Impact of time and work:rest ratio matched sprint interval training programmes on performance: A randomised controlled trial

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6 Abstract

- 7 Objectives
- 8 The aim of this study was to examine the effects of a short training intervention using two
- 9 repeated sprint protocols matched for total sprint duration and work:rest ratio.
- 10 Design
- 11 Randomised-controlled trial
- 12 Methods
- 13 Thirty physically active males were randomly allocated to one of two sprint training groups: a
- 14 6 second group, a 30 second group or a non-exercising control. The training groups were
- 15 matched for work:rest ratio and total sprint time per session, and completed 6 training
- 16 sessions over a 2-week period. Before and after the 2 week training period, participants
- 17 completed a VO_{2max} test and a 10km time trial on a cycle ergometer.
- 18 Results
- 19 Time trial performance increased significantly by 5.1% in 6 sec (630 \pm 115 sec to 598 \pm 92
- 20 sec; p<0.05) and 6.2% in 30 sec (579 ± 68 sec to 543 ± 85 sec; p<0.05) from baseline
- testing, but there was no significant change in the control group (p>0.05), and no significant
- difference between exercise groups (p>0.05). The 6 sec group increased peak power output
- by 9.0% (from 1092 ± 263 W to 1181 ± 248 W; p<0.05) from sprint session 1 to 6, and the 30
- 24 sec group by 20.0% (1041 \pm 161 W to 1237 \pm 159 W; p<0.05).
- 25 Conclusions
- 26 This study indicates that both 6 and 30 second bouts of repeated sprint exercise, matched
- 27 for total sprint duration and W:R can improve athletic performance.
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 35 Keywords: HIIT; Time-trial; Cycling; Exercise; Athletic

37 Introduction

There has been renewed interest in the use of sprint interval training (SIT) as an exercise 38 intervention in athletic, recreational and sedentary populations.¹⁻² Despite total exercise time 39 being considerably lower in comparison with traditional endurance training approaches, 40 similar increases in VO_{2max} (defined as the maximum rate at which an individual can take up 41 and utilise oxygen), muscle oxidative capacity and exercise performance have been 42 observed.³⁻⁷ This training approach is characterised by repeated bouts of relatively brief 43 44 intermittent exercise, with an 'all-out' effort⁸, and may be more enjoyable than prolonged endurance training.9-10 45

Many studies looking into the effect of SIT have used a 30 second supramaximal exercise 46 sprint bout with 4 minutes recovery (1:8, work to rest ratio (W:R)), and have reported a range 47 of central adaptations, such as increased cardiac output and stroke volume, and peripheral 48 adaptations including increases in a range of enzymatic concentrations, in both trained an 49 untrained individuals.¹¹⁻¹⁵ It has been hypothesised that some of the adaptations to this type 50 of training are associated with the metabolic demands and signalling responses which occur 51 52 in the early stages of a sprint. Studies have therefore also investigated shorter exercise bouts to determine whether adaptations similar to those observed following 30 second 53 sprints can be elicited.¹⁶⁻¹⁸ 54

Taylor et al.¹⁹ investigated the acute effects of SIT on cell signalling responses in matched 55 56 duration interval and continuous protocols, reporting no difference between the two distinct 57 bouts of exercise despite differences in total work done, which suggests that total work is not 58 necessarily a crucial factor when monitoring adaptations to such protocols. Chia-Lun et al.³ 59 studied the more chronic effect of high intensity training when matched for total time. Although they reported significant improvements in VO_{2max} , neither of the interval training 60 groups were supramaximal, nor did they utilise any form of performance test. Jakeman et 61 al.¹⁶ reported an improvement in time trial performance and time to exhaustion following 2 62 weeks of SIT consisting of 10x6 second sprints, with the improvements in time trial 63 performance being associated with a prolonged time to reach the onset of blood lactate 64 65 accumulation (OBLA – defined¹⁶ as a blood lactate concentration of 4 mmol.L⁻¹). Such enhancements in time trial performance have also been reported following a 30 seconds SIT 66 programme.^{7,20} While these, and other studies¹⁷⁻¹⁸ have reported similar adaptations, there 67 68 are frequent differences between training protocols and consequently, the importance of aspects such as work:rest ratio and the duration of each individual sprint is unclear. Despite 69 the range of SIT paradigms which have been used, there is very little information to compare 70 very short duration (<10 sec) sprint training directly with the more traditional 30 sec SIT 71 approach. It is therefore unknown if the adaptive mechanisms causing improvements in 72 73 performance following SIT will generate similar adaptations with different sprint durations. 74 Therefore, the purpose of this study was to investigate the impact of individual sprint duration on time trial performance and VO_{2max} when W:R and total sprint duration are 75 matched. It was hypothesised that the shorter, 6 second SIT would be at least as effective 76 77 as the 30 second SIT intervention in improving these parameters.

