

Research article

Open Access

## The effects of four weeks of creatine supplementation and high-intensity interval training on cardiorespiratory fitness: a randomized controlled trial

Jennifer L Graef<sup>†</sup>, Abbie E Smith<sup>†</sup>, Kristina L Kendall<sup>†</sup>, David H Fukuda<sup>†</sup>, Jordan R Moon<sup>†</sup>, Travis W Beck<sup>†</sup>, Joel T Cramer<sup>†</sup> and Jeffrey R Stout<sup>\*†</sup>

Address: Department of Health and Exercise Science, University of Oklahoma, Huston Huffman Center, 1401 Asp Avenue, Norman, OK 73019, USA

Email: Jennifer L Graef - Jennifer.L.Graef-1@ou.edu; Abbie E Smith - abbiesmith@ou.edu; Kristina L Kendall - krissykendall@ou.edu; David H Fukuda - david.fukuda@ou.edu; Jordan R Moon - jordanmoon@ou.edu; Travis W Beck - tbeck@ou.edu; Joel T Cramer - jcramer@ou.edu; Jeffrey R Stout\* - jrout@ou.edu

\* Corresponding author †Equal contributors

Published: 12 November 2009

Received: 11 June 2009

Accepted: 12 November 2009

*Journal of the International Society of Sports Nutrition* 2009, **6**:18 doi:10.1186/1550-2783-6-18

This article is available from: <http://www.jissn.com/content/6/1/18>

© 2009 Graef et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

### Abstract

**Background:** High-intensity interval training has been shown to be a time-efficient way to induce physiological adaptations similar to those of traditional endurance training. Creatine supplementation may enhance high-intensity interval training, leading to even greater physiological adaptations. The purpose of this study was to determine the effects of high-intensity interval training (HIIT) and creatine supplementation on cardiorespiratory fitness and endurance performance (maximal oxygen consumption ( $VO_{2PEAK}$ ), time-to-exhaustion ( $VO_{2PEAK}$ TTE), ventilatory threshold (VT), and total work done (TWD)) in college-aged men.

**Methods:** Forty-three recreationally active men completed a graded exercise test to determine  $VO_{2PEAK}$ ,  $VO_{2PEAK}$ TTE, and VT. In addition, participants completed a time to exhaustion (TTE) ride at 110% of the maximum workload reached during the graded exercise test to determine TWD (TTE (sec)  $\times$  W = J). Following testing, participants were randomly assigned to one of three groups: creatine (creatine citrate) (Cr;  $n = 16$ ), placebo (PL;  $n = 17$ ), or control ( $n = 10$ ) groups. The Cr and PL groups completed four weeks of HIIT prior to post-testing.

**Results:** Significant improvements in  $VO_{2PEAK}$  and  $VO_{2PEAK}$ TTE occurred in both training groups. Only the Cr group significantly improved VT (16% vs. 10% improvement in PL). No changes occurred in TWD in any group.

**Conclusion:** In conclusion, HIIT is an effective and time-efficient way to improve maximal endurance performance. The addition of Cr improved VT, but did not increase TWD. Therefore, 10 g of Cr per day for five days per week for four weeks does not seem to further augment maximal oxygen consumption, greater than HIIT alone; however, Cr supplementation may improve submaximal exercise performance.

## Background

Traditional endurance training has been shown to improve aerobic capacity, such as the ability to sustain a given submaximal workload for an extended period of time, or to produce a higher average power output over a fixed distance or time [1,2]. Physiological adaptations from training, resulting from an increase in mitochondrial density, include changes in skeletal muscle substrate utilization and improved respiratory control sensitivity [3]. High-intensity interval training (HIIT) is a time-efficient way to induce similar adaptations, such as increased maximal mitochondrial enzyme activity [4] and a reduction in glycogen utilization and lactate accumulation [5,6]. In addition, HIIT may be more effective than conventional endurance training at improving muscle buffering capacity [7,8]. HIIT consists of repeated bouts of short to moderate duration exercise completed at intensities greater than the anaerobic threshold, interspersed with brief periods of low-intensity or passive rest. HIIT is designed to repeatedly stress the body, physiologically, resulting in chronic adaptations and improving metabolic and energy efficiency [9,10]. Helgerud et al. [11] found that HIIT significantly augmented maximal oxygen consumption ( $VO_{2PEAK}$ ) and time to exhaustion (TTE) greater than a traditional training program with moderately-trained males. The velocity at which ventilatory threshold (VT) occurred increased as well, which may signify a higher training capacity and, therefore, should also represent an improvement in endurance performance. It was determined during this study that different protocols of HIIT, matched for frequency and total work done, provided similar results [11]. In support, Burke et al. [12] examined the effects of two different interval training protocols on  $VO_{2PEAK}$ , VT, and lactate threshold in a group of untrained women, demonstrating that both interval-training protocols significantly improved all performance variables. Similarly, an increase in  $VO_{2PEAK}$  and VT was found in three groups of well-trained cyclists following three different HIIT protocols of varying intensities and work-to-rest ratios [9].

