



Caffeine Ingestion Increases Estimated Glycolytic Metabolism during Taekwondo Combat Simulation but Does Not Improve Performance or Parasympathetic Reactivation

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OPEN ACCESS

Citation: Lopes-Silva JP, Silva Santos JFd, Branco BHM, Abad CCC, Oliveira LFd, Loturco I, et al. (2015) Caffeine Ingestion Increases Estimated Glycolytic Metabolism during Taekwondo Combat Simulation but Does Not Improve Performance or Parasympathetic Reactivation. PLoS ONE 10(11): e0142078. doi:10.1371/journal.pone.0142078

Editor: Nir Eynon, Victoria University, AUSTRALIA

Received: August 19, 2015

Accepted: October 16, 2015

Published: November 5, 2015

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Data Availability Statement: All relevant data are within the paper.

Funding: The authors have no support or funding to report.

Competing Interests: The authors have declared that no competing interests exist.

Abstract

Objectives

The aim of this study was to evaluate the effect of caffeine ingestion on performance and estimated energy system contribution during simulated taekwondo combat and on post-exercise parasympathetic reactivation.

Methods

Ten taekwondo athletes completed two experimental sessions separated by at least 48 hours. Athletes consumed a capsule containing either caffeine (5 mg·kg $^{-1}$) or placebo (cellulose) one hour before the combat simulation (3 rounds of 2 min separated by 1 min passive recovery), in a double-blind, randomized, repeated-measures crossover design. All simulated combat was filmed to quantify the time spent fighting in each round. Lactate concentration and rating of perceived exertion were measured before and after each round, while heart rate (HR) and the estimated contribution of the oxidative (W_{AER}), ATP-PCr (W_{PCR}), and glycolytic ($W_{[La-]}$) systems were calculated during the combat simulation. Furthermore, parasympathetic reactivation after the combat simulation was evaluated through 1) taking absolute difference between the final HR observed at the end of third round and the HR recorded 60-s after (HRR $_{60s}$), 2) taking the time constant of HR decay obtained by fitting the 6-min post-exercise HRR into a first-order exponential decay curve (HRR $_{\tau}$), or by 3) analyzing the first 30-s via logarithmic regression analysis (T30).



Results

Caffeine ingestion increased estimated glycolytic energy contribution in relation to placebo (12.5 \pm 1.7 kJ and 8.9 \pm 1.2 kJ, P = 0.04). However, caffeine did not improve performance as measured by attack number (CAF: 26. 7 \pm 1.9; PLA: 27.3 \pm 2.1, P = 0.48) or attack time (CAF: 33.8 \pm 1.9 s; PLA: 36.6 \pm 4.5 s, P = 0.58). Similarly, RPE (CAF: 11.7 \pm 0.4 a.u.; PLA: 11.5 \pm 0.3 a.u., P = 0.62), HR (CAF: 170 \pm 3.5 bpm; PLA: 174.2 bpm, P = 0.12), oxidative (CAF: 109.3 \pm 4.5 kJ; PLA: 107.9 kJ, P = 0.61) and ATP-PCr energy contributions (CAF: 45.3 \pm 3.4 kJ; PLA: 46.8 \pm 3.6 kJ, P = 0.72) during the combat simulation were unaffected. Furthermore, T30 (CAF: 869.1 \pm 323.2 s; PLA: 735.5 \pm 232.2 s, P = 0.58), HRR_{60s} (CAF: 34 \pm 8 bpm; PLA: 38 \pm 9 bpm, P = 0.44), HRR τ (CAF: 182.9 \pm 40.5 s, PLA: 160.3 \pm 62.2 s, P = 0.23) and HRR_{amp} (CAF: 70.2 \pm 17.4 bpm; PLA: 79.2 \pm 17.4 bpm, P = 0.16) were not affected by caffeine ingestion.

Conclusions

Caffeine ingestion increased the estimated glycolytic contribution during taekwondo combat simulation, but this did not result in any changes in performance, perceived exertion or parasympathetic reactivation.

Introduction

Oxidative metabolism is the main metabolic pathway to provide energy during taekwondo combat [1], although the ability to maintain high-intensity actions via the anaerobic system is considered a further determining factor during a taekwondo match [2]. Campos et al. [1] demonstrated that high-intensity actions were maintained by ATP-PCr metabolism during taekwondo combat simulation, in which glycolytic metabolism has been considered important to perform numerous high-intensity actions [3]. Thus, nutritional strategies that could improve glycolytic metabolism, such as caffeine supplementation, could improve performance during simulated taekwondo combat. However, the effect of caffeine ingestion on energy cost during taekwondo combat is currently unexplored in the literature.

Rapid recovery between rounds is crucial to taekwondo success since official matches are composed of 3 rounds of 2 min with 1 min intervals [2]. Higher maximal aerobic power (VO_{2max}) and, consequently, a faster parasympathetic reactivation during the intervals between rounds could help the athletes to improve taekwondo performance during either isolated (training) or consecutive combats (competition). Heart rate variability (HRV) and heart rate recovery (HRR) are non-invasive tools to provide information regarding the cardiac sympathetic and parasympathetic modulations during and after exercise [4]. Goldberger et al. [5] proposed a simple temporal time-varying parasympathetic index to evaluate parasympathetic reactivation after exercise, which is derived from the time-course of the root mean square of successive differences between R-R intervals (RMSSD) measured during successive 30-s segments (RMSSD_{30s}). HRR can be assessed by the number of heart beats recovered within 60s after the cessation of exercise [6], fitting post-exercise HR recovery (HRR) to a first-order exponential decay [7, 8], and analyzing the time constant of 30s short-time recovery. Buchheit et al. [4] showed that all HRR indexes were significantly correlated with glycolytic metabolism during high-intensity exercise. Caffeine ingestion has previously been shown to impair parasympathetic reactivation [9], though it could be hypothesized that caffeine supplementation would



prolong sympathetic activity that occurs during exercise, increasing both adrenergic factors and local metabolites (i.e. epinephrine, lactate) during recovery. Thus, caffeine may actual slow parasympathetic reactivation after combat simulation.

