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Pre-participation cardiovascular evaluation in Pacific Island athletes

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Abstract

**Background:** Pre-participation cardiovascular evaluation (PPE) aims to detect cardiac disease with sudden cardiac death (SCD) risk. No study has focused on Pacific Island athletes.

**Methods:** A total of 2,281 Pacific Island athletes were studied with (i) a questionnaire on family, personal history and symptoms, (ii) a physical examination and (iii) a 12-lead ECG.

**Results:** 85% presented a normal history and examination. A positive family history was 1.4-1.9 fold higher in Melanesians, Polynesians and Métis than in Caucasians, while a positive personal history, abnormal symptoms and abnormal examination was 1.3 fold higher in Melanesians and Métis than in others. Neither gender nor training level had a bearing on these results. Melanesians had higher T wave inversions (TWIs) in V2-V4 leads but had no CV abnormalities. Lateral or infero-lateral TWIs were found in 6 male and in 5 highly trained athletes and cardiomyopathies were diagnosed in 3/6 athletes. Overall, 3.9% athletes were found to have a CV abnormality and 0.8% had a risk of SCD. Polynesians and males were more at risk than the others while the level of training made no difference. In athletes at risk of SCD, the main detected CV diseases were cardiomyopathies, Wolff-Parkinson-White (WPW) and severe valve lesions of rheumatoid origin.

**Conclusions:** PPE revealed that 3.9% presented CV abnormalities. A risk of SCD was found in 0.8% with cardiomyopathies, WPW, and severe valve lesions of rheumatoid
origin. Melanesians, Polynesians and male of high level of training were more at risk than others.

Keywords: Pre-participation cardiovascular evaluation; Cardiomyopathy; Sudden cardiac death; T wave inversion; Melanesians; Polynesians
1 – Introduction

Sport provides health benefits, but sudden cardiac death (SCD) can reveal hitherto undetected cardiovascular (CV) disease. The estimated incidence rate of SCD is 1-2/100 000 athletes/year [1] and can be as high as 1/17 000 or 1/3 300 in some specific sports and ethnicities like Afro-Caribbean basketball players [2]. The proposition of a pre-participation cardiovascular evaluation (PPE) is presented as justifiable on medical grounds [3] with the aim to detect a CV disease and/or a disease with SCD risk. As about 80% of these diseases are asymptomatic, a resting 12-lead electrocardiogram (ECG) although controversial [4], is recommended as a valid, cost effective means of athlete screening [5].

Athlete’s ethnicity has a significant impact on the CV response to a high level of physical training as on the SCD risk. In comparison with Caucasians, heart specificities have been reported in Afro-Caribbean [6, 7], Asian [8] or West-Asian athletes [9]. Moreover, it has been reported that Afro-Caribbean athletes are more susceptible to SCD than Caucasians [10]. No study has focused on Pacific Island athletes.

The aim of this study was to present the results of a PPE carried out on 2 281 Pacific Island athletes. A second aim was to specify the differences related to the ethnicity of Melanesians, Polynesians, Caucasians and Métis, to their gender and to their level of training.

2 – Methods
The study design was approved from the University and Sports Medicine Hospital ethics committee of Saint-Etienne.

2.1 - Population

From 2012 to 2015, 2,281 athletes, 716 females, were studied and written informed consent. They were regularly involved in competition and registered in 40 sports federations. Football (392), volley ball (303), martial arts (286) and rugby (213) accounted for 52% of the athletes. They were also involved in 2 Pacific-specific sports Va’a, namely, sea canoe, (119) and sailing (93).

Ethnicity

Athletes were recruited from 14 Pacific Islands: New Caledonia (1450), French Polynesia (489), Fiji (108), Wallis and Futuna (69), Tonga (38), Tuvalu (36), Vanuatu (31), Papua New Guinea (28), 2 Samoas (15), Cook, Salomon and Palau islands (17).

Attention was focused on Melanesians (n = 702, 278 females), Polynesians (n = 653, 165 females), Caucasians (n = 469, 138 females) and Métis (n = 457, 135 females). Almost all of them, including Caucasians were native to these Pacific Islands. The term “Métis” was used when at least one parent or one of the 4 grand-parents was not from the same ethnicity as the others.

Level of training

Athletes were divided into two groups, those training more than 6 h/week (HT, n = 1336, 363 females) and those training less than 6 h/week (LT, n = 945, 353 females).

2.2 - Pre-participation cardiovascular evaluation
All athletes underwent the same PPE that included family and personal history, a physical examination and a 12-lead resting ECG. When athletes had several PPEs, only the first PPE was recorded. **In New Caledonia and French Polynesia, as in France, the PPE is routinely performed. Concerning the other Islands, the PPE was performed as a part of the research project during the 2013 Mini Pacific Games in Wallis.**

*History and physical examination.*

All athletes completed a questionnaire including the main items from the AHA 12-elements [11]. The medical history addressed present and past complaints as well as family history [12].

The physical examination was based on the Consensus Statement from the European Society of Cardiology sports group [13, 14].

*Resting 12-Lead Electrocardiography (ECG)*

After resting for 5-10 min, in the supine position, a 12-Lead ECG (Mac 1 200, GE, USA or Nihon Kohden, Japan) was recorded. The ECG analysis was initially carried out using The Seattle criteria [15]. At the end of the study, they were reviewed using the international classification proposed in 2017 [3].

