

ORIGINAL ARTICLE

Outcomes of Cardiac Screening in Adolescent Soccer Players

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ABSTRACT

BACKGROUND

Reports on the incidence and causes of sudden cardiac death among young athletes have relied largely on estimated rates of participation and varied methods of reporting. We sought to investigate the incidence and causes of sudden cardiac death among adolescent soccer players in the United Kingdom.

METHODS

From 1996 through 2016, we screened 11,168 adolescent athletes with a mean (\pm SD) age of 16.4 \pm 1.2 years (95% of whom were male) in the English Football Association (FA) cardiac screening program, which consisted of a health questionnaire, physical examination, electrocardiography, and echocardiography. The FA registry was interrogated to identify sudden cardiac deaths, which were confirmed with autopsy reports.

RESULTS

During screening, 42 athletes (0.38%) were found to have cardiac disorders that are associated with sudden cardiac death. A further 225 athletes (2%) with congenital or valvular abnormalities were identified. After screening, there were 23 deaths from any cause, of which 8 (35%) were sudden deaths attributed to cardiac disease. Cardiomyopathy accounted for 7 of 8 sudden cardiac deaths (88%). Six athletes (75%) with sudden cardiac death had had normal cardiac screening results. The mean time between screening and sudden cardiac death was 6.8 years. On the basis of a total of 118,351 person-years, the incidence of sudden cardiac death among previously screened adolescent soccer players was 1 per 14,794 person-years (6.8 per 100,000 athletes).

CONCLUSIONS

Diseases that are associated with sudden cardiac death were identified in 0.38% of adolescent soccer players in a cohort that underwent cardiovascular screening. The incidence of sudden cardiac death was 1 per 14,794 person-years, or 6.8 per 100,000 athletes; most of these deaths were due to cardiomyopathies that had not been detected on screening. (Funded by the English Football Association and others.)

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SUDDEN CARDIAC DEATH IN A SEEMINGLY healthy athlete invariably prompts discussion about possible preventative screening strategies. The absence of a systematic registry of deaths in young athletes (<35 years old) has been a major obstacle to ascertaining the precise incidence of sudden cardiac death in this group; estimates vary between 0.5 and 13 deaths per 100,000 athletes,^{1,2} depending on the population studied and methods of data collection used.³ Furthermore, data on outcomes in adolescent athletes who have been screened for cardiovascular disease in a well-defined cohort are lacking.

The English Football Association (FA) has run a mandatory cardiac screening program for adolescent athletes in the United Kingdom since 1997, with a total of more than 11,000 athletes screened since its inception. The aim of this study was to determine the incidence and causes of sudden cardiac death in this well-defined population of previously screened soccer players.

METHODS

SCREENING PROGRAM

Between January 1, 1996, and December 31, 2016, a total of 11,168 soccer players at clubs affiliated with the FA underwent mandatory cardiovascular screening. The program encompassed all youth academy players (15 to 17 years of age) across the 92 professional clubs in the soccer league system who had excelled within the preceding 5 years. All such high-ranking players were offered a formal remunerated scholar contract (usually at the age of 16 years) with a view to progressing to a professional senior career. Written informed consent for screening was obtained from each player by the team doctor. Written informed consent from a parent or guardian was required for athletes younger than 16 years of age, in accordance with the FA governance department.

Athletes underwent assessment with a health questionnaire (Fig. S1 in the Supplementary Appendix, available with the full text of this article at NEJM.org), physical examination, 12-lead electrocardiography (ECG), and echocardiography. Mobile screening units staffed by accredited sonographers visited the clubs to conduct the evaluations, and the results were reviewed by an expert regional cardiologist who was a member of the FA-approved cardiology consensus panel.

A formal report was sent to the FA medical department, in which each athlete was classified in one of three categories on the basis of the evaluation: normal; further evaluation needed, if an abnormality was detected that required further investigation to confirm or refute the presence of cardiac disease; or cardiac disease detected. The last category was subclassified into disorders that are associated with sudden cardiac death or those encompassing congenital septal and valvular conditions.

