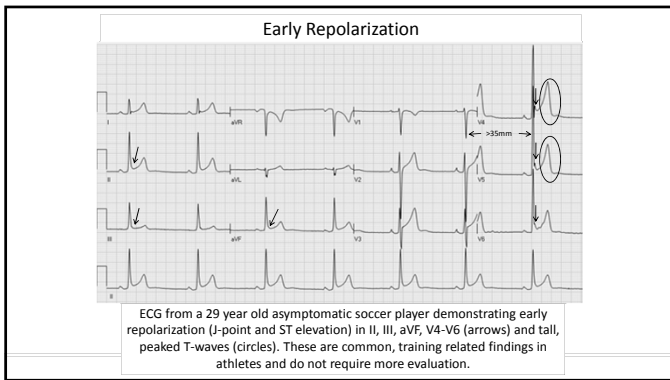
International Criteria for ECG Interpretation in Athletes

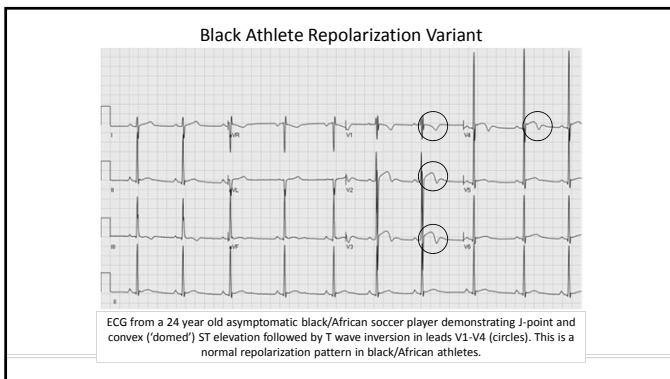


Isolated Increased QRS Voltage

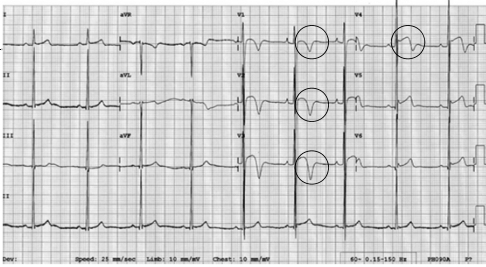






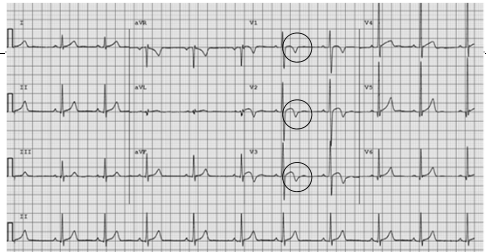


**Black Athlete Repolarization Variant:
Confined to Leads V1-V4**



ECG from a black/African athlete demonstrating voltage criterion for LVH, J-point elevation and convex ('domed') ST segment elevation followed by T wave inversion in V1-V4 (circles). This is a normal repolarization pattern in black athletes.

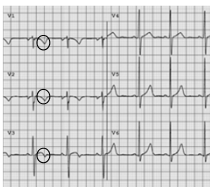
Juvenile T Wave Inversion



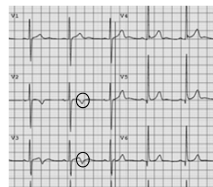
ECG from a 12 year old asymptomatic Caucasian female soccer player demonstrating the juvenile pattern of T wave inversion in leads V1-V3 (circles). This is a normal finding in athletes <16 years of age.

Juvenile T Wave Inversion

Age <16 yo; Independent of race; TWI in V1-V3; Does not extend to V4



13 yo Caucasian female



15 yo Asian female

No further evaluation needed

Junctional Escape Rhythm

A 28 year old Caucasian male demonstrating a junctional escape rhythm (red arrows). Note the constant RR interval between beats.

1° Atrioventricular Block

ECG shows 1° AV block (PR interval >200 ms). The PR interval is measured from the beginning of the P wave to the beginning of the QRS complex. In this ECG tracing, the PR interval is constant from beat to beat and measures 300 ms.

