Sudden Cardiac Death in Young Athletes

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Disclosure

Christopher M. Rausch, MD

No relevant financial disclosures with ineligible companies to report.



Overview

1 Case Presentation

2 Sudden cardiac death discussion

3 Screening and prevention



Case Presentation

History of Present Illness

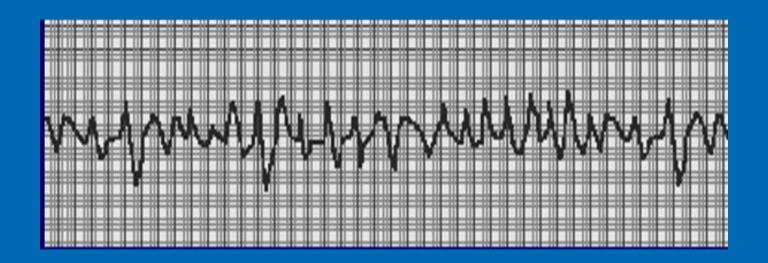
- 13-year-old male collapsed and became cyanotic while playing basketball
- Bystander unable to find pulse and began CPR
- EMS arrives and places defibrillator patches on the chest



Google image, June 2022



Ventricular Fibrillation (V Fib)





Case Presentation

History of Present Illness

- Defibrillated with 100 joules causing successful conversion to sinus rhythm
- Bag and mask ventilation provided with improvement in mental status within 10-15 minutes
- Transferred to ICU



Case Presentation

Past Medical History

- Episode of syncope 1 year prior
- No other major illnesses or hospitalizations

Family History

- Mother with episodes of lightheadedness and palpitations
- No history of sudden death
- No known inheritable diseases



What is the most common cardiac cause of sudden death in athletes?

- 1. Tetralogy of Fallot
- 2. Commotio cordis
- 3. Hypertrophic cardiomyopathy
- 4. Long QT syndrome



Sudden Cardiac Death

Incidence

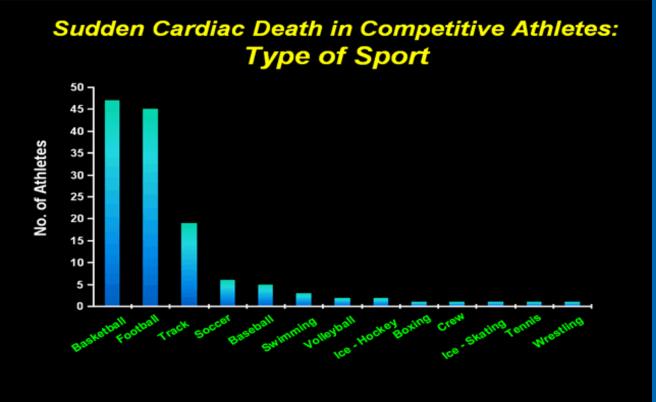
- 1/40,000 1/300,000
 - AHA: 350,000 Americans each year (7,000 are children)
- Male>Female
- Occurs most commonly in the most common sports
- Two-thirds of the deaths occur during exercise, making SCA the leading cause of death in young athletes

Table 2 Incidence of sudden death stratified by athletic or general population						
Country	Reference	Population	Years	Incidence per 100,000 person-year		
Italy	Corrado [18]	Athletes	1980-1981	3.6		
Italy	Corrado [18]	Athletes	2007-2008	0.4		
Israel	Steinvil [39]	Athletes	1985-1997	2.54		
Israel	Steinvil [39]	Athletes	1998-2009	2.66		
USA	VanCamp [8]	Athletes	1983-1993	0.33		
USA	Maron [40]	Athletes	1985-2006	0.44		
Denmark	Holst [27]	Athletes	2000-2006	1.21		
Denmark	Holst [27]	All children	2000-2006	3.76		
Japan	Tanaka [38]	All children	1989-1997	1.32		
USA	Eckart [50]	Military recruits	1997-2001	13		
USA and Canada	Atkins [3]	All children ages 1-11	2005-2007	3.73		
USA and Canada	Atkins [3]	All children ages 12-19	2005-2007	6.37		

