Reliability of heart rate variability threshold and parasympathetic reactivation after a submaximal exercise test

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Abstract — The objective of this study was to evaluate reproducibility of heart rate variability threshold (HRVT) and parasympathetic reactivation in physically active men (n = 16, 24.3 ± 5.1 years). During the test, HRVT was assessed by SD1 and r-MSSD dynamics. Immediately after exercise, r-MSSD was analyzed in segments of 60 seconds for a period of five minutes. High absolute and relatively reproducible analysis of HRVT were observed, as assessed by SD1 and r-MSSD dynamics (ICC = 0.92, CV = 10.8, SEM = 5.8). During the recovery phase, a moderate to high reproducibility was observed for r-MSSD from the first to the fifth minute (ICC = 0.69–0.95, CV = 7.5–14.2, SEM = 0.07–1.35). We conclude that HRVT and r-MSSD analysis after a submaximal stress test are highly reproducible measures that might be used to assess the acute and chronic effects of exercise training on cardiac autonomic modulation during and/or after a submaximal stress test.

Keywords: reproducibility, anaerobic threshold, heart rate variability, vagal reactivation.

Introduction

Assessment of cardiac autonomic nervous system modulation during an incremental stress test or during the recovery phase has been considered a promising strategy for risk stratification and exercise prescription for different populations. As established, during exercise there is reciprocity in heart rate modulation between sympathetic hyperactivity and parasympathetic inhibition; short-term, post-exercise cardiovascular adaptation is associated with a simultaneous rapid increase in parasympathetic and decrease in sympathetic activity.

Time-domain analysis and Poincaré method plotting of spontaneous heart rate variability (HRV) based on an R-R interval (i-RR) series is a feasible, non-invasive tool for evaluating cardiac autonomic modulation in different functional conditions. These allow an indirect inference about cardiac autonomic nervous system control on the sinus node. During incremental stress tests, an exponential reduction in the root mean square of successive differences between the adjacent normal R-R intervals (r-MSSD, temporal index) and the standard deviation of instantaneous beat-to-beat variability (SD1, Poincaré index) are expected consequences of reduced parasympathetic activity. The point where the values in these two parasympathetic indices are stabilized is called the heart rate variability threshold (HRVT) and is considered an indicator of parasympathetic deactivation during an incremental exercise test.

Recent publications demonstrated that HRVT coincided with the first ventilatory threshold and lactate threshold, suggesting that HRVT might be a practical and attractive alternative to identify adequate workload for exercise prescription. Conversely, analysis of vagal activity immediately post-exercise has been considered a complementary alternative for monitoring training status and control of internal load.

Therefore, considering the clinical and functional implication of vagal assessment during physical stress and its recovery, previous studies have assessed the validity of HRV indexes to represent vagal activity under these conditions. However, the reproducibility of HRV analysis in the exercise test (HRVT) and in the recovery phase are not fully established.

Considering that both HRVT and parasympathetic reactivation are autonomic phenomena assessed in non-stationary conditions, it is essential to evaluate their reproducibility during exercise test and its recovery period, since a poor reproducibility may lead to erroneous inference about the subject’s physiological status. Therefore, we aimed to evaluate the reproducibility of HRVT and parasympathetic reactivation immediately after a submaximal stress test in young, physically active men.
Methods

Participants

We evaluated 16 healthy, physically active, male non-athlete subjects, aged 24.3 ± 5.1 years (range: 20–32 years), with body mass index (BMI) of 25.1 ± 3.7 kg/m² (range: 22.8–29.7 kg/m²). They underwent exercise testing during the afternoon, between 2:00 and 5:00 pm, and had previously been instructed to abstain from stimulants, alcoholic beverages, medicines, and physical activity for at least 48 h prior. The volunteers were oriented to the experimental protocol before the beginning of the tests, and all signed an informed consent to participate, with the approval of the Ethical Committee on Human Research of the UNIEURO University Center in compliance with the Brazilian National Research Ethics System Guidelines and Declarations of Helsinki.

First, we obtained anthropometrical, clinical and basic physiological data, and information on lifestyle, habits, and physical activity level using the International Physical Activity Questionnaire (IPAQ)17.

The basic physiological data, blood pressure and HR (HRbaseline) were measured after 10 minutes in the supine position according to a protocol standardized in our laboratory18. All measures were performed in a quiet, clinical laboratory room at controlled ambient temperature (21–24°C).