Mobitz Type I (Wenckebach) 2° AV Block

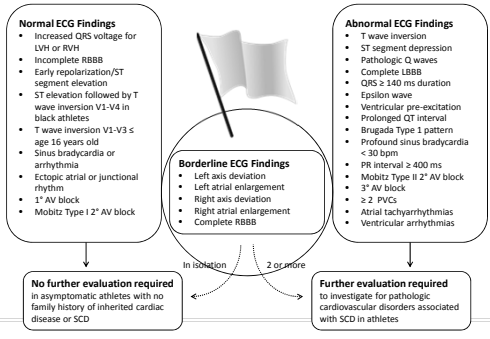
PR 140 ms PR 190 ms PR 200 ms P PR 140 ms PR 190 ms

- Mobitz Type I (Wenckebach) 2° AV block is demonstrated by progressively longer PR intervals until there is a non-conducted P-wave and no QRS.
- The first PR interval after the dropped beat is shorter than the last conducted PR interval prior to the dropped beat

Mobitz Type I (Wenckebach) 2° AV Block



International Criteria for ECG Interpretation in Athletes



European Heart Journal (2013) 34, 1041–1048
doi:10.1093/eurheartj/ehs399

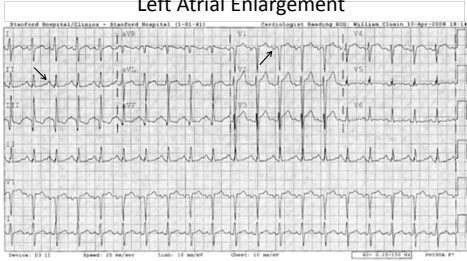
CLINICAL RESEARCH
Sports cardiology

Should axis deviation or atrial enlargement be categorised as abnormal in young athletes? The athlete's electrocardiogram: time for re-appraisal of markers of pathology 2013

Sabira Gazi^{1,2}, Nabeel Sheikh¹, Saqib Ghani¹, Abbas Zaidi¹, Mathew Wilson¹, Harsharan Raju¹, Andrew Cox¹, Matt Reed¹, Michael Papadakis¹, and Sanjay Sharma^{1,3*}

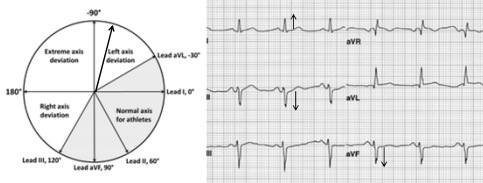
- N=2533 athletes: no athlete with isolated left or right axis deviation or atrial enlargement showed evidence of cardiomyopathy
- N=171 patients with HCM: co-existing ECG abnormalities present in 89% with axis deviation or atrial enlargement

Left Atrial Enlargement



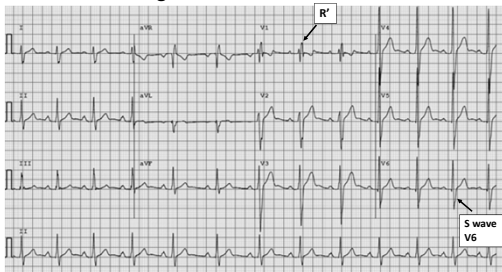
ECG demonstrates left atrial enlargement, defined as a prolonged P wave duration of >120 ms in leads I or II with negative portion of the P wave ≥ 1 mm in depth and ≥ 40 ms in duration in lead V1

Left Axis Deviation



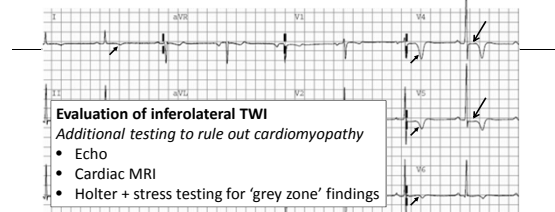
ECG demonstrates abnormal left axis deviation defined as frontal plane QRS axis of less than -30° . The QRS is positive in lead I and negative in aVR and lead II. The QRS axis shown here is about -70° .

Right Bundle Branch Block



- 19 yo Caucasian male athlete with complete RBBB. The QRS duration is ≥ 120 ms with rSR' pattern in V1 and S wave wider than R wave in V6.
- When found in isolation without other borderline or abnormal findings, and without other clinical markers of concern, complete RBBB does not require more investigation.

Inferolateral T Wave Inversion and ST Depression



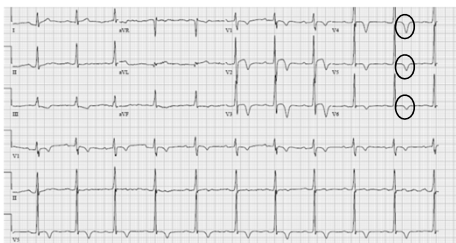
Evaluation of inferolateral TWI

Additional testing to rule out cardiomyopathy

- Echo
- Cardiac MRI
- Holter + stress testing for 'grey zone' findings

Abnormal ECG from a patient with hypertrophic cardiomyopathy. Note T wave inversions in I, aVL, and V4-V6 (red arrows), as well as ST segment depression in V4-V5 (black arrows).