*Further evaluation*

All athletes presenting abnormal findings were offered further complementary cardiac screening. As the burden of rheumatoid heart disease (RHD) is significant in the Pacific region, echocardiography was systematically offered to athletes with a family and/or personal history of Acute Rheumatic Fever (ARF) and/or who presented with a heart murmur at the physical examination. The WHF 2012 criteria [16] were applied.
Athletes with normal history and physical examination and with normal ECG did not undergo any other investigation.

It must be pointed out that in some low-income Pacific Islands some investigations, although highly recommended, like transthoracic echocardiography (TTE), 24h Holter recording, stress ECG, Magnetic Resonance Imaging (MRI), could not always be performed either for economic reasons, or the lack of sports cardiologists or specialized medical facilities.

2.3 - Statistical analysis

Multivariate Anova Analysis and one-way Anova analysis were carried out between ECG criteria, age, ethnicity, gender, and training level. A Fisher and Scheffe post hoc procedure was used to determine the location of the differences. A $P$ value of 0.05 was chosen as the level of statistical significance. The Stata 11 program was used for the statistical analysis.

3 – Results

3.1 - Athletes’ demographics

The 2 281 athletes were on average 19.4 ± 6.5 years old (age range 10-40) and trained 8.6 ± 5.0 h/week (Table 1 and 2). Male Polynesians were significantly older, taller, and heavier than the other ethnic groups ($P<0.05$). Regarding female athletes, Melanesians were younger and smaller than Polynesians and Métis and trained less than Polynesians.
(P<0.05), while female LT athletes were younger and smaller than female HT athletes (P<0.05).

3.2 - History and physical examination

85.4% of all athletes presented a normal history and physical examination, whereas 14.6% presented an abnormal history and/or physical examination (Figure 1).

The family history was positive for at least one item in 6.8% of athletes. The most frequent item was death of CV origin (n = 52, 1.3%), father n = 28, mother n = 17, brothers and sisters (n = 5) with 31 (0.8%) of family members younger than the age of 50. Other items were coronary artery diseases, valvular diseases, aneurysms, inter atrial or ventricular communications, pace-makers, hypertrophic cardiomyopathy, arrhythmias and high blood pressure.

The incidence of a positive family history was 1.4-1.9 fold higher (P<0.05) in Melanesians (6.3%), Polynesians (8.6%) and Métis (7.2%) when compared to Caucasians (4.5%). There was no statistical difference between males and females or between LT and HT athletes.

The personal history was positive in at least 1 item in 3.2% athletes and 2.0% athletes complained about at least 1 abnormal symptom during exercise. The most frequent item was RHD history (n = 47) followed by abnormal dyspnea (n = 23), while the most frequent symptoms were palpitations (n = 13), syncope or near syncope (n = 8) and thoracic pain (n = 7).

The physical examination was abnormal in 4.2% of athletes including 91 cardiac murmurs and 6 high blood pressure values.
The incidence of a positive personal history, abnormal symptom and abnormal examination was 1.3 fold higher \((P<0.05)\) in Melanesians (9.8\%) and Métis (12.9\%) when compared to Polynesians (6.3\%) and Caucasians (8.5\%). Gender did not have an impact on these findings nor did the level of training.

### 3.3 - ECG analysis

Most of the ECG variables were independently related to age, ethnicity, gender and duration of training.

In the total population, heart rate (HR) was < 40 bpm in 10 (0.4\%) athletes and < 50 bpm in 140 (6.1\%) athletes. No HR lower than 30 bpm was observed.

Electrical left atrial enlargement (LAE) and right atrial enlargement (RAE) were observed respectively in 2.2\% and 0.6\% athletes. They did not reveal any abnormalities.

Atrioventricular block (AVB) type 1 was observed in 2.5\% and a PR interval duration of \(\geq 220\) ms in 0.9\% and \(\geq 300\) ms in 0.2\% athletes. No PR duration longer than 400 ms was observed. A PR interval duration \(<120\) ms with pre-excitation was found in 6 athletes (0.3\%).

Isolated complete right bundle branch block (RBBB) was observed in only 1 athlete. Isolated QRS duration > 140 ms was found in 9 athletes (0.4\%). Isolated QRS complex axis deviation > 120° was found in 7 (0.3\%) male athletes and < -30° in 5 (0.2\%) athletes (4 males and 1 female). Isolated electrical left ventricular hypertrophy, (LVH), Sokolow-Lyon index, was found in 14\% of athletes, Melanesians (10.7\%), Polynesians (15.3\%), Caucasians (18.1\%) and Métis (12.9\%). Isolated electrical right ventricular hypertrophy
(RVH) was found in 7.4% athletes, Melanesians (4.3%), Polynesians (12.1%), Caucasians (6.4%) and Métis (6.3%).

Q waves steeper than 3 mm were found in 0.4% athletes, 9 males and 1 female. Of these athletes, only 1 had a Q/R ratio > 25%, none had a Q wave duration > 40 ms. They did not reveal any abnormalities.

T wave inversions (TWIs) were observed in 45 athletes (2.0%). TWIs concerned V2-V4 leads in 39 athletes, 27 (69%) were ≤ 16 years old. The incidence was 1.4-2.0 fold higher ($P < 0.05$) in Melanesians (2.6%) than in Polynesians (1.2%), Caucasians (1.5%) and Métis (1.3%). In most cases (31/39 = 79%), TWIs were associated with a J point elevation with convex ST segment with a 2-fold higher incidence in Melanesians (2.1%) than in Polynesians (1.1%), Caucasians (0.9%) and Métis (1.1%).

Lateral or infero-lateral TWIs were found in 6 athletes, 3 Melanesians, 1 Caucasian and 2 Métis. They were all males and 5 were HT athletes. HCM was diagnosed in the only 3 athletes that could be further investigated.