J Interv Card Electrophysical (2013) 36: 167-175

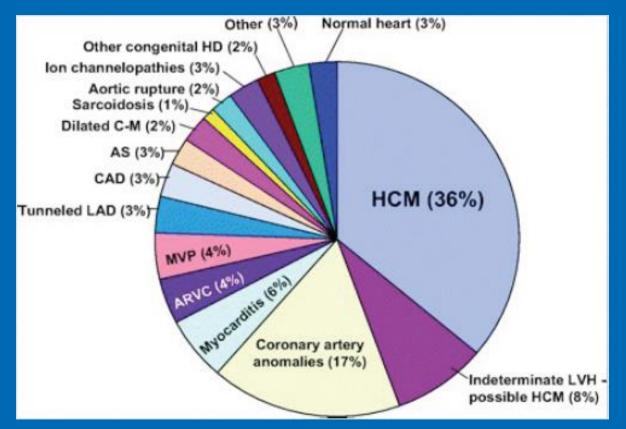


Sudden Cardiac Death





Sudden Cardiac Death





Cardiomyopathies

Hypertrophic Cardiomyopathy (HCM)

Arrhythmogenic Right Ventricular Dysplasia/Cardiomyopathy (ARVD)

Dilated Cardiomyopathy

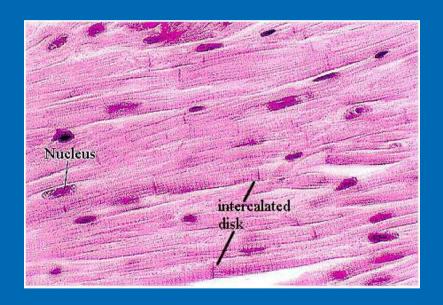


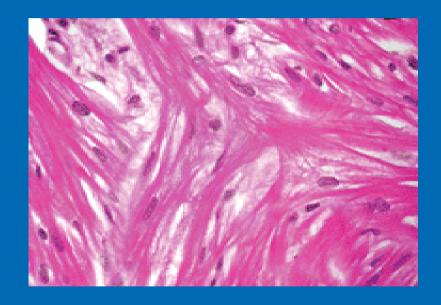
Cardiomyopathies: Hypertrophic Cardiomyopathy

- Prevalence 1:500
- Left ventricular hypertrophy
 - May be concentric or variable pattern
 - Disorganized myocardial architecture
 - Predisposition for ventricular arrhythmias
- Autosomal dominant
 - Any of 12 different cardiac sarcomere genes
 - Over 400+ mutations identified



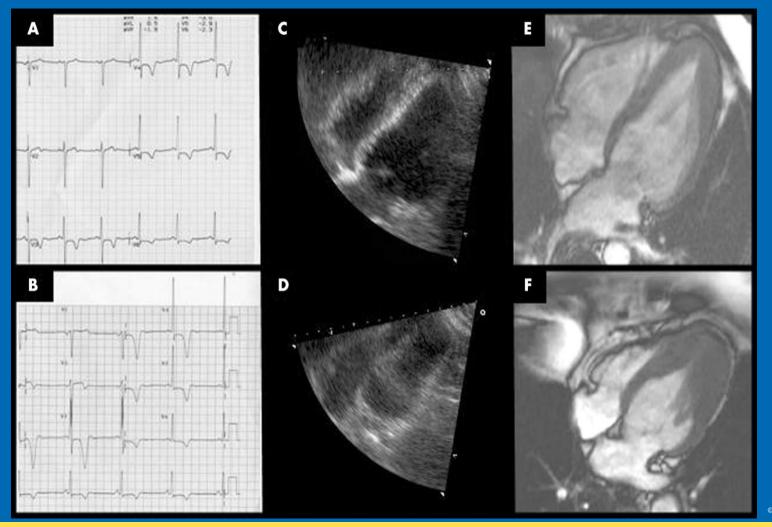
Hypertrophic Cardiomyopathy





AHA, Circulation Research; 2017



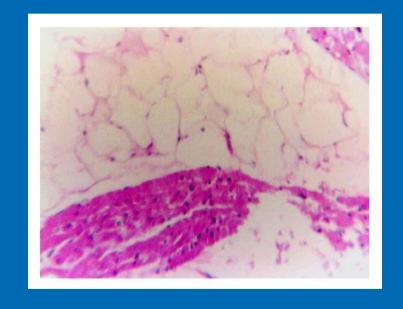


Moon, Heart; 2004

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Cardiomyopathies: Arrhythmogenic Ventricular Cardiomyopathy