Further investigations were performed at regional specialist centers. Athletes with T-wave inversion, abnormally enlarged cardiac dimensions, or a low ejection fraction and those fulfilling echocardiographic criteria for left ventricular noncompaction underwent maximal exercise stress testing, 24-hour Holter monitoring, and cardiovascular magnetic resonance imaging (MRI). Athletes with a prolonged QT interval underwent exercise stress testing and 24-hour Holter monitoring, and athletes with a Wolff–Parkinson–White ECG pattern were risk-stratified on the basis of an exercise test and an electrophysiological study. From 2012 onward, there was sufficient evidence indicating that among black athletes, anterior T-wave inversion was a common variant that was not associated with structural heart disease or serious cardiac arrhythmias⁴; therefore, asymptomatic black athletes with this repolarization abnormality did not undergo further investigation unless the echocardiogram was abnormal.⁵ In 2013, we ceased to investigate athletes who fulfilled echocardiographic criteria for left ventricular noncompaction but had a normal ECG and normal cardiac function.⁶ The results of all the investigations were reported back to the FA headquarters and reviewed independently by the first and last authors.

DIAGNOSIS OF DISORDERS ASSOCIATED WITH SUDDEN CARDIAC DEATH

The diagnosis of hypertrophic cardiomyopathy was based on a left ventricular wall thickness of 15 mm or greater in any myocardial segment on echocardiography or cardiovascular MRI in the absence of another condition capable of producing left ventricular hypertrophy.⁷⁻⁹ The diagnosis of dilated cardiomyopathy was considered if the patient had a dilated left ventricle (a left ventricular end-diastolic dimension of >59 mm in males and >53 mm in females) and a reduced ejection

fraction (<52% in males and <54% in females).^{10,11} The diagnosis of arrhythmogenic right ventricular cardiomyopathy was based on published criteria.^{12,13} The diagnosis of long-QT syndrome was based on a corrected QT (QTc) interval of 500 msec or greater or of 470 to 490 msec in association with notched T waves in at least three leads, paradoxical prolongation of the QT interval with exercise, or a positive genetic test.^{14,15} Identification of the Wolff–Parkinson–White ECG pattern was based on findings of a short PR interval and a slurred upstroke to the QRS complex.

RECOMMENDATIONS FOR FOLLOW-UP ASSESSMENTS OR EXCLUSION FROM PLAY

Some of the screened athletes had abnormal T-wave inversion but structurally normal hearts, and others had borderline-abnormal cardiac dimensions but no other features to support the diagnosis of cardiomyopathy. These athletes were investigated with ECG and echocardiography performed every year and cardiovascular MRI performed every 2 years. Athletes with congenital valvular abnormalities or septal defects were monitored with annual ECG and echocardiography.

Athletes with cardiac disorders that are associated with sudden cardiac death were advised not to compete and were discharged into the care of the National Health Service. Decisions to disqualify such athletes were made by the FA cardiology consensus panel after discussions in accordance with current exercise recommendations of the European Society of Cardiology and the American Heart Association.¹⁶⁻¹⁸ The decision was relayed to the player in the presence of the player's parent or guardian and the club doctor by the regional cardiologist.

OUTCOMES

Calculation of the follow-up period per athlete was based on the number of years of competition within the FA, which was determined from the FA registry of players. Deaths among athletes were ascertained through the development of a database that was compiled from voluntary reports to the FA. A second method to ascertain the number of deaths was through a secure survey that was sent to health professionals at each of the 92 FA-affiliated clubs, asking specifically about deaths from any cause. In addition, regular Internet searches had been performed since

2005, with the use of three different search engines (Google, Yahoo, and MSN search [Bing]) and at least 16 keywords (student, athlete, collapsed, died, death, heart, cardiac, arrest, attack, soccer, running, school, unknown, college, defibrillator, and saved).