Phosphocreatine (PCr), a high-energy storage molecule within skeletal muscle, provides immediate replenishment of ATP during intense exercise [13]. Multiple HIIT bouts are designed to deplete PCr stores in the working skeletal muscle, reducing power output. It has been reported that it takes more than six minutes to fully recover depleted PCr stores after exercise-induced PCr depletion [14]. Therefore, if recovery intervals during HIIT bouts are less than six minutes, PCr may not be fully replenished, resulting in a reduced ability to meet the demands of cellular ATP resynthesis and a reduced performance [13]. Supplementing with creatine (Cr) has been demonstrated to effectively augment muscle phosphocreatine (PCr) stores [15]. Specifically, one study

showed a 20% increase in muscle creatine with ingestion of 20 g of Cr per day for just 5 days [16].

It has been suggested that increases in skeletal muscle PCr concentration may improve muscle buffering capacity and moderate glycolysis [17,18]. In addition, Cr supplementation may increase the rate of PCr resynthesis between HIIT exercise bouts and enhance mitochondrial shuttling of ATP into the cytoplasm, providing significant physiological adaptations [15,16].

Current research suggests that Cr supplementation, when combined with training, has been shown to significantly augment performance [19]. Moreover, the combination of Cr supplementation and HIIT may lead to greater improvements in  $VO_{2PEAK}$ , VT, and TTE than previously reported with HIIT or Cr supplementation alone. While Cr is known to improve anaerobic performance, its use in aerobic performance has been under-researched. To date, no one has examined the concurrent effects of Cr supplementation and HIIT. Therefore, we propose that Cr supplementation may increase training capacity during HIIT, resulting in improved endurance performance as measured by  $VO_{2PEAK}$ , VT, and TTE, beyond what has been demonstrated for HIIT alone. The purpose of this study was to determine the combined effects of four weeks of HIIT and Cr supplementation on  $VO_{2PEAK}$ , VT, and TTE in recreationally active college-aged men.

## Methods

Forty-three recreationally active (1-5 hours of regimented exercise per week) college-aged men (mean  $\pm$  SD, Age:  $22.6 \pm 4.9$  years; Ht:  $178.1 \pm 7.1$  cm; Wt:  $83.0 \pm 13.8$  kg) volunteered to participate in this study. Participants were screened for health conditions that would have prevented them from participating in the study, including heart and joint conditions. Any participants who had taken sports supplements, including any form of Creatine, in the three months prior to the beginning of the study were excluded. Participants kept a food diary, and none of the participants consumed a vegetarian diet. Participants were asked not to change training or dietary habits for the duration of the study. This study was approved by the University's Institutional Review Board for Human Subjects and informed consent was obtained from each participant prior to enrollment.

### **Determination of $VO_{2PEAK}$ , ventilatory threshold, and total work done**

A maximal graded exercise test (GXT) on a cycle ergometer (Corival 400, Groningen, The Netherlands) was completed by all participants to determine maximal oxygen consumption ( $VO_{2PEAK}$ ). Participants began pedaling at a cadence of 60-80 revolutions per minute (RPM) at a work-

load of 20 watts (W). The workload increased 1 W every 3 seconds (a total of 20 W every minute). This continued until the subject could no longer maintain 60-80 RPM or until volitional exhaustion, despite verbal encouragement. Respiratory gases were monitored and continuously analyzed with open-circuit spirometry using a calibrated metabolic cart (True One 2400®, Parvo-Medics, Inc., Provo, UT). Data were averaged over 15-second intervals, with the highest 15-second oxygen consumption and heart rate recorded as the peak oxygen consumption ( $VO_{2PEAK}$ ) and maximum heart rate (HR<sub>max</sub>), respectively. Time to exhaustion for the GXT ( $VO_{2PEAK}TTE$ ) was recorded. In addition, ventilatory threshold (VT) was measured during this test. VT was determined from a plot of ventilation ( $V_E$ ) against  $VO_2$  as described previously [20]. Two linear regression lines were fit to the lower and upper portions of the  $V_E$  vs.  $VO_2$  curve, before and after the break points, respectively. The intersection of these two lines was defined as VT.

At least 24-48 hours following the  $VO_{2PEAK}$  test, participants returned to determine time to exhaustion (TTE) at 110% of the maximum workload reached during the graded exercise test. Participants completed a warm-up consisting of a five-minute cycle at a workload of 50 W. Following warm-up, participants pedaled at 110% of the maximum workload achieved during their  $VO_{2PEAK}$  test. Keeping a cadence of 70 RPM, they pedaled until volitional exhaustion. Time was recorded in seconds, and total work done (TWD) was reported in kilojoules, determined by multiplying the workload in watts and the time to exhaustion in seconds.