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79 <u>Methods</u>

- 80 Thirty physically active (minimum of 5 hours week⁻¹ in a range of sports) males volunteered
- to participate in this study, which received ethical approval from the local university ethics
- 82 committee, and was carried out in accordance with the declaration of Helsinki. Participants
- 83 were randomly allocated, by blind draw, to one of three groups and completed either two
- 84 weeks training (3 sessions week⁻¹ with a minimum of 24hours between sessions) or were
- 85 asked to follow their normal training programme. Participants in the treatment groups could
- 86 continue exercising outside of the experimental conditions, however all participants were
- informed that this must not be exhaustive exercise. Outcome measures were a VO_{2max} test
- and a 10km time trial; both completed on a cycle ergometer and were assessed before and
- 89 after two weeks of SIT or normal training (control group).

90 Participant characteristics

- 91 On the first visit to the laboratory, basic anthropometric measures were taken, height was
- 92 measured to the nearest 0.1cm using a stadiometer (Holtain, Crosswell, Wales), weight and
- 93 body composition were measured using bioelectrical impedance analysis (BIA) (Tanita, BC-
- 94 418MA, Amsterdam, The Netherlands) to the nearest 0.1kg (table 1).
- 95 Insert table 1 here

96 VO_{2max} test and Time Trial

- 97 During visits one and nine, participants completed an incremental VO_{2max} test, on a Lode
- 98 Excaliber cycle ergometer. Following a 5-minute warm up against a 50Watt (W) load,
- 99 participants cycled against a progressively increasing resistance (25W min) until volitional
- 100 exhaustion. During the VO_{2max} test, heart rate (Polar, FT1, England) and RPE (Borg scale 6-
- 101 20) were recorded every minute, with respiratory variables monitored continuously (Cortex
- 102 Metalyzer 3B, Leipzig, Germany). Mean VO₂ during the final 30 seconds of each maximal 103 test was recorded, and the highest value within 2SD of this mean was taken as recorded as
- the VO_{2max} . Prior to all sessions, gas analysers were calibrated using a gas standard and the
- 105 volume transducer was calibrated with a 3L syringe following manufacturers guidelines.
- 24-48 hours following the VO_{2max} test (session 2 and 10), participants completed a selfpaced 10km time trial on a Lode Excaliber cycle ergometer. The ergometer was set in linear
 mode, and the linear factor was calculated according to the participants' average cadence
 and maximum work rate from the VO_{2max} test. Participants were aware of the distance cycled
 but were blinded to time.
- 111

112 Training intervention

113 Volunteers in the treatment groups were randomly assigned to one of the sprint training

programmes, with both set at a resistance of 7.5% of body weight on a Lode Excaliber cycle

ergometer (LEM Software, Lode, The Netherlands). Both protocols used a W:R of 1:8, such

that all participants completed 2 minutes of sprint work. Each volunteer completed 6 sprint

sessions spread over 14 days, with a minimum of 24 hours rest in between each session.