Lateral T Wave Inversion



Markedly abnormal ECG showing TWI ≥ 2 mm in V4-V6. Note that the ST segment preceding TWI in V4-6 is flat or downsloping.

Table 2 Evaluation of ECG abnormalities in athletes

ECG abnormality	Potential cardiac diagnosis*	Recommended evaluation†	Considerations
T wave inversion in the lateral or inferolateral leads	HCM DCM LSCM APVC (with predominant left ventricular involvement) Myocarditis	Echocardiogram Cardiac MRI Exercise ECG test Minimum 24-hour ECG monitor	Lateral or inferolateral T wave inversion is common in primary myocardial disease. Cardiac MRI should be a routine diagnostic test for this ECG phenotype and is superior to echocardiography for detecting apical HCM, left ventricular hypertrophy localized to the free lateral wall, APVC, with predominant left ventricular involvement and myocarditis. If cardiac MRI is not available, echocardiography with contrast should be considered as an alternative investigation for apical HCM in patients with deep T wave inversion in leads V4-V6. Consider family evaluation if available and genetic screening. Annual follow-up testing is recommended for athletes with lateral T wave inversion in leads V4-V6.
T wave inversion isolated to the inferior leads	HCM LSCM Myocarditis	Echocardiogram	Consider cardiac MRI based on echocardiogram findings or clinical suspicion.
T wave inversion in the anterior leads‡	APVC DCM	Echocardiogram Cardiac MRI Exercise ECG test Signal-averaged ECG	The extent of investigations may vary based on clinical suspicion for APVC, and results from initial testing.
ST segment depression	HCM DCM LSCM APVC Myocarditis	Echocardiogram	Consider cardiac MRI and additional testing based on echocardiogram findings or clinical suspicion.
Pathological Q waves	HCM DCM LSCM Myocarditis Prior myocardial infarction	Echocardiogram Coronary artery disease risk factor assessment Repeat ECG for repeat (V1-V2) Q waves; done investigations recommended if repeat Q waves are persistent	Consider cardiac MRI (with perfusion study if available) based on echocardiogram findings or clinical suspicion. In the absence of cardiac MRI, consider exercise stress testing, dobutamine stress echocardiogram or a myocardial perfusion scan for evaluation of coronary artery disease in athletes with suspicion of prior myocardial infarction or multiple risk factors for coronary artery disease.
Complete left bundle branch block	DCM HCM LSCM Sarcoidosis Myocarditis	Echocardiogram Cardiac MRI (with stress perfusion study)†	A comprehensive cardiac evaluation to rule out myocardial disease should be considered.

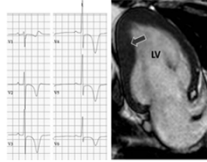
Evaluation of Lateral or Inferolateral TWI

Comprehensive evaluation to r/o cardiomyopathy

Echocardiogram

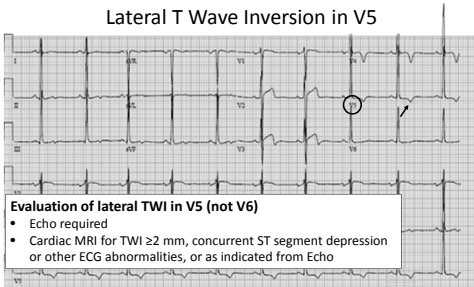
Cardiac MRI should be a routine diagnostic test for this ECG phenotype

24 hour ECG monitor + stress testing for 'grey zone' findings



Apical HCM

Lateral T Wave Inversion in V5

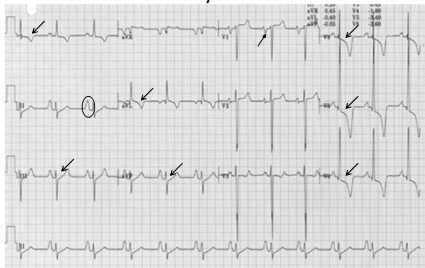


Evaluation of lateral TWI in V5 (not V6)