Caffeine (1,3,7 –trimethylxanthine) ingestion (3–9 mg·kg⁻¹) has been shown to significantly improve performance during endurance tasks (activities lasting greater than 30 min) [10], although its effect on short-term (< 30 min) high-intensity exercise is less clear [11]. Caffeine has been shown to increase anaerobic contribution [11] and reduce the rating of perceived exertion [12, 13] during exercise. However, little attention has been given to the effect of caffeine ingestion on combat sports, and only one study investigated its effect on tackwondo combat. Santos et al. [14] showed improved performance with 5 mg·kg⁻¹ caffeine supplementation on performance during simulated tackwondo combat. The results of this study suggested caffeine supplementation allowed individuals to maintain an increased exercise intensity during the latter rounds as indicated by increased blood lactate concentrations following the second and third rounds. However, no study to date has evaluated the effect of caffeine ingestion on the contribution of different energy systems during, or parasympathetic reactivation after, tackwondo combat simulation.

Therefore, the aim subsequent rmance feine on performance? You need to put this in order. My suggestion is below:easure following the second and thof this study was to evaluate the effect of caffeine ingestion on estimated energy systems contribution during simulated taekwondo combat and its subsequent effects on performance. Furthermore, we determined whether caffeine supplementation influenced parasympathetic indexes measured following the combat simulation. We hypothesized that caffeine supplementation would increase glycolytic metabolism, and thus lead to performance improvements, during the combat simulation. Furthermore, it was supposed that the higher glycolytic metabolism contribution with caffeine would delay cardiac autonomic recovery.

Material and Methods

Subjects

Ten male taekwondo athletes (age: 21 ± 4 years, body mass: 71.0 ± 12.9 kg, height: 1.80 ± 0.08 m), who were black belt holders with a minimum of 9 years' experience took part in the present study. All of the subjects were active competitors in national and international championships. The athletes were at the competitive phase of their preparation and engaged in 10 sessions of training per week; approximately 1.5 h was dedicated to technical workouts, 12 h to tactical training and the remaining time to strength and flexibility (2h). The sample was representative of high-level male taekwondo athletes from all Olympic weight divisions: <58 kg (n = 3), 58–68 kg (n = 3), 68–70 kg (n = 2), >80 kg (n = 3). Prior to testing, the athletes were informed of the procedures, including the possible risks involved, and signed an informed consent form. This study was conducted in accordance with the International Ethical Guidelines and Declaration of Helsinki and all procedures were approved by the Ethics and Research Committee of the University of São Paulo.

A priori sample-size calculations, based on previously published data with similar participant characteristics [14], determined that 8 subjects would be necessary to detect a statistical difference given an estimated effect size of 0.5, a 1- β error probability of 0.8 and a P value significance level less than 0.05 for lactate concentration, which was the primary measurement used to estimate energy contribution during exercise. To account for athlete drop out, we aimed to recruit eleven participants. However, one participant was excluded from the study due to technical problems with the gas analyzer during collection and the athlete's lack of time to repeat all procedures. Thus, statistical analysis was performed on ten participants.



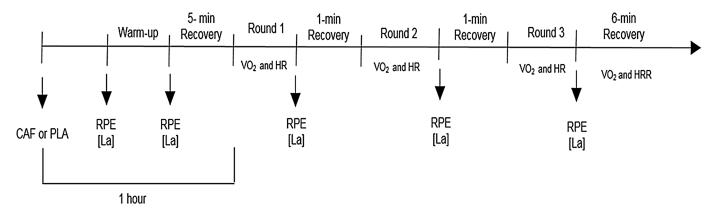


Fig 1. Experimental Design. CAF: Caffeine; PLA: Placebo; RPE: Rating of perceived exertion; [La⁻]: Plasma lactate concentration; HR: Heart rate; HRV: Heart rate variability; VO₂: Oxygen uptake.

Experimental Design

Athletes attended two sessions in which they were required to consume one capsule containing either 5 mg·kg⁻¹ of pure caffeine (CAF; Formula Ativa Pharmacy, São Paulo, São Paulo, Brazil) or cellulose (placebo; PLA) alongside 200 ml water, in a double-blind, randomized, repeated-measures crossover design (Fig 1). Immediately following combat simulation, athletes answered a questionnaire to evaluate any side effects (e.g., increased urine output, gastrointestinal problems, tachycardia, muscle soreness, or headache) experienced by participants during the combat simulation [15]. The two visits were separated by a 7-day washout period [14, 16, 17]. Following supplementation, athletes remained seated for 50-min and then performed a 5-min, self-selected warm-up. The self-selected warm-up was recorded and repeated in the subsequent experimental session. Thereafter, a 5-min rest was allowed and the combat was performed a full 1-hr after supplementation (Fig 1).

All tests were conducted at the same time of day (1 pm– 5 pm) to minimize circadian variance. To avoid the influence of diet on performance, athletes filled-out a 24-h food diary on the day preceding the first trial and were instructed to repeat the same diet on the day preceding the second test. Furthermore, during combat simulation the athletes did not ingest water or any food during the experiments. The athletes were also instructed not to ingest alcohol during the study and to refrain from consuming caffeine-containing substances (i.e., coffee, chocolate, and soft drinks) for 24-h before the experimental tests. Athletes were also required not to be performing any rapid weight loss in order to make weight during the period in which experiments were conducted. Habitual caffeine intake was assessed via a questionnaire with only two participants reporting a high-consumption level for caffeine (corresponding to more than six cups of coffee per day), while the remaining participants all reported a low-consumption level (less than two cups of coffee per day). The athletes performed the tests hydrated and fed, with the last meal ingested two hours before the start of the supplementation protocol. The temperature and relative humidity in CAF and PLA were 27.7°C, 13.4% and 29.5°C, 13.4%, respectively.