118 Those in the 6 sec group completed 20x6 second sprints with a 48s recovery, and the 30

119 sec group completed 4x30 second sprints with a 4-minute recovery to replicate commonly 120 used SIT protocols. Participants were given a 10 second countdown before each sprint and were instructed to increase the cadence so they were at their maximal sprint velocity at the 121 start of each sprint. Throughout each sprint the participants were given encouragement to 122 ensure an 'all-out' effort. Power output was recorded throughout sprints using LEM software, 123 124 at a sampling frequency of 5Hz. Peak power output was calculated as the highest recorded power output per sprint. Total work per session was calculated from the mean power output 125 126 per sprint, multiplied by sprint time. The sum of the four or 20 sprints was then converted to 127 kJ.

128 Data Analysis

Data were analysed using SPSS v21.0, and are expressed as mean ± standard deviation, 129 130 unless otherwise stated. Outcome measures were analysed using a repeated measures (RM) ANOVA. VO_{2max} and time trial performance were analysed using a 3 x 2 (group x time) 131 RM ANOVA, and power output data were analysed using a 2 x 6 (group x time) RM ANOVA. 132 The Mauchly sphericity test was used to assess the assumption of sphericity, with the 133 Greenhouse-Geisser correction used for violations. Statistical significance level was set at 134 p≤0.05, and the Scheffé post hoc test was used where appropriate. Cohens D effect sizes 135 136 were calculated, with 0.2, 0.5 and 0.8 being considered a small, medium or large effect size

137 respectively.²¹

138 <u>Results</u>

139 There were no significant differences between groups in participant characteristics,10km 140 time trial, VO_{2max} and peak power output at baseline.

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There was a main effect of time for time trial performance, which improved significantly in both training groups (Figure 1), by 5.1% in 6 sec (d=0.31), 6.2% in 30 sec (d=0.47) (p<0.05), but there was no change in the control group (-1.0%; p>0.05). An interaction effect was also observed, with post hoc analysis showing no significant difference between the 6 sec and 30 sec groups (p>0.05). There was a significant difference between the 30 sec and control groups (p<0.05), but no statistically significant difference between the 6 sec and control qroups (p>0.05). There was no significant main effect for time in VO_{2max} in either

140 groups (p > 0.03). There was no significant main effect for time in \sqrt{O}_{2max} in 140 intervention, and there was no groups time interaction (n > 0.05).

intervention, and there was no group*time interaction (p>0.05).

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151 Insert Figure 1 here

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Peak power output was achieved on either the first or second sprint of each training session. While there was no significant difference in peak power output between training groups (p>0.05), there was a significant time and group*time interaction effect for peak power in both conditions (p<0.05; Figure 2a). Peak power output in 6 sec increased significantly by

9% from session 1 to session 6 (d=0.3), and by 20% in 30 sec (d=1.2) from session 1 to the

final sprint session (Figure 2a). There was a significant group and group*time interaction for

total work done (kJ; p<0.05), with the 6 sec group doing significantly more work than the 30 sec group (d=2.1), however, there was no main effect of time for total work done (Figure 2b).

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162 Insert Figure 2 here

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164 **Discussion**

The main finding of this study is that both a 6 and 30 second repeated sprint intervention for 2 weeks, that were matched for total sprint time and W:R, resulted in similar improvement in time trial performance compared to a control group. There were, however, no effects of either training protocol on VO_{2max} .

Time trial (TT) performance improved significantly in both the 30 sec and 6 sec training 169 group, and remained unchanged in the control group. The improvement in performance of 170 the 30 sec group (6.2%) is similar to that reported by Burgomaster et al.²⁰, who used the 171 same 30 sec training protocol, reporting a 9.6% improvement. TT performance in the 6 sec 172 group also improved (5.1%). In a training study paper by Taylor *et al.*²³ their training control 173 174 group followed a similar programme as to the 30 sec group in this current study. 175 Interestingly, they reported no improvement in time trial performance following the training period. The difference in the findings of this work and the current study could be firstly that 176 their work was conducted on trained cyclists and also that the time trial distance was of a 177 longer duration. SIT seems to be a potent method for improving performance over shorter 178

179 time trials.