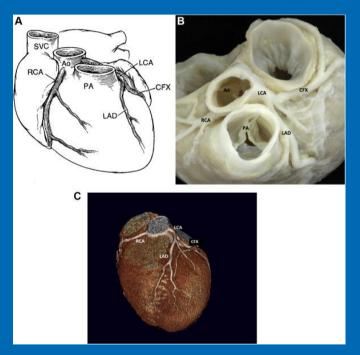
- Incidence 1:5000
- Autosomal dominant primarily (some recessive)
 - Multiple different genetic variants identified (at least 17)
 - Right and Left ventricular involvement
- Linked to disorder of desmosomes and adherens in the majority of cases
- Fibrofatty replacement of myocardium





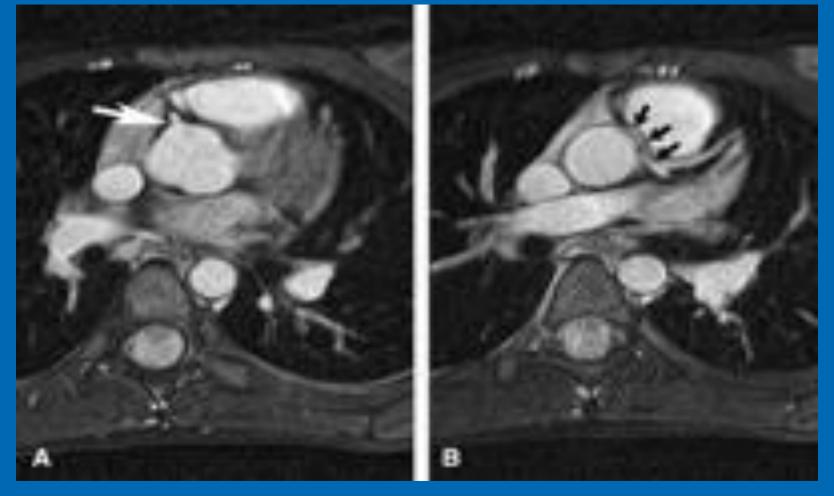
Frances, Int J of Cardiol; 2006

Congenital Coronary Artery Anomalies

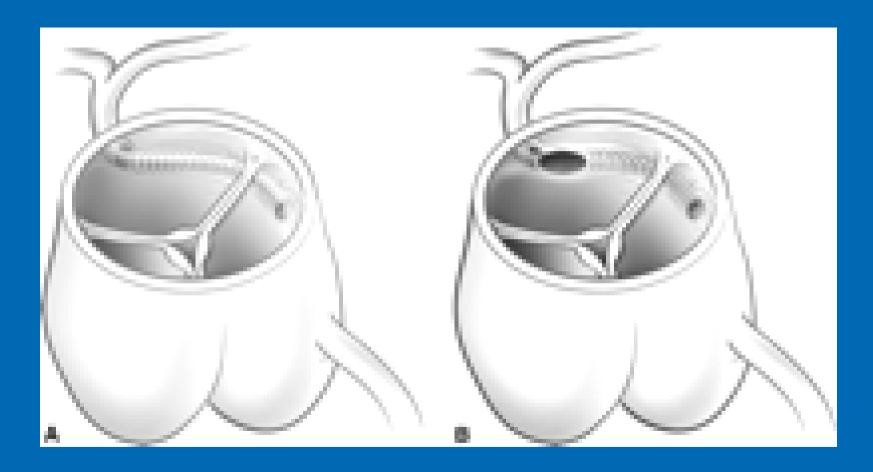


Frommelt, JASE: 2020

- Coronary originating from the opposite sinus of Valsalva
 - Acute angle takeoff
 - Intramural coronary
- Myocardial bridge

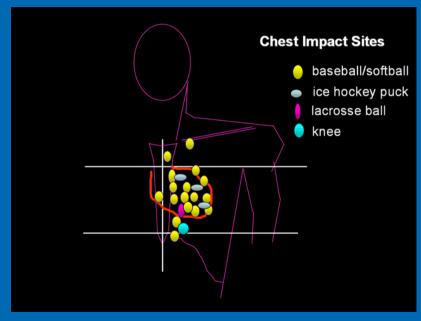


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Commotio Cordis

- Blunt, low energy precordial blow
 - 15-30 ms before Twave peak
 - VF
- No structural disease
- Mean age 13 yrs