Simulated Taekwondo Combat

Simulated combat was performed according to the World Taekwondo Federation's (WTF) official rules and followed the Olympic weight categories. Briefly, each combat consisted of three 2-min rounds with 1-min intervals, and took place within an official area (8 x 8 m). A black belt in taekwondo who was blinded to the capsule content that the athletes had ingested refereed the matches. Although new taekwondo rules establish the use of electronic body protectors to



protect the trunk of the athlete and at the same time allow more reliable and accurate score identification [18], we did not use this system because it would cause interference with the gas analyzer. The gas analyzer was placed on the athlete's back and no attack to this area was allowed. Each athlete competed in a one-on-one match, and only one athlete was evaluated per match (CAF or PLA vs. non-supplemented athlete); the opponent was the same in both conditions. Furthermore, although coaches and others athletes were present during the experiments, no verbal encouragement was provided to the athletes. These procedures were similar to those used in studies analyzing karate [19] and taekwondo [1]. A video camera (Sony® DCR-SX40, China) was used to record each combat for subsequent performance analysis.

Physiological and Psychological Responses

Twenty-five microliters (μL) of blood was collected from the athletes' earlobes to measure lactate concentration, analyzed by the electrochemical method, using a YSI Model 231 Lactate Analyzer (Yellow Sprints, US) which had been calibrated according to manufacturer instructions. The blood sample collections took place immediately before and after the warm-up, immediately after each round, and 1, 3 and 5-min after the last round.

Rating of perceived exertion (RPE) was measured immediately before and after the warm-up, and immediately before and after each round using the 6–20 Borg scale [20]. The subjects were asked to report RPE using cues derived from all sensations experienced during the exercise. Oxygen consumption was measured breath-by-breath during combat and rest using the K4b² (Cosmed, Italy), which has previously been validated [21]. The athlete remained with the equipment during the entire combat and for 6-min after the activity ended. Mean oxygen consumption ($\rm VO_{\rm 2MEAN}$) for each round was calculated as the average $\rm VO_2$ during the entire time of the round. Since the gas analyzer uses telemetry, the start and finish of all combat phases were marked by one researcher who made these markers during all matches procedures. Thus, $\rm VO_2$ between rounds ($\rm VO_{\rm 2INTERVAL}$) also was recorded.

Estimated Energy System Contribution

Estimates of the contributions of the oxidative, glycolytic and ATP-PCr systems during exercise were calculated using the measurements of oxygen consumption during activity, peak blood lactate concentration, and the fast phase of excess oxygen consumption after exercise (EPOC_{FAST}). Oxidative metabolism (W_{AER}) was estimated by subtracting resting VO₂ (VO_{2R}-EST; measured during a 5-min period before the warm-up) from VO_{2MEAN} during the rounds using the trapezoidal method [22]. The ATP-PCr system (W_{PCR}) contribution was estimated using oxygen consumption during the interval between rounds and the EPOC_{FAST} after the third round [19, 22, 23]. In the present study, we fitted the kinetics of post-match oxygen consumption to bi- and mono-exponential models and, since the slow component of the bi-exponential model was negligible, post-match breath-by-breath VO2 data were fitted to a monoexponential model and W_{PCR} was obtained by calculating the integral of the exponential part (Origin 6.0, Microcal, Massachusetts, USA). The contribution of the glycolytic system (W_{[La}-]) was calculated using lactate concentration after combat, assuming that the accumulation of 1 mmol·L⁻¹ [La⁻] is equivalent to 3 ml O₂ kg⁻¹ of body mass [24]. The caloric quotient of 20.92 kJ [25] was used in all three different estimated energy systems. The delta change in lactate $(\Delta_{[La^{-}]})$ was calculated as lactate concentration after each round, minus the lactate concentration at the beginning of the round.

Total metabolic work (W_{TOTAL}) was calculated as the sum of the three estimated energy systems ($W_{AER} + W_{PCR} + W_{[La}]$). Additionally, the contribution of the three estimated energy systems was also expressed as a percentage relative to total energy expenditure.



Heart Rate and post-exercise Heart Rate Recovery (HRR) assessment

Athletes remained in a seated position during the intervals between rounds. HR was measured continuously throughout the match simulation using a HR monitor (Polar S810i, Finland). Mean (HR_{mean}) and peak (HR_{peak}) HR were determined during each round. Parasympathetic function was assessed during the 6-min period after the third round. At the end of third round, all athletes immediately sat passively on a chair. Time between the end of exercise and sitting was less than 5-s. Particular attention to this detail was made because differences in body posture have been shown to result in different absolute HR recovery values. HRR was calculated by 1) taking the absolute difference between the final HR at the end of third round (mean of 5-s) and the HR recorded 60-s later (HRR_{60s}: mean of 5-s), 2) taking the time constant of HR decay obtained by fitting the 6-min post-exercise HRR into a first-order exponential decay curve (HRR $_{\tau}$), or by 3) analyzing the first 30-s (from the 10th to the 40th seconds) via logarithmic regression analysis (T30) [4, 7].

Time-varying vagal-related Heart Rate Variability (HRV) index

A progressive increase in the R-R interval is generally observed during the initial 5-min of HRR, although on shorter scales (i.e., 15-60-s), the curve is piecewise linear with superimposed oscillations. Thus, a time-varying vagal-related index (the root mean square of successive differences in the R-R intervals; RMSSD) was calculated for each of the subsequent 30-s segments of recovery (RMSSD $_{30s}$). To smooth any transient outliers in the HRV plots (HRV vs recovery time), a median filter operation was performed in which each value was replaced with the median of the value as well as the preceding and following values. The first and last values were not median filtered [5].

Video Analyses

The recorded images were analyzed using the Kinovea software (0.8.15), according to the criteria established by Santos et al. [14, 26]. Time-motion analyses were conducted in an attempt to calculate work:rest ratios for each match. The events were measured in tenths of a second, using the marking tool in the analysis software. Attack time (AT), stepping time (ST) and pause time (PT) were recorded separately for each round. The AT was the total time during which the athlete attacked or tried to attack his opponent. Specifically, we recorded the time elapsed between the moment that the athlete began to move his foot or hand in the direction of the opponent and the moment when the athlete finished the attack movement. The ST was defined as the total time in which there was no attempt to attack and the PT was characterized by time-outs determined by the referees.