The incremental exercise test on the treadmill was applied immediately after the basic physiological data assessment. The experimental design consisted of three visits to our laboratory at intervals between 48 and 72 hours; at each visit the participants were subjected to a standardized exercise test.

Heart rate variability analysis

The R–R interval series was obtained by the model RS800CX Polar cardiac monitor® at a sampling frequency of 1,000 HZ19,20. Then, each series was transferred to a microcomputer for offline data processing and analysis of R–R interval variability, employing a Kubios HRV analyzer (Matlab TM version 2.0 beta, Kuopio, Finland). Before processing the HRV data, all R–R interval series were visually verified on a beat-to-beat basis to validate sinus rhythm and identify ectopic beats, artifacts, and signal reliability. When present, spurious beats were deleted from the series with their previous and following intervals, without adding new intervals.

The time-domain index measured was the root mean square of successive differences between the adjacent normal R–R intervals (r-MSSD). The Poincaré plot was measured as the instantaneous beat-to-beat variability of the date (SD1). Both measures reflect the cardiac parasympathetic modulation.

Exercise test and recovery

A submaximal incremental exercise test on a treadmill (Centurion, Micromed) was conducted. The test protocol included a two-minute preliminary warm-up period at a constant 3.0 km/h and 2.5% slope for adaptation. Immediately after the warm-up, the effective exercise test protocol was started at an initial velocity of 3.0 km/h, which was increased by 1.0 km/h each minute until the volunteer reached 85% of his maximum heart rate (HRpeak), predicted by the formula HRpeak = 208–(0.7 × age)21. The incline of the treadmill was set at 2.5% throughout the test.

After achieving 85% of HRpeak (HRpeak), the exercise protocol was interrupted and an active 5-minute recovery phase was immediately started, with the speed of treadmill reduced to 2.4 km/h but keeping the incline at 2.5%1.

Determination of heart rate variability threshold and parasympathetic reactivation

Before HRVT determination, artifacts were removed by visual inspection when necessary (<3%). We adopted this analysis because <3% is within the standards commonly used in the literature to proceed this analysis22.

The 60 seconds of i-RR analysis in each stage were performed using the Kubios HRV analyzer software previously mentioned. HRVT was considered the load (m/min) corresponding to the point of stabilization at which there was no further significant decline in the values of the r-MSSD (HRVT_r-MSSD) and SD1 (HRVT_SD1) variables during the incremental test by means of visual graphic evaluation1,23. The HRVT analysis was performed by two independent evaluators with considerable experience in this analysis, and in case of disagreement between them, a third evaluator would be consulted to examine the graph. All data were analyzed by consensus between the two evaluators; thus, the third evaluator was not necessary.

Heart rate variability after submaximal exercise test

While a progressive increase in the R–R interval was generally observed during initial recovery, the r-MSSD index was calculated each minute for five minutes after exercise cessation using time-varying analysis8. The r-MSSD is widely used to analyze the parasympathetic reactivation after maximal and submaximal exercise24,25,26,27.

Statistical analysis

Data were analyzed using SPSS v20 (SPSS Inc., USA). Data normality was examined using the Shapiro-Wilk test. Post-exercise r-MSSD was transformed by taking the natural logarithm to allow parametric statistical comparisons that assume a normal distribution. Comparison between the three trials was made using ANOVA for repeated measures, and differences were identified by a Bonferroni post-hoc test. Comparisons were analyzed considering sphericity correction by a Green-House test. Relative reliability was assessed with intra-class correlation coefficient (ICC) (two-way mixed), and absolute reliability was assessed with the standard error of measurement (SEM) and the intra-subject coefficient of variation (CV). CV is a measure of discrepancy and is expressed as a percentage.
Reliability of R-R interval variability

of the mean (CV = standard deviation/mean × 100). SEM quantifies the precision of individual scores on a test and was calculated as follows:

\[
\text{SEM} = \sqrt{\text{MSE}},
\]

where MSE is the mean-squared error.