Link, Circ; 1999



Myocarditis

- Inflammatory disease of myocardium
- Most common etiology in North America is viral
 - Coxsackie A and B, ECHO viruses, and Influenza viruses
 - Drugs and toxins
- Inflammation, focal necrosis, and replacement fibrosis predisposes to ventricular arrhythmias



COVID-19 Clinical Pathways

Cardiac Evaluation for Post COVID-19 (SARS-CoV-2) Infection Return to Play in Children and Adolescents Return to Play Clinical Pathway

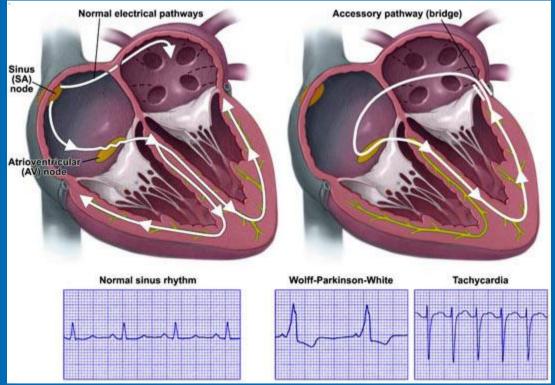


Arrhythmias

- Wolff Parkinson White (WPW)
- Long QT syndrome
- Catecholaminergic Polymorphic Ventricular Tachycardia (CPVT)
- Brugada syndrome

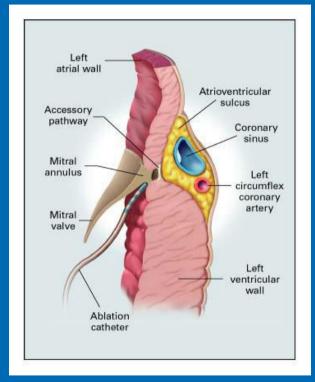


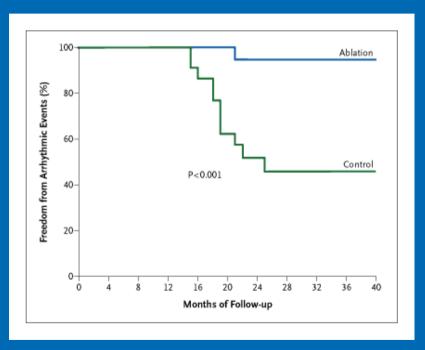
Wolff-Parkinson-White





Wolff-Parkinson-White









Morady, NEJM; 1999

- Inherited mutations of cardiac ion channels
 - Autosomal dominant and recessive forms
 - 13 distinct genetic loci identified
 - Affects primarily K and Na channels
- Abnormal ion channel function results in delayed repolarization

Table 1. LQTS Genes							
Gene	Syndrome	Frequency	Locus	Protein (Functional Effect)			
KCNQ1 (LQT1)	RWS, JLNS	40–55	11p15.5	Kv7.1 (↓)			
KCNH2 (LQT2)	RWS	30-45	7q35-36	Kv11.1 (↓)			
SCN5A (LQT3)	RWS	5-10	3p21-p24	NaV1.5 (†)			
ANKB (LQT4)	RWS	<1%	4q25-q27	Ankyrin B (↓)			
KCNE1 (LQT5)	RWS, JLNS	<1%	21q22.1	MinK (↓)			
KCNE2 (LQT6)	RWS	<1%	21q22.1	MiRP1 (↓)			
KCNJ2 (LQT7)	AS	<1%	17q23	Kir2.1 (↓)			
CACNA1C (LQT8)	TS	<1%	12p13.3	L-type calcium channel (†)			
CAV3 (LQT9)	RWS	<1%	3p25	Caveolin 3 (↓)			
SCN4B (LQT10)	RWS	<1%	11q23.3	Sodium channel-β4 (↓)			
AKAP9 (LQT11)	RWS	<1%	7q21-q22	Yotiao (↓)			
SNTA1 (LQT12)	RWS	<1%	20q11.2	Syntrophin α 1 (\downarrow)			
KCNJ5 (LQT13)	RWS	<1%	11q24	Kir3.4 (↓)			

