Left Atrial Enlargement in Young High-Level Endurance Athletes – Another Sign of Athlete’s Heart?

by

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Enlargement of the left atrium is perceived as a part of athlete’s heart syndrome, despite the lack of evidence. So far, left atrial size has not been assessed in the context of exercise capacity. The hypothesis of the present study was that LA enlargement in athletes was physiological and fitness-related condition. In addition, we tried to assess the feasibility and normal values of left atrial strain parameters and their relationship with other signs of athlete’s heart. The study group consisted of 114 international-level rowers (17.5 ± 1.5 years old; 46.5% women). All participants underwent a cardio-pulmonary exercise test and resting transthoracic echocardiography. Beside standard echocardiographic measurements, two dimensional speckle tracking echocardiography was used to assess average peak atrial longitudinal strain, peak atrial contraction strain and early left atrial diastolic longitudinal strain. Mild, moderate and severe left atrial enlargement was present in 27.2%, 11.4% and 4.4% athletes, respectively. There were no significant differences between subgroups with different range of left atrial enlargement in any of echocardiographic parameters of the left ventricle diastolic function, filling pressure or hypertrophy. A significant correlation was found between the left atrial volume index and maximal aerobic capacity (R > 0.3; p < 0.001). Left atrial strain parameters were independent of atrial size, left ventricle hypertrophy and left ventricle filling pressure. Decreased peak atrial longitudinal strain was observed in 4 individuals (3.5%). We concluded that LA enlargement was common in healthy, young athletes participating in endurance sport disciplines with a high level of static exertion and was strictly correlated with exercise capacity, therefore, could be perceived as another sign of athlete’s heart.

Key words: speckle tracking echocardiography, left atrium, maximal aerobic capacity, strain.

Introduction

Athlete’s heart as a result of intensive, prolonged exercise has been investigated for many years. It has been suggested that training, especially in endurance sports, can lead to both left and right atrial enlargement (D’Andrea et al., 2010; Pagourelias et al., 2013; Pelliccia et al., 2005). Many previous studies have proved that left ventricular remodeling due to physical activity is physiological as the diastolic and systolic function remains normal or even supernormal. Left atrium (LA) enlargement is perceived as physiological as well by extrapolation, even though data confirming this statement is lacking. LA is of particular interest owing to its involvement in
pathophysiology of atrial fibrillation (AF) which is arrhythmia more often occurring in athletes than in sedentary controls (Baldesberger et al., 2008).

Deformation (strain and strain rate) analysis, assessed with two-dimensional speckle tracking echocardiography (2D STE), is a new echocardiographic method that allows a non-invasive evaluation of the atrial function. Decreased LA strain and the strain rate enable to predict the onset of atrial fibrillation (Cameli et al., 2014; Shaikh et al., 2012; Spethmann et al., 2014; Yoon et al., 2012). Moreover, strain value correlates with atrial fibrosis in paroxysmal and persistent AF (Kuppahally et al., 2010). Left atrial deformation assessment has been previously carried out in male and female athletes (D’Andrea et al., 2008; D’Ascenzi et al., 2014, 2015; Sun et al., 2009). However, to our knowledge, there is no study that has determined normal values of LA strain in a large sample of elite athletes.

Even though increased exercise capacity is one of the most obvious results of intensive training, cardiopulmonary exercise test results were seldom correlated with echocardiography in the athletic population (Yamazaki et al., 2000). It is known that exercise capacity is correlated with left ventricle size (Whyte et al., 2000; Yamazaki et al., 2000). Whether LA size also correlates with fitness has not been studied before.

Among all sports, those which combine high static and dynamic exertion induce the morphologic adaptation of the heart to the highest degree. Therefore, the biggest LA enlargement can be expected in those groups of athletes (Pelliccia et al., 2005).

The hypothesis of the study was that LA enlargement in athletes was physiologic, fitness-related condition in both male and female athletes. In addition, we tried to assess the feasibility of left atrial strain parameters and their relation with other signs of athletes’ heart observed in echocardiography. Last of all, we aimed to establish the normal values of 2D STE left atrial strain in young, healthy athletes.

