Estimation of anaerobic threshold by cardiac repolarization instability: a prospective validation study

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Abstract

Background: Assessing lactate (LT) or anaerobic thresholds (AT) in athletes is an important tool to control training intensities and to estimate individual performance levels. Previously we demonstrated that ECG-based assessment of cardiac repolarization instability during exercise testing allows non-invasive estimation of AT in recreational athletes. Here, we validate this method in professional and amateur team sports athletes.

Methods: We included 65 team sports athletes (32 professionals and 33 amateur athletes; 51 men, 14 women, mean age 22.3 ± 5.2 years) undergoing a standardized incremental cycle exercise test. During exercise testing a high-resolution ECG (1000 Hz) was recorded in Frank-leads configuration and beat-to-beat vector changes of cardiac repolarization ($dT°$) were assessed by previously established technologies. Repolarization-based AT ($AT_{dT°}$) was estimated by its typical $dT°$-signal pattern. Additionally, LT was detected in accordance to methods established by Mader (LT Mader) and Dickhuth (LT Dickhuth).

Results: All athletes performed exercise testing until exhaustion with a mean maximum workload of 262.3 ± 60.8 W (241.8 ± 64.4 W for amateur athletes and 283.4 ± 49.5 W for professional athletes). Athletes showed $AT_{dT°}$ at 187.6 ± 44.4 W, LT Dickhuth at 181.1 ± 45.6 W and LT Mader at 184.3 ± 52.4 W. $AT_{dT°}$ correlated highly significantly with LT Dickhuth ($r = 0.96$, $p < 0.001$) and LT Mader ($r = 0.98$, $p < 0.001$) in the entire cohort of athletes as well as in the subgroups of professional and amateur athletes ($p < 0.001$ for all).

Conclusions: $AT_{dT°}$, defined by the maximal discordance between $dT°$ and heart rate, can be assessed reliably and non-invasively via the use of a high-resolution ECG in professional and amateur athletes.

Keywords: Cardiac repolarization, Autonomic nervous system, Anaerobic threshold, Professional athlete

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Background
Controlling exercise intensities has been shown to be one of the key measures to improve endurance capacity and performance: The concept of using submaximal workload parameters such as lactate (LT) or ventilatory thresholds (VT) to determine individual cardio-respiratory fitness and to schedule training intensities is commonly accepted in this context and has been demonstrated in endurance as well as team-sports athletes [1–3].

Assessment of parameters of the autonomic nervous system (ANS) has gained pronounced attention. The concept of testing biomarkers of the ANS for estimating thresholds is based on the idea that the ANS exerts distinct influences on the cardiorespiratory system during exercise to regulate heart rate, cardiac contractility and blood pressure. Recently, different study groups implemented the evaluation of autonomic biomarkers to control training intensities and to detect training-induced states of fatigue [4]. Measuring ANS biomarkers is promising as it is non-invasive and cost-efficient. Heart rate variability (HRV)-derived parameters reflecting ANS activity showed an association with LT in healthy and diseased cohorts [5–8]. Despite different methods to assess AT including metabolic markers (lactate, glucose), ventilatory responses, autonomic markers (e.g. HRV-derived markers such as RMSSD) or neuroendocrine markers (catecholamines), they all determine AT within close ranges [9]. However, as the physiological downstream is faster in some systems than in others, these different methods assessing the anaerobic threshold may show slightly different workloads at the AT. Nevertheless, as all these methods differ in invasiveness, costs, time-consumption and easiness of determination, new methods to indicate AT may be helpful.

Sympathetic-activity associated periodic repolarization dynamics (PRD) is a novel ECG-based parameter, that reflects influences of efferent cardiac sympathetic activity on the ventricular myocardium during repolarization [10, 11]. The non-invasive assessment via high-resolution ECG is based on beat-to-beat changes of the T wave vector (dT°) with periodic components of repolarization in the low-frequency range (≤ 0.1 Hz). Large clinical trials demonstrated that increased levels of PRD are strong predictors of sudden cardiac death (SCD) in patients with ischemic and non-ischemic cardiomyopathy [11–13].