Schwartz et al ,Circ Arr; 2012



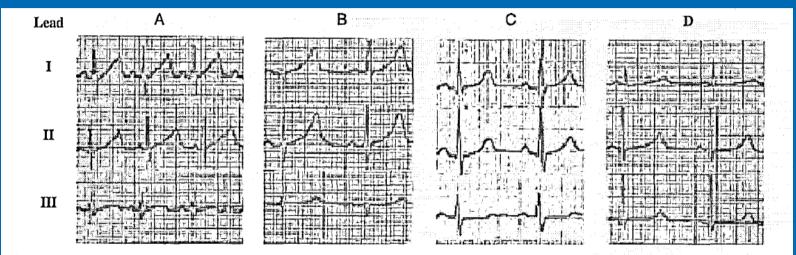


FIGURE 1—Four typical LQT1 patterns. A. Infantile pattern. B. Broad-based T-wave pattern. C. Normal-appearing T-wave pattern. D. Late-onset normal-appearing T-wave pattern. (Reproduced with permission from Zhang et al. (47)).



Torsades de pointes





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Probability of LQTS

 \leq 1 pt = Low

2-3 pts = Intermediate

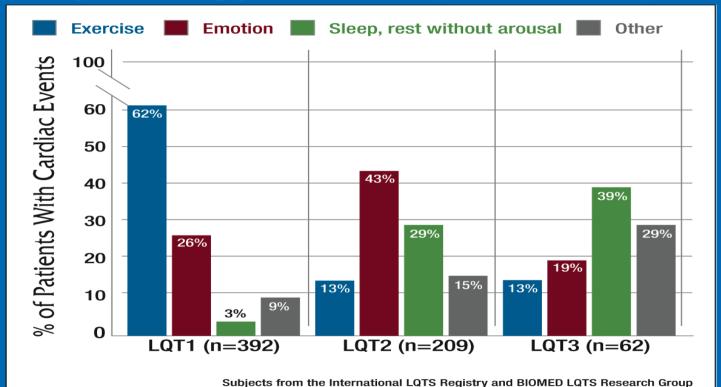
≥ 4 pts = High

	Points
ECG findings ^a	
A. QTc ^b	
> 480 ms	3
460–470 ms	2
450 ms (in males)	1
B. Torsade de pointes	2
C. T-wave alternans	1
D. Notched T wave in three leads	1
E. Low HR for age ^c	0.5
Clinical history	
A. Syncope ^d	
With stress	2
Without stress	1
B. Congenital deafness	0.5
Family history	
A. Family members with definite LQTS'	1
 B. Unexplained sudden cardiac death at younger than age 30 among immediate family members 	0.5

Schwartz, Circ; 1993

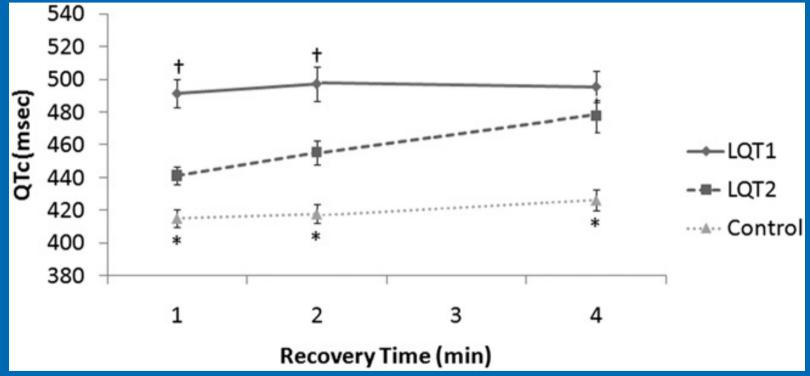


Arrhythmogenic Triggers Differ by LQTS Subtype





Prolongation of QTc with exercise





Chattha, et al. Heart Rhythm 2010

Catecholaminergic Polymorphic Ventricular Tachycardia (CPVT)

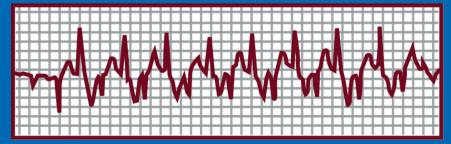
Inherited arrhythmogenic disorder characterized by ventricular ectopy induced by exercise or emotional stress.