Death certificates were obtained from the U.K. government for all deceased persons in the cohort to ascertain the causes of death, which were categorized broadly as accidental, suicide, drug-related, cancer, or cardiac causes. Autopsy data were available in all cases of sudden cardiac death, and diagnoses were based on previously established pathological criteria in conjunction with consultation with an expert cardiac pathologist.¹⁹

Data on survival status during the screening program over the 20-year period among athletes who had diagnoses of cardiac disorders that are associated with sudden cardiac death were obtained from attending cardiologists, most of whom were part of the expert FA consensus panel.

RESULTS

FURTHER EVALUATION AFTER SCREENING

The mean (\pm SD) age of the 11,168 soccer players who underwent cardiovascular screening was 16.4 ± 1.2 years; 10,581 (95%) of the athletes were male. In the entire cohort, 830 athletes (7%) underwent further investigation after preliminary assessment. Among these athletes, 104 (0.9%) reported symptoms that were deemed noncardiac in origin after assessment by a cardiologist. Among the remaining 726 athletes, 292 (3% of the total cohort) underwent investigation for T-wave inversion (anterior [153], inferior or lateral [114], or widespread [25]). A total of 25 athletes (0.2%) underwent investigation for a prolonged QTc interval. A further 409 athletes (4%) underwent cardiovascular MRI because of ventricular remodeling consistent with cardiomyopathy.⁹ Specifically, 229 athletes (2%) had a left ventricular wall thickness of 13 mm or greater; 106 (0.9%) had an enlarged left ventricular cavity with a borderline-low ejection fraction (of 50 to 52%)¹⁰; 80 athletes (0.7%) had a suspected right ventricular regional wall-motion abnormality, and 19 (0.2%) had increased left ventricular trabeculation and a mildly reduced ejection fraction (of $\leq 50\%$).⁶ Twenty-five athletes had abnormal T-wave inversion and enlarged cardiac dimensions.