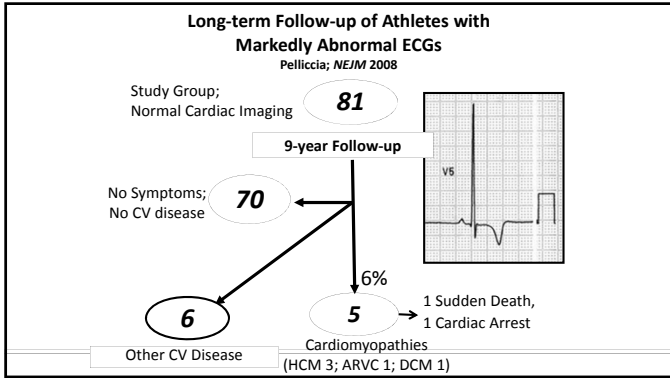
- Echo required
- Cardiac MRI for TWI ≥ 2 mm, concurrent ST segment depression or other ECG abnormalities, or as indicated from Echo

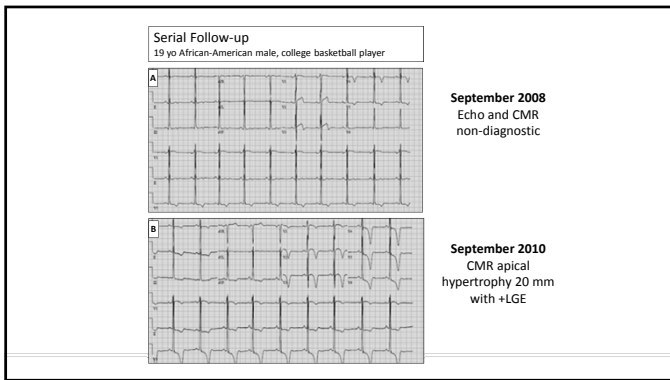
ECG in a 18 yo African-American male. TWI extending to V5 is considered abnormal. Only one lead of TWI required in V5 or V6.

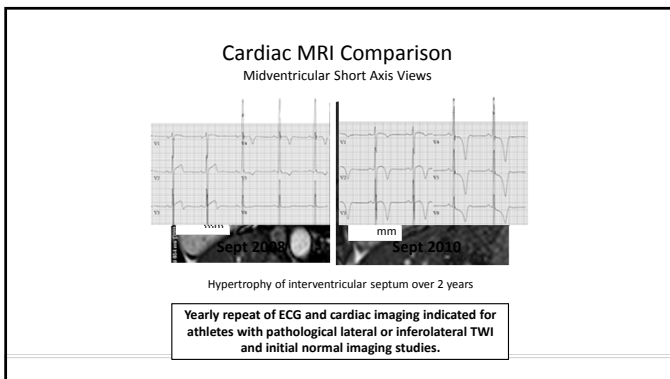
The "Markedly Abnormal" ECG



ECG from a patient with HCM demonstrating QRS voltage criterion for LVH in association with deep T wave inversion and ST segment depression predominantly in the lateral leads (I, aVL, V4-V6), voltage criterion for left and right atrial enlargement, and left axis deviation.







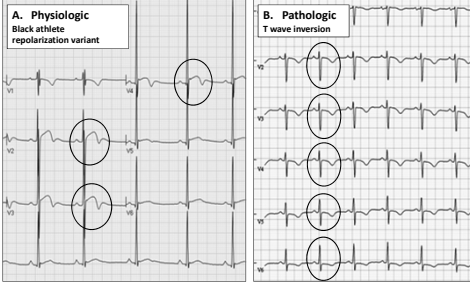
Normal or Abnormal?



Evaluation of Inferolateral TWI

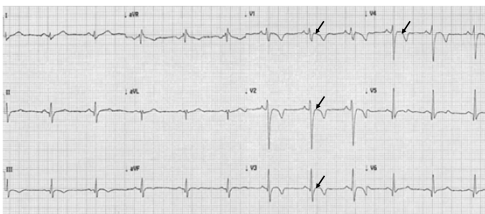
Additional testing to rule out cardiomyopathy

- Echo
- Cardiac MRI
- Holter + stress testing for 'grey zone' findings
- If initial studies are non-diagnostic → serial (annual) follow-up with ECG + Echo (at minimum); cardiac MRI for changes in ECG or Echo



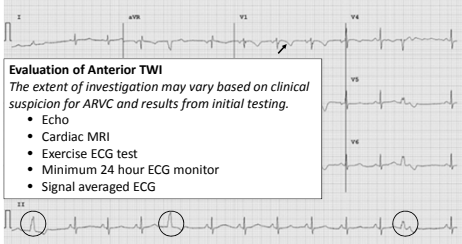
Physiologic (A) and pathologic T wave inversion (B). Panel A demonstrates physiologic repolarization in a black athlete with TWI in V1-V4 preceded by J-point and convex 'domed' ST segment elevation (green circles). Panel B demonstrates pathologic TWI in V1-V6 with absent J-point elevation and a downsloping ST segment (red circles).