180 Although not directly measured in the current study, an increase in mitochondrial enzymes including citrate synthase activity have previously been reported following 30 second SIT 181 protocols^{6,15,20,24}, and changes such as these may have improved the oxidative potential of 182 the muscle and subsequent exercise performance during the current study in both training 183 groups. Although still not clear, the increased flux between rest and exercise may have 184 caused greater perturbations to the muscle milieu during the 6s supramaximal efforts is a 185 possible factor for the adaptations reported following such training bouts.¹⁹ Recent work of 186 Taylor et al.¹⁹ evaluating duration matched interval and continuous exercise demonstrated 187 188 that, despite completing significantly more work in their interval training group, the magnitude of AMPK phosphorylation did not differ between groups. This work supports that of the 189 current study in that a major determinant for adaptation stems from the ability to achieve 190 191 repeated peak power outputs during the intervals rather than complete more work as demonstrated in the 6s training group. It has previously been suggested that the major 192 drivers of performance improvements may occur in the first 6-10 seconds of SIT, with a 6 193 194 second training approach being sufficient to elicit significant performance benefits.¹⁶ While there was a slightly greater time trial improvement in the 30 sec group, the lack of significant 195 performance differences between groups would suggest that a 6 second protocol can be as 196 197 equally beneficial method to elicit performance adaptations when matched for total sprint time, and W:R. Peak power output is typically observed within the early portion of the sprint, 198 and as similar adaptations occurred following both sprint protocols, achieving peak power 199 may be an important feature of performance related adaptations.¹⁸ In the current study, 200 participants accelerated to their top speed at the start of the sprint, ensuring that peak power 201

output was achieved and sustained for as long as possible. Zelt *et al.*²⁵ compared a 30 and
 15 second sprint, and found performance adaptations including an increased VO_{2max} and

critical power, but found no difference between conditions. It may therefore be that

205 repeatedly reaching peak power, rather than sprint duration or total work completed, is the

206 determining factor for improvements in exercise performance. To our knowledge, this current

study is the first to attempt to investigate this by controlling for the work:rest ratio in this way.

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Following two weeks of SIT, peak power output per session significantly increased in both 209 the 30 sec (+20%) and the 6 sec groups (+9%) (Figure 2a). Burgomaster *et al.*²⁰ reported an 210 increase in peak power output by 5.4% following 6 sessions of 30 sec sprints in 2 weeks, 211 however the increase in the current study is more similar to that of Burgomaster et al.⁷, who 212 reported a 17% increase in peak power output following 6 weeks of the 30 sec SIT protocol. 213 Improvement in peak power output has also been reported in studies utilizing shorter (<10 214 seconds) supramaximal bouts, including repeated 5 second and 6 second sprints^{17,16}, which 215 may be linked with an increase in the activity of glycolytic enzymes phosphofructokinase 216 (PFK) and Hexokinase (Hex)²⁶⁻²⁷ and the improved resynthesis of PCr during the recovery 217 period.²⁸ While not assessed in the current study, increases in PFK have been shown to 218 219 occur as pH increases and accelerates the rate of glycolysis, fuelling the initial 5-10 seconds of sprinting.²⁹ Despite the significant increase in peak power, analysis of total work done 220 during the training sessions indicated that there was no significant change in either training 221 group. This therefore indicates a poorer fatigue index following training, particularly in the 30 222 sec group, and although participants were able to achieve similar peak power outputs, these 223 data suggest that restoration of mean power output was slower.²⁸ In addition, those in the 6 224 sec group did significantly more work than those in the 30 sec group, which is likely to reflect 225 the fact that the shorter sprint resulted in less depletion in stored glycogen, and an ability to 226 better resynthesize PCr needed to achieve repeated peak performance. This may also have 227 resulted in the process of glycolysis becoming the dominant driver of exercise¹⁸, potentially 228 increasing glycolytic enzyme activity including Hex and PFK to fuel the sprint during the 229 230 latter sprints. This again may indicate that total work done is not necessarily the main driver of adaptation. It should be noted, that while the differences in work done between the two 231 training groups was significant, individual pacing strategies may have contributed to a 232 portion of this difference. We did not specifically look to protect against pacing, aside from 233 giving strong verbal encouragement, and this is a potential limitation of this study, which 234 future designs may wish to consider guarding against. 235