The QTc interval duration was on average 10 ms longer in females than in males. Four athletes presented a prolonged QTc duration between 470 and 510 ms, 2 males, 2 females, 2 Polynesians, 1 Melanesian and 1 Métis.

More than 2 premature ventricular beats were observed in 6 athletes, 4 Polynesians, 1 Melanesian and 1 Métis, 4 males and 2 females, 5 HT and 1 LT athlete. Supra ventricular tachycardia was observed in only 1 male HT Polynesian.

### 3.4 - Results of the pre-participation evaluation

The results of the PPE are summarized in Figure 1 and Table 3.
History and physical examination abnormalities concerned 14.6% athletes of which 1.6% presented abnormalities in both their history and examination. Of these athletes, an associated abnormal ECG was observed in 45 (2.0%). In these 45 athletes, 7% had a positive family history, 42% a positive personal history, 22% were symptomatic and 56% had an abnormal physical examination.

Isolated abnormal ECG results concerned 1% athletes.

Athletes with an abnormal ECG, and/or abnormal history, symptoms and physical examination were invited to undergo complementary CV evaluations: 71 underwent all the CV evaluations. Very few athletes refused to undergo further investigation. However in the case of some low-income athletes/islands, for economic reasons, 17 did not undergo either the TTE (n = 5) or the MRI scan (n = 12). Overall 88 athletes (3.9%) presented a CV abnormality.

A group of 21 athletes presented both abnormal examination and normal ECG. In 18, the CV abnormalities were diagnosed by echocardiography. In the 21 athletes, 14 RHDs had an abnormal personal history (n = 11) and or heart murmur (n = 6): 4 inter auricular or ventricular communication had a heart murmur: 3 cases of high blood pressure an abnormal examination.

The incidence of CV abnormalities was 1.4 fold higher ($P < 0.05$) in Polynesians (5.1%), when compared to Melanesians (3.4%), Caucasians (3.4%), or Métis (3.3%). The incidence was higher ($P < 0.05$) in males (4.2%) than in females (3.1%) but was almost the same between LT and HT athletes (3.8% as opposed to 3.9%).

Globally a CV disease associated with a potential risk of SCD was identified in 18 (0.8%) athletes. Familial and personal history was abnormal in 5 athletes, personal
history in 7, abnormal symptom in 6: syncope (n = 4), palpitations (n=1) and thoracic pain (n=1). Lastly the disease was detected by isolated abnormal ECG in 6 athletes.

The incidence was 2 to 6-fold higher (P<0.05) in Polynesians (1.7%) than in Melanesians (0.6%), Caucasians (0.2%) or Métis (0.4%), 3-fold higher in males than in females (1.1% vs. 0.3%, P < 0.01) and 4-fold higher in HT than in LT athletes (1.1% vs. 0.3% P<0.01). There were 6 (0.3%) cardiomyopathies detected in 3 Polynesians, 2 Melanesians and 1 Métis that included 4 HCMs, 1 left and 1 right uncompacted ventricle, all had abnormal ECGs, however, only 3/6 were symptomatic; 6 cases (0.3%) of Wolff-Parkinson-White in 4 Polynesians and 2 Melanesians; 5 severe valve lesions (0.2%), in 3 Polynesians, 1 Caucasian and 1 long QT syndrome (LQT1 type) in a Polynesian.

Fifty-three other CV diseases not linked to a risk of SCD were identified. They included 30 low grade valve diseases, 8 mild inter-auricular or ventricular communications, 8 low-risk arrhythmias, 6 cases of high blood pressure, and 1 low-risk coronary artery malformation. The incidence was 1.3-1.4 fold higher (P < 0.05) in Melanesians and Caucasians (2.3%-2.8%) than in Polynesians and Métis (2.1%-2.0%). In the severe or low grade valve diseases, a rheumatoid origin was identified in 60% athletes.

Other abnormal ECGs (n = 17) included 10 abnormal TWIs that could not be investigated by an MRI scan for economic reasons; 4 prolonged QTc, 2 were considered as normal after an expert opinion and 2 could not be investigated further. The 3/17 last abnormal ECGs were 1 abnormal Q wave and 2 abnormal QRS in right leads associated
with axis deviation, complete RBBB and RVH. No right cardiomyopathy was confirmed by complementary examination. Regular monitoring was recommended.

3.4 - 2012-2017, follow-up

A 5-year follow up could be carried out only on the 1 450 New Caledonian studied. Indeed PPE was not regularly organized in the 13 other Pacific Islands. Despite the PPE, 3 athletes died of a SCD. The first one was an 18 year old male Melanesian football player diagnosed with HCM. Although medically advised against participation in intensive sports, he continued to play and died during a football match. The second was a female Polynesian who died at rest. ECG and echocardiography carried out previously were normal. The diagnosis of prolonged QT was suspected because her untrained sister presented a prolonged QTc > 480 ms. Neither an autopsy nor a genetic test could be carried out. The third athlete was a Caucasian male triathlete with normal ECG and echocardiography. He died during a swimming session. The autopsy, without genetic or toxicology tests, was concluded as negative. In first-degree relatives, ECG, echocardiography and 24h Holter ECG recordings were normal. PPE could not prevent 2 if not 3 SCDs in 5 years in 1 450 athletes per year. It corresponded to 1 SCD per 2 416 or 3 625 athletes/year.

4 – Discussion
The first aim of this study was to present the results of a PPE carried out on 2,281 athletes from Pacific Islands. The PPE performed was fully normal for more than 85%. This value is close to the 90% found by Fuller et al. in 5,615 high school athletes [17].