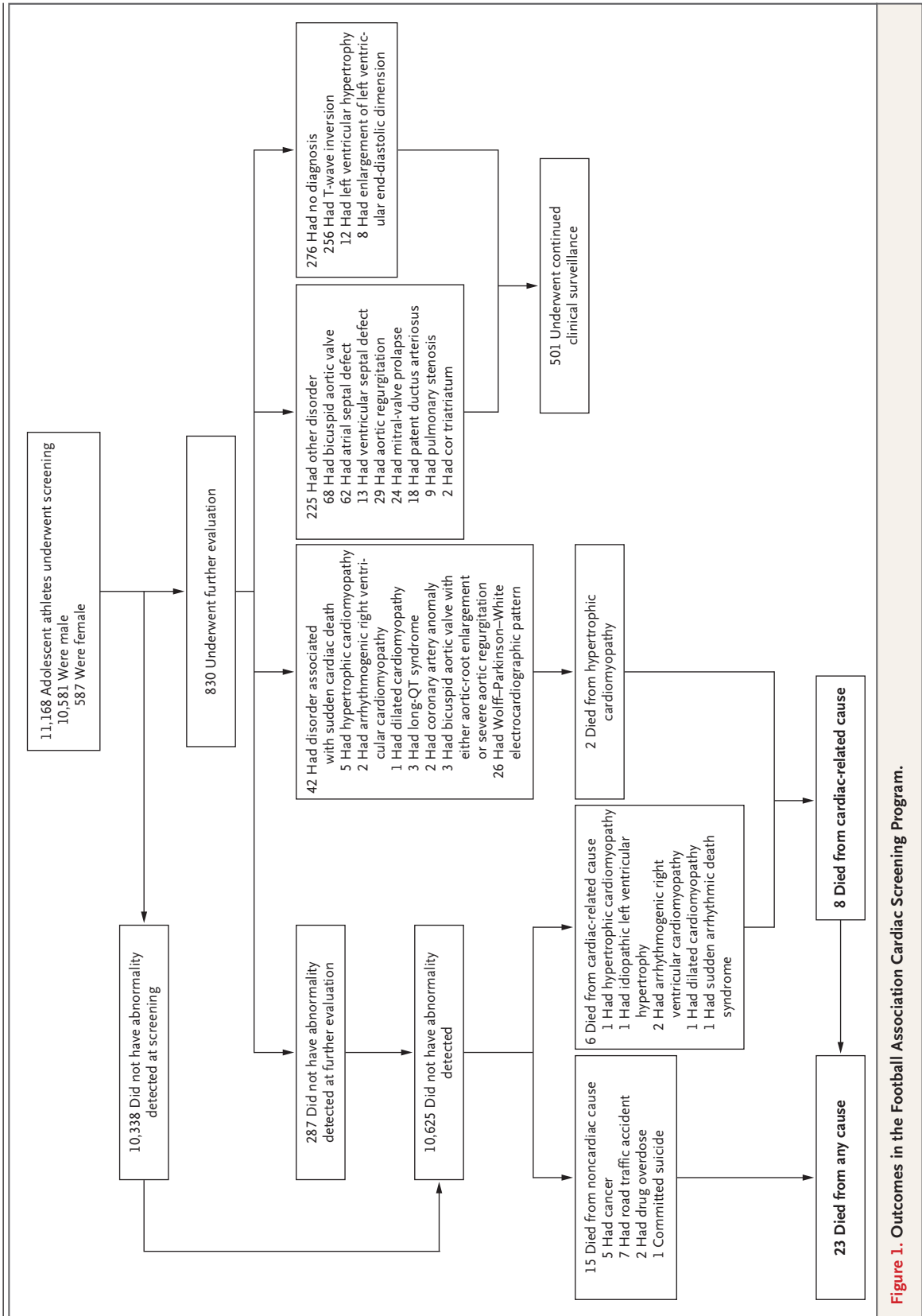


Figure 1. Outcomes in the Football Association Cardiac Screening Program.

Table 1. Clinical Features and Outcomes in Athletes with Cardiac Conditions That Are Associated with Sudden Cardiac Death.*

Condition, Sex, and Age	Race†	History and Examination	ECG Result	Echocardiography Result	LGE on Cardiac MRI	Exercise Test Result	Genetic Test Result‡	Outcome
HCM								
M, 16 yr	White	Negative	TWI (leads II, aVF, V2–V6)	LVTW 16 mm (asymmetric septal hypertrophy)	Yes	Normal	MYBPC3 mutation	Advised not to play
M, 15 yr	White	Negative	TWI (leads V2–V6), LAD, isoelectric ST segments	LVTW 15 mm (asymmetric septal hypertrophy)	No	Normal	MYBPC3 mutation	Advised not to play
M, 16 yr	White	Negative	TWI (leads II, III, aVF), ST depression	Apical hypertrophy	No	Normal	MYH7 mutation	Advised not to play
M, 16 yr	Black	Negative	TWI (leads V1–V5), ST depression	LVTW 16 mm (asymmetric septal hypertrophy)	Yes	Normal	Negative	Advised not to play
M, 16 yr	Mixed	Negative	TWI (leads V4–V6), isoelectric ST segments	Apical hypertrophy	No	Normal	Negative	Advised not to play
ARVC								
M, 16 yr	White	Palpitations	TWI (leads V1–V3)	Reduced LV systolic function; dilated and aneurysmal RV	Yes	Ventricular ectopy of LBBB morphology	Negative	Advised not to play
M, 17 yr	White	Negative	Normal	Aneurysmal RV with hypokinetic free wall	No	Ventricular ectopy of LBBB morphology	PKP2 mutation	Advised not to play
DCM								
M, 16 yr	White	Dyspnea	TWI (leads V1–V4), ST depression	LVEDD, 61 mm; EF, 45%	Yes	LV ejection fraction did not increase with exercise	Negative	Advised not to play
LQTS								
F, 16 yr	White	Negative	QTc, 510 msec	Normal	NA	QTc, >500 msec	KCNQ1 mutation	Advised not to play
M, 15 yr	White	Negative	QTc, 503 msec	Normal	NA	QTc, >500 msec	Negative	Advised not to play
M, 16 yr	White	Negative	QTc, 490 msec	Normal	NA	Paradoxical increase in QTc during recovery	KCNQ1 mutation	Advised not to play
CAA								
M, 16 yr	White	Negative	Normal	Left coronary artery arising from right sinus of Valsalva	NA	Positive for myocardial ischemia	NA	Underwent corrective surgery and returned to play
M, 15 yr	White	Negative	Normal	Right coronary artery arising from left sinus of Valsalva with adverse course.	NA	Normal	NA	Underwent corrective surgery and returned to play