Similar to previous studies, which have implemented a 2-week SIT programme²⁰, there was 236 no significant change in VO_{2max} in either group. Previous research has indicated that 237 changes in the activity of oxidative enzymes associated with improvements in VO_{2max}, such 238 as citrate synthase, can take up to 6 weeks to reach a higher steady state.³⁰ Additionally, 239 240 central adaptations that influence VO_{2max} may take longer to occur than the 2-week intervention used in the present study.³¹ While there were no statistically significant 241 improvements observed in the current study, it would be of interest to further investigate 242 243 responses to SIT over a more prolonged period.

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Conclusion

This study found that two-weeks of SIT comprising either a 6 or 30 second repeated bouts of exercise which were matched for total sprint time and work:rest ratio elicited similar changes in performance. In comparison with a control group, there were significant improvements in time trial performance, and sprint power output significantly increased for both groups. Adaptations due to the shorter sprint bout may be due to the greater amount and quality of work that can be completed during the 6 sec protocol. This study is the first to match duration and work:rest ratio in this way, and provides interesting insight into adaptations to this type of training. **Practical applications** Two-weeks of SIT using either a 6 or 30 second repeated bouts significantly • improved athletic performance in comparison with a control group on a 10km TT As long as work:rest ratio and total sprint duration are matched, either 6 or 30 second SIT programme is beneficial for performance adaptations **Acknowledgements** We would like to thank the volunteers who participated in this study. No financial assistance was provided for the project, and the authors report no conflict of interest

278		<u>References</u>
279 280 281	1.	Babraj, J., Vollaard, N., Keast, C. <i>et al.</i> (2009). Extremely short duration high intensity interval training substantially improves insulin action in young health males. <i>BMC Endocrine Disorders,</i> 9(3).
282 283	2.	Adamson, S., Lorimer, R., Cobley, J. <i>et al.</i> (2014). High intensity training improves health and physical function in middle aged adults. <i>Biol</i> , 3, 333-344.
284 285 286	3.	Chia-Lun, L., Wei-Chieh, H., Ching-Feng, C. (2016). Physiological Adaptations to Sprint Interval Training with Matched Exercise Volume. <i>Med Sci Sports Exerc</i> , <i>in press</i> , DOI: 10.1249/MSS.0000000000001083.
287 288 289	4.	Nybo, L., Sundstrup, E., Jakobsen, M.D. <i>et al.</i> (2010). High-Intensity Training versus Traditional Exercise Interventions for Promoting Health. <i>Med Sci Sports Exerc.</i> 42 (10), 1951-1958.
290 291 292	5.	Helgerud, J., Høydal, K., Wang, E. <i>et al.</i> (2007). Aerobic High-Intensity Intervals Improve VO _{2max} More Than Moderate Training. <i>Med Sci Sports Exerc</i> . 39 (4), 665- 671.
293 294 295	6.	Gibala, M.J., Little, J.P., Essen, M.V. <i>et al.</i> (2006). Short-term Sprint Interval versus Traditional Endurance Training: Similar Initial Adaptations in Human Skeletal Muscle and Exercise Performance. <i>J Physiol.</i> 575 (3), 901-911.
296 297 298	7.	Burgomaster, K.A., Howarth, K.R., Phillips, S.M. <i>et al.</i> (2008). Similar Metabolic Adaptations During Exercise After Low Volume Sprint Interval and Traditional Endurance Training in Humans. <i>J Physiol</i> . 586 (1), 151-160.
299 300 301	8.	Gibala, M.J. & McGee, S.L. (2008). Metabolic Adaptations to Short-term High- Intensity Interval Training: A Little Pain for a Lot of Gain? <i>Exerc Sport Sci Reviews</i> . 36 (2), 58-63.
302 303 304	9.	Bartlett, J.D., Close, G.L., Maclaren, D.P.M. <i>et al.</i> (2011). High-Intensity Interval Running is Perceived to be More Enjoyable than Moderate-Intensity Continuous Exercise: Implications for Exercise Adherence. <i>J Sport Sci.</i> 29 (6), 547-553.
305 306 307 308	10.	Kong, Z., Fan, X., Sun, S. <i>et al.</i> (2016). Comparison of high-intensity interval training and moderate-to-vigorous continuous training for cardiometabolic health and exercise enjoyment in obese young women: A randomized controlled trial. <i>PLoS ONE</i> . 11(7), e0158589. doi: 10.1371/journal.pone.0158589.
309 310	11.	Laursen, P.B. & Jenkins, D.G. (2002). The Scientific Basis for High Intensity Interval Training. <i>Sports Med</i> . 32 (1), 53-73.
311 312 313 314	12.	Bayati, M., Farzad, B., Gharakhanlou, R. <i>et al.</i> (2011). A Practical Model of Low- Volume High-Intensity Interval Training Induces Performance and Metabolic Adaptations that Resemble 'all-out' Sprint Interval Training. <i>J Sport Sci Med.</i> 10 (1), 571-576.