**Material and Methods**

The study group was comprised of 117 young, elite international-level rowers, average age of 17.5 years (15-21), participating in a routine annual medical check-up in the Institute of Sport (Warsaw, Poland) between January and March, 2013. The study was conducted during the preseason preparation period, in the middle of the intensive training program. The medical assessment was performed in the morning (at least 12 hours from the last training session) and consisted of consecutive stages: clinical and physical examination, resting ECG, transthoracic echocardiography and an exercise cardiopulmonary test. The incremental exercise test (3 min exercise bouts interspersed with 30 s rest) until exhaustion was performed on a Concept-II ergonometer (Morrisville, USA). The respiratory exchange parameters were measured with the MetaLyser 3B and MetaMax 3B system (Cortex, Germany). The exact methodology of the cardiopulmonary test had been previously described (Klusiewicz et al., 2014). Maximal aerobic capacity (VO2max) was expressed in l/min and in ml/kg/min. Exclusion criteria were hypertension, valvular heart disease, cardiomyopathy, absence of sinus rhythm, pulmonary disease, endocrine disease and/or renal failure. For final analysis, only participants with acceptable quality of echocardiographic examination were included. All athletes gave informed consent. The Research Ethics Committee of the Institute of Sport (Warsaw, Poland) approved of the study protocol.

Echocardiographic examination was performed by two experienced physicians, using a Vivid 7 machine (GE, Vingmed Ultrasound, Boston, USA), with a M4S 1.5 to 4.0 MHz transducer. ECG was continuously registered. Attention was given to adjust the ECG curve (lead selection, gain control) in order to present a well defined P wave and QRS complex. The examination was performed at rest in the left-lateral decubitus position using standard parasternal and apical views. All echocardiographic acquisitions were digitally stored and off-line analysis from three consecutive cardiac cycles was performed using dedicated software (EchoPac ver. 112 GE, USA). The chamber size measurements were performed according to recommendations (Lang et al., 2015; Rudski et al., 2010). In order to decrease the influence of preload-dependent parameters and to increase the quality of recorded data, all acquisitions were made during shallow end-expiration. Body surface area (BSA) was calculated according to the Mosteller’s formula.
LA measurements were performed at left ventricular end-systole, in atrial maximal size. The LA area was measured in a 4-chamber view. LA volume (LAV) was calculated by the Simpson’s method and then indexed to the body surface area as a LA volume index (LAVI). During LA size assessment the LA appendage and pulmonary veins were excluded and mitral valve annulus plane was used as the inferior border. LA enlargement was defined as follows (Lang et al., 2015): Mild $\geq 34$ml/m² and <42ml/m², moderate $\geq 42$ml/m² and <48ml/m², severe $>48$ml/m².

STE derived LA strain analysis was performed offline by one physician. The methodology was adopted to comply with methodology used in the most experienced echo-labs involved in athletes’ assessment which showed good reproducibility (D’Ascenzi et al., 2012). The ultrasound focus position was set on the level of LA to optimize the image quality, the frame rate was set between 60-80 Hz. The commercially available 2D semi-automated software was used (Echo Pac, ver. 112). The strain measurement was performed on generated curves. The scale was adjusted to set 0 level at the beginning of the QRS complex. The average (from all accepted segments) peak atrial longitudinal strain (PALS) and peak atrial contraction strain (PACS) were measured, as parameters of atrial reservoir and atrial pump (buster), respectively. The difference between PALS and PACS was early LA diastolic longitudinal strain (ELAS) and represented passive conduit function. PACS was defined as maximal longitudinal strain before the peak of the P wave (Figure 1).

To distinguish athletes with low PALS, normal values of PALS for the general healthy population were used from the study of Cameli et al. (2009) using the same methodology, echo system and software vendor.