- Most commonly caused by mutations of the cardiac ryanodine receptor gene (RYR2).
- ~1%-2% of CPVT is caused by recessive mutations of the calsequestrin (CASQ2) gene.
- If left untreated, CPVT is lethal in 30%-50% of patients.
- The onset of CPVT symptoms typically occurs in childhood and adolescence.



CPVT

- CPVT cannot be diagnosed on the basis of a resting ECG.
- Exercise stress testing is an important part of a CPVT workup.
 - However, in as many as 20% of CPVT patients, formal exercise stress testing will not produce ventricular ectopy.
- During exercise stress testing, bidirectional VT with a beat-to-beat 180-degree rotation of the QRS complex is often observed.





CPVT





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Brugada Syndrome

- Prevalence estimates vary from 5 66 / 10,000
- Increased prevalence in Asia, where identified as sudden unexplained nocturnal death syndrome (SUNDS)
- Male predominance in its phenotype (~8:1)
- Average age at presentation is 40 yr, but cases diagnosed from infancy into late 70s
- As in long QT syndrome Type 3, arrhythmia and death common during sleep and bradycardia



Brugada Syndrome

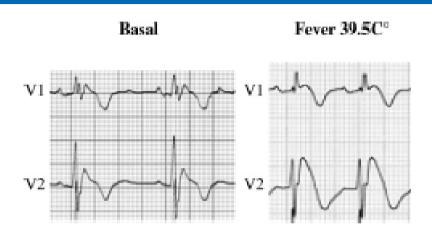


Figure 2. Representation of the right precordial leads V_1 and V_2 of a basal ECG (left) and an ECG taken during a febrile episode in a 5-year-old child. During fever, ST-segment elevation was accentuated, and coved-type morphology of the ST segment (>2 mm) occurred in lead V_2 .



Case Conclusion

Work-up

- Echocardiogram demonstrated normal structure
- Cardiac catheterization confirmed normal structure
- EKG demonstrated sinus rhythm with corrected QT interval of 440-450 milliseconds
- Mother's EKG showed a corrected QT interval of 480 milliseconds
- Started on Beta-blocker



Case Conclusion

- Electrophysiology study
 - No inducible arrhythmias
 - QT prolongation with epinephrine challenge test
- Long QT syndrome diagnosis
- Dual chamber cardioverter/defibrillator placed

What is the average survival for an out of hospital sudden cardiac arrest?

- 1. 79%
- 2. 52%
- 3. 37%
- 4. 7%
- 5. 0%



Case Conclusion – What Worked?

- Average survival for out of hospital SCA = 5-10%
 - Improved survival if presenting rhythm is VF
- Immediate CPR
 - Survival decreases 10% per minute if no CPR
 - CRP prolongs VF and delays asystole
- Prompt defibrillation
 - CPR alone is unlikely to restore a perfusing rhythm





Screening

Preparticipation Evaluation

Electrocardiogram

Echocardiogram



Electrocardiograms

Many causes of SCD produce EKG changes

- Abnormal in up to 95% of HCM
- Abnormal in >90% of ARVC



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Electrocardiograms

ECG may pick up:

- HCM
- ARVC
- Dilated cardiomyopathy
- Pre-excitation syndromes
- Conduction diseases
- Long/short QT syndrome

ECG will not pick-up:

- CPVT
- Commotio cordis
- Marfan syndrome

- Valvular Heart Disease
- Congenital anomalies of coronary arteries
- Myocarditis
- Congenital heart disease

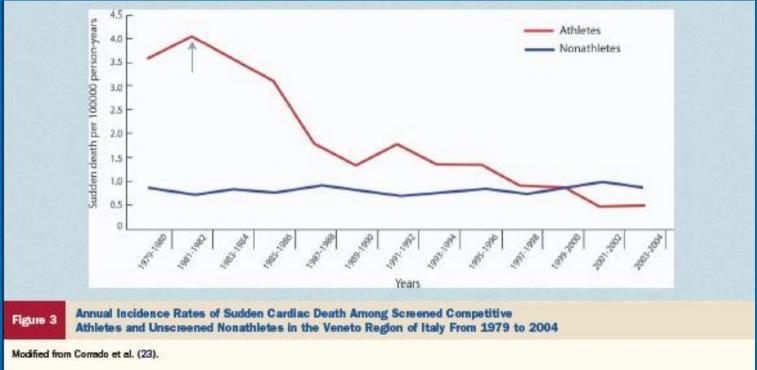


Electrocardiograms - pro

- Italian study showed 10 fold decrease in SCD over a 20-year period (Corrado, JAMA, 2006)
 - >40,000 athletes aged 12-35 in Veneto, Italy from 1979-2004
 - Rate of SCD dropped 89% (P<0.001) from all causes
 - HCM diagnosed on ECG (82%) vs H/P (23%)