- 315 13. Harmer, A.R., McKenna, M.J., Sutton, J.R. *et al.* (2000). Skeletal Muscle Metabolic
 and Ionic Adaptations During Intense Exercise Following Sprint Training in Humans.
 317 *J Appl Physiol.* 89 (1), 1793-1803.
- 14. Creer, A.R., Ricard, M.D., Conlee, R.K. *et al.* (2004). Neural, Metabolic, and
 Performance Adaptations to Four Weeks of High Intensity Sprint-Interval Training in
 Trained Cyclists. *Int J Sports Med.* 25 (1), 92-98.
- 15. Burgomaster, K.A., Hughes, S.C., Heigenhauser, G.J.F. *et al.* (2005). Six Sessions of
 Sprint Interval Training Increases Muscle Oxidative Potential and Cycle Endurance
 Capacity in Humans. *J Appl Physiol.* 98 (1), 1985-1990.
- 16. Jakeman, J., Adamson, S. & Babraj, J. (2012). Extremely short duration high
 intensity training substantially improves endurance performance in triathletes. *Appl Physiol, Nut Metabol.* 37 (5), 976-981.
- 17. Linossier, M.T., Denis, C., Dormois, D. *et al.* (1993). Ergometric and Metabolic
 Adaptation to a 5-s Sprint Training Programme. *Eur J Physiol*. 67 (1), 408-414.
- 18. Hazell, T.J., MacPherson, R.E.K., Gravelle, B.M.R. *et al.* (2010). 10 or 30-s Sprint
 Interval Training Bouts Enhance Both Aerobic and Anaerobic Performance. *Eur J Appl Physiol Occ Physiol.* 110 (1), 153-160.
- 19. Taylor, C.W., Ingham, S.A., Hunt, J.E.A. *et al.* (2016a). Exercise duration-matched
 interval and continuous sprint cycling induce similar increases in AMPK
 phosphorylation, PGC-1α and VEGF mRNA expression in trained individuals. *Eur J Physiol*, 116, 1445-1454.
- 20. Burgomaster, K.A., Heigenhauser, G.J.F. & Gibala, .M.J. (2006). Effect of Short Term Sprint Interval Training on Human Skeletal Muscle Carbohydrate Metabolism
 During Exercise and Time-Trial Performance. *J Appl Physiol*. 100 (1), 2041-2047.
- 21. Cohen, J (1988). *Statistical Power Analysis for the Behavioural Sci*. 2nd ed. Hillsdale:
 NJ: Lawrence Erlbaum.
- 22. Paton, C.D. & Hopkins, W.G. (2006). Variation in Performance of Elite Cyclists from
 Race to Race. *Eur J Sport Sci.* 6 (1), 25-31.
- Taylor, C.W., Ingham, S.A., and Ferguson, R.A. (2016b). Acute and chronic effect of
 sprint interval training combined with postexercise blood-flow restriction in trained
 individuals. *Exp Physiol*, 101.1, 143-154.
- 24. Little, J.P., Safdar, A., Wilkin, G.P. *et al.* (2010). A practical model of low-volume
 high-intensity interval training induces mitochondrial biogenesis in human skeletal
 muscle: potential mechanisms. *J Physiol.* 588 (6), 1011-1022.
- 25. Zelt, J.G.E., Hankinson, P.B., Foster, W.S. *et al.* (2014). Reducing the Volume of
 Sprint Interval Training Does Not Diminish Maximal and Submaximal Performance
 Gains in Healthy Men. *Eur J Physiol*. 114 (11), 2427-2436.