As exercise is known to affect ventricular repolarization crucially [4, 14] we previously tested repolarization instability (dT° signal) during exercise testing [15]: We demonstrated that the dT° signal shows a characteristic three-phasic pattern that allows a reliable and non-invasive estimation of the anaerobic threshold (we called it AT_{dT°}) in healthy recreational athletes. This pattern at the anaerobic threshold is characterized by a maximal discordance of dT° and heart rate and this point highly significantly correlated with lactate thresholds measured by the methods of Mader and Dickhuth [15].

In the present study we validated this non-invasive ECG-based assessment of the anaerobic threshold by our previously described methods in a cohort of 65 team sport athletes to check if this method can be transferred to professional athletes as well as well-trained amateur athletes who are able to achieve markedly higher maximal workloads during exercise and have increased workloads at lactate thresholds.

Methods
Study population
We included 65 healthy team sport athletes (14 women, 51 men, mean age 22.3 ± 5.2 year. (standard deviation), minimum age 14 year, maximum age 36 year.) who underwent a graded cycle ergometer test until maximal exhaustion. Our study cohort consisted of 32 professional athletes (1st league and 2nd league European football clubs and 1st league basketball club) and 33 amateur athletes (4th and 6th league European football clubs). Exclusion criteria were acute or chronic infections, presence of pacemakers or implantable cardioverter defibrillators (ICDs), history of cardiovascular diseases or risk factors and other contraindications for performing exercise testing [16].

All individuals gave written informed consent. For participants under 16 years old, written informed consent was obtained from a parent or guardian. This study was approved by the local ethics committee (Ethikkommission der Medizinischen Fakultät der LMU München) and was conducted in accordance to the Declaration of Helsinki.

Exercise testing
All subjects performed a standardized graded cycle exercise test [17] (starting at 90 W workload) until fatigue which we defined as not being capable to maintain pedal cadence above 70 rpm. Increases of 30 W took place every 3 min while pedal cadence was kept constantly at 70–90 rpm.

Assessment of ECG-based cardiac repolarization instability and detection of anaerobic threshold via dT° (AT_{dT°})
Determination of AT via dT° signals was carried out analogously to a previous study by our group [15]. For details, we thus refer to this publication. In brief, we analyzed high-resolution data from Frank's orthogonal lead ECG (1000 Hz, Schiller medilog AR4 plus, Schiller diagnostics, CH) which was recorded throughout the entire exercise test including a 5 min resting phase prior and
after the end of each cycle test with SMARTlab computer and R peak and T wave detection algorithms [17, 18]. In this process, the obtained spatiotemporal properties of each T wave are then used to assess the angle \( \theta \) between two successive repolarization (T wave) vectors [11, 13, 15]. When plotted over time \( \theta \) displays a variability with typical underlying oscillations in the low-frequency range (\( \leq 0.1 \text{ Hz} \)) [11, 13, 15]. During exercise this \( \theta \) signal shows a characteristic three-phasic pattern and \( \text{AT}_{\theta} \) is defined as the point of maximal discordance between \( \theta \) signal and heart rate [15]. Figure 1 illustrates an exemplary \( \theta \) signal and corresponding heart rate signal during graded exercise test and shows the moment of maximal discordance between these signals defined as \( \text{AT}_{\theta} \). This point was converted into power output (W) assuming a linear increase in exercise increments and the corresponding heart rate was extracted from the ECG signal.

Detection of lactate thresholds via methods by Mader and Dickhuth
Capillary blood samples were obtained from earlobes and lactate concentrations (in mmol/l) were measured (lactate Scout+, EKF Diagnostics, Cardiff, GB) before exercise at rest, during exercise at the end of each incremental step and after exercise. The individual lactate threshold was calculated using a standardized computer software (winlactat V 5.2.1.6., Mesics, Münster, Germany). The calculation of LT was determined according to the methods by Mader (fixed threshold at 4 mmol/l) and Dickhuth [1, 19]. Dickhuth defined the LT as the lactate concentration 1.5 mmol/l above the lactate equivalent (i.e. the lowest value of the lactate-performance ratio marking the onset of the lactate increase during exercise) [19].