The Italian Experience





Corrado, 2004

Is ECG Screening Cost Effective?

- 12 lead ECG is most cost effective: \$44,000 per year of life saved
 - H&P would have to be 2x more sensitive for similar cost-effectiveness
 - Echocardiogram would have to decrease in cost 4.5x for similar cost-effectiveness

(Fuller, Med Sci Sports & Ex; 2000)

- Model based on Corrado estimates of sudden death
 - Adding ECG to H&P
 - Saves 2.06 life-years per 1000 athletes
 - increased cost \$89 per athlete
 - Extra \$42,900 (\$21,200-\$71,300) per life-year saved
 - H&P and ECG Versus NO SCREENING
 - Saves 2.66 life-years per 1000 athletes
 - \$199 per athlete
 - \$76,100 (\$62,400-\$130,000) per life-year saved



Electrocardiograms - con

- Italian study not applicable to US population
 - Large population of ARVC
- US screening unreasonable
 - 10-12 million young athletes
 - Athletes' EKGs frequently abnormal
 - 1.5 million abnormal EKG but only 35 true positives
 - \$250,000 to detect one case of HCM (Maron, Circulation, 1996)
 - Medical legal (liability of PCP to read ECG)
 - Ethical (player disqualification)
 - Feasibility (Infrastructure)

Maron, Circulation, 1996



Electrocardiograms - con

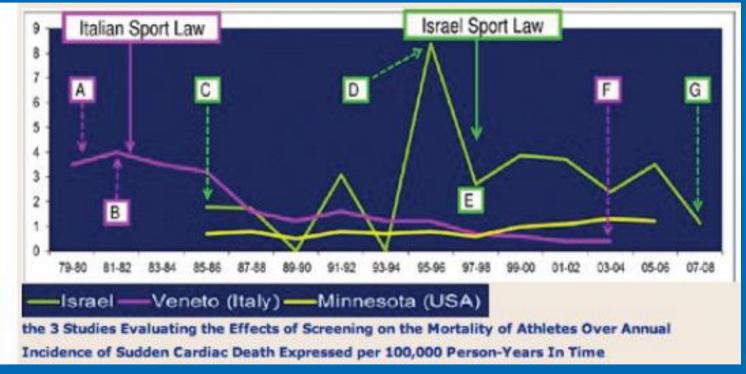
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Pediatric cardiologists in the U.S. = 1,521
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# of athletes to be screened = 12 million
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of exams/ECGs to be performed = 7,890 / year by each pediatric cardiologist = 658 / month = 150 / week



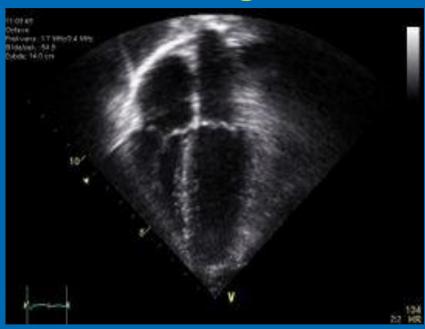
Does it make a difference?





Viskin, JACC; 2011

Echocardiograms



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- Identify structural abnormalities
 - valvular disease, cardiomyopathy, coronary abnormalities, aortic root dilation, etc.
- Identify functional abnormalities
 - cardiomyopathy, myocarditis
- May not identify
 - arrhythmia syndromes, coronary abnormalities, early cardiomyopathy



Screening

\$1.8 billion per year to screen US athletes

\$30 million to detect one athlete at risk for sudden cardiac death





Moss and Adams'; 7th ed

Screening

The Preparticipation evaluation

- Goal: identify clinically relevant cardiovascular abnormalities
 - Many causes of SCD are inheritable
 - Many patients with early symptoms
 - Identify noncardiac concerns





AHAJOKES.com, 2022

PREPARTICIPATION PHYSICAL EVALUATION

5th Edition

American Academy of Family Physicians

American Academy of Pediatrics

American College of Sports Medicine

American Medical Society for Sports Medicine

American Orthopaedic Society for Sports Medicine

American Osteopathic Academy of Sports Medicine

American Academy of Pediatrics



DEDICATED TO THE HEALTH OF ALL CHILDREN®



American Academy of Pediatrics, 2021

See References slide for additional information.