BAV	Black	Dyspnea, diastolic murmur	Left axis deviation	Fusion of right and left coronary cusps and severe aortic regurgitation; LVEDD, 60 mm	No	Terminated prematurely because of fatigue	NA	Underwent corrective surgery and returned to play
M, 16 yr	Black	Negative	Normal	Fusion of right and noncoronary cusps with mixed aortic valve disease; diameter of aortic root at sinuses of Valsalva, 53 mm	No	Normal	NA	Underwent corrective surgery and returned to play
M, 17 yr	White	Negative	Normal	Fusion of right and left coronary cusps and severe aortic regurgitation; LVEDD, 63 mm	No	Normal	NA	Underwent corrective surgery and returned to play

* ARVC denotes arrhythmogenic right ventricular cardiomyopathy, BAV bicuspid aortic valve, CAA coronary-artery anomaly, DCM dilated cardiomyopathy, ECG electrocardiogram, EF ejection fraction, HCM hypertrophic cardiomyopathy, LAD left axis deviation, LBBB left bundle-branch block, LGE late gadolinium enhancement, LQTS long-QT syndrome, LV left ventricular, LVEDD left ventricular end-diastolic diameter, LVWT left ventricular wall thickness, NA not applicable, QTc corrected QT interval, RV right ventricle, and TWI T-wave inversion.
 † Race was reported by the athlete or the parent or guardian.
 ‡ Genes that are associated with cardiac disease were tested for mutations; among the athletes with cardiac conditions that are associated with sudden cardiac death, mutations were found in the genes encoding potassium voltage-gated channel subfamily Q member 1 (KCNQ1), myosin binding protein C, cardiac (MYBPC3), myosin heavy chain 7 (MYH7), and plakophilin 2 (PKP2).

DETECTION OF CARDIAC DISEASE

A total of 42 athletes (0.38%) were found to have cardiac disorders that are associated with sudden cardiac death (Fig. 1 and Table 1). Five (0.04%) had a diagnosis of hypertrophic cardiomyopathy, 2 (0.02%) of arrhythmogenic right ventricular cardiomyopathy,¹² and 1 (0.01%) of dilated cardiomyopathy. Three athletes (0.03%) had a diagnosis of long-QT syndrome. Two athletes (0.02%) had a diagnosis of an anomalous origin of a coronary artery, and 3 (0.03%) were found to have a bicuspid aortic valve associated with either aortic-root enlargement of 50 mm or greater (1) or severe aortic regurgitation (2). A total of 26 (0.23%) athletes had the Wolff–Parkinson–White ECG pattern (Table S1 in the Supplementary Appendix).

Among the 42 athletes with cardiac disorders that are associated with sudden cardiac death, 2 (5%) had symptoms, 1 (2%) had an abnormality detected on cardiac examination, and 1 (2%) had both symptoms and an abnormality detected on examination (Table 2). Abnormal ECGs were obtained in 36 of these athletes (86%), and abnormal echocardiograms in 12 (29%).

There were 225 athletes (2%) with other cardiac disorders, including congenital septal and valvular abnormalities (Fig. 1). The diagnoses in all of these athletes were made with the use of echocardiography; 48 of the athletes (21%) had an abnormal ECG, and 74 (33%) had an abnormality detected on examination (Table 2).