**Statistical analysis**

For statistical analysis, commercially available software STATISTICA ver. 10 (StatSoft, Tulsa, USA) was used. Continuous variables are presented as mean ± standard deviation (SD), categorical variables as percentages. Normal distribution of all continuous variables was examined using the Shapiro-Wilk test. The unpaired t-test and the Mann-Whitney u-test were used according to data distribution to assess between-groups differences in continuous variables and chi-square for categorical variables. The correlations between continuous variables were assessed with the Pearson correlation coefficient (r) and between categorical variables with the Spearman rank correlation coefficient, correlations with R$>0.3$ with p<0.05 were reported. In order to assess significance of differences between multiple subgroups analysis of variance (ANOVA) was used. The post hoc Scheffe test was performed to distinguish the intergroup differences. The level of significance was set at p < 0.05.

**Results**

**Clinical characteristics and conventional echocardiography**

General characteristics of the examined group are presented in Table 1. From primarily examined 117 athletes, 114 were finally included in the analysis. All exclusions were due to poor image quality. No cardiovascular disorder was found. From the analyzed 114 athletes 53 were female (46.5%). Standard echocardiographic and Doppler parameters are summarized in Table 2. The parameters of LA size are presented in Table 3. Left atrium enlargement according to LAVI was present in nearly half of the athletes (43%) and was more frequent in man then in women (52.5% vs 32.1%; p <0.05). Mild, moderate or severe LA enlargement was present in 27.2%, 11.4% and 4.4% athletes, respectively. Following ANOVA results, there were no significant differences between subgroups with different ranges of LA enlargement with regard to left ventricular parameters of the diastolic function (mitral E and A waves velocities, E/A ratio), filling pressure (E/e’ ) and hypertrophy (LVMi, IVSd).

**Cardiopulmonary exercise test**

Cardiopulmonary exercise test parameters are presented in Table 3. Maximal aerobic capacity was, as expected, higher in males, both in absolute (l/min) and relative (ml/min/kg) values. Resting and maximal exercise heart rates were similar in males and females. VO2max correlated significantly with most LA size parameters (except LA area indexed to BSA).

**Left atrial strain analysis**

From the primary group of 117 athletes, 114 (97%) had acceptable image quality for STE analysis. From this group the assessment of 97.7%
of LA segments (668/684) was feasible. In 10 athletes (9%) one segment and in 3 athletes (2.6%) two segments could not be traced properly. The LA roof and interatrial septum were the regions of poor tracking quality in 6 athletes and 5 athletes, respectively (see technical consideration). STE derived LA strain parameters are presented in Table 4 with 5-95 percentile as suggested by normal ranges. Linear correlation between LA strain and parameters of: diastolic function (mitral inflow E velocity, E deceleration time, A velocity, E/A ratio, TDI mitral velocities: E', A', E/E'), LV size and hypertrophy (LVM, LVMi, LVEDd, IVSd), exercise capacity (VO2max) and LA size (LAV, LAVI, LA area, LA diameter) were assessed. No significant correlation was found between LA functional parameters and LV and LA morphology. No significant correlation was noted between LA strain and parameters of exercise capacity. According to normal value for general population, decreased LA PALS (<29%) was present in 4 athletes (3.5%). The only significant differences in athletes with reduced PALS were a larger intraventricular septum diameter (1.15 vs 1.07cm), a lower mitral E wave velocity (78cm/s vs 98cm/s) and consequently higher mitral E/E' (6.0 vs 5.0).

**Figure 1**

Example of left atrial strain (%) curves with selected measured parameters.

PALS- Peak atrial longitudinal strain; PACS – peak atrial contraction strain, ELAS - Early atrial longitudinal strain;

**Table 1**

| Subjects’ characteristics | All (N = 114) | Women (n = 53) | Men (n = 61) | p |
|--------------------------|--------------|----------------|-------------|---|
| Age (yrs)                | 17.5 ± 1.5   | 17.5 ± 1.39    | 17.4 ± 1.5  | NS|
| Training experience (yrs)| 4.3 ± 1.7    | 4.37 ± 1.9     | 4.3 ± 1.6   | NS|
| Body mass (kg)           | 74.2 ± 9.6   | 67.8 ± 7.5     | 79.9 ± 7.5  | <0.001|
| Height (cm)              | 180.8 ± 8.8  | 173.4 ± 5.6    | 187.2 ± 5.5 | <0.001|
| BMI (kg/m²)              | 22.7 ± 2.1   | 22.5 ± 2.3     | 22.8 ± 1.8  | <0.001|
| BSA (m²)                 | 1.9 ± 0.16   | 1.8 ± 0.12     | 2.04 ± 0.11 | <0.001|
| Heart rate (bpm)         | 65.4 ± 12.0  | 64.2 ± 12      | 66.4 ± 11   | NS|

BMI – body mass index, BSA – body surface area, NS – no significant difference.