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Screening - History

- Passed out during exercise?
- Passed out after exercise?
- Chest pain/discomfort during exercise?
- Heart race or skip beats with exercise?
- Test for heart?

- HTN, high cholesterol, heart murmur or infection?
- Sudden death in family?
- FHx heart condition?
- FHx death before 50?
- FHx Marfan Syndrome?



Screening - Exam

- Normal PE
- Blood pressure
 - F/U if abnormal
- Palpate pulses
- Cardiac auscultation
 - Supine and standing
- Marfan's stigmata



The Pre-Participation Evaluation



Current practice makes an unclear picture

- Multiple venues
- Multiple practitioners with variable knowledge base (Glover, AJC; 2007)

The Pre-Participation Evaluation

- Retrospectively, only 3% athletes who died of SCD were suspected to have predisposing cardiac disease by routine history and physical (Maron, JAMA; 1996)
- 40% of SCD events have unrecognized or underappreciated warning symptoms (Campbell, Pediatrics; 2006)

Other interventions (AED, Mandatory CPR)?

Effectiveness of Emergency Response Planning for Sudden Cardiac Arrest in United States High Schools With Automated External Defibrillators

Jonathan A. Drezner, MD; Ashwin L. Rao, MD; Justin Heistand, MD; Megan K. Bloomingdale; Kimberly G. Harmon, MD

Table 1. Proportion of US High Schools With AED Programs
That Train School Staff in CPR or AED Use and Include Staff in
Annual Practice of the EAP for SCA

	CPR Trained, %	AED Trained, %	Practice EAP, %
Coaches	80	72	34
School nurses	75	71	22
Administrators	64	63	30
Teachers	49	45	22

Table 3. Comparison of Case Details of SCA in High School Student Athletes and Older Nonstudents

SCA Victims	Age (Range), y	Witnessed Collapse, n/N (%)	Selzure-Like Activity After Collapse, n/N (%)	Bystander CPR, n/N (%)	Time From Arrest to CPR, Mean; Median; Range, min	Shock Deployed, n/N (%)	Time From Arrest to First Shock, Mean; Median; Range, min	Survival to Hospital Discharge, n/N (%)
High school student athletes (n=14)	16 (14–17)	14/14 (100)	7/12 Reported (58)	13/14 (93)	1.5; 0.5; 0-5.75	13/14 (93)	3.6; 2.4; 0.75-11.5	9/14 (64)
Older nonstudents (n=22)	57 (42-71)	21/22 (95)	5/17 Reported (29)	21/22 (95)	0.8; 0.75; 0-1.75	19/22 (86)	1.8; 1.75; 0.5–3.25	14/22 (64)

Conclusions—School-based AED programs provide a high survival rate for both student athletes and older nonstudents who suffer SCA on school grounds. High schools are strongly encouraged to implement onsite AED programs as part of a comprehensive emergency response plan to SCA. (Circulation. 2009;120:518-525.)

Is your school prepared for a cardiac emergency?

Development and Implementation of Heart Safe Schools in Your District

https://www.childrenscolorado.org/doctors-and-departments/departments/heart/programs-and-clinics/project-adam/



Conclusions

- SCD a tragedy that affects society's "models of health"
- Many causes of SCD include personal or family history
- Currently, PPE is the most accessible and affordable screening tool
- Use of adequate PPE forms needed
- Prompt CPR and defibrillation is critical to survival of SCA events



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Google Image (June, 2022)https://www.google.com/url?sa=i&url=https%3A%2F%2Fwww.union-bulletin.com%2Fsports%2Fsports_columns%2Fcolumn-inside-the-amazing-convoluted-journey-marjon-beauchamp-took-from-yakima-to-the-nba-draft



Thank You