In the 15% athletes that had to undergo further investigations, 3.9% CV abnormalities were found. They included 3.1% proven CV diseases and 0.7% suspected CV diseases as the CV investigations could not be carried out fully.

CV diseases linked to a risk of SCD were limited to 0.8% athletes.

The second aim of the study was to identify ECG pattern related to ethnicity, gender and training level. Melanesian athletes presented with frequent TWIs in V2-V4 similar to those described in Afro-Caribbean athletes. Most of infero-lateral TWIs were associated with cardiomyopathies. ECG findings were different between males and females and between low and high level trained athletes confirming that the 2017 international recommendations [3] are applicable for Pacific Island athletes.

4.1 - History and physical examination

Positive family, personal history, symptoms and physical examination were present in 14.6% of the athletes. Of them, only 20% presented with a proven or suspected cardiac disease. From these data, it must not be concluded that history and physical examination are of minor importance in athletes’ PPE. Indeed, in the 3.9% athletes having a cardiac abnormality, 50% athletes had one or more positive item. These data are in contrast with previous studies showing that history and physical examination led to an accurate diagnosis in only 1 to 3% of athletes [18, 19]. Effectiveness to detect diseases is debated
as many cardiovascular diseases remain silent and difficult to diagnose as some warning symptoms may be misinterpreted by medical providers [20].

These data confirm that history and physical examination remain of major importance in athletes’ PPE [21].

4.2 - ECG analysis

Of the 13 studied ECG criteria, 9 were related to ethnicity, 13 to gender and 9 to the training level.

Early repolarization concerned around 50% of athletes. This ECG pattern that has classically been viewed as benign [14] confirmed previous studies concerning Afro-Caribbean athletes [22].

Melanesians presented more frequent abnormal TWIs than other ethnicities. Most of TWIs were found in athletes under the age of 16 and TWIs were located in V2 to V4 leads confirming previous observations in young Caucasians [23]. They were often associated with a J point elevation and a domed ST segment. As none of them presented any CV disease, they were considered as early repolarization as in the case of Afro-Caribbean athletes [3].

The relationship observed between ECG criteria and training level confirms that the classifications [15, 19, 24] proposed for the interpretation of athletes’ ECG and especially the 2017 international recommendations [3] are applicable for Pacific Island athletes.

4.3 - Results of the pre-participation evaluation
Out of the athletes, 3.9% presented with a CV disease that needed a regular medical follow-up, 0.8% were identified with a cardiac disease associated with a potential risk of SCD. These values are close to those reported in a review of 12 studies grouping 55,396 athletes [1].

Athletes with SCD risk were contraindicated to competition, training being allowed only at a low intensity. Unfortunately, not all followed these recommendations as at least one died during a football match. For WPW, athletes were allowed to train and to compete again after successful ablation treatment.

The 2.6/1,000 athlete prevalence of HCM found in the present study is also close to the prevalence found by Wilson et al. [9] i.e. 3.2/1,000 in West Asian Athletes. In these 2 studies, Afro-Caribbeans or Melanesians were highly present 31% as opposed to 25%.

This data highlights that cardiomyopathies concern in the same way Melanesians and Polynesians. Only 50% were symptomatic, while all had an abnormal ECG confirming the value of this exam [3, 5, 25].

Specific attention was given to the 45 athletes with deep TWIs. In this population, 4 HCMs were diagnosed. However, because of economic reasons, all athletes did not undergo echocardiography and cardiac MRI. Thus, the diagnosis of cardiomyopathy was probably underestimated as MRI alone increases by 35% the diagnosis of HCM in athletes with ITWs and a normal echocardiography [26].

In the present study, the incidence of WPW was high and close to the total incidence of the cardiomyopathies, confirming previous screening studies [1, 7].

A major observation resulting from this study was the high prevalence (1.5%) of valvular diseases of rheumatoid origin. The prevalence was 3.9 fold higher than those
reported in athletes of Western and Middle East countries [1], confirming that RHD remains highly prevalent in the Pacific region [27, 28].

4.4 - Incidence rate of sudden cardiac death.

An estimate of the efficacy of the PPE was somewhat difficult. Indeed, 18 athletes were identified with a cardiac disease associated with a potential risk of SCD. Clearly, PPE could not detect nor prevent 2 SCDs in the New Caledonian population confirming previous observation in athletes [17, 29, 30]. In the present study, the observed SCD rate was very high when compared with previous ones. However, strength of the study was the sources of data that were reliable. Indeed, New Caledonia is a small country easy to survey. It is also pointed out that these values were close to values found by Harmon et al. [31] in black African-American basketball players.

4.5 - Limitations of the study

A major limitation of this study was the lack of control groups. However, it seemed very hard to recruit a large group of healthy volunteers from the populations studied. As we compared data observed in LT and HT levels athletes, it was thought that the LT level subjects could be considered close to the 2-3 h/week average of school, college or adult untrained people.

As emphasised above, a detailed and exhaustive CV evaluation deemed necessary after the initial examination could not be carried out because of economic reasons and/or lack of medical staff and facilities.

5 - Conclusion
This study gives for new information concerning family and personal history, symptoms, physical examination, ECG findings and cardiovascular abnormalities detected after a PPE in a large population of Pacific Island athletes.

A normal PPE was recorded for 85% of athletes. ECG findings were different between males and females and between low and high level trained athletes confirming that the 2017 international recommendations [3] are applicable for Pacific Island athletes.

Melanesians presented with frequent TWIs in V2-V4 similar to those described in Afro-Caribbean athletes. Most of other TWIs were associated with cardiomyopathies.