Data expressed as mean ± SD
Table 2

| Variable                  | All (N = 114) | Women (n = 53) | Men (n = 61) | p     |
|---------------------------|---------------|----------------|--------------|-------|
| RVOTprox (cm)            | 2.92 ± 0.3    | 2.80 ± 0.31    | 3.02 ± 0.23  | <0.001|
| LVDd (cm)                | 5.02 ± 0.4    | 4.77 ± 0.40    | 5.24 ± 0.32  | <0.001|
| LVDD/BSA (cm/m²)         | 2.61 ± 0.22   | 2.65 ± 0.26    | 2.58 ± 0.17  | NS    |
| IVSd (cm)                | 1.08 ± 0.1    | 1.02 ± 0.10    | 1.13 ± 0.11  | <0.001|
| PWDd (cm)                | 1.05 ± 0.1    | 0.99 ± 0.12    | 1.10 ± 0.12  | <0.001|
| LVM (g)                  | 246 ± 55      | 206 ± 35       | 281 ± 45     | <0.001|
| LVMi (g/m²)              | 127 ± 22      | 115 ± 19       | 138 ± 18     | <0.001|
| Ao (root) (cm)           | 3.00 ± 0.3    | 2.83 ± 0.26    | 3.14 ± 0.34  | <0.001|
| Aoo (root) (cm²/m²)      | 1.56 ± 0.15   | 1.57 ± 0.14    | 1.54 ± 0.17  | NS    |
| LV EF (%)                | 68.4 ± 2.3    | 68.1 ± 2.0     | 69.8 ± 2.5   | NS    |
| TAPSE (mm)               | 23.7 ± 3.0    | 23.2 ± 2.9     | 24.2 ± 3.1   | NS    |
| MV E (m/s)               | 0.94 ± 0.2    | 0.96 ± 0.17    | 0.92 ± 0.15  | NS    |
| MV A (m/s)               | 0.46 ± 0.1    | 0.45 ± 0.12    | 0.46 ± 0.13  | NS    |
| MV E/A                   | 2.20 ± 0.6    | 2.26 ± 0.55    | 2.15 ± 0.57  | NS    |
| MV E DecT (ms)           | 214.3 ± 53    | 212 ± 48       | 216 ± 56     | NS    |
| MV e' (cm/s)             | 19.2 ± 2.2    | 19.5 ± 2.1     | 18.9 ± 2.2   | NS    |
| MV a' (cm/s)             | 8.3 ± 1.6     | 8.42 ± 1.8     | 8.23 ± 1.4   | NS    |
| MV s' (cm/s)             | 11.1 ± 1.7    | 10.7 ± 1.6     | 11.35 ± 1.7  | 0.04  |
| MV E/e'                  | 4.96 ± 0.9    | 4.99 ± 0.93    | 4.93 ± 0.85  | NS    |

RVOT – right ventricle outflow tract, d – diameter, 
LVED – left ventricular end-diastolic diameter, IVS – interventricular septum and-diastolic diameter, PWD – posterior wall end-diastolic diameter, 
Ao – aortic root diameter, EF – ejection fraction, TAPSE – tricuspid annular plane systolic excursion, 
MV– mitral valve, E – early inflow velocity, A – atrial inflow velocity, DecT – deceleration time, e’ – early diastolic velocity, a’ – atrial diastolic velocity, s’– systolic velocity, NS – no significant difference.
Data expressed as mean ± SD – standard deviation