The following indexes were calculated from these measurements: AT for each round; ST for each round; sum of AT for each round; sum of ST for each round; total attack number (AN) for each round; and average AT:ST ratio for each round [14, 26]. To verify the reliability of analysis, all matches were analyzed twice by the same researcher and a reliability of analyses was performed using an Intraclass Coefficient Correlation (ICC). The ICC revealed that the time-motion analyses conducted in this study was highly reliable for all 5 variables investigated, with values ranging from 0.95 to 0.99 (P < 0.05): attack time (ICC: 0.98; P < 0.01), total attack number for each round (ICC: 0.95; P < 0.01), sum of attack time for each round (ICC: 0.99; P < 0.01), stepping time (ICC: 0.97; P < 0.01), sum of stepping time for each round (ICC: 0.99; P < 0.01).

Statistical Analysis

The Kolmogorov-Smirnov test was applied to determine if the data met parametric assumptions (normality). The data were tested for homogeneity using Levene's test. After confirmation



of normality and homogeneity, the variables were compared using a two-way (condition and time of measurement) analysis of variance (ANOVA) with repeated measurements in the second factor (condition x time). Where necessary, a Bonferroni multiple comparison test was used in order to identify possible differences between conditions and time. When the assumption of sphericity was violated, as indicated by the Mauchley test, the critical value of F was adjusted using the Greenhouse-Geisser epsilon value. All indexes of HRR were compared using a Student's t-test in both CAF and PLA. Effect sizes were calculated using eta squared (η^2). A η^2 of 0.2, 0.5 and 0.8 were considered as small, moderate and large effects [27]. The results of descriptive statistics are reported as the mean \pm SD. Significance was defined as P < 0.05. All analyses were performed using SPSS software (version 17.0; IBM, Chicago, IL, USA).

Results

Success of the blinding and Pre-Test Recommendations

Of the ten athletes investigated, six athletes correctly distinguished which capsule had been ingested based on the effects they felt after taking it. CAF produced an increase in the sensations of tachycardia (20% of athletes), anxiety/nervousness (10% of athletes) and increases of vigor/activeness (60% of athletes). Furthermore, following pre-test instructions, all athletes confirmed that they did not drink alcohol, caffeine, or perform vigorous exercise during the 24 h before the tests.

Physiological Measurements

The HR and oxygen consumption results are presented in Table 1. There was no main effect of condition for HR_{mean} ($F_{1,8} = 3.02$; P = 0.12; $\eta^2 = 0.27$, [small]), but there was a main effect of round ($F_{2,16} = 20.31$, P = 0.01; $\eta^2 = 0.71$, [moderate]), with lower values in the first round compared to the second (P = 0.01) and third (P = 0.01) rounds; similarly, there were lower values in the second round than in the third round (P = 0.04). However, there was no condition by round interaction ($F_{2,16} = 0.46$; P = 0.63; $\eta^2 = 0.05$, [small]). There was no main effect of condition for HR_{peak} ($F_{1,8} = 3.35$; P = 0.10; $\eta^2 = 0.29$, [small]), but there was a main effect of round ($F_{2,16} = 12.15$, P = 0.01; $\eta^2 = 0.60$, [moderate]), with lower values in the first round compared to those in the second (P = 0.02) and third (P = 0.02) rounds, and lower values in the second round than in the third round (P = 0.05). However, there was no condition by round interaction ($F_{2,16} = 2.32$; P = 0.13; $\eta^2 = 0.22$, [small]) (Table 1).

There was no main effect of condition for VO_{2mean} ($F_{1,9} = 4.33$; P = 0.06; $\eta^2 = 0.32$, [moderate]), but there was a main effect of round ($F_{2,18} = 20.74$; P = 0.01 $\eta^2 = 0.69$, [large]), with lower values in the first round than in the second (P = 0.01) and third (P = 0.05) rounds, and lower

Table 1. Physiological responses during a taekwondo combat.

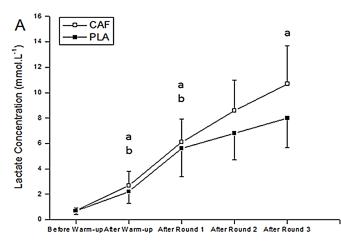
	Round 1		Round 2		Round 3	
	CAF	PLA	CAF	PLA	CAF	PLA
HR _{mean} (bpm)	170 ± 8 ^{a,b}	167 ± 13 ^{a,b}	175 ± 6 ^b	173 ± 10 ^b	178 ± 6	177 ± 10
HR _{peak} (bpm)	181 ± 10 ^{a,b}	177 ± 12 ^{a,b}	185 ± 7 ^b	183 ± 10 ^b	188 ± 8	188 ± 10
VO ₂ (ml.kg ⁻¹ .min ⁻¹)	$37.9 \pm 4.2^{a,b}$	$34.5 \pm 5.5^{a,b}$	43.2 ± 5.9 ^b	41.35 ± 3.6 ^b	41.4 ± 5.1	38.4 ± 6.1

Data are reported as Mean ± SD. CAF: caffeine; PLA: placebo. HR_{peak}: peak heart rate; HR_{mean}: mean heart rate; VO_{2mean} = mean oxygen uptake.

^a different from round 2 (P < 0.05);

doi:10.1371/journal.pone.0142078.t001

^b different from round 3 (P < 0.05).



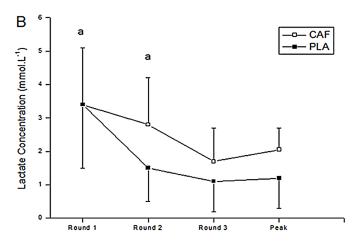


Fig 2. a. Peak blood lactate concentration before and after warm-up, and after rounds 1, 2 and 3; CAF = Caffeine; PLA = Placebo. ^asignificantly higher than the preceding value in PLA; ^bsignificantly higher than the preceding value in CAF **Fig 2b.** Delta blood lactate concentration after the warm up minus blood lactate at the beginning of rounds 1, 2 and 3. ^asignificantly higher than round 3 (n = 10; values are mean ± SD).

values in the second compared to the third round (P = 0.05). However, there was no condition by round interaction ($F_{2.18} = 0.65$; P = 0.55; $\eta^2 = 0.06$, [small]) (Table 1).