INTERVENTIONS AND OUTCOMES

Athletes with a diagnosis of cardiomyopathy (8) or long-QT syndrome (3) were advised against participation in competitive soccer. None of the athletes with cardiomyopathy were deemed to have a sufficiently high-risk profile to warrant a prophylactic implantable cardioverter–defibrillator.^{7,20,21} The 3 athletes with long-QT syndrome began treatment with beta-blockers. Both athletes with anomalous coronary-artery origins underwent corrective surgery and returned to play. The 3 athletes with bicuspid aortic valves underwent surgical intervention for aortic-root dilatation (1) or severe aortic regurgitation (2) and returned to play (Table 1). All 26 athletes with the Wolff–Parkinson–White ECG pattern underwent risk stratification, and 24 underwent ablation before returning to play. The remaining 2 athletes with the Wolff–Parkinson–White pattern were deemed

Table 2. Summary of Cardiac Conditions Detected According to Screening Tool.

Condition	No. of Athletes	No. of Athletes with Abnormal Result			
		History	Examination	ECG	Echocardiography
Any cardiac condition	267	6	76	84	237
Condition associated with sudden cardiac death	42	3	2	36	12
Hypertrophic cardiomyopathy	5	0	0	5	5
Arrhythmogenic right ventricular cardiomyopathy	2	1	0	1	2
Dilated cardiomyopathy	1	1	0	1	1
Coronary-artery anomalies	2	0	0	0	2
Bicuspid aortic valve–associated disease*	3	1	2	0	3
Long-QT syndrome	3	0	0	3	0
Wolff–Parkinson–White ECG pattern	26	0	0	26	0
Other cardiac condition	225	3	74	48	225
Bicuspid aortic valve	68	1	32	15	68
Atrial septal defect	62	1	6	26	62
Aortic regurgitation	29	0	16	2	29
Mitral-valve prolapse	24	0	12	3	24
Patent ductus arteriosus	18	0	1	1	18
Ventricular septal defect	13	0	3	1	13
Pulmonary stenosis	9	1	4	0	9
Cor triatriatum	2	0	0	0	2

* Bicuspid aortic valve–associated disease includes bicuspid aortic valve with either aortic-root enlargement or severe aortic regurgitation.

to have low-risk accessory conduction pathways that were not ablated, and they continued to compete (Table S1 in the Supplementary Appendix).

Of the 42 athletes who, during screening, received diagnoses of cardiac disorders that are associated with sudden cardiac death, 40 (95%) were alive at the end of the study period. Two athletes with a diagnosis of hypertrophic cardiomyopathy continued to compete despite medical advice and died subsequently during intensive exercise.

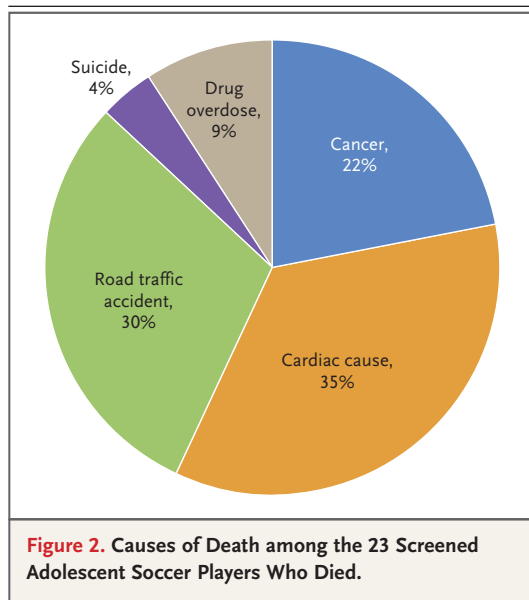
Among the 225 athletes with other cardiac disorders, 7 with an atrial septal defect and 1 with a ventricular septal defect had hemodynamic indications to warrant percutaneous closure before returning to play. The remaining 217 athletes (97%) had mild disease and were permitted to compete. We could not establish a cardiac diagnosis in 276 athletes (2%) who had abnormal T-wave inversion or borderline abnormal cardiac dimensions. All these athletes continued to compete but remained under clinical surveillance.

ALL-CAUSE MORTALITY

Of the 92 professional clubs, 86 (93%) responded to the questionnaire regarding all-cause mortality. During a mean follow-up period of 10.6±8.3 years, there were 23 deaths. The causes of death included road traffic accidents (7 deaths [30%]), cancer (5 [22%]), drug overdose (2 [9%]), and suicide (1 [4%]). Cardiac disorders accounted for 8 deaths (35%), all of which were sudden and occurred during exercise (Fig. 2).