Reliability of  $VO_{2PEAK}$ , VT and TWD was determined using a subsample of subjects ( $n = 10$ ) measured during each scheduled testing week. The test-retest intraclass correlation coefficient ( $R$ ) was 0.96 ( $SE \pm 0.1$  L), 0.67 ( $SE \pm 0.3$  L), and 0.79 ( $SE \pm 4.8$  kJ), respectively, for the three measurement variables.

A total of three testing sessions occurred throughout a nine-week period--familiarization (week 1), baseline (week 4), and post (week 9). Familiarization testing was implemented to reduce any learning effect--possibly influencing the dependent variables as well as the training intensity--from the initial  $VO_{2PEAK}$  testing.

### Supplementation

Following familiarization testing, participants were randomly assigned, in a double-blind fashion to either a Cr ( $n = 16$ ) or a Pl ( $n = 17$ ) group. A control group (CON;  $n = 10$ ), neither supplemented nor completed the high-intensity interval training, and instead only completed the testing measurements during each of the scheduled testing weeks. Participants supplemented for a total of 30

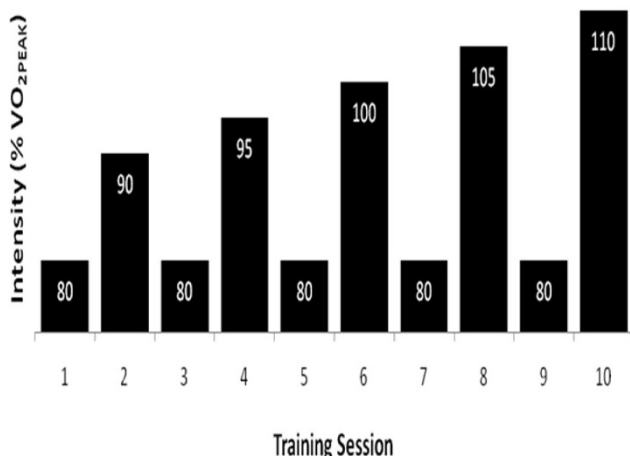
days (10 days of familiarization period followed by an additional 20 days of supplementing and training) at a dose of 10 g per day, taken in two doses--one dose 30 minutes prior to and one dose immediately following training. Participants only supplemented on training days (5 days/week) under the supervision of the researchers, to monitor compliance. Participants in the Cr group consumed 5 g of creatine citrate mixed with 15 g dextrose per packet (Creatine Edge, FSI Nutrition, Omaha, NE), dissolved in 4-8 ounces of water. Similarly, participants in the Pl group consumed 20 g of dextrose per packet dissolved in 4-8 ounces of water. Both drinks were identical in appearance and taste.

### High-intensity interval training (HIIT)

Training began at least 24-48 hours following the TTE test. Participants were required to visit the lab five days per week, for six weeks, to perform the HIIT. A two-week familiarization training period was implemented before taking baseline testing measurements. Due to the effectiveness of the training, and to the generally untrained population, a familiarization period was implemented to allow for all participants to quickly adapt to the high-intensity protocol. Previous research has shown significant improvements in performance with just two weeks of HIIT [21]. Furthermore, in a previous study from our lab in which a familiarization period was not used, the large adaptations from training may have masked any effects from supplementation [22]. After the two-week familiarization period, participants re-tested their  $VO_{2PEAK}$  and TTE. Following baseline testing, participants completed four additional weeks of training, in which the intensities were re-evaluated based on baseline  $VO_{2PEAK}$  power output values. Three of the five days per week of training consisted of training at progressively increasing workloads, determined as a percentage of the participant's baseline  $VO_{2PEAK}$  max workload. One recovery day (two days per week) occurred between each of the three difficult training sessions. During these recovery days, participants completed a training session at 80% of their  $VO_{2PEAK}$  max workload. Difficult training days increased in intensity each session beginning at 90% of their  $VO_{2PEAK}$  max workload and progressing up to 120% of their  $VO_{2PEAK}$  max workload (Figure 1). Each training session began with a five-minute warm up at 50 W, followed by a protocol of five sets of two-minute exercise bouts, with one minute of passive rest in between exercise bouts.

### Statistical analysis

Descriptive statistics were evaluated to determine group demographics. A mixed factorial ANOVA (group [Cr vs. Pl vs. Con]  $\times$  time [pre vs. post]) was evaluated, looking for any significant differences ( $P \leq 0.05$ ) between treatment groups and across time for each variable measured. If a significant interaction occurred, the statistical model was



**Figure 1**  
**HIIT protocol.** Represents the first two weeks of the HIIT protocol. Training intensity eventually reached 120% of the VO<sub>2PEAK</sub> maximum workload.

decomposed and the simple main effects were examined using separate one-way repeated measures ANOVAs for each group. If the result was a simple main effect, Bonferroni *post-hoc* comparisons were performed among groups, while dependent-samples t-tests with Bonferroni corrections were performed across time. If no interactions occurred, the main effects were analyzed by collapsing across the non-interacting variables and analyzed in the same approach as described for the simple main effect.