There was a main effect of condition for $[La^{-1}]_{peak}$ ($F_{1,9} = 8.90$; P = 0.01; $\eta^2 = 0.5$, [small]) with higher values in CAF compared to PLA (P = 0.03). Furthermore, $[La^{-1}]_{peak}$ increased significantly with time in all conditions ($F_{4,36} = 120.27$; P = 0.01; $\eta^2 = 0.93$, [large]). Additionally, there was a condition by round interaction ($F_{4,36} = 3.41$; P < 0.01; $\eta^2 = 0.16$, [small]), with values after the first round higher than before (P < 0.01) and after (P < 0.01) warm-up in CAF and PLA. Moreover, $[La^{-1}]_{peak}$ after the third round (P < 0.01) was significantly higher than following the second round in CAF (P < 0.01) (Fig 2a).

There was no main effect of condition for Δ [La⁻] (F_{1,9} = 2.64; P = 0.13; η^2 = 0.22, [small]), but there was a main effect of round (F_{2,18} = 13.13; P = 0.01; η_p^2 = 0.6, [small]) with Δ [La⁻] being higher after the first round in relation to the second (P = 0.05) and third (P = 0.01) rounds. However, there was no condition by round interaction (F_{2,18} = 1.33; P = 0.28; η_p^2 = 0.12, [small]) (Fig 2b).

There was no main effect of condition on absolute estimated aerobic energy cost $(F_{1,9}=0.27, P=0.61, \eta^2=0.02, [small])$, but there was a main effect of round $(F_{2,18}=13.81; P=0.01; \eta^2=0.60, [moderate])$, with higher estimated absolute aerobic cost during the second round than the first (P=0.01) and third rounds (P=0.02). No condition by round interaction was found $(F_{2,18}=1.87; P=0.18; \eta^2=0.17, [small])$. Similarly, there was no main effect of condition on the estimated relative aerobic cost $(F_{1,9}=0.12; P=0.91; \eta^2=0.01, [small])$, but there was a main effect of round $(F_{2,18}=15.72; P=0.01; \eta^2=0.63, [moderate])$, with higher estimated relative aerobic cost in the second round than in the first (P=0.01) and third (P=0.01) rounds. However, there was no condition by round interaction $(F_{2,18}=0.97; P=0.39; \eta^2=0.1, [small])$ (Table 2).

There was a main effect of condition on estimated absolute glycolytic contribution ($F_{1,9} = 5.60$; P = 0.04; $\eta^2 = 0.38$, [moderate]), with higher values in CAF compared to PLA (P = 0.04). Furthermore, there was a main effect of round ($F_{2,18} = 10.47$; P = 0.01; $\eta^2 = 0.53$, [moderate]), with higher estimated absolute glycolytic contribution in the first round compared to the second (P = 0.04) and third (P = 0.01) rounds, but there was no condition by round interaction ($F_{2,18} = 0.24$; P = 0.78; $\eta^2 = 0.02$, [small]). Similarly, for estimated relative glycolytic



Table 2. Estimated metabolic responses on the three rounds of taekwondo combat simulation in placebo and caffeine conditions.

	Rou	Round 1		Round 2		Round 3	
	CAF	PLA	CAF	PLA	CAF	PLA	
Oxidative							
W _{AER} (kJ)	103.2 ± 11.8 ^b	97.1 ± 7.8 ^b	117.3 ± 15.2	116.8 ± 15.2	107.5 ± 20.8 ^b	109.8 ± 16.4 ^b	
Relative (%)	66 ± 6 ^b	63 ± 5 ^b	66 ± 6	70 ± 3	65 ± 6 ^b	64 ± 9 ^b	
ATP-PCr							
W _[PCR] (kJ)	37.8 ± 10.0	43.4 ± 7.4	43.1 ± 10.2	42.6 ± 8.7	55.3 ± 13.7 ^{ab}	54.7 ± 24.2 ^{ab}	
Relative (%)	24 ± 4	27 ± 3	24 ± 3	26 ± 2	31 ± 4 ^{ab}	33± 9 ^{ab}	
Glycolytic							
W _[La] (kJ)	17.7 ± 9.94 ^c	13.3 ± 6.9	11.8 ± 7.9 ^{ac}	7.5 ± 5.5 ^a	8.1 ± 4.3 ^{ac}	5.8 ± 3.3 ^a	
Relative (%)	10 ± 5	9 ± 5	7 ± 4 ^a	4 ± 3 ^a	4 ± 2 ^a	3 ± 2 ^a	
W _{Total} (kJ)	158.4 ± 22.7 ^b	154.2 ± 17.0 ^b	175.3 ± 26.3	163.9 ± 27.1	173.6 ± 28.0	167.6 ± 38.3	

Data are reported as mean and standard deviation. CAF: Caffeine; PLA: Placebo. Waer = estimated oxidative energy, $W_{[La-]}$ = estimated glycolytic energy, W_{PCR} = estimated ATP-PCr energy, W_{Total} = estimated total metabolic work ($W_{AER} + W_{[La-]} + W_{PCR}$).

doi:10.1371/journal.pone.0142078.t002

contribution there was a main effect of condition ($F_{1,9} = 5.70$; P = 0.04; $\eta^2 = 0.38$, [moderate]), with higher values in CAF compared to PLA. Furthermore, there was a main effect of round ($F_{2,18} = 13.10$; P = 0.01; $\eta^2 = 0.60$, [moderate]) with higher estimated relative glycolytic contribution in the first round compared to second (P = 0.02) and third (P = 0.01) rounds. However, there was no condition by round interaction ($F_{2,18} = 1.32$; P = 0.29; $\eta^2 = 0.12$, [small]) (Table 2).