SUDDEN CARDIAC DEATHS

Autopsy data were available for all sudden cardiac deaths. Among the athletes who died from cardiac disorders, the mean time between screening and sudden death was 6.8 years (range, 0.1 to 13.2). Cardiomyopathies were the most common cause of death and accounted for 7 of 8 (88%) sudden cardiac deaths (Table 3). There were a total of 118,351 person-years of follow-up over the 20-year study period. The resulting overall incidence of sudden cardiac death among ado-



lescent soccer players was 1 per 14,794 person-years, or 6.8 per 100,000 athletes.

Of the 8 sudden cardiac deaths, 6 (75%) were in athletes who had normal findings during preliminary screening (Fig. 1 and Table 3). Variation in test interpretation could have accounted for the normal findings. Therefore, 50 deidentified ECGs and corresponding echocardiograms from the cohort were analyzed by two independent experts in inherited cardiac diseases who were unaware of the outcomes. The ECGs and echocardiograms were from 45 athletes whose results had been interpreted as normal, including the 6 decedents, and 5 athletes with abnormal results, including 1 athlete each with a diagnosis of arrhythmogenic right ventricular cardiomyopathy, dilated cardiomyopathy, hypertrophic cardiomyopathy, long QT interval, and the Wolff–Parkinson–White ECG pattern. Blind reading showed 100% agreement between the two reviewers for the 6 decedents with normal results (Table 3).

COST OF DIAGNOSIS

On the basis of the U.K. National Health Service tariffs,²² the cost of preliminary investigation with consultation (£160 [\$213 in U.S. dollars]), ECG (£25 [\$33]), and echocardiography (£72 [\$96]) would amount to £257 (\$342) per athlete, resulting in an initial cost of screening 11,168 athletes of £2,870,176 (\$3,817,334). The cost of further investigating 830 athletes was £375,587

(\$499,531), with a total estimated outlay of £3,245,763 (\$4,316,865). The cost to detect serious cardiac disease associated with sudden cardiac death (42 athletes) was £77,280 (\$102,782) per case, and the cost to identify any cardiac disorder (267 athletes) was £12,156 (\$16,167) per case.

DISCUSSION

We report the outcomes of cardiovascular screening of adolescent soccer players, determined with the use of data from the FA in the United Kingdom. The prevalence of disorders associated with sudden cardiac death in young athletes was 0.38%, which is similar to that reported in other screening programs.^{23–25} Congenital septal and minor valvular disorders were detected in an additional 2% of the athletes, leading to an overall prevalence of 2.4% for all cardiac conditions.

In this cohort, electrical diseases accounted for 29 (69%) of the 42 cases of cardiac disorders that are associated with sudden cardiac death, whereas the primary cardiomyopathies accounted for only 8 (19%) of these cases. Anomalous coronary-artery origins accounted for 2 (5%) of the cases but were almost certainly underrepresented because of the limitations of echocardiography in detecting this type of disorder. The remaining 3 (7%) of the 42 cases were due to advanced aortic-valve disease. History, findings on physical examination, and ECG were abnormal in 7%, 5%, and 86%, respectively, of athletes with cardiac disorders associated with sudden cardiac death.

Hypertrophic cardiomyopathy was the most commonly detected cardiomyopathy, and all 5 athletes in whom this condition was detected had an abnormal ECG and echocardiogram. The prevalence of hypertrophic cardiomyopathy among these elite adolescent soccer players was 1 in 1861 athletes (including 1 case not detected by screening), which is considerably lower than that reported in the general population, raising the possibility that athletes with a more advanced phenotype at this age may have been selected out of competition because of reduced cardiorespiratory capacity.^{26,27} Age-related penetrance of hypertrophic cardiomyopathy is also an important consideration, since in many affected persons the disease is not expressed during adolescence.^{28–33} The prevalence of arrhythmogenic right ventric-

Table 3. Characteristics of Athletes with Sudden Cardiac Death.