Overall, 3.9% of athletes presented a CV disease requiring an annual CV follow-up and 0.8% athletes were disqualified from competitive sport due to the risk of SCD. A high prevalence of RHD 1.5% was found. All these data confirm the importance of a PPE including ECG for Pacific Island athletes.

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Contributors

JCC had full access to all data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. JCC, FE, JCB and FC had the idea for and designed the study. All authors had a substantial contribution to the conception and design and to the collection of the data. JCC and FE provided statistical analysis. JCC, FE, AG, FR, KI, FC were responsible for interpretation of the data. JCC, FE and FC drafted the manuscript and submitted the paper for publication. All authors approved the final version.

Findings
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Authors have no conflict of interest to declare.
References

Figures legend

**Figure 1**: Number (%) of diagnosis elements found in the 88 cardiovascular (CV) abnormalities of the 2 281 studied athletes after questionnaire concerning family history, personal history, symptoms and physical examination and after ECG and complementary investigations. CVD: cardiovascular disease.

**Figure 2**: Cardiovascular abnormalities, n=88, with sudden cardiac death risk, n=18, (A), without any sudden cardiac death risk, n=53, (B), with isolated abnormal ECG, n=17, (C). QT : Prolonged QT; CV mal: cardiovascular malformation; Inter AV: inter atrio-ventricular communication.

**Figure 3**: ECG of the 3 athletes that died of a sudden cardiac death. (A) was a Caucasian male triathlete with normal ECG. He died during a swimming session. (B) was a female Polynesian who died at rest. ECG was normal. The diagnosis of prolonged QT was suspected because her untrained sister presented a prolonged QTc > 480 ms. (C) was an 18 year old male Melanesian football player diagnosed with HCM that died during a football match.
Table 1. Mean ± SD characteristics of the 1565 males in relation to ethnicity and training level.

<table>
<thead>
<tr>
<th>Ethnicity</th>
<th>Melanesians n = 424</th>
<th>Polynesians n = 488</th>
<th>Caucasians n = 331</th>
<th>Métis n = 322</th>
<th>LT n = 592</th>
<th>HT n = 973</th>
<th>Total n = 1565</th>
</tr>
</thead>
<tbody>
<tr>
<td>Training (h/w) SD</td>
<td>8.6 ± 5.7</td>
<td>9.2 ± 6.0</td>
<td>9.0 ± 4.0</td>
<td>8.2 ± 3.4</td>
<td>4.8* ± 1.2</td>
<td>11.2 ± 5.0</td>
<td>8.8* ± 5.1</td>
</tr>
<tr>
<td>Age (years) SD</td>
<td>18.9 ± 6.6</td>
<td>21.0 ± 5.6</td>
<td>19.4 ± 7.1</td>
<td>19.5 ± 6.4</td>
<td>19.3* ± 6.7</td>
<td>20.1 ± 6.3</td>
<td>19.8* ± 6.4</td>
</tr>
<tr>