Anterior T Wave Inversion



21 yo Caucasian male with ECG demonstrating anterior T wave inversion (V1-V4) preceded by a non-elevated J-point and ST segment. Delayed S wave upstroke in V2 and low voltage (<5 mm) QRS complexes in limb leads I and aVL suggest possible ARVC.

Anterior T Wave Inversion



Evaluation of Anterior TWI
 The extent of investigation may vary based on clinical suspicion for ARVC and results from initial testing.

- Echo
- Cardiac MRI
- Exercise ECG test
- Minimum 24 hour ECG monitor
- Signal averaged ECG

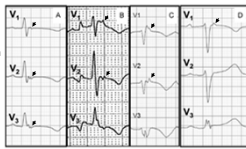
ECG from a patient with ARVC. Note pathological TWI in V1-V3 (arrows) preceded by a flat or downsloping ST segment and without J-point elevation. PVCs also present (circles).

Epsilon Wave

Definition

2010 ARVC/D Task Force Criteria:
 "Reproducible low amplitude signal between end of QRS complex to onset of the T wave in the right precordial leads (V1 to V3)."

2017 Athlete 'International Criteria':
 "Distinct low amplitude signal (small positive deflection or notch) between the end of the QRS complex and onset of the T wave in leads V1-V3."



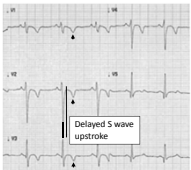
Platonov et al.
Heart Rhythm, 2016

Epsilon Wave

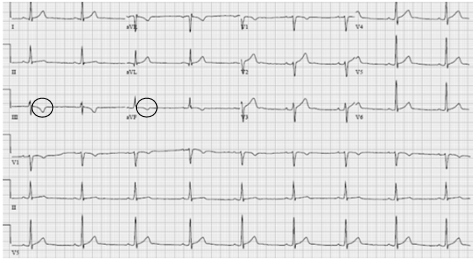
Epsilon waves are typically a manifestation of more advanced disease and unlikely to be an isolated ECG finding

In patients with ARVC that express an epsilon wave:

- 89% also manifest TWI in the right precordial leads
- 100% have a delayed S wave upstroke (prolonged terminal activation duration) >55 ms from the nadir of the S wave to the end of the QRS complex
- Thus, a suspected epsilon wave should prompt evaluation for other ECG abnormalities suggestive of ARVC (TWI; delayed S wave upstroke; low limb lead voltage).

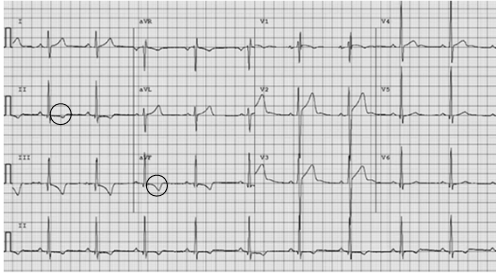


Inferior T Wave Inversion Normal or Abnormal?



18 yo Caucasian male with TWI in leads III and aVF. Lead III is excluded (need 2 contiguous leads). In the absence of symptoms or other clinical markers of concern, this is a normal ECG and no further evaluation is needed.

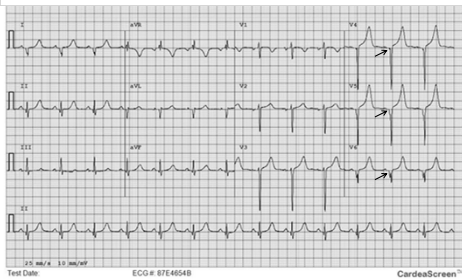
Inferior T Wave Inversion



ECG demonstrates TWI in the inferior leads II and aVF. This is an abnormal ECG and requires further evaluation (echocardiogram).