Table 3

| Variable                  | All (N = 114) | Women (n = 53) | Men (n = 61) | p     |
|---------------------------|---------------|----------------|--------------|-------|
| LA (cm)                   | 3.65 ± 0.40   | 3.50 ± 0.3     | 3.77 ± 0.44  | <0.001|
| LAA (cm²)                 | 22.6 ±3.4     | 20.1 ±2.6      | 24.0 ± 3.4   | <0.001|
| RAA (cm²)                 | 18.0 ±3.1     | 16.6 ±2.7      | 19.3 ± 2.8   | <0.001|
| LA/BSA (cm²/m²)           | 1.90 ±0.20    | 1.95 ±0.17     | 1.86 ± 0.22  | 0.02  |
| RAA/BSA (cm³)             | 9.35 ±1.4     | 9.20 ±1.53     | 9.48 ± 1.27  | NS    |
| LAA/BSA (cm³/m²)          | 11.7 ±1.6     | 11.7 ±1.5      | 11.8 ±1.6    | NS    |
| LAVI (cm³/m²)             | 34.3 ±6.9     | 31.9 ±6.1      | 36.3 ±6.9    | <0.001|
| VO₂max (l/min)            | 4.20 ±0.9     | 3.36 ±0.4      | 4.94 ± 0.6   | <0.001|
| VO₂max (ml/kg/min)        | 56.4 ±8.2     | 49.9 ±5.1      | 62.1 ±5.8    | <0.001|
| HRmax (/min)              | 195 ±8.2      | 194 ±7.6       | 195 ±8.7     | NS    |

LA – left atrial diameter, LAA – left atrial area, RAA – right atrial area, 
BSA – body surface area, LAV – left atrial volume, LAVI – left atrial volume index, 
VO₂max – maximal aerobic capacity, HRmax – maximal heart rate.
Data expressed as mean ± SD – standard deviation

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### Table 4

Mean values of two dimensional speckle tracking echocardiography left atrial strain parameters assessed in 4 – chamber view, with 5-95 percentile as normal value.

|                  | All (N = 114) | 5 - 95 Percentile | Women (n = 53) | Men (n = 61) | p (Men vs Women) |
|------------------|---------------|-------------------|----------------|--------------|-----------------|
| PALS (%)         | 42.3 ± 8.1    | 30.0 – 58.1       | 42.9 ± 0.1     | 41.7 ± 0.1   | NS              |
| PACS (%)         | 12.1 ± 3.6    | 6.75 –19.0        | 12.2 ± 3.6     | 12.1 ± 3.7   | NS              |
| ELAS (%)         | 30.2 ± 7.2    | 20.6 - 44.4       | 30.7 ± 7.4     | 29.7 ± 7.0   | NS              |

PALS – Peak atrial longitudinal strain, PACS – peak atrial contraction strain, ELAS – Early atrial longitudinal strain, TPALS – time to peak atrial strain, TPACS – time to peak atrial contraction strain, RR – time between consecutive heart cycles.

Data expressed as mean ± SD – standard deviation

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**Figure 2**

Correlation between VO2max and LAV ($r=0.35; p<0.001$)

Correlation between left atrial volume and maximal aerobic capacity.

LAV – left atrial volume, VO2max – maximal aerobic capacity
Discussion

In our study we demonstrated that left atrial enlargement was common condition in young, healthy rowers. LA dilatation was more often observed in males, and was not related with LV hypertrophy or LV diastolic function what confirmed observations of D’Andrea et al. (2010). The most often postulated mechanism of physiological LA dilatation in athletes is due to an increased volume load in contrast to increased LV filling pressure observed in pathologic LV hypertrophy (Pelliccia et al., 2005). In our study, even though the selected group of athletes participated in one of the most demanding endurance sports, the degree of left atrial enlargement was surprisingly high, especially given the young age of participants. Observed LA volumes were comparable to those reported by Baggish et al. (2010) in senior, elite and sub-elite rowers. Despite the fact that in the updated Recommendation for Chamber Quantification published by Lang et al. (2015) the upper reference value of the LA volume index was increased, still more than 40% of young, healthy athletes exceeded the norms. This observation shows the need for establishing normal values of LAVI for this specific population what has been already postulated by other authors (Galderisi et al., 2015).