<td>Height (cm) SD</td>
<td>168.3 ± 14.2</td>
<td>179.8 ± 8.1</td>
<td>173.1 ± 11.6</td>
<td>174.7 ± 11.5</td>
<td>172.5* ± 13.1</td>
<td>175.3 ± 11.6</td>
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<tr>
<td>Weight (kg) SD</td>
<td>66.7 ± 20.7</td>
<td>82.0 ± 16.5</td>
<td>65.1 ± 16.1</td>
<td>72.2 ± 18.4</td>
<td>71.0* ± 20.4</td>
<td>73.0 ± 18.6</td>
<td>72* ± 19</td>
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<tr>
<td>HR (bpm) SD</td>
<td>66.3 ± 12.4</td>
<td>66.2 ± 11.5</td>
<td>62.4 ± 11.8</td>
<td>63.0 ± 11.4</td>
<td>67.4* ± 12.3</td>
<td>63.2 ± 11.3</td>
<td>64.8* ± 11.9</td>
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<td>PR (ms) SD</td>
<td>152 ± 23</td>
<td>156 ± 22</td>
<td>147 ± 22</td>
<td>152 ± 25</td>
<td>150* ± 23</td>
<td>154 ± 23</td>
<td>152* ± 23</td>
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<tr>
<td>LAE (%) SD</td>
<td>2.8 ± 0.2</td>
<td>0.2 ± 0.3</td>
<td>4.5 ± 0.6</td>
<td>3.0 ± 0.6</td>
<td>0.2* ± 0.2</td>
<td>0.7 ± 0.4</td>
<td>0.5* ± 0.3</td>
</tr>
<tr>
<td>RAE (%) SD</td>
<td>0.2 ± 0.2</td>
<td>0.6 ± 0.6</td>
<td>0.6 ± 0.6</td>
<td>0.6 ± 0.6</td>
<td>0.2* ± 0.2</td>
<td>0.7 ± 0.4</td>
<td>0.5* ± 0.3</td>
</tr>
<tr>
<td>QRS (ms) SD</td>
<td>96 ± 12</td>
<td>98 ± 11</td>
<td>97 ± 12</td>
<td>98 ± 11</td>
<td>95* ± 12</td>
<td>98 ± 11</td>
<td>97* ± 11</td>
</tr>
<tr>
<td>LVH (%) SD</td>
<td>15.6 ± 19.9</td>
<td>25.1 ± 11</td>
<td>25.1 ± 11</td>
<td>17.1 ± 12</td>
<td>15.2* ± 12</td>
<td>21.7 ± 11</td>
<td>19.2* ± 11</td>
</tr>
<tr>
<td>RVH (%) SD</td>
<td>5.4 ± 13.9</td>
<td>7.9 ± 0.3</td>
<td>7.9 ± 0.6</td>
<td>8.1 ± 0.2</td>
<td>7.3* ± 0.2</td>
<td>10.3 ± 0.4</td>
<td>9.1* ± 0.3</td>
</tr>
<tr>
<td>IRBBB % SD</td>
<td>56.4 ± 0.7</td>
<td>45.9 ± 1.2</td>
<td>67.4 ± 1.2</td>
<td>60.9 ± 1.2</td>
<td>54.7 ± 1.2</td>
<td>57.3 ± 1.2</td>
<td>56.4* ± 1.3</td>
</tr>
<tr>
<td>CRBBB (%) SD</td>
<td>0.2 ± 0.2</td>
<td>0.2 ± 0.2</td>
<td>0.3 ± 0.3</td>
<td>0.6 ± 0.6</td>
<td>0.2 ± 0.2</td>
<td>0.4 ± 0.4</td>
<td>0.3* ± 0.3</td>
</tr>
<tr>
<td>ILBBB (%) SD</td>
<td>0.7 ± 1.2</td>
<td>1.2 ± 0.6</td>
<td>1.2 ± 0.6</td>
<td>1.2 ± 0.6</td>
<td>1.2 ± 0.6</td>
<td>0.8 ± 0.6</td>
<td>11.0* ± 0.8</td>
</tr>
<tr>
<td>ERP ≥2 % SD</td>
<td>60.6 ± 26</td>
<td>63.7 ± 23</td>
<td>58.6 ± 23</td>
<td>65.5 ± 23</td>
<td>56.4* ± 23</td>
<td>65.7 ± 23</td>
<td>62.2* ± 23</td>
</tr>
<tr>
<td>TWI (%) SD</td>
<td>2.6 ± 1.2</td>
<td>1.2 ± 1.2</td>
<td>1.5 ± 1.5</td>
<td>1.2 ± 1.2</td>
<td>1.5* ± 1.2</td>
<td>2.1 ± 1.2</td>
<td>1.9* ± 1.2</td>
</tr>
<tr>
<td>QTc (ms) SD</td>
<td>413 ± 22</td>
<td>413 ± 23</td>
<td>407 ± 23</td>
<td>409 ± 23</td>
<td>411 ± 22</td>
<td>410 ± 23</td>
<td>411* ± 23</td>
</tr>
</tbody>
</table>
SD standard deviation; $P<0.05$