**Results**

Separate one-way ANOVAs indicated no differences between groups in any of the variables at baseline measurement. In addition, there was no change measured in the Con group over time in any of the variables.

**Body Weight (BW)**

There was no change in BW from baseline to post measurement in the Cr (84.0 ± 12.5 kg and 84.4 ± 12.3 kg, respectively) or Pl (82.9 ± 15.2 kg and 83.2 ± 15.0 kg, respectively) groups.

**Maximal Oxygen Consumption (VO<sub>2PEAK</sub>) and Time to Exhaustion (VO<sub>2PEAK</sub>TTE)**

A significant two-way interaction (time × treatment, p < 0.001) for VO<sub>2PEAK</sub> occurred, and a post hoc Bonferroni analysis indicated no significant differences between groups at post measurements. However, a main effect for time (p < 0.001) occurred due to a change in VO<sub>2PEAK</sub> over time in the Cr (p = 0.002) and Pl (p = 0.001) groups, as indicated by separate Bonferroni-adjusted (p < 0.017) dependent-samples t-tests (Table 1).

There was a significant two-way interaction (time × treatment, p < 0.001) for VO<sub>2PEAK</sub>TTE; however, a post hoc Bonferroni analysis indicated no significant differences between groups at post measurement. A main effect for time (p < 0.001) occurred, and separate Bonferroni-adjusted (p < 0.017) dependent-samples t-tests indicated a significant change over time in the Cr (p < 0.001) and Pl (p < 0.001) groups.

**Ventilatory Threshold (VT)**

A significant two-way interaction (time × treatment, p = 0.040) occurred for VT (l·min<sup>-1</sup>). A post hoc Bonferroni analysis indicated no difference between Cr and Pl (Table 1). Separate Bonferroni-adjusted (p < 0.017) dependent-samples t-test indicated a change over time for Cr (p = 0.001), but not for Pl (p = 0.040) (Figure 2).

**Total Work Done (TWD)**

Table 2 summarizes the mean changes in TWD at 110% of the VO<sub>2PEAK</sub> maximum workload within the three treatment groups. There was no interaction and no main effect for time for either group.

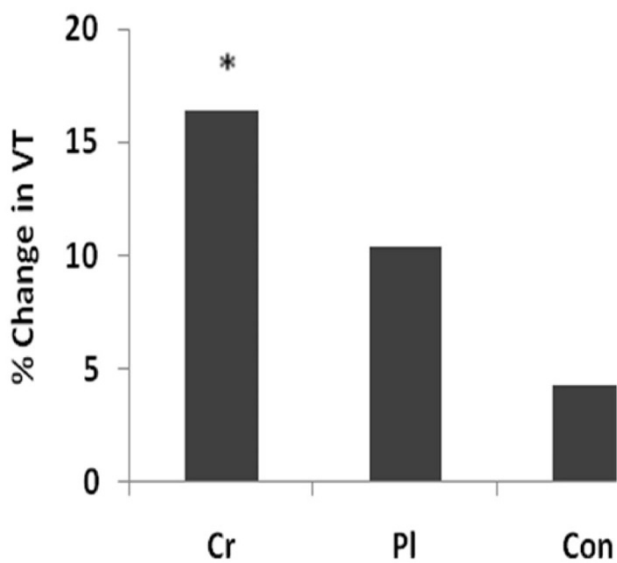
**Discussion**

High-intensity interval training has been shown to be an effective method for improving endurance performance [7,12,23-26]. The results of the present study are in agreement with many studies demonstrating an increase in VO<sub>2PEAK</sub> after HIIT [12,27-29]. In addition, time to exhaustion during the graded exercise test was also improved. However, few studies have examined the concurrent effects of HIIT with Cr supplementation on endurance performance. The current study demonstrated no

**Table 1: Mean ± SD of maximal oxygen consumption (VO<sub>2PEAK</sub>), time to exhaustion (VO<sub>2PEAK</sub>TTE), and ventilatory threshold (VT) at baseline and following four weeks of treatment**

	VO <sub>2PEAK</sub> (l·min <sup>-1</sup> )		VO <sub>2PEAK</sub> TTE (seconds)		VT (l·min <sup>-1</sup> )	
	Baseline	Post	Baseline	Post	Baseline	Post
<b>Cr (n = 16)</b>	3.60 ± 0.55	3.87 ± 0.47*	818.3 ± 127.2	869.3 ± 130.0*	2.14 ± 0.53	2.49 ± 0.57*
<b>Pl (n = 17)</b>	3.65 ± 0.59	4.00 ± 0.59*	837.7 ± 130.1	899.4 ± 127.9*	2.30 ± 0.51	2.54 ± 0.48
<b>Con (n = 10)</b>	3.67 ± 0.71	3.54 ± 0.71	802.8 ± 148.9	781.9 ± 151.2	2.08 ± 0.70	1.99 ± 0.48

\*Indicates a significant (p ≤ 0.01) change over time within treatment groups.



