Comments from the editors and reviewers:

Reviewer 0

General

comments

<u>Reviewer</u>: This study examines changes in various factors when the 4X4 HIIT regime is performed in normoxic and hypoxic conditions. This is novel, but the authors need to make a better case for the importance of this topic, and need to better evaluate text found in the Discussion to increase the impact of these findings.

Authors' response: We thank the reviewer for their careful consideration of our work.

Specific

comments

Reviewer: Please consider these comments re. your paper-thank you.

Authors' response: We have responded to the comments individually raised below.

<u>Reviewer</u>: If word count allows in Abstract, please list the age and VO2max of these runners; thank you.

<u>Authors' response</u>: We thank the reviewers for this comment and have subsequently added the age of the runners who participated in this study. An explanation regarding the absence of a VO_{2MAX} assessment is provided below.

P2 L4 – Nineteen <u>trained</u> runners (33.4 \pm 9.1 years) completed a...

<u>Reviewer</u>: Line 2: HIIR is not a common abbreviation and should be deleted or revised. And because this set of text is referring to benefits of high intensity interval training, why not just use this abbreviation (HIIT) which is so well-recognized?

<u>Authors' response</u>: We acknowledge the reviewers comment that 'HIIT' is a more conventional acryonym that 'HIIR', and have replaced the latter with the former throughout the manuscript.

<u>Reviewer</u>: Line 9; be careful..this 4X4 model that you use is not time efficient vs. MICT (the bout takes 27 min not including warmup and cooldown), unless you are referring to the actual duration of interval exercise which is 16 min. Please consider revising this text.

<u>Authors' response</u>: We thank the reviewer for highlighting this comment and have subsequently added in further information regarding the time efficiency of HIIT compared to MICT.

P4 L5-8 – Compared to moderate-intensity continuous running, <u>HIIT</u> leads to similar improvements in cardiorespiratory fitness <u>that is achieved with a shorter effective exercise</u> <u>duration</u> per session (2). Due to the reduced time-commitment and <u>exercise</u> training volume...

<u>Reviewer</u>: Line 25: and it could be argued that these perceptually regulated bouts better mirror how exercisers actually choose to modify intensity during acute exercise.

<u>Authors' response</u>: We thank the author for suggesting this important point, and have subsequently added this information to the manuscript.

P5 L25-26 – ... offer a viable solution, and is perhaps more reflective of how exercisers modify intensity during acute exercise.

<u>Reviewer</u>: Lines 32-34: I am not familiar with this study, but exercise at 75 %HRmax does not meet the criteria of Weston et al. (2014) designating HIIT of > 85 %HRmax.

<u>Authors' response</u>: We thank the reviewer for highlighting the required percentage of maximal heart rate for exercise to be considered as a high-intensity. According to Weston et al. (2014), this does not classify the exercise employed by Chacaroun et al. (2018) as high-intensity the reviewer suggests. However, Chacaroun et al. (2018) compared continuous (30 mins) and interval (1 min on, 1 min off) cycling in hypoxic (clamped SpO₂ = 75%) and normoxic conditions at a similar absolute heart rate (75% of max). The authors make reference to a position stand from the ACSM (1998), highlighting that 75% of maximal heart rate is recommended for continuous exercise. Therefore, although this intensity may not be aligned with the more recent work of Weston et al. (2014), the intensity employed and findings of Chacaroun et al. (2018) support our rationale. As such, we have not made any changes to the current manuscript.

ACSM. The recommended quantity and quality of exercise for developing and maintaining cardiorespiratory and muscular fitness, and flexibility in healthy adults. MSSE. 1998;30(6):975-991.

Chacaroun, S., Gonzalez, I. V. E., Flore, P., Doutreleau, S. and Verges, S. Physiological responses to hypoxic constant-load and high-intensity interval exercise sessions in healthy subjects. Euro J App Phys. 2018:1-12.

Weston, K. S., Wisløff, U., & Coombes, J. S. High-intensity interval training in patients with lifestyle-induced cardiometabolic disease: a systematic review and meta-analysis. Br J Sports Med. 2014;48(16):1227-1234.

<u>Reviewer</u>: Line 40: remember that the Bartlett et al. study that you cite here was in men with VO2max well above 55 mL/kg/min, so these results poorly generalize to the typical adult.

<u>Authors' response</u>: We thank the reviewer for highlighting the work we cite by Bartlett et al. (2011) includes a participant cohort with a relatively high VO_{2MAX}, which may not be representative of adults within the general population. As such, we have since replaced this reference with a study comparing HIIT and continuous exercise (Thum et al., 2017). These authors conclude that HIIT is more enjoyable in active males and females with a more representative VO2_{MAX} of the general population ($41.4 \pm 4.1 \text{ mL/kg/min}$) (Thum et al., 2017).

P26 L447-449 – <u>Thum, J., Parsons, G., Whittle, T., & Astorino, T. (2017). High-intensity</u> <u>interval training elicits higher enjoyment than moderate intensity continuous exercise.</u> <u>PloS one. 2017;12(1):e0166299.</u>

Bartlett, J. D., Close, G. L., MacLaren, D. P. M., Gregson, W., Drust, B., Morton, J. P. High intensity-interval running is perceived to be more enjoyable than moderate intensity continuous exercise: implications for exercise adherence. J Sports Sci. 2011; 29:547–553.

<u>Reviewer</u>: Please fix the typos in lines 44-45; thank you. Also, this text does not lay a solid or clear foundation for this text referring to cognitive function, breathlessness, reaction time, etc. as currently written. This information just appears here with little transition text to relate this particular topic to the current study—please rewrite this text here to better denote the importance or relevance of cognitive function, breathlessness, motivation, etc.