$^a$ Melanesians vs. Polynesians; $^b$ Melanesians vs. Caucasians; $^c$ Melanesians vs. Mètis
$^d$ Melanesians vs. Caucasians and Mètis; $^e$ Polynesian vs. Caucasian; $^f$ Polynesian vs. Mètis
$^g$ Polynesian vs. Caucasian and Mètis; $^h$ Caucasian vs. Mètis;

* LT vs. HT; $^*$ males vs. females see table 1b
Table 2. Mean ± SD characteristics of the 716 females in relation to ethnicity and training level.

<table>
<thead>
<tr>
<th>Ethnicity</th>
<th>Melanesians n = 278</th>
<th>Polynesians n = 165</th>
<th>Caucasians n = 138</th>
<th>Métis n = 135</th>
<th>LT n = 353</th>
<th>HT n = 363</th>
<th>Total n = 716</th>
</tr>
</thead>
<tbody>
<tr>
<td>Training (h/w) SD</td>
<td>7.6 a, 5.3</td>
<td>9.4, 5.4</td>
<td>7.9, 3.5</td>
<td>8.4, 4.5</td>
<td>4.6*, 1.3</td>
<td>11.7, 4.6</td>
<td>8.2*, 4.9</td>
</tr>
<tr>
<td>Age (years) SD</td>
<td>17.7 ac, 6.4</td>
<td>19.7, 6.2</td>
<td>18.0, 6.4</td>
<td>19.3, 7.0</td>
<td>17.5*, 6.3</td>
<td>19.6, 6.6</td>
<td>18.6*, 6.5</td>
</tr>
<tr>
<td>Height (cm) SD</td>
<td>161.0 ac, 9.6</td>
<td>170.5 g, 7.7</td>
<td>164.3, 8.6</td>
<td>166.5, 8.1</td>
<td>163*, 9.4</td>
<td>167, 9.0</td>
<td>165*, 9.5</td>
</tr>
<tr>
<td>Weight (kg) SD</td>
<td>59.0 ac, 14.2</td>
<td>70.2 g, 14.3</td>
<td>57.0 h, 12.6</td>
<td>61.8, 12.1</td>
<td>60*, 14.8</td>
<td>64, 13.6</td>
<td>62*, 14.4</td>
</tr>
<tr>
<td>HR (bpm) SD</td>
<td>71.0 ad, 14.1</td>
<td>67.0, 12.5</td>
<td>66.7, 12.8</td>
<td>67.7, 12.4</td>
<td>70*, 13.7</td>
<td>67, 12.8</td>
<td>69*, 13.3</td>
</tr>
<tr>
<td>PR (ms) SD</td>
<td>144 a, 21</td>
<td>150 e, 21</td>
<td>141 h, 20</td>
<td>147, 20</td>
<td>143*, 22</td>
<td>147, 20</td>
<td>145*, 21</td>
</tr>
<tr>
<td>LAE (%)</td>
<td>1.8 ad, 0.6 c</td>
<td>0.6 e, 0.7</td>
<td>4.3 h, 0.7</td>
<td>0.7, 2.8</td>
<td>2.3*, 1.4</td>
<td>1.4, 0.8</td>
<td>1.8*, 0.8</td>
</tr>
<tr>
<td>RAE (%)</td>
<td>0</td>
<td>2.4</td>
<td>0.7</td>
<td>0.6* 1.2</td>
<td>0.6, 0.3</td>
<td>0.1, 0.1</td>
<td>0.8*, 0.1</td>
</tr>
<tr>
<td>QRS (ms) SD</td>
<td>89, 9</td>
<td>89</td>
<td>89</td>
<td>90, 11</td>
<td>88*, 9.2</td>
<td>90, 10.6</td>
<td>89*, 9.9</td>
</tr>
<tr>
<td>LVH (%)</td>
<td>3.2 ab, 1.4</td>
<td>4.2 e, 1.4</td>
<td>1.8, 1.1</td>
<td>3.7, 2.8*</td>
<td>4.1, 3.2</td>
<td>4.1, 3.8</td>
<td>4.1* 3.2</td>
</tr>
<tr>
<td>RVH (%)</td>
<td>2.5 a, 6.7 g</td>
<td>2.9</td>
<td>2.9</td>
<td>2.2, 3.1*</td>
<td>3.9, 3.5</td>
<td>3.9, 3.5</td>
<td>3.9, 3.5</td>
</tr>
<tr>
<td>IRBBB %</td>
<td>37.8 ad, 27.9 g</td>
<td>52.2, 14.3 h</td>
<td>48.1, 13.3</td>
<td>39.9, 9.3*</td>
<td>40.5, 15.4</td>
<td>40.2, 12.4</td>
<td>40.2, 12.4</td>
</tr>
<tr>
<td>CRBBB (%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0, 0.3</td>
<td>0.3, 0.1*</td>
<td>0.3, 0.1*</td>
<td>0.3, 0.1*</td>
</tr>
<tr>
<td>ILBBB (%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0, 0.3</td>
<td>0.3, 0.1*</td>
<td>0.3, 0.1*</td>
<td>0.3, 0.1*</td>
</tr>
<tr>
<td>ERP ≥2 %</td>
<td>12.6 ad, 9.7 g</td>
<td>14.3 h, 13.3</td>
<td>13.3, 9.3</td>
<td>15.4, 12.4*</td>
<td>15.4, 12.4</td>
<td>15.4, 12.4</td>
<td>15.4, 12.4</td>
</tr>
<tr>
<td>TWI (%)</td>
<td>3.6 ad, 1.2 e</td>
<td>2.2 h, 0.7</td>
<td>2.8* 1.7</td>
<td>1.7, 2.2*</td>
<td>1.7, 2.2</td>
<td>1.7, 2.2</td>
<td>1.7, 2.2</td>
</tr>
<tr>
<td>QTc (ms) SD</td>
<td>422 b, 24</td>
<td>425 e, 22</td>
<td>416 h, 25</td>
<td>422, 22</td>
<td>421, 25</td>
<td>423, 25</td>
<td>422*, 23</td>
</tr>
</tbody>
</table>
SD standard deviation; P<0.05
a Melanesians vs. Polynesians; b Melanesians vs. Caucasians; c Melanesians vs. Métis
d Melanesians vs. Caucasians and Métis; e Polynesian vs. Caucasian; f Polynesian vs. Métis
g Polynesian vs. Caucasian and Métis; h Caucasian vs. Métis.
*LT vs. HT; *males vs. females see table 1a
Table 3. Number (%) of cardiovascular (CV) abnormalities (1) with risk of sudden cardiac death (SCD), (2) without any risk of SCD, (3) isolated abnormal ECGs without any diagnosis and (4) number of SCDs over the 5-year study period in the 2 281 athletes grouped by ethnicity, gender and training level.