Pathologic Q Waves

New Criteria: Q/R ratio ≥ 0.25 or ≥ 40 ms in duration



Test Date: ECG #: 67E46548 CardeaScreen™

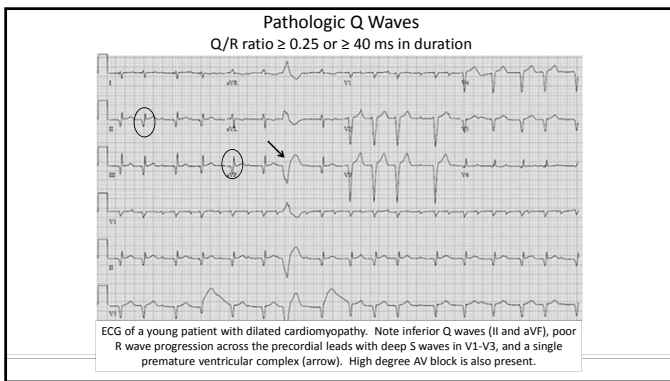


Table 2 Evaluation of ECG abnormalities in athletes

ECG abnormality	Potential cardiac disease*	Recommended evaluation†	Considerations
T wave inversion in the lateral or inferior leads	HCM DCM LINC APVC (with predominant left ventricular involvement) Myocarditis	Echocardiogram Cardiac MRI Exercise ECG test Minimum 24-hour ECG monitor	Lateral or inferior T wave inversion is common in primary myocardial disease. Cardiac MRI should be a routine diagnostic test for this ECG phenotype and is superior to echocardiography for detecting apical HCM. Left ventricular hypertrophy localized to the free lateral wall (AVC) with predominant left ventricular involvement and myocarditis. If cardiac MRI is not available, echocardiography with contrast should be considered as an alternative investigation for apical HCM in patients with deep T wave inversion in leads V5-V6. Consider family evaluation if available and genetic screening. Annual follow-up testing is recommended for regular athletic career in athletes with normal results.
T wave inversion isolated to the inferior leads	DCM LINC Myocarditis	Echocardiogram	Consider cardiac MRI based on echocardiogram findings or clinical suspicion.
T wave inversion in the anterior leads*	APVC DCM	Echocardiogram Cardiac MRI Exercise ECG test Minimum 24-hour ECG monitor Signal averaged ECG	The extent of investigations may vary based on clinical suspicion for APVC and results from initial testing.
ST segment depression	HCM DCM LINC APVC Myocarditis	Echocardiogram	Consider cardiac MRI and additional testing based on echocardiogram findings or clinical suspicion.
Pathological Q waves	DCM LINC Myocarditis Prior myocardial infarction	Echocardiogram Coronary artery disease risk factor assessment Repeat ECG for septal (V1-V2) QS pattern; above investigations recommended if septal Q waves are persistent	Consider cardiac MRI (with perfusion study if available) based on echocardiogram findings or clinical suspicion. In the absence of cardiac MRI, consider exercise stress testing, dobutamine stress echocardiogram or a myocardial perfusion scan for evaluation of coronary artery disease in athletes with suspicion of prior myocardial infarction or multiple risk factors for coronary artery disease.
Complete left bundle branch block	DCM LINC Myocarditis	Echocardiogram Cardiac MRI (with stress perfusion study)†	A comprehensive cardiac evaluation to rule out myocardial disease should be considered.

Evaluation of Pathologic Q Waves

Echocardiogram

- Consider cardiac MRI (with perfusion study) based on echocardiogram findings and clinical suspicion

Coronary artery disease risk factor assessment

- Consider exercise stress testing, dobutamine echo, or myocardial perfusion scan in athletes >30 yo or if multiple risk factors for CAD are present

Repeat ECG for septal (V1-V2) QS pattern

- "pseudo-septal" infarct pattern from high lead misplacement

Anterior Q Waves V1-V2 / QS Pattern

Evaluation of Anterior Q waves
 1) Repeat ECG
 2) If Q waves persist → Echo
 3) CAD risk assessment

- Stress testing for multiple risk factors or age >30 years

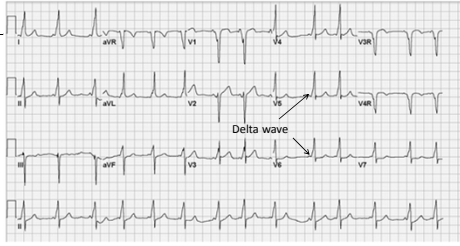
ECG in a 22 yo African-American male. Anterior Q waves can be from incorrect high lead placement of V1-V2 (ie "pseudo-septal" infarct).

Anterior Q Waves?

Repeat ECG in same athlete shows RS in V2 (arrows). This suggests QS pattern was lead placement issue. No further evaluation needed.