Results

Considering the hemodynamic variables evaluated at rest condition, no significant differences were observed between the three trials (p = 0.30–0.62, Table 1).

| Variables          | Trial 01       | Trial 02       | Trial 03       | p     |
|--------------------|----------------|----------------|----------------|-------|
| HR_{baseline} (bpm)| 61.1 ± 9.9     | 63.3 ± 12.4    | 63.3 ± 10.7    | 0.34  |
| SBP (mmHg)         | 115.2 ± 7.8    | 116.1 ± 9.3    | 117.3 ± 10.3   | 0.30  |
| DBP (mmHg)         | 74.6 ± 8.8     | 74.7 ± 5.9     | 72.9 ± 10.3    | 0.62  |

ANOVA for repeated measures. HR_{baseline}: resting heart rate; SBP: systolic blood pressure; DBP: Diastolic blood pressure.

Similarly, no differences were observed in HRVT, as evaluated by SD1 (p = 0.86) and r-MSSD (p = 0.86). Additionally, all parasympathetic reactivation measures during recovery (r-MSSD1 to r-MSSD5) were similar between the three tests (p = 0.21–0.94, Table 2).

| Variables          | Trial 01       | Trial 02       | Trial 03       | p     |
|--------------------|----------------|----------------|----------------|-------|
| HRVT_{SD1} (m/sec)| 128.6 ± 23.8   | 129.6 ± 24.4   | 130.7 ± 25.6   | 0.86  |
| HRVT_{r-MSSD} (m/sec)| 128.6 ± 23.8  | 129.6 ± 24.4   | 130.7 ± 25.6   | 0.86  |
| r-MSSD1            | 1.4 ± 0.45     | 1.4 ± 0.56     | 1.4 ± 0.53     | 0.94  |
| r-MSSD2            | 1.9 ± 0.55     | 1.8 ± 0.74     | 1.9 ± 0.60     | 0.75  |
| r-MSSD3            | 1.9 ± 0.61     | 1.9 ± 0.73     | 1.8 ± 0.61     | 0.66  |
| r-MSSD4            | 2.0 ± 0.73     | 1.9 ± 0.65     | 1.9 ± 0.69     | 0.29  |
| r-MSSD5            | 2.2 ± 0.92     | 1.8 ± 0.66     | 1.8 ± 0.65     | 0.21  |

ANOVA for repeated measures. HRVT_{SD1}: heart rate variability threshold evaluated by SD1; HRVT_{r-MSSD}: heart rate variability threshold evaluated by r-MSSD; r-MSSD1 to r-MSSD5: r-MSSD analysis from first to fifth minute of recovery.

Table 1: The mean and standard deviation of resting hemodynamics variables assessed on three occasions.

Table 2: The mean and standard deviation of heart rate variability threshold and post-exercise r-MSSD analysis assessed on three occasions.

As shown in Table 3, higher relative and absolute reproducibility were observed in HRVT_{SD1} (ICC = 0.92, CV = 10.8, SEM = 5.8) and HRVT_{r-MSSD} (ICC = 0.92, CV = 10.8, SEM = 5.8). Similar absolute and relative reproducibility observed in HRVT_{SD} and HRVT_{r-MSSD} can be explained by the physiological nature of the markers, both representing parasympathetic activity, which results in a similar kinetic during an incremental stress test (Figure-1).

Table 3: Absolute and relative reproducibility of heart rate variability threshold and post-exercise r-MSSD analysis.

| Variables          | ICC (95% confidence) | CV% (95% confidence) | SEM |
|--------------------|-----------------------|----------------------|-----|
| HRVT_{SD1} (m/min) | 0.92 (0.82-0.97)      | 10.8 (0.61-21.1)     | 5.8 |
| HRVT_{r-MSSD} (m/min)| 0.92 (0.82-0.97)    | 10.8 (0.61-21.1)     | 5.8 |
| r-MSSD1            | 0.87 (0.72-0.95)      | 14.2 (17.3, 38.2)    | 0.07|
| r-MSSD2            | 0.92 (0.83-0.97)      | 8.9 (-2.5, 18.7)     | 0.20|
| r-MSSD3            | 0.91 (0.80-0.96)      | 9.2 (-6.8, 25.4)     | 0.25|
| r-MSSD4            | 0.95 (0.90-0.98)      | 8.4 (-1.8, 18.8)     | 0.37|
| r-MSSD5            | 0.69 (0.28-0.88)      | 7.5 (-4.5, 19.6)     | 1.05|

ICC: intraclass correlation coefficient; CV: coefficient of variation; SEM: standard error of measurement; HRVT_{SD1}: heart rate variability threshold evaluated by SD1; HRVT_{r-MSSD}: heart rate variability threshold evaluated by r-MSSD; r-MSSD1 to r-MSSD5: r-MSSD analysis from first to fifth minute of recovery.