**Figure 2**  
**Effect of Creatine and HIIT on VT.** Percent change in VT over time for each group.

additional improvements in  $VO_{2PEAK}$  when combining Cr supplementation and HIIT. However, when measuring VT, improvements were only demonstrated in the Cr group. Interestingly, in contrast to previous reports of significant increases in TWD with Cr supplementation or HIIT alone, no change in TWD was observed [5,28,30-33].

Endurance performance is commonly assessed using a measure of aerobic capacity,  $VO_{2PEAK}$ . HIIT has been reported to be effective in improving  $VO_{2PEAK}$  5-15% [12,27-29,34-40]. In the current study, a 9% increase in  $VO_{2PEAK}$  was observed. Short recovery periods between intense exercise bouts during HIIT may create a greater reliance on aerobic metabolism, due to the importance of aerobic metabolism in the removal of lactic acid and for the resynthesis of phosphocreatine [41]. HIIT may also induce up-regulation of glycolytic and oxidative enzymes, a possible mechanism influencing the improvements in  $VO_{2PEAK}$  [34]. In addition, an increase in stroke volume

following HIIT [11] may contribute to an increase in  $VO_{2PEAK}$ .

While the HIIT program was effective in improving  $VO_{2PEAK}$  by 9%, creatine supplementation had no further influence on aerobic capacity. These results are in agreement with the few studies that have examined the effects of Cr supplementation on  $VO_{2PEAK}$  [30,42-44]. Cr has been shown to be effective in improving short-duration, intense activities, but few studies have examined the effects of Cr on longer duration, endurance-type activities. Due to the intensity and time duration (two minutes) of the interval work periods, it was hypothesized that Cr would provide for a greater training capacity, and, therefore, the Cr group would show greater improvements in the testing measurements. McConell and colleagues [45] found that Cr improved the maintenance of energy balance in the muscle during intense aerobic exercise; however, performance was not improved, which is in agreement with the current study.

Ventilatory threshold (VT) may be another useful predictor of endurance performance. The VT has been suggested as an indicator of the ability of the cardiovascular system to adequately supply oxygen to the working muscles, preventing muscle anaerobiosis [46]. Performing exercise at intensities greater than VT commonly result in an inadequate supply of oxygen to the working muscles, quickly leading to fatigue [47]. Therefore, improvements in VT may correspond to an augmented time to exhaustion and a greater threshold for fatigue. Additionally, it has been proposed that training at intensities greater than VT, much like the HIIT protocol of the current study, may enhance the efficiency of the body to supply oxygen to the working muscles (i.e. VT) [12,48-50]. Furthermore, a concomitant rise in muscle lactate levels and a drop in pH at high intensities of exercise may signal arterial chemoreceptors, altering ventilatory regulating mechanisms. Therefore, improvements in cardiovascular fitness may also coincide with a decrease in lactate accumulation resulting in an improvement in VT. However, in the current study, significant improvements in VT were only observed in the Cr group (16%), although the Pl group demonstrated a trend for improved VT (10%). The increased VT in the Cr group is in agreement with previous studies that demonstrated improved VT following Cr supplementation but without training [30,42,44]. Improved exercise efficiency at submaximal workloads following Cr supplementation may be attributed to an increase in muscle PCr levels, which may augment the ratio of ATP/ADP, stimulating mitochondrial respiration [42] and delaying reliance on anaerobic glycolysis [51,52]. In addition to acting as an energy buffer, PCr also acts as a proton ( $H^+$ ) buffer when the creatine kinase reaction favors regenerating ATP [53]. This utilization of  $H^+$  may delay the decrease in skeletal muscle

**Table 2: Mean  $\pm$  SD of total work done (TWD) at 110% of  $VO_{2PEAK}$  maximum workload at baseline and following four weeks of treatment**

	TWD (kJ)	
	Baseline	Post
<b>Cr (n = 16)</b>	42.3 $\pm$ 8.0	40.5 $\pm$ 9.4
<b>Pl (n = 17)</b>	47.5 $\pm$ 14.1	43.3 $\pm$ 10.0
<b>Con (n = 10)</b>	37.7 $\pm$ 9.1	39.0 $\pm$ 11.6

pH, possibly signaling arterial chemoreceptors and augmenting the ventilatory response. Future studies, however, are needed to validate our results.

Both HIIT and Cr have been reported to improve total work done [5,28,30-33]. However, no improvements were observed in TWD during a ride to exhaustion at 110% of the maximum workload reached during the GXT in the present study. One reason for the lack of improvement in TWD in the current study may be participant motivation. The 110% workload during which TWD was measured was the first ride to exhaustion in a session of 3 rides to exhaustion. Therefore, participants may have quit early in order to save energy for the subsequent work bouts. In addition, studies that observed improvements in TWD following Cr supplementation implemented a loading phase (20 g/d for 5-7 days) into the supplementation protocol [30-33]. A loading phase was not used in the current study, so it may be possible that muscle PCr levels were not increased enough to aid in improving TWD.