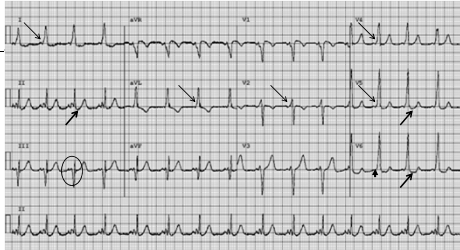
ECG abnormality	Definition
T wave inversion	<ul style="list-style-type: none"> ▶ Asymetric <ul style="list-style-type: none"> ▶ V2-V4 <ul style="list-style-type: none"> - excludes black athletes with J point elevation and convex ST segment elevation followed by TW in V2-V4; athletes < age 16 with TW in V1-V2 and biphasic T waves in V3-V4 ▶ I and aVL, V5 and/or V6 (only one lead of TW required in V5 or V6) ▶ II and aVF, V5-V6, I and aVL ▶ II and aVF ▶ Symetric <ul style="list-style-type: none"> ▶ I and aVL, V5 and/or V6 (only one lead of TW required in V5 or V6) ▶ II and aVF, V5-V6, I and aVL ▶ II and aVF
ST segment depression	<ul style="list-style-type: none"> ▶ Asymetric <ul style="list-style-type: none"> ▶ ≥0.5mm in depth in two or more contiguous leads ▶ Symetric <ul style="list-style-type: none"> ▶ ≥0.5mm in depth in two or more contiguous leads
Pathological Q waves	<ul style="list-style-type: none"> ▶ QRS ratio ≥25% or others in duration in two or more leads (including III and aVF)
Complete left bundle branch block	<ul style="list-style-type: none"> ▶ QRS >120ms, predominantly negative QRS complex in lead V1 (QS or rS) and upright notched or slurred S wave in leads I and V6
Profound non-specific intraventricular conduction delay	<ul style="list-style-type: none"> ▶ Any QRS duration >140ms
Episodic wave	<ul style="list-style-type: none"> ▶ Distinct low amplitude signal (small positive deflection or notch) between the end of the QRS complex and onset of the T wave in leads V3-V5
Ventricular pre-excitation	<ul style="list-style-type: none"> ▶ PR interval <120ms with a delta wave (slowed upstroke in the QRS complex) and with QRS >120ms
Prolonged QT interval*	<ul style="list-style-type: none"> ▶ QTc >470ms (male) ▶ QTc >480ms (female) ▶ QTc >490ms (black/CP)
Brugada type 1 pattern	<ul style="list-style-type: none"> ▶ Coved pattern: initial ST elevation ≥2mm high take-off with downsloping ST segment elevation followed by a negative symmetric T wave in leads V1-V3
Profound sinus bradycardia	<ul style="list-style-type: none"> ▶ <40bpm per minute or sinus pauses >2s
Profound 1° atrioventricular block	<ul style="list-style-type: none"> ▶ >400ms
Mild to moderate 2° atrioventricular block	<ul style="list-style-type: none"> ▶ Intermittently non-conducted P waves with a fixed PR interval
3° atrioventricular block	<ul style="list-style-type: none"> ▶ Complete heart block
Atrial tachyarrhythmias	<ul style="list-style-type: none"> ▶ Sinus bradycardia, sinus tachycardia, sinus flutter, atrial flutter
Premature ventricular contractions	<ul style="list-style-type: none"> ▶ ≥ premature ventricular contractions per 15s testing
Ventricular arrhythmias	<ul style="list-style-type: none"> ▶ Couplets, triplets and non-sustained ventricular tachycardia

Ventricular Pre-excitation / Wolff-Parkinson-White



ECG demonstrating the classic findings of WPW with a short PR interval (<120 ms), delta wave (slurred QRS upstroke), and prolonged QRS (>120 ms).

Ventricular Pre-excitation / WPW Pattern



17 yo asymptomatic female with negative family history. ECG demonstrates a short PR interval and delta waves (black arrows). Other findings suggestive of WPW include a large Q wave in lead III (red circle), absence of a Q wave in V6 (blue arrowhead), and ST segment depression in V5-V6 and lead II (red arrows).