352 353 354	 Rodas, G., Ventura, J.L., Cadefau, J.A. <i>et al.</i> (2000). A Short Training Programme fo the Rapid Improvement of Both Aerobic and Anaerobic Metabolism. <i>Eur J Physiol</i>. 82 (5-6), 480-486.
355 356 357	 MacDougall, J.D., Hicks, A.L., MacDonald, J.R. <i>et al.</i> (1998). Muscle Performance and Enzymatic Adaptations to Sprint Interval Training. <i>J Appl Physiol</i>. 84 (6), 2138- 2142.
358 359 360	 Bogdanis, G.C., Nevill, M.E., Boobis, L.H. <i>et al.</i> (1995). Recovery of power output and muscle metabolites following 30 s of maximal sprint cycling in man. <i>J Appl Physiol</i>, 482(2), 467-480.
361 362	29. Beneke, R., Pollman, C., Bleif, I. <i>et al</i> . (2002). How Anaerobic is the Wingate Anaerobic Test for Humans? <i>Eur J Physiol</i> . 87 (4-5), 388-392.
363 364	 Hood, D.A. (2001). Invited Review: Contractile Activity-Induced Mitochondrial Biogenesis in Skeletal Muscle. J Appl Physiol. 90 (3), 1137-1157.
365 366 367	 Gist, N.H., Freese, E.C. & Cureton, K.J. (2014). Comparison of Responses to Two High-Intensity Intermittent Exercise Protocols. J Strength Cond Res. 28 (11), 3033- 3040.
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373	Figure 1: 10km time trial performance; * denotes a significant difference from baseline
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375 376 377	Figure 2: Panel a) Peak power output; * denotes a significant increase from session 1 to session 6; Panel b) Total work done; * denotes a significant difference between groups

Group	Age (yrs)	Height (m)	Body Mass (kg)	Body Fat (%)
6 sec (n=10)	21 ± 4	1.78 ± 0.06	75.7 ± 13.9	14.0 ± 5.9
30 sec (n=10)	21 ± 4	1.84 ± 0.06	83.0 ± 10.2	14.0 ± 3.3
Con (n=10)	23 ± 3	1.82 ± 0.07	82.4 ± 7.6	14.9 ± 3.6

Table 1. Participant anthropometric characteristics

Variable	Condition		Pre	Post	d
	6 sec	Mean ± SD	630 ± 115	598 ± 92*	0.31
		95% CI	559-701	541-655	
	30 sec	Mean ± SD	579 ± 68	543 ± 85*	0.47
TT (Seconds)		95% Cl	537- 621	490-596	0.47
	Con	Mean ± SD	631± 104	634 ± 99	0.03
		95% Cl	567- 695	573-695	0.03
	6 sec	Mean ± SD	57 ± 8	59 ± 10	0.00
		95% Cl	52-62	53-65	0.22
	30 sec	Mean ± SD	57 ± 6	58 ± 9	0.40
VO _{2max} (ml.kg.min ')		95% CI	53-61	52-64	0.13
	Con	Mean ± SD	52 ± 9	52 ± 6	0 12
		95% CI	47-59	48-56	0.13

Table 2: Time trial and VO_{2max} data

*Denotes a significant difference from baseline (p<0.05).



Fig 1. 10km time trial performance; * denotes a significant difference from baseline.



Fig 2. Panel (a) Peak power output; * denotes a significant increase from session 1 to session 6; Panel (b) Total work done; * denotes a significant difference between groups.