|                | Ethnicity | Melanesians | Polynesians | Caucasians | Métis | Males | Females | LT 945 | HT 1336 | Total 2 281
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
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<th></th>
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<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>(1) 18 CV</td>
<td>CM</td>
<td>2</td>
<td>3</td>
<td>0</td>
<td>1</td>
<td>6</td>
<td>0</td>
<td>1</td>
<td>5</td>
<td>6</td>
</tr>
<tr>
<td>abnormalities</td>
<td>WPW</td>
<td>2</td>
<td>4</td>
<td>0</td>
<td>0</td>
<td>5</td>
<td>0</td>
<td>2</td>
<td>4</td>
<td>6</td>
</tr>
<tr>
<td>with SCD risk</td>
<td>S. Val. lesion</td>
<td>0</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>4</td>
<td>0</td>
<td>0</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>Prolonged QT</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>Sub-total (%)</td>
<td>4 (0.6)</td>
<td>11 (1.7)</td>
<td>1 (0.2)</td>
<td>2 (0.4)</td>
<td>16 (1.0)</td>
<td>2 (0.3)*</td>
<td>3 (0.3)*</td>
<td>15 (1.1)</td>
<td>18 (0.8)</td>
</tr>
<tr>
<td>(2) 53 CV</td>
<td>Other valve lesion</td>
<td>11</td>
<td>6</td>
<td>8</td>
<td>5</td>
<td>20</td>
<td>10</td>
<td>14</td>
<td>16</td>
<td>30</td>
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<tr>
<td>abnormalities</td>
<td>Inter AV com.</td>
<td>4</td>
<td>0</td>
<td>3</td>
<td>1</td>
<td>5</td>
<td>0</td>
<td>5</td>
<td>3</td>
<td>8</td>
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<tr>
<td>with regular</td>
<td>Arrhythmia</td>
<td>1</td>
<td>4</td>
<td>1</td>
<td>2</td>
<td>6</td>
<td>0</td>
<td>4</td>
<td>4</td>
<td>8</td>
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<tr>
<td>follow up</td>
<td>HBP</td>
<td>0</td>
<td>5</td>
<td>0</td>
<td>1</td>
<td>6</td>
<td>0</td>
<td>1</td>
<td>5</td>
<td>6</td>
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<tr>
<td></td>
<td>CVM</td>
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<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>1</td>
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<tr>
<td></td>
<td>Sub-total (%)</td>
<td>16 (2.3)</td>
<td>15 (2.3)</td>
<td>13 (2.8)</td>
<td>9 (2.0)</td>
<td>37 (2.4)</td>
<td>16 (2.2)</td>
<td>24 (2.5)</td>
<td>29 (2.2)</td>
<td>53 (2.3)</td>
</tr>
<tr>
<td>(1)+(2)</td>
<td>Total (%)</td>
<td>20 (2.8)</td>
<td>26 (4.0)</td>
<td>14 (3.0)</td>
<td>11 (2.4)</td>
<td>52 (3.3)*</td>
<td>18 (2.5)</td>
<td>27 (2.9)</td>
<td>43 (3.2)</td>
<td>71 (3.1)</td>
</tr>
<tr>
<td>(3) 17 isolated</td>
<td>T wave inversion</td>
<td>3</td>
<td>5</td>
<td>2</td>
<td>0</td>
<td>8</td>
<td>2</td>
<td>7</td>
<td>3</td>
<td>10</td>
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<tr>
<td>abnormal ECG</td>
<td>Prolonged QT</td>
<td>1</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>2</td>
<td>2</td>
<td>2</td>
<td>2</td>
<td>4</td>
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<td>CM suspicion</td>
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<td>0</td>
<td>0</td>
<td>2</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td>Q wave</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td>Sub-total (%)</td>
<td>4 (0.6)</td>
<td>7 (1.1)</td>
<td>2 (0.4)</td>
<td>4 (0.9)</td>
<td>14 (0.9)</td>
<td>3 (0.4)</td>
<td>9 (1.0)*</td>
<td>8 (0.6)</td>
<td>17 (0.7)</td>
</tr>
<tr>
<td>(1)+(2)+(3)</td>
<td>Total (%)</td>
<td>24 (3.4)</td>
<td>33 (5.1)</td>
<td>16 (3.4)</td>
<td>15 (3.3)</td>
<td>66 (4.2)*</td>
<td>22 (3.1)</td>
<td>36 (3.8)</td>
<td>52 (3.9)</td>
<td>88 (3.9)</td>
</tr>
<tr>
<td>(4) SCD over</td>
<td></td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>5yrs</td>
<td></td>
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</tbody>
</table>

CM: cardiomyopathy, WPW: Wolff-Parkinson-White, S. Val.: Severe valve, Vent: ventricular, Inter AV com: inter atrial or ventricular communication, HBP: high blood pressure, CVM: cardio vascular malformation, R: Right. LT, HT: low and high level of training, P<0.05
Melanesians vs. Polynesians; Melanesians vs. Caucasians; Melanesians vs. Métis
Melanesians vs. Caucasians and Métis; Polynesian vs. Caucasian; Polynesian vs. Métis; Polynesian vs. Caucasian and Métis
Caucasian vs. Métis; LT vs. HT; males vs. females
Highlights: What are the new findings?

 ► This study provides new information on the prevalence of cardiovascular disease and abnormalities in male and female Melanesians, Polynesian, Caucasians and Métis Pacific Island athletes and in low and highly trained athletes.
 ► Melanesians had a higher rate of T waves inversion in V2-V4 than others and were similar to those in Afro-Caribbean athletes.
 ► The prevalence of valvular diseases of rheumatoid origin was clearly higher than in Western and Middle East countries.
Figure 1

2,281 athletes

History and physical examination

Normal 1,947 (85.4)

Abnormal 334 (14.6)

36 (1.6) athletes had several answers

Family history 154 (6.8)

Personal history 73 (3.2)

Symptoms 46 (2.0)

Physical exam 97 (4.2)

Normal ECG 1,925 (84.4)

- Normal ECG 148 (6.5)
  - Proven CVD 2
  - Suspected CVD 1

- Abnormal ECG 6 (0.3)
  - Proven 5
  - Suspected CVD 1

- Normal ECG 54 (2.4)
  - Proven CVD 12
  - Suspected CVD 1

- Abnormal ECG 19 (0.8)
  - Proven 18
  - Suspected CVD 1

- Normal ECG 36 (1.6)
  - Proven CVD 10
  - Suspected CVD 1

- Abnormal ECG 10 (0.4)
  - Proven CVD 14
  - Suspected CVD 5

Abnormal ECG 22 (1.0)

- Normal ECG 310 (13.6)
  - Proven CV disease 21 (2.6)
  - Suspected CV disease 7 (0.3)

- Abnormal ECG 45 (2.0)
  - Proven CV disease 38 (1.7)
  - Suspected CV disease 7 (0.3)
Figure 2

A
- Severe valve lesion: 5
- Cardio myopathy: 6
- Wolff Parkinson White: 6

B
- Other valve lesion: 30
- HBP: 6
- Rhythm trouble: 8
- Inter AV communication: 8
- CV malf: 1

C
- Inverted T waves: 10
- Q waves: 1
- Cardio myopathy suspicion: 2
- Prolonged QT: 4