Figure 1. Heart rate variability dynamics during a submaximal exercise test.

Higher ICC values were also observed in r-MSSD from minutes 1–4 in the recovery phase (ICC > 0.87, CV < 14.7, SEM < 0.37). In the fifth minute of recovery, the ICC values were somewhat lower, whereas SEM values were higher (ICC = 0.69, CV = 7.5, SEM = 1.05).
Discussion

The major finding of this study is the higher relative and absolute reliability observed in HRVT SD1 and HRVT r-MSSD measurements identified by visual inspection. Additionally, the analysis of r-MSSD in the initial phase of recovery (5 minutes) also showed moderate to high reproducibility.

HRVT is a valid indicator of parasympathetic withdrawal during exercise. An increasing amount of evidence shows that the HRVT point coincides with the ventilatory and lactate threshold, which are considered simple and practical measures for estimating the aerobic/anerobic transition. Therefore, our results suggest that, beyond being a valid indicator of parasympathetic withdrawal, HRVT is a reproducible measure in a submaximal exercise test.

Relative reliability is the degree to which individuals maintain their position in a sample with repeated measurements, and it was considered very high when ICC was higher than 0.90, high at 0.70–0.89 and moderate at 0.50–0.69. The results were considered unreliable when ICC was lower than 0.50. Corroborating our findings, Candido et al. found very high relative reproducibility for analysis of HRVT analyzed by visual inspection of SD1 (ICC = 0.81) and r-MSSD (ICC = 0.81) dynamics in six incremental tests on a cycle simulator. Dourado and Guerra also observed excellent relative reliability of HRVT during the incremental shuttle-walk test in healthy subjects (ICC = 0.92).

However, to our knowledge, this is the first study that investigates the reproducibility of HRVT measure in an incremental submaximal exercise test on a treadmill in healthy male subjects. Although the reliability of the measure has been tested in previous studies, it needs to be tested in different protocols and populations for more comprehensive understanding and for practical applications.

Absolute reliability represents the degree to which repeated measurements vary for individuals. Good absolute reliability was also observed in HRVT and in post-exercise r-MSSD analysis. The CV and SEM of HRVT SD1 and HRVT r-MSSD indicate low variation of these markers, occurring with a magnitude inferior to the load increment for each stage (16.6 min). Similar results are observed by Candido et al., who found CV values ranged from 11.4 to 13.1% for HRVT SD1 and HRVT r-MSSD, respectively, values slightly higher than those found in this study (CV = 10.8% for HRVT r-MSSD and HRVT SD1).

In practical terms, when small variations in HRVT are observed in different tests, an interval not greater than 16.6 m/min (1 km/h) should be considered when evaluating the training effects of this variable. For example, an increase of 1 km/h in HRVT occurrence after an exercise training program can be the result of random error and not an adaptation to the training program. Additionally, in a protocol adopted in this study, HRVT measured by SD1 and r-MSSD were observed in similar effort intensities in most cases.

Immediately after a stress test (5 min), good relative and absolute reproducibility were observed in r-MSSD values obtained in 60-second segments during active recovery, especially in the first four minutes of the recovery phase. Dupuy et al. observed a larger range in ICC for HRV analysis after a maximal short analysis of r-MSSD. Good absolute reliability was also observed in HRVT and in post-exercise r-MSSD analysis. The CV and SEM of HRVT SD1 and HRVT r-MSSD indicate low variation of these markers, occurring with a magnitude inferior to the load increment for each stage (16.6 min). Similar results are observed by Candido et al., who found CV values ranged from 11.4 to 13.1% for HRVT SD1 and HRVT r-MSSD, respectively, values slightly higher than those found in this study (CV = 10.8% for HRVT r-MSSD and HRVT SD1).

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Conclusion

We concluded that HRVT, analyzed by visual inspection during an incremental submaximal treadmill stress test, and r-MSSD analysis during recovery are highly reproducible measures. Our results suggest that these indices might be useful tools for assessing the acute and chronic effects of exercise training on cardiac autonomic modulation and practical parameters for exercise prescription.

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