Statistics
All results are represented as mean ± standard deviation. For statistical analyses and graphical illustration, we used CRAN “R” 3.6.3. Kruskall-Wallis test was performed to detect statistical differences between mean determined thresholds (\( \text{AT}_{\theta}, \text{LT}_{\text{Mader}} \) and \( \text{LT}_{\text{Dickhuth}} \)). The relationship between the three different methods was assessed using the Pearson correlation coefficient test. Intra class correlation (ICC) was tested for these three methods. Bland-Altman plots were performed to visualize the differences between the methods and the respective average.

Results
Baseline characteristics of study participants
Table 1 shows the baseline characteristics of all study participants (A) as well as of professional athletes (B) and amateur athletes (C) as subgroups. Altogether 65 healthy team sport athletes (14 women, 51 men, mean age 22.3 ± 5.2 yrs.) were included in this study. Mean BMI was 22.7 ± 1.8 kg×m\(^{-2}\), mean maximum workload was 262.3 ± 60.8 W (241.8 ± 64.4 W for amateur athletes and 283.4 ± 49.5 W for professional athletes).
participants finished exercise testing until exhaustion. Table 2 shows performance parameters, heart rates and lactate thresholds assessed via methods by Mader and Dickhuth as well as AT\textsuperscript{dT°} for all participants (A) and professional (B) and amateur athletes (C) alone. To exclude that maximal performance was not reached during the test, we calculated theoretical maximal heart rate using the formula 208-(age x 0.7) as described elsewhere [20] and calculated %HR\textsubscript{max} expected for all thresholds.

**dT° signal pattern during exercise and determination of AT\textsuperscript{dT°}**

We were able to identify the typical, previously described three-phasic dT° pattern [15] in all participating athletes during cycle exercise test: We found a low dT° signal at rest (first 15 min) which immediately increased concurrently to the heart rate with the beginning of exercising. At AT\textsuperscript{dT°}, dT° and heart rate showed minimal correlation. Then dT° transiently declines before increasing again until the end of the exercise test (exemplary signal: see Fig. 1). During recovery the dT° signal drops but remains higher than baseline levels. AT\textsuperscript{dT°} was determined as previously described [15].

**Correlation of AT\textsuperscript{dT°} with LTs by Mader and Dickhuth**

AT\textsuperscript{dT°} occurred at 187.6 ± 44.4 W, LT\textsubscript{Dickhuth} at 181.1 ± 45.6 W and LT\textsubscript{Mader} at 184.3 ± 52.4 W. Mean heart rate at AT\textsuperscript{dT°} was 154.0 ± 14.2 bpm, at LT\textsubscript{Dickhuth} 152.8 ± 14.9 bpm and at LT\textsubscript{Mader} 153.1 ± 14.9 bpm (Table 2).

AT\textsuperscript{dT°} highly significantly correlated with LT\textsubscript{Dickhuth} (R = 0.96, R\textsuperscript{2} = 0.92, p < 0.001) and LT\textsubscript{Mader} (R = 0.98, R\textsuperscript{2} = 0.96, p < 0.001) (Fig. 2A and C, respectively) investigating power output. Similar results were detectable correlating heart rates at AT\textsuperscript{dT°} with LT\textsubscript{Dickhuth} (R = 0.97, R\textsuperscript{2} = 0.94, p < 0.001) and with LT\textsubscript{Mader} (R = 0.92, R\textsuperscript{2} = 0.85, p < 0.001). Intra class correlation for these three methods was excellent with intraclass correlation coefficients (ICC) of 0.95 (power output) and 0.93 (heart rate). Bland-Altman plots illustrate close concordance between AT\textsuperscript{dT°} and LT\textsubscript{Dickhuth} (Fig. 2B) as well as between AT\textsuperscript{dT°} and LT\textsubscript{Mader} (Fig. 2D) with power output showing a mean difference of 6.6 W between AT\textsuperscript{dT°} and LT\textsubscript{Dickhuth}.