## Conclusion

In conclusion, the current study supports previous evidence that HIIT is an efficient way to induce cardiorespiratory improvements [7,12,23-26]. However, although Cr supplementation has been shown to improve intense exercise [54,55], no apparent benefits were observed in the present study. Furthermore, while improvements in VT were observed following Cr supplementation, it did not lead to an increase in TWD. A Cr loading phase followed by a maintenance phase might improve HIIT more than the low-dose supplementation used in the current study. However, Jager et al. found improvements in interval exercise performance using a similar dose of creatine citrate (5 g/day for 28 days) [56]. Due to the possibility that any benefits of low-dose creatine supplementation were masked by the effectiveness of HIIT alone, a longer training period may be implemented in future studies to determine whether low-dose Cr supplementation will induce further improvements when results from training begin to plateau.

## Competing interests

The authors declare that they have no competing interests.

## Authors' contributions

JG, AS, KK and DF contributed in writing and editing the manuscript along with concept and design, data acquisition, and data analysis and interpretation. JM, TB, JC, and JS contributed in writing and editing the manuscript, as well as concept and design. All authors have read and approved the final manuscript.

## References

1. Coyle EF: **Integration of the physiological factors determining endurance performance ability.** *Exerc Sport Sci Rev* 1995, **23(25-63)**.
2. Hawley JA: **Adaptations of skeletal muscle to prolonged, intense endurance training.** *Clin Exp Pharmacol Physiol* 2002, **29(3)**:218-22.
3. Holloszy JO, Coyle EF: **Adaptations of skeletal muscle to endurance exercise and their metabolic consequences.** *J Appl Physiol* 1984, **56(4)**:831-8.
4. Burgomaster KA, Hughes SC, Heigenhauser GJ, Bradwell SN, Gibala MJ: **Six sessions of sprint interval training increases muscle oxidative potential and cycle endurance capacity in humans.** *J Appl Physiol* 2005, **98(6)**:1985-90.
5. Harmer AR, McKenna MJ, Sutton JR, Snow RJ, Ruell PA, Booth J, Thompson MW, Mackay NA, Stathis CG, Cramer RM, Carey MF, Eager DM: **Skeletal muscle metabolic and ionic adaptations during intense exercise following sprint training in humans.** *J Appl Physiol* 2000, **89(5)**:1793-803.
6. Burgomaster KA, Heigenhauser GJ, Gibala MJ: **Effect of short-term sprint interval training on human skeletal muscle carbohydrate metabolism during exercise and time-trial performance.** *J Appl Physiol* 2006, **100(6)**:2041-7.
7. Weston AR, Myburgh KH, Lindsay FH, Dennis SC, Noakes TD, Hawley JA: **Skeletal muscle buffering capacity and endurance performance after high-intensity interval training by well-trained cyclists.** *Eur J Appl Physiol Occup Physiol* 1997, **75(1)**:7-13.
8. Edge J, Bishop D, Goodman C: **The effects of training intensity on muscle buffer capacity in females.** *Eur J Appl Physiol* 2006, **96(1)**:97-105.
9. Laursen PB, Shing CM, Peake JM, Coombes JS, Jenkins DG: **Influence of high-intensity interval training on adaptations in well-trained cyclists.** *J Strength Cond Res* 2005, **19(3)**:527-33.
10. Jenkins DG, Quigley BM: **The influence of high-intensity exercise training on the Wlim-Tlim relationship.** *Med Sci Sports Exerc* 1993, **25(2)**:275-82.
11. Helgerud J, Hoydal K, Wang E, Karlsen T, Berg P, Bjerkaas M, Simonsen T, Helgesen C, Hjørth N, Bach R, Hoff J: **Aerobic high-intensity intervals improve VO2max more than moderate training.** *Med Sci Sports Exerc* 2007, **39(4)**:665-71.
12. Burke J, Thayer R, Belcamino M: **Comparison of effects of two interval-training programmes on lactate and ventilatory thresholds.** *Br J Sports Med* 1994, **28(1)**:18-21.
13. Cottrell GT, Coast JR, Herb RA: **Effect of recovery interval on multiple-bout sprint cycling performance after acute creatine supplementation.** *J Strength Cond Res* 2002, **16(1)**:109-16.
14. Bogdanis GC, Nevill ME, Boobis LH, Lakomy HK, Nevill AM: **Recovery of power output and muscle metabolites following 30 s of maximal sprint cycling in man.** *J Physiol* 1995, **482(Pt 2)**:467-80.
15. Hultman E, Soderlund K, Timmons JA, Cederblad G, Greenhaff PL: **Muscle creatine loading in men.** *J Appl Physiol* 1996, **81(1)**:232-7.
16. Harris RC, Soderlund K, Hultman E: **Elevation of creatine in resting and exercised muscle of normal subjects by creatine supplementation.** *Clin Sci (Lond)* 1992, **83(3)**:367-74.
17. Stout J, Eckerson J, Ebersole K, Moore G, Perry S, Housh T, Bull A, Cramer J, Batheja A: **Effect of creatine loading on neuromuscular fatigue threshold.** *J Appl Physiol* 2000, **88(1)**:109-12.
18. Volek JS, Kraemer WJ: **Creatine Supplementation: Its effect on human muscular performance and body composition.** *J Strength Cond Res* 1996, **10(200-210)**.
19. Derave W, Eijnde BO, Verbessem P, Ramaekers M, Van Leemputte M, Richter EA, Hespel P: **Combined creatine and protein supplementation in conjunction with resistance training promotes muscle GLUT-4 content and glucose tolerance in humans.** *J Appl Physiol* 2003, **94(5)**:1910-6.
20. Orr GW, Green HJ, Hughson RL, Bennett GW: **A computer linear regression model to determine ventilatory anaerobic threshold.** *J Appl Physiol* 1982, **52(5)**:1349-52.
21. Talanian JL, Galloway SD, Heigenhauser GJ, Bonen A, Spriet LL: **Two weeks of high-intensity aerobic interval training increases the capacity for fat oxidation during exercise in women.** *J Appl Physiol* 2007, **102(4)**:1439-47.