There was no main effect of condition on estimated absolute ATP-PCr contribution ($F_{1,9} = 0.13$; P = 0.72; $\eta^2 = 0.15$, [small]), but there was a main effect of round on estimated absolute ATP-PCr contribution ($F_{2,18} = 16.34$; P = 0.01; $\eta^2 = 0.64$, [moderate]), with higher values in the third round compared to the first (P = 0.01) and second (P = 0.01) rounds. However, there was no condition by round interaction ($F_{2,18} = 0.62$; P = 0.54; $\eta^2 = 0.06$, [small]). Similarly, for estimated relative ATP-PCr contribution there was no main effect of condition ($F_{1,9} = 1.83$; P = 0.20; $\eta^2 = 0.16$, [small]). However, there was a main effect of round for estimated relative ATP-PCr contribution ($F_{2,18} = 19.41$; P = 0.01; $\eta^2 = 0.68$, [moderate]), with higher estimated relative ATP-PCr contribution in the third round when compared to the first (P = 0.01) and second (P = 0.02) rounds. In addition, no condition by round interaction was shown ($F_{2,18} = 0.55$; P = 0.94; $\eta^2 = 0.01$, [small]) (Table 2).

There was no main effect of condition on estimated absolute energy expenditure ($F_{1,9}$ = 2,24; P = 0.16; η^2 = 0.20, [small]), but there was a main effect of round ($F_{2,18}$ = 5.04; P = 0.01; η^2 = 0.35, [moderate]), with higher estimated absolute energy expenditure in the second round compared to the first round (P = 0.01). Furthermore, there was no condition by round interaction ($F_{2,18}$ = 0.46; P = 0.63; η^2 = 0.04, [small]) (<u>Table 2</u>).

There was no main effect of condition for RPE ($F_{1,9} = 0.27$; P = 0.62; $\eta^2 = 0.03$, [small]), but there was a main effect of round ($F_{4,36} = 105.82$; P = 0.01; $\eta^2 = 0.92$, [large]), with values increasing significantly throughout the combat in both CAF and PLA conditions (P = 0.01). However, there was no condition by round interaction ($F_{3,27} = 1.24$; P = 0.31; $\eta^2 = 0.12$, [small]) (Fig 3).

^aDifferent than round 1;

^bDifferent than round 2;

^cDiferent than PLA condition.

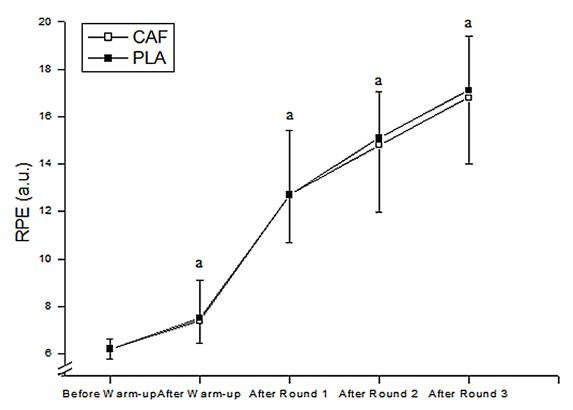


Fig 3. Rating of perceived exertion (RPE) before and after the warm-up and after each round in CAF and PLA. ^asignificantly higher than preceding values in CAF and PLA. CAF = Caffeine; PLA = Placebo.

Time-Motion Analyses

Time-motion analyses are presented in Table 3. There was no main effect of condition ($F_{1,9} = 0.31$; P = 0.58; $\eta^2 = 0.03$, [small]), round ($F_{2,18} = 0.40$; P = 0.67; $\eta^2 = 0.04$, [small]) or condition by round interaction ($F_{2,18} = 0.70$; P = 0.50; P = 0.07, [small]) on AT. Similarly, there was no main effect of condition ($F_{1,9} = 0.25$; P = 0.62; P = 0.07, [small]), round ($P_{2,18} = 0.17$; P = 0.08; P = 0.17, [small]) or condition by round interaction ($P_{2,18} = 0.49$; P = 0.61; P = 0.05, [small]) for ST (Table 3).

Table 3. Attack Time, Stepping Time, total time of Attack Time, Stepping Time, ratio between Attack Time and Stepping Time, and ratio between summed time of the Attack Time + Stepping Time and Pause Time during a taekwondo combat simulation in both CAF and PLA conditions.

	Round 1		Round 2		Round 3	
	CAF	PLA	CAF	PLA	CAF	PLA
Attack Time (s)	1.3 ± 0.4	1.4 ± 0.8	1.3 ± 0.2	1.2 ± 0.3	1.2 ± 0.1	1.3 ± 0.2
Stepping Time (s)	3.0 ± 1.0	3.0 ± 0.8	3.3 ± 1.0	3.7 ± 1.1	3.8 ± 2.6	4.0 ± 1.6
Attack Time sum (s)	35.2 ± 8.8	45 ± 2,9	33.6 ± 6.5	32.0 ± 7.3	32.8 ± 8.1	33.0 ± 10.0
Stepping Time sum (s)	73.5 ± 18.1	76.0 ± 14.0	79.2 ± 11	79.7 ± 14.5	79.3 ± 9.7	78.5 ± 14.1
Attack Number (times)	28.0 ± 7.0	30.6 ± 7.0	25.1 ± 4.6	25.8 ± 7	27.2 ± 83	25.6 ± 8.5
Attack Time/Stepping Time ratio	0.5 ± 0.2	0.6 ± 0.5	0.4 ± 0.1	0.4 ± 0.2	0.4 ± 0.1	0.4 ± 0.18

CAF: Caffeine; PLA: Placebo.

doi:10.1371/journal.pone.0142078.t003



Table 4. Heart rate variability indexes of parasympathetic reactivation in both CAF and PLA
conditions.

	CAF	PLA
HRR _{60s} , bpm	34 ± 8	38 ± 9
T ₃₀ , s	869.1 ± 323.2	735.5 ± 232.2
HRR_{τ} , s	182.9 ± 40.5	160.3 ± 62.2
HRR _{amp} , bpm	70.2 ± 17.4	79.2 ± 17.4

 HRR_{60s} , number of heart rate beat recovered in 60-s after exercise cessation; T_{30} , time constant of short-time heart rate recovery; HRR_{τ} , time constant of HR during the 6-min recovery. CAF = Caffeine; PLA = Placebo.

doi:10.1371/journal.pone.0142078.t004

There was no main effect of condition ($F_{1,9}=0.66$; P=0.43; $\eta^2=0.06$, [small]), round ($F_{2,18}=2.45$; P=0.11; $\eta^2=0.21$, [small]) or condition by time interaction ($F_{2,18}=2.20$; P=0.13; $\eta^2=0.19$, [small] on the sum of attack time. Furthermore, there was no effect of condition ($F_{1,9}=0.10$; P=0.75; $\eta^2=0.01$, [small]), round ($F_{2,18}=2.82$; P=0.08; $\eta^2=0.23$, [small]) or condition by round interaction for the sum of stepping time ($F_{2,18}=0.27$; P=0.76; $\eta^2=0.02$, [small]) (Table 3).