Table 2 Evaluation of ECG abnormalities in athletes

ECG abnormality	Potential cardiac disease*	Recommended evaluation†	Considerations
Multiple premature ventricular contractions	HCM DCM LVNC ARVC Myocarditis Sarcoidosis	Echocardiogram 24-hour ECG monitor Exercise ECG test	If >2000 PVCs or non-sustained ventricular tachycardia are present on initial testing, comprehensive cardiac testing (include of cardiac MRI) is warranted to investigate for myocardial disease. Consider signal-averaged ECG.
Ventricular pre-excitation	Wolff-Parkinson-White	Exercise ECG test Echocardiogram	Absent cessation of the delta wave (pre-excitation) on exercise ECG denotes a low-risk pathway. Electrophysiological study for risk assessment should be considered if a low-risk pathway cannot be confirmed by non-invasive testing. Consider electrophysiological study for moderate to high intensity sports.
Prolonged QTc	Long QT syndrome	Repeat resting ECG on separate day Review for QT-prolonging medication Acquire ECG of first-degree AV block if possible	Consider exercise ECG test, laboratory electrolyte screening, family screening and genetic testing when clinical suspicion is high. Consider direct referral to a heart rhythm specialist or sports cardiologist for a QTc >500 ms.
Brugada type 1 pattern	Brugada syndrome	Refer to cardiologist or heart rhythm specialist	Consider high precardial lead ECG with leads V1 and V2 in second intercostal space or sodium channel blockade if Brugada pattern is indeterminate. Consider genetic testing and family screening.
Profound sinus bradycardia <30 beats per minute	Myocardial or electrical disease	Repeat ECG after mild aerobic activity	Consider additional testing based on clinical suspicion.

Evaluation of Ventricular Pre-excitation

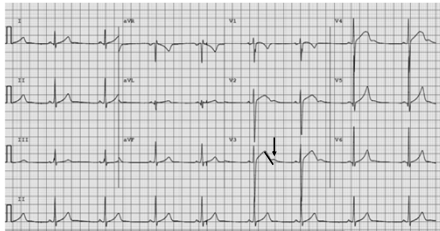
Exercise ECG test

- Abrupt cessation of the delta wave (pre-excitation) denotes a low risk pathway
- EP study should be considered if a low risk accessory pathway cannot be confirmed by non-invasive testing
- Consider EP study for moderate to high intensity sports

Echocardiogram

- Association of pre-excitation with Ebstein's anomaly and cardiomyopathy

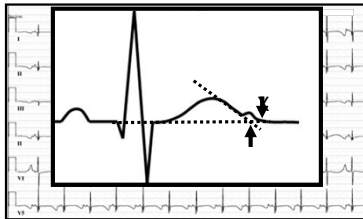
Long QT Syndrome?



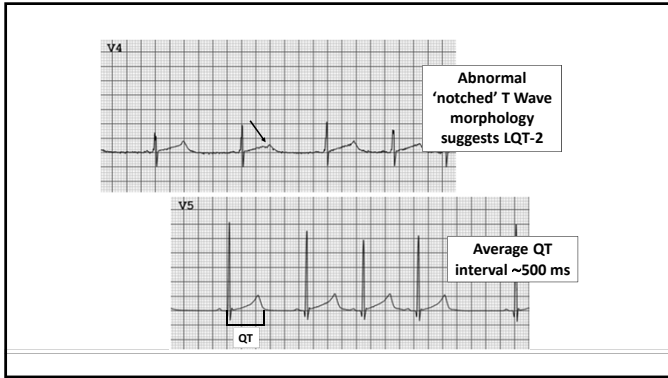
Normal ECG

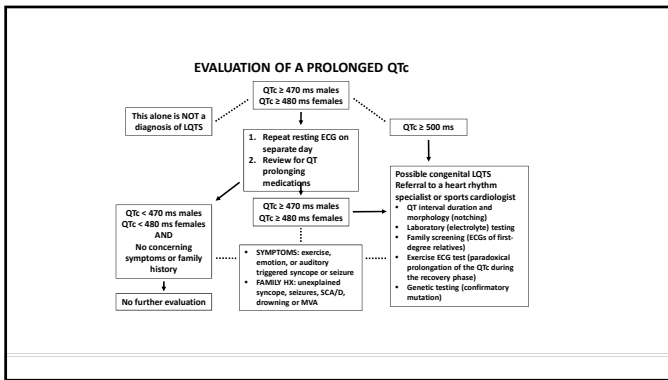
- QTc is normal
- Don't include the U wave in anterior precordial leads!
 - "Teach-the-tangent" or "Avoid-the-tail" method for manual measurement of the QT interval

No further evaluation needed



This figure illustrates the "Teach-the-tangent" or "Avoid-the-tail" method for manual measurement of the QT interval. A straight line is drawn on the downslope of the T wave to the point of intersection with the isoelectric line. The U wave is not included.





Abnormal ECG = Temporary Restriction?

Temporary restriction from athletic activity should be considered for athletes with abnormal ECGs, especially when there is high clinical suspicion for pathologic cardiac disease, until secondary investigations are completed.

Conditional clearance for sports participation pending further evaluation can be considered on a case-by-case basis.

Thank you!

- Questions or clarifications?

Please feel free to email me:
siebert@uw.edu