22. Smith AE, Moon JR, Kendall KL, Graef JL, Lockwood CM, Walter AA, Beck TW, Cramer JT, Stout JR: **The effects of beta-alanine supplementation and high-intensity interval training on neuromuscular fatigue and muscle function.** *Eur J Appl Physiol* 2009, **105(3)**:357-63.
23. Daniels JT, Yarbrough RA, Foster C: **Changes in VO<sub>2</sub> max and running performance with training.** *Eur J Appl Physiol Occup Physiol* 1978, **39(4)**:249-54.
24. Dolgener FA, Brooks WB: **The effects of interval and continuous training on VO<sub>2</sub> max and performance in the mile run.** *J Sports Med Phys Fitness* 1978, **18(4)**:345-52.
25. Thomas TR, Adeniran SB, Etheridge GL: **Effects of different running programs on VO<sub>2</sub> max, percent fat, and plasma lipids.** *Can J Appl Sport Sci* 1984, **9(2)**:55-62.
26. Westgarth-Taylor C, Hawley JA, Rickard S, Myburgh KH, Noakes TD, Dennis SC: **Metabolic and performance adaptations to interval training in endurance-trained cyclists.** *Eur J Appl Physiol Occup Physiol* 1997, **75(4)**:298-304.
27. Burgomaster KA, Howarth KR, Phillips SM, Rakobowchuk M, Macdonald MJ, McGee SL, Gibala MJ: **Similar metabolic adaptations during exercise after low volume sprint interval and traditional endurance training in humans.** *J Physiol* 2008, **586(1)**:151-60.
28. Edge J, Bishop D, Goodman C, Dawson B: **Effects of high- and moderate-intensity training on metabolism and repeated sprints.** *Med Sci Sports Exerc* 2005, **37(11)**:1975-82.
29. Gross M, Swensen T, King D: **Nonconsecutive- versus consecutive-day high-intensity interval training in cyclists.** *Med Sci Sports Exerc* 2007, **39(9)**:1666-71.
30. Zoeller RF, Stout JR, O'Kroy JA, Torok DJ, Mielke M: **Effects of 28 days of beta-alanine and creatine monohydrate supplementation on aerobic power, ventilatory and lactate thresholds, and time to exhaustion.** *Amino Acids* 2007, **33(3)**:505-10.
31. Preen D, Dawson B, Goodman C, Lawrence S, Beilby J, Ching S: **Effect of creatine loading on long-term sprint exercise performance and metabolism.** *Med Sci Sports Exerc* 2001, **33(5)**:814-21.
32. van Loon LJ, Oosterlaar AM, Hartgens F, Hesselink MK, Snow RJ, Wagenmakers AJ: **Effects of creatine loading and prolonged creatine supplementation on body composition, fuel selection, sprint and endurance performance in humans.** *Clin Sci (Lond)* 2003, **104(2)**:153-62.
33. McNaughton LR, Dalton B, Tarr J: **The effects of creatine supplementation on high-intensity exercise performance in elite performers.** *Eur J Appl Physiol Occup Physiol* 1998, **78(3)**:236-40.
34. MacDougall JD, Hicks AL, MacDonald JR, McKelvie RS, Green HJ, Smith KM: **Muscle performance and enzymatic adaptations to sprint interval training.** *J Appl Physiol* 1998, **84(6)**:2138-42.
35. Hickson RC, Bomze HA, Holloszy JO: **Linear increase in aerobic power induced by a strenuous program of endurance exercise.** *J Appl Physiol* 1977, **42(3)**:372-6.
36. Keith SP, Jacobs I, McLellan TM: **Adaptations to training at the individual anaerobic threshold.** *Eur J Appl Physiol Occup Physiol* 1992, **65(4)**:316-23.
37. Rodas G, Ventura JL, Cadefau JA, Cusso R, Parra J: **A short training programme for the rapid improvement of both aerobic and anaerobic metabolism.** *Eur J Appl Physiol* 2000, **82(5-6)**:480-6.
38. Tabata I, Nishimura K, Kouzaki M, Hirai Y, Ogita F, Miyachi M, Yamamoto K: **Effects of moderate-intensity endurance and high-intensity intermittent training on anaerobic capacity and VO<sub>2</sub>max.** *Med Sci Sports Exerc* 1996, **28(10)**:1327-30.