There was no effect of condition ($F_{1,9}=0.04$; P=0.83; $\eta^2=0.01$, [small]), round ($F_{2,18}=3.05$; P=0.07; $\eta^2=0.25$, [small]) or condition by time interaction on number of attacks ($F_{2,18}=1,27$; P=0.30; $\eta^2=0.12$, [small]). Additionally, there was no effect of condition ($F_{1,9}=1.02$; P=0.33; $\eta^2=0.10$, [small]), round ($F_{2,18}=0.38$; P=0.68; $\eta^2=0.04$, [small]) or condition by time interaction on attack time/stepping time ratio ($F_{2,18}=1.18$; P=0.33; $\eta^2=0.11$, [small]) (Table 3).

HRV indexes are presented in Table 4. There were no significant differences in HRR $_{60}$ (t(8) = 1.17, P > 0.05, effect size = 0.38, [moderate], 95% CI = -3.96–12.1), HRR $_{\tau}$ (t(8) = -1.18, P > 0.05, effect size = 0.38, [moderate], 95% CI = -66.7–21.4), T $_{30}$ (t(8) = -1.12, P > 0.05, effect size = 0.36, [moderate], 95% CI = -407.3–140.1) or HR $_{amp}$ (t(8) = 1.57, P > 0.05, effect size = 0.48, [moderate], 95% CI = -4.2–22.3) between CAF and PLA.

The time course of the R-R interval after the third round is shown in Fig 4. There was no main effect of condition on RMSSD_{30s} ($F_{1,8} = 0.67$; P = 0.67; P = 0.07, [small]), time ($F_{11,88} = 1.92$; P = 0.16; $P_{11,88} = 1.96$; P = 0.18; $P_{11,88} = 1.96$; P = 0.18; $P_{11,88} = 1.96$; $P_{11,88} = 1.96$;

The time course of HRR after the third round is illustrated in Fig 5. There was no main effect of condition for HRR ($F_{1,8} = 2.04$; P > 0.05; $\eta^2 = 0.18$, [small]). However, there was a main effect of time for HRR ($F_{11,99} = 200.11$; P < 0.05; $\eta^2 = 0.01$, [small]), with values systematically higher until the 150s after the third round in relation to values after 180s of the recovery. Furthermore, there was no condition by time interaction ($F_{11,99} = 1.63$; P > 0.05; $\eta^2 = 0.15$, [small]).

Discussion

The main findings of the present study were that caffeine ingestion increases absolute and relative estimated glycolytic energy contribution during simulated taekwondo combat, although this did not translate into any improvements in performance. Furthermore, caffeine ingestion did not affect autonomic control after the combat.

In this study, caffeine supplementation did not change any time-motion performance variable during simulated taekwondo combat. Previous studies have reported benefits after caffeine

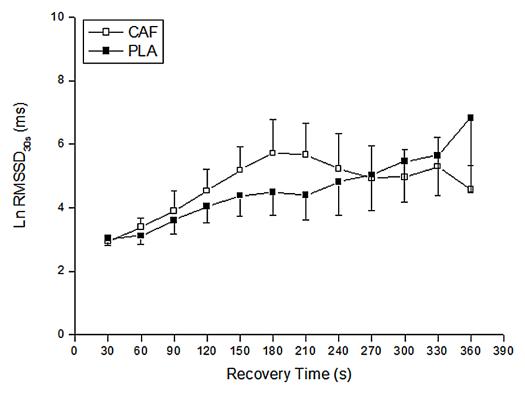


Fig 4. Root mean square of successive differences in the R-R intervals measured on successive 30-s segments (RMSSD_{30s}) during the 6-min recovery period after the third round in both CAF and PLA.

ingestion on high-intensity exercise [10, 11, 16, 28], as well as during taekwondo combat simulation [14]. However, not all studies have shown a positive effect of caffeine with no performance changes shown during successive maximal cycling bouts [29], wrestling [30], and judo [17]. Similarly to our results, Lopes-Silva et al. [17] showed that caffeine ingestion did not improve performance (number of throws) during the Special Judo Fitness Test after a 5-day weight loss period. More recently, Felippe et al. [31] also reported that caffeine ingestion did not improve performance (number of throws) during the Special Judo Fitness Test. Conversely, Santos et al. [14] showed that caffeine supplementation reduced the sum of time-outs enforced by the referee in the first and second rounds of a taekwondo combat simulation. The reasons for these differences are not clear, but may be related to different performance protocols that used different durations, intensities and type of exercise. In the present study, we evaluated simulated taekwondo combat and during this activity performance is determined by technical, tactical, physiological and physical characteristics of the athletes [2]. Despite an increased estimated glycolytic contribution with caffeine this did not alter the pattern of technical actions during the simulation. However, in the study conducted by Santos et al. [14] the combats were performed between supplemented athletes (i.e. caffeine versus caffeine), which suggests that improvement in performance may have been caused by the fact that both athletes were supplemented. In contrast to the study of Santos et al. [14], athletes in the present study were evaluated individually (supplemented versus non-supplemented athletes). Thus, when only one athlete is assessed (i.e., the supplemented athlete) there was no improvement in performance variables following caffeine supplementation during simulated taekwondo combat.