**Table 1** Shows baseline characteristics for all participants (A) and subgroups of professional athletes and amateur athletes. All data presented as mean ± standard deviation. BMI body mass index

|                      | (A) all participants (n = 65) | (B) professional athletes (n = 32) | (C) amateur athletes (n = 33) |
|----------------------|-------------------------------|-----------------------------------|-----------------------------|
| female (n)           | 14                            | 0                                 | 14                          |
| male (n)             | 51                            | 32                                | 19                          |
| age (yr)             | 223 ± 5.2                     | 23.1 ± 5.5                        | 21.6 ± 4.9                  |
| weight (kg)          | 73.9 ± 10.5                   | 78.5 ± 8.4                        | 69.4 ± 10.5                 |
| height (cm)          | 180.2 ± 9.6                   | 185.4 ± 7.5                       | 175.1 ± 8.6                 |
| BMI (kg x m\textsuperscript{-2}) | 22.7 ± 1.8                  | 22.8 ± 1.3                        | 22.5 ± 2.1                  |

**Table 2** Shows parameters of performance and heart rates in means ± standard deviation. LTs calculated via method by Mader and Dickhuth and AT assessed via dT°. HR heart rate, bpm beats per minute, PO power output, W Watt, LT lactate threshold, AT anaerobic threshold. Maximal expected heart rate calculated using the formula HR\textsubscript{max(expected)}(bpm) = 208-(age x 0.7)

|                      | (A) all participants (n = 65) | (B) professional athletes (n = 32) | (C) amateur athletes (n = 33) |
|----------------------|-------------------------------|-----------------------------------|-----------------------------|
| PO\textsubscript{max} (W) | 262.3 ± 60.8                  | 283.4 ± 49.5                      | 241.8 ± 64.4                |
| LT\textsubscript{Dickhuth}: PO (W) | 181.1 ± 45.6                | 196.2 ± 37.0                      | 166.5 ± 48.8                |
| LT\textsubscript{Dickhuth}: %PO\textsubscript{max} | 69.0 ± 6.4                   | 69.2 ± 5.6                        | 68.9 ± 7.2                  |
| LT\textsubscript{Mader}: PO (W) | 184.3 ± 52.4                  | 198.6 ± 41.5                      | 170.4 ± 58.6                |
| LT\textsubscript{Mader}: %PO\textsubscript{max} | 69.7 ± 8.7                   | 70.0 ± 8.0                        | 69.5 ± 9.5                  |
| AT\textsuperscript{dT°}: PO (W) | 187.6 ± 44.4                  | 204.1 ± 33.9                      | 171.7 ± 47.9                |
| AT\textsuperscript{dT°}: %PO\textsubscript{max} | 71.8 ± 6.8                   | 72.3 ± 5.2                        | 71.3 ± 8.0                  |
| LT\textsubscript{Dickhuth}: HR (bpm) | 152.8 ± 14.9                  | 151.0 ± 15.7                      | 154.3 ± 14.4                |
| LT\textsubscript{Mader}: HR (bpm) | 153.1 ± 14.9                  | 151.3 ± 15.8                      | 154.6 ± 14.1                |
| AT\textsuperscript{dT°}: HR (bpm) | 154.0 ± 14.2                  | 151.7 ± 14.3                      | 156.0 ± 14.1                |
| HR\textsubscript{max(expected)}(bpm) | 192.4 ± 3.7                   | 191.9 ± 3.8                       | 192.9 ± 3.5                 |
| LT\textsubscript{Dickhuth}: % HR\textsubscript{max(expected)} | 794 ± 7.2                    | 787.7 ± 7.3                       | 80.0 ± 7.2                  |
| LT\textsubscript{Mader}: % HR\textsubscript{max(expected)} | 796 ± 7.3                    | 789.7 ± 7.4                       | 80.2 ± 7.3                  |
| AT\textsuperscript{dT°}: % HR\textsubscript{max(expected)} | 800 ± 6.9                    | 791.1 ± 6.6                       | 80.9 ± 7.1                  |
of 3.3 W between AT_{dT} and LT_Mader and of 3.2 W between LT_Dickhuth and LT_Mader.