39. Ray CA: **Sympathetic adaptations to one-legged training.** *J Appl Physiol* 1999, **86(5)**:1583-7.
40. Hoogeveen AR: **The effect of endurance training on the ventilatory response to exercise in elite cyclists.** *Eur J Appl Physiol* 2000, **82(1-2)**:45-51.
41. Linossier MT, Denis C, Dormois D, Geysant A, Lacour JR: **Ergometric and metabolic adaptation to a 5-s sprint training programme.** *Eur J Appl Physiol Occup Physiol* 1993, **67(5)**:408-14.
42. Nelson AG, Day R, Glickman-Weiss EL, Hegsted M, Kokkonen J, Sampson B: **Creatine supplementation alters the response to a graded cycle ergometer test.** *Eur J Appl Physiol* 2000, **83(1)**:89-94.
43. Reardon TF, Ruell PA, Fiatarone Singh MA, Thompson CH, Rooney KB: **Creatine supplementation does not enhance submaximal aerobic training adaptations in healthy young men and women.** *Eur J Appl Physiol* 2006, **98(3)**:234-41.
44. Murphy AJ, Watsford ML, Coutts AJ, Richards DA: **Effects of creatine supplementation on aerobic power and cardiovascular structure and function.** *J Sci Med Sport* 2005, **8(3)**:305-13.
45. McConell GK, Shinewell J, Stephens TJ, Stathis CG, Canny BJ, Snow RJ: **Creatine supplementation reduces muscle inosine monophosphate during endurance exercise in humans.** *Med Sci Sports Exerc* 2005, **37(12)**:2054-61.
46. Wasserman K, Beaver WL, Whipp BJ: **Gas exchange theory and the lactic acidosis (anaerobic) threshold.** *Circulation* 1990, **81(1 Suppl)**:114-30.
47. Wasserman K, Whipp BJ, Koyle SN, Beaver WL: **Anaerobic thresholds and respiratory gas exchange during exercise.** *J Appl Physiol* 1973, **35(2)**:236-43.
48. Poole DC, Gaesser GA: **Response of ventilatory and lactate thresholds to continuous and interval training.** *J Appl Physiol* 1985, **58(4)**:1115-21.
49. Acevedo EO, Goldfarb AH: **Increased training intensity effects on plasma lactate, ventilatory threshold, and endurance.** *Med Sci Sports Exerc* 1989, **21(5)**:563-8.
50. Laursen PB, Blanchard MA, Jenkins DG: **Acute high-intensity interval training improves Tvent and peak power output in highly trained males.** *Can J Appl Physiol* 2002, **27(4)**:336-48.
51. Balsom PD, Soderlund K, Sjodin B, Ekblom B: **Skeletal muscle metabolism during short duration high-intensity exercise: influence of creatine supplementation.** *Acta Physiol Scand* 1995, **154(3)**:303-10.
52. Febbraio MA, Flanagan TR, Snow RJ, Zhao S, Carey MF: **Effect of creatine supplementation on intramuscular TCr, metabolism and performance during intermittent, supramaximal exercise in humans.** *Acta Physiol Scand* 1995, **155(4)**:387-95.
53. Volek JS, Kraemer WJ, Bush JA, Boetes M, Incledon T, Clark KL, Lynch JM: **Creatine supplementation enhances muscular performance during high-intensity resistance exercise.** *J Am Diet Assoc* 1997, **97(7)**:765-70.
54. Casey A, Constantin-Teodosiu D, Howell S, Hultman E, Greenhaff PL: **Creatine ingestion favorably affects performance and muscle metabolism during maximal exercise in humans.** *Am J Physiol* 1996, **271(1 Pt 1)**:E31-7.
55. Tarnopolsky MA, MacLennan DP: **Creatine monohydrate supplementation enhances high-intensity exercise performance in males and females.** *Int J Sport Nutr Exerc Metab* 2000, **10(4)**:452-63.
56. Jager R, Metzger J, Lautmann K, Shushakov V, Purpura M, Geiss KR, Maassen N: **The effects of creatine pyruvate and creatine citrate on performance during high intensity exercise.** *J Int Soc Sports Nutr* 2008, **5(4)**:

Publish with **BioMed Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:  
http://www.biomedcentral.com/info/publishing\_adv.asp