Our results show that there was no significant difference in HR_{mean} and HR_{peak} during simulated taekwondo combat after caffeine ingestion. Our data corroborate with Santos et al. [14], who

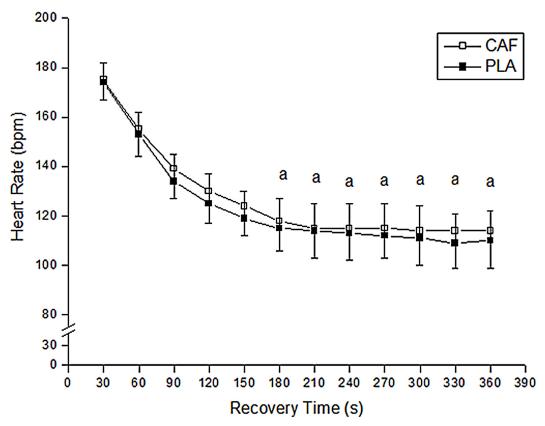


Fig 5. Heart Rate Recovery during the 6-min recovery period after the third round in both CAF and PLA. CAF = Caffeine; PLA = Placebo. ^asignificantly lower in relation to times 30, 60, 90, 120 and 150-s after the third round.

did not report significant differences in HR responses during taekwondo combat simulation after caffeine supplementation in relation to placebo. However, there was an increase in HR values over the course of the combat. This corroborates with other studies, which have shown an increase in cardiovascular demand as the number of rounds continues in taekwondo athletes [1, 14, 32].

The results of our study showed a predominant estimated energy contribution of oxidative metabolism (65%) during simulated taekwondo combat, followed by ATP-PCr (27%) and glycolytic (8%) contribution. These results are in agreement with Campos et al. [1], who reported that energy contribution during a taekwondo combat simulation was predominantly supplied from aerobic sources, with lesser contribution from both the ATP-PCr and glycolytic energy systems. Furthermore, although caffeine supplementation had no effect on the ATP-PCr contribution, in the present study, there was an increase in ATP-PCr contribution in the third round in relation to the first and second rounds. This difference may be due to the method used to calculate ATP-PCr metabolism; a 1-min break between rounds is insufficient to determine VO_2 offset, while the fast component of EPOC is considered for determination of energy contribution during the third round.

Ratings of perceived exertion increased linearly across rounds, although no differences were shown between caffeine and placebo. These results are in accordance with Santos et al. [14], who showed that caffeine supplementation did not reduce RPE during simulated taekwondo combat. However, Doherty and Smith [13] did show that caffeine ingestion reduced perceived exertion; 29% of the variance explaining the ergogenic effect of caffeine on performance was due to decreased RPE during the exercise. As opposed to endurance exercise, taekwondo



combat is acyclic and athletes distribute the technical and tactical actions through the combat. Thus, our results showed that caffeine supplementation does not affect RPE during simulated taekwondo combat.

Caffeine ingestion resulted in increased peak lactate concentration following simulated taekwondo combat. This result is supported by other studies, which showed an increase in blood lactate concentration after caffeine ingestion [14, 17, 33]. For example, Santos et al. [14] showed that caffeine ingestion resulted in increased lactate concentrations during simulated taekwondo combat when compared to placebo. Furthermore, Lopes-Silva et al. [16] showed that caffeine ingestion following a 5-day weight loss period resulted in increased lactate concentrations after successive bouts of the Special Judo Fitness Test. Thus, the increase in the lactate concentration after caffeine ingestion corroborates with the results reported in the literature.

Our results showed that caffeine ingestion increased estimated glycolytic contribution during simulated taekwondo combat when compared to placebo. Although no study in the literature has reported that CAF supplementation increases glycolytic contribution during a combat taekwondo simulation, it is well established that caffeine increases anaerobic contribution during cycling exercise [16, 28]. Santos et al. [28] showed that caffeine ingestion increased anaerobic contribution during a 4-km cycling time-trial and two mechanisms have been proposed to explain the increase in blood lactate concentration. Firstly, CAF may increase anaerobic activity through its antagonistic action on peripheral adenosine receptors, which could prevent the inhibitory effects of adenosine on phosphofructokinase activity in skeletal muscle [34]. Secondly, caffeine may promote catecholamine release to facilitate the conversion of phosphorylase *b* to its more active *a* form, accelerating muscle glycogenolysis [33]. Although previous studies showed a decrease in glycolytic contribution over subsequent rounds during simulated taekwondo combat [1], our data show that caffeine supplementation actually increased estimated glycolytic contribution during simulated taekwondo combat simulation in relation to placebo. However, this did not translate across into any changes in technical actions performed by the athletes.

Caffeine supplementation did not affect post-exercise parasympathetic reactivation indexes after the taekwondo combat simulation. In contrast to our results, Bunsawat et al. [9] demonstrated that HR was increased 2-min after incremental treadmill exercise with caffeine in relation to placebo. However, this delay in HR was accompanied by an increase in time to exhaustion after caffeine supplementation. Thus, the delay in autonomic recovery after incremental exercise could have been caused by a combination of caffeine-induced sympathetic activation along with increased performance. Furthermore, although parasympathetic indexes after repeated sprint exercise have been correlated with anaerobic contribution, possibly through a high elevation of adrenergic factors and local metabolites during recovery [4], the increase in glycolytic metabolism after caffeine ingestion did not influence the parasympathetic indexes after simulated taekwondo combat.

Practical Applications

Although time-motion performance variables during simulated taekwondo combat were not changed following supplementation, the results of the present study suggest that caffeine supplementation might be an effective strategy to increase glycolytic energy metabolism in taekwondo athletes, and could be an effective pharmacological strategy during training sessions to improve anaerobic capacity.

Conclusion

In summary, caffeine supplementation increased estimated glycolytic metabolism contribution during simulated taekwondo combat, although this did not translate into improvements in



performance. Furthermore, caffeine supplementation did not influence post-exercise parasympathetic reactivation following a taekwondo simulation.

Acknowledgments

The authors are grateful to Dr. Bryan Saunders (FAPESP grant number: 2011/19513-2) for the proofreading of the manuscript.

Author Contributions

Conceived and designed the experiments: JPLS JFSS BHMB CCCA LFO IL EF. Performed the experiments: JPLS JFSS BHMB CCCA LFO IL EF. Analyzed the data: JPLS JFSS BHMB CCCA LFO IL EF. Contributed reagents/materials/analysis tools: JPLS JFSS BHMB CCCA LFO IL EF. Wrote the paper: JPLS JFSS BHMB CCCA LFO IL EF.

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