Left and right ventricle size is generally larger in male athletes, however, indexation with a body surface area can decrease observed differences (Konopka et al., 2015; Krol et al., 2011). A similar observation can be made with regard to LA enlargement in our examined group. In absolute numbers all LA size parameters (dimension, area and estimated volume) were significantly greater in males (Table 3). After indexation with the body surface area, differences of some measurements (atrial area) lose significance, but the most recommended parameter (LAVI) does not. The between gender differences in LAVI have been reported by other authors and discussed in the context of a lower incidence of AF in women (D’Andrea et al., 2010; Mont et al., 2009).

The relationship between results of cardiopulmonary tests and heart morphology in athletes has not been extensively studied. It was shown that VO2max correlated with the LV diameter in athletes, but not in obese controls (Yamazaki et al., 2000). Also the LV diameter and LV posterior wall thickness correlated positively with cycling performance in triathletes (Mondillo et al., 2011). To our knowledge, this study is the first to show a similar relationship of left atrial size with exercise capacity. This observation gives another evidence of physiological nature of LA enlargement due to intensive training.

The image quality which is a main limitation of 2D STE is very important in reliable quantitative strain assessment. Athletes, having in general a very good acoustic window, seem to be an appropriate population for the wider introduction of STE into clinical practice. In our study, the functional assessment of LA was feasible in nearly all examined athletes. The important information is that LA STE derived strain was independent of LA size, which had been postulated before based on smaller sample groups (N < 25) (D’Ascenzi et al., 2012; Gabrielli et al., 2014). LA systolic and diastolic strains in young athletes are also independent of the parameters of left ventricular hypertrophy and left ventricle filling pressure. In this regard, LA strain analysis can be used in differentiating pathological and physiological left ventricular hypertrophy in hypertrophic cardiomiopathy and in hypertension what has been investigated before (D’Andrea et al., 2008; Sun et al., 2009).

Although LA enlargement is common condition in athletes, only few data exist on the long term clinical consequences of this phenomenon. It has been shown that LA enlargement is more prominent in athletes with electrocardiographic changes such as early repolarisation (Konopka et al., 2016). There are theories suggesting that LA enlargement and remodeling may contribute to paroxysmal supraventricular arrhythmias in athletes (Benito et al., 2011; Guasch et al., 2013). Benito et al. (2011) on an animal mice model found that intensive, prolonged physical training led to increased LA dilatation and fibrosis accompanied with increased arrhythmia inducibility. Nevertheless, some studies have shown that despite high prevalence of LA dilatation in athletes participating in IIIC sports, AF develops in a small percentage of them (Pelliccia et al., 2005). Even though LA dilatation may be present in more than ¼ of endurance athletes, the highest
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prevalence of AF reported by Baldesberger et al. (2008) was “only” 10% in the 7th decade of life (Baldesberger et al., 2008). These data suggest that LA enlargement alone is not a good indicator of the risk for AF. Therefore, other predictors of AF should be sought. Functional atrial assessment seems to be one of possibilities.

In our study, we applied the most often cited methodology from the center experienced in sports cardiology (D’Ascenzi et al., 2012, 2014, 2015). In this setting, LA strain parameters assessed in 2D STE were very similar to other publications evaluating different groups of athletes, which strengthens the reproducibility of the method. The significance of the observed decreased (<29%) PALS in 4 athletes (3.5%), described as a normal for the general population by Camelli et al. (2009), needs further observation. So far the medical follow-up has not revealed any pathologies.

In summary, our study provides new information about LA morphologic and functional remodeling in athletes, which confirms that LA enlargement may be legitimately perceived as another sign of athlete’s heart.

Conclusions:

1. LA enlargement is common in healthy, young athletes participating in endurance sport disciplines with a high level of static exertion and is strictly correlated with exercise capacity, therefore, can be perceived as another sign of athlete’s heart.

2. In endurance athletes with LA enlargement, there are no echocardiographic signs of an impaired LV diastolic function or increased LV filling pressure.

3. The magnitude of left atrial strains assessed in STE is not correlated with LA enlargement, LV hypertrophy or LV diastolic function in the population of healthy, young competitive athletes.

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