We further investigated thresholds for subgroups of professional athletes (n = 32) and amateur athletes (n = 33). In professional athletes mean AT_{dT} was at 204.1 ± 33.9 W, LT_Dickhuth at 196.2 ± 37.0 W and LT_Mader at 196.2 ± 37.0 W (ICC = 0.91). Elite athletes had a mean heart rate of 151.7 ± 14.3 bpm at AT_{dT} of 151.0 ± 15.7 bpm at LT_Dickhuth and of 151.3 ± 15.8 bpm at LT_Mader (ICC = 0.94). In amateur athletes we detected AT_{dT} at 171.7 ± 47.9 W, LT_Dickhuth at 166.5 ± 48.8 W and LT_Mader at 170.4 ± 58.6 W (ICC = 0.96). Amateur athletes had a mean heart rate of 156.0 ± 14.1 bpm at AT_{dT} of 154.3 ± 14.4 at LT_Dickhuth and of 154.6 ± 14.1 at LT_Mader (ICC = 0.93). Figure 3 visualizes the strong correlation between AT_{dT} and LT_Dickhuth for both subgroups of amateur (R = 0.97, R^2 = 0.94, p < 0.001, Fig. 3A and B) and professional athletes (R = 0.93, R^2 = 0.86, p < 0.001, Fig. 3C and D) regarding power output. Figure 4A shows box plots for power outputs at AT_{dT}, LT_Mader and LT_Dickhuth with no significant differences between methods of threshold determination. Intergroup comparison revealed no significant differences as checked by Kruskal-Wallis test (p = 0.73). Figure 4B shows box plots for heart rates at AT_{dT}, LT_Mader and LT_Dickhuth with no significant differences between methods of threshold determination. Kruskal-Wallis test again detected no significant differences between assessment methods (p = 0.91).

Discussion

In the present study we were able to validate a non-invasive cardiac repolarization-based method [15] to determine AT in a large cohort of professional and amateur team sport athletes while performing a standardized incremental cycle exercise test. We confirmed the characteristic three-phasic pattern in all study participants who showed a gradual increase concordantly to the heart rate at the start of exercise, a sudden drop in repolarization instability (here at a power output of mean 187.6 ± 44.4 W) discordantly to the heart rate and a new rise of the dT° signal at the end of the workout. The moment of minimal concordance of dT° and heart rate (AT_{dT}) correlated highly significantly with lactate thresholds by Mader and Dickhuth.

Therefore, this pattern has been shown to be characteristic in both young and healthy average-trained athletes as well as professional athletes and very well-trained amateur athletes undergoing standardized incremental cycle exercise tests. It is thus valid and reproducible in both athletes who are only able to perform...
exercise until lower maximal workloads and in athletes capable to continue exercise tests up to very high maximum workloads. This determination of AT is similar to results previously published by Milagro et al. who also found changes in the profile of ventricular repolarization instability as well as of oscillations in the low frequency spectrum once reaching the anaerobic threshold [21]. It is noteworthy that these changes in repolarization instability cannot be provoked by increased heart rate or by fixed atrial pacing [11, 15]. The dT° signal occurs independently of breathing rates and is not associated with heart rate variability [11]. Emerging data validly attributes the dT° signal to efferent sympathetic cardiac nerve activity at the ventricular myocardium [11, 22].