Athlete No.	Sex and Age	Race*	Years from Screening to Death	Diagnosis	Initial Screening Result	Blind Reading (Reviewer 1)	Blind Reading (Reviewer 2)
1	M, 16.8 yr	Black	0.1	Idiopathic left ventricular hypertrophy	Negative	Negative	Negative
2	M, 16.6 yr	Mixed	1.0	Hypertrophic cardiomyopathy	Abnormal ECG and echocardiogram	NA	NA
3	M, 16.6 yr	Black	3.3	Hypertrophic cardiomyopathy	Negative	Negative	Negative
4	M, 16.3 yr	Black	7.7	Dilated cardiomyopathy	Negative	Negative	Negative
5	M, 17.0 yr	White	7.9	Arrhythmogenic right ventricular cardiomyopathy	Negative	Negative	Negative
6	M, 17.2 yr	White	9.7	Arrhythmogenic right ventricular cardiomyopathy	Negative	Negative	Negative
7	M, 15.7 yr	White	11.5	Hypertrophic cardiomyopathy	Abnormal ECG and echocardiogram	NA	NA
8	M, 16.8 yr	White	13.2	Sudden arrhythmic death syndrome	Negative	Negative	Negative

* Race was reported by the athlete or the parent or guardian.

ular cardiomyopathy was 1 in 2792 athletes; this disorder has a highly variable clinical course, and the mean age at presentation is 31 ± 13 years.³⁴ Exercise has been shown to accelerate the phenotypic manifestations of this condition,^{35,36} and years of intensive training regimes may well contribute to the unmasking of phenotypes that could not be detected during adolescence.

Of the seven athletes who died suddenly from cardiomyopathy, five (71%) had a normal ECG and echocardiogram at a mean age of 16 years. Screening at this age seems logistically appropriate, given that most people will be postpubertal and will have overt evidence of any electrical or structural cardiac abnormalities. However, this study shows that screening during late adolescence will fail to detect a substantial proportion of athletes who have or will eventually have a cardiomyopathy, either because the disease is not yet manifest or because ECG and echocardiography are not sensitive enough to detect early disease in some adolescents.

This systematic study revealed that the incidence of sudden cardiac death among screened 16-year-old soccer players was approximately 1 per 14,800 person-years, or 6.8 per 100,000 athletes. This figure is considerably higher than previous estimates among athletes who have been screened with the use of history and physical examination alone or who have not undergone cardiac screening.^{1-3,37-41} Sudden cardiac death may

be more common in some sports than in others; these results in adolescent soccer players are similar to findings previously reported in male basketball players.³ The National Collegiate Athletic Association in the United States adopted a policy in 2014 requiring all affiliated institutions to report cases of catastrophic injury or death. Such mandatory reporting initiatives across other sporting organizations would offer an important source of prospective data in already well-defined populations.

Several limitations of our study should be noted. For our data, we relied on voluntary reporting of sudden cardiac deaths by clubs and on retrospective recall. Although we confirmed all sudden cardiac deaths with death certificates and autopsy reports, it is possible that we did not capture all cases. Our results therefore represent the minimum incidence of sudden cardiac death among screened adolescent soccer players. The end point of our study was sudden death, and therefore we are unable to comment on the number of athletes in whom a quiescent cardiomyopathy may have developed after the initial screen or who survived a sudden cardiac arrest. Finally, our study included only adolescent soccer players of the highest ability and may underestimate the burden of cardiac disorders or prevalence of sudden cardiac death among non-elite or older players.

In conclusion, we investigated the results of

cardiovascular screening of a large cohort of adolescent soccer players in the United Kingdom. Diseases associated with sudden cardiac death, the majority of them electrical cardiac disorders, were identified in 0.38% of participants. The incidence of sudden cardiac death among these previously screened athletes was approximately 1 per 14,800 person-years, or 6.8 per 100,000

athletes. Most of these deaths were due to cardiomyopathies that were not detected on screening.

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APPENDIX

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