Studies have broadly investigated the alterations within the autonomic nervous system during exercise and the mechanisms found are dynamic, complex and still remain incompletely understood: It is established that the activity of the sympathetic ANS gradually increases with greater workloads shifting from an approximately 4:1 vagal-sympathetic balance to a 4:1 sympa-tho-vagal balance in the course of an exercise [23]. The change in autonomic cardiac modulation during increasing workload subsequently results in altered HRV-derived parameters and this has been used to assess AT in athletes [24, 25]. Our present study, studies assessing AT via HRV-derived parameters [24, 25] and previous studies investigating the connection of changes in repolarization instability and AT [15, 21] thus suggest that the autonomic control of the electrical activity of the myocardium, especially on the level of the ventricular myocardium seems to change once reaching the anaerobic threshold.

While the exact physiological mechanisms underlying AT_{dT°} remain unclear, our study provides several relevant practical implications for future sports research and training physiology: (i) We were able to present a method which is non-invasive and does not require any puncture to gain blood samples. (ii) Furthermore, as it is ECG-based it is less cost-intensive as point-of-care lactate measurements. (iii) A link between HRV indices and blood lactate levels was found during resistance exercise [6–8] in endurance athletes [5] and team sports athletes [24, 25]. Of note, HRV indices are dynamic and rather sensitive to environmental conditions [4]. dT° was proven to be independent of heart rate and breathing rate [11]. Nevertheless, up to date there has no study been conducted comparing the assessment of thresholds based on HRV indices with repolarization patterns. (iv) As threshold-based training has been shown to lead to

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**Fig. 3** Pearson correlation coefficient test for power output between AT_{dT°} and LT_{Dickhuth} for amateur athletes (A) and professional athletes (C). Bland-Altman plots indicate concordance between AT_{dT°} and LT_{Dickhuth} (B + D).
performance benefits in endurance as well as team-sports athletes [1–3] the investigation of AT<sub>dT°</sub>-based training intensities in athletes might be of high interest and should be investigated in future studies.

Our study has some limitations. First, we investigated only team sport athletes and the group of professional athletes contained no female athletes. Whether determination of LTs via AT<sub>dT°</sub> can be validly transferred to endurance athletes has to be elucidated in future studies. Secondly, we only investigated the dT° signal on a cycle ergometer and not with different methods such as treadmill running. Third, AT<sub>dT°</sub> was only correlated to LT and not to ventilatory thresholds as assessed during cardiopulmonary exercise testing. This should be performed in future studies. Additionally, we performed graded exercise tests. However, the gold standard to determine lactate thresholds is the assessment of the maximal steady state during multiple rectangular exercise protocols. Finally, our study included healthy individuals without any known underlying cardiovascular diseases. Whether the assessment of AT<sub>dT°</sub> can be performed in patients with cardiovascular disease or on medication influencing the ANS has to be investigated in future studies.

**Conclusions**

We demonstrated that AT<sub>dT°</sub> is a reliable and non-invasive measurement to assess AT. It correlates with established methods of LT assessment in a large cohort of professional and well-trained amateur athletes. The results of this validation study indicate that AT<sub>dT°</sub> might represent a promising tool for future routine application.

**Abbreviations**

ANS: Autonomic nervous system; AT: Anaerobic threshold; BMI: Body mass index; ECG: Electrocardiogram; HRV: Heart rate variability; ICD: Implantable
cardioverter defibrillator; LT: Lactate threshold; PRD: Periodic repolarization dynamics; SCD: Sudden cardiac death; VT: Ventilatory threshold

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Authors’ contributions
DS prepared the manuscript and analysed data. SK performed the experiments. LVs, LS and AB revised the manuscript. WH performed experiments and analysed data. SB analysed data, revised the manuscript, had the idea for the study and was responsible for conducting the study. All authors gave final approval and agree to be accountable for all aspects of work ensuring integrity and accuracy.

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Availability of data and materials
All data can be obtained by request from the corresponding author.

Declarations

Ethics approval and consent to participate
The study was performed in accordance with the Declaration of Helsinki. The ethics committee of the Ludwig-Maximilians University (LMU) of Munich, Germany approved the study. All participants gave written informed consent prior to our study. For participants under 16 years old written informed consent was obtained from a parent or guardian. The recorded data was irrevocably anonymised.

Consent for publication
Not applicable.

Competing interests
The authors declare no conflict of interest.

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