<u>Authors' response</u>: We thank the reviewer for pointing out the spelling errors in the manuscript and these have been corrected accordingly. Cognitive function is usually decreased when assessed in hypoxic conditions, or shortly after exercise at a relative, fixed-inensity in hypoxia, compared to normoxia (McMorris et al., 2017). Similarly, exercise-related sensations are at a premium during maximal repeated sprints in hypoxia compared to normoxia (Brocherie et al., 2017). Therefore, the importance of our work relates to attempting to incorporate a perceptually-regulated exercise intensity to compensate for the negative influence hypoxia has on fixed-intensity exercise versus normoxia, denoted via defects in cognitive function and exercise-related sensations. We have re-written the section highlighted by the reviewer to better explain the transition into exercise-related sensations and cognitive function.

P5 L47-P6 L51 – ... to <u>maintain</u> exercise-related sensations <u>contributing</u> to RPE (16) in hypoxia and normoxia. <u>Cyling continuously for 10 min at a fixed-intensity (corresponding</u>

to 50% VO_{2Max}) in hypoxia *versus* normoxia negatively impacts cognitive function (17). Slower self-selected running velocities may assist <u>with mitigating</u> hypoxic-induced negative cognitive function compared to normoxia (18).

Brocherie, F., Millet, G. P., & Girard, O. (2017). Psychophysiological responses to repeatedsprint training in normobaric hypoxia and normoxia. International Journal of Sports Physiology and Performance, 12(1), 115-123.

McMorris, T., Hale, B. J., Barwood, M., Costello, J., & Corbett, J. (2017). Effect of acute hypoxia on cognition: A systematic review and meta-regression analysis. Neuroscience & Biobehavioral Reviews, 74, 225-232.

<u>Reviewer</u>: Please list a rationale for this study aim—why does this really matter or to whom do these data apply? Is this for training purposes or something else?

<u>Authors' response</u>: We thank the reviewer for suggesting to list a short rationale after the aim of the study. We have subsequently added this into the manuscript which can also be found below.

P6 L59-61 – <u>Decreasing external load with matched internal load during perceptually-</u> regulated HIIT in hypoxia compared to normoxia may benefit athletes during heavy <u>training blocks prior to competition.</u>

<u>Reviewer</u>: Is it wise to have only 3 women in the study? Please comment.

Authors' response: In response to the reviewers comment, our aim was not to investigate the effect of gender in response to perceptually-regulated interval running in hypoxia and normoxia. Rather our intention was to investigate the effect of perceptually-regulated interval running in hypoxia and normoxia on adjustments in running velocity and aoociated exerciserelated sensations of trained runners. During our recruitment period, three eligible females volunteered for the study who matched the inclusion criteria (P6 L67-69). We have calculated groups means \pm SD and re-analysed the primary outcomes of our study (velocity, heart rate, SpO_2 and exercise-related sensations) only with the 16 male participants (excluding the three females). As shown at the bottom of this response document, this analysis indicates that including only three females does not lead to different groups means \pm SD, probability values and effect sizes compared to a dataset of 16 males. Consequently, including the 3 female participants in our final sample of 19 participants does not change the overall message of the study. Therefore, we believe that keeping the three females within the current participant total (n = 19) is warranted since our power calculation indicated 21 participants are required to yield sufficient power in the statistical tests carried out (P6 L73-P7 L76). This would not be achieved if the data from the three females were removed (n = 16).

Reviewer: Is there a reason why a VO2max test was not undertaken?

<u>Authors' response</u>: We thank the reviewer for raising this comment. Although identifying and presenting VO_{2MAX} values of the participants in our study would have been informative, we didn't feel it was necessary. This is because we used a modified, and validated, method (Martin et al., 1992) to determine the individual velocity required for each individual to run at an RPE of 16, which is presented (P17 L268-269) and the main purpose of our work. As such, of the ceiling value (VO_{2MAX}) did not need to be identified in the current study.

Martin, P. E., Rothstein, D. E., Larish, D. D. Effects of age and physical activity status on the speed-aerobic demand relationship of walking. J Appl Phys 1992;73:200–206.

<u>Reviewer</u>: Were participants informed about the study aim e.g. to test changes in your measures in hypoxia vs. normoxia?

<u>Authors' response</u>: We appreciate the reviewer raising this important comment. Prior to enrolling onto the study, participants were informed that they will perform two HIIT protocols, randomly in hypoxic and normoxic conditions. We explained the measures that we would be performing, but did not directly outline that we would be looking at the comparison between hypoxic and normoxic conditions to the participants. Naturally, this may have impacted their perceptions or lead to social desirability bias in response to perceptual scales and assessment of attention and executive function. Further, as outlined in the manuscript (P9 L132-135), participants were blinded to the environmental condition during the HIIT protocol as we removed the hypoxic generator from their view, and simulated 100 m (machine switched on) during normoxia. We have subsequently added in some further information regarding this comment to the manuscript as outlined below.

P7 L86-88 – <u>To minimise the impact of social desirability bias, participants were made</u> <u>aware of the purpose of the study but were naïve to experimental hypotheses.</u>

Adams, S. A., Matthews, C. E., Ebbeling, C. B., Moore, C. G., Cunningham, J. E., Fulton, J., & Hebert, J. R. The effect of social desirability and social approval on self-reports of physical activity. Am J Epidem, 2005;161(4):389-398.

<u>Reviewer</u>: I think the Results would be better structured by describing the fidelity of HIIT first, by denoting the HR and velocity data. This confirms that these bouts actually represent HIIT based on 85 %HRmax. Then, follow this with your other outcomes.

<u>Authors' response</u>: We acknowledge and agree with the reviewer that the results structure would be better when beginning the first section with velocity and heart rate data. The heading of this section has been renamed, and only includes velocity and heart rate data.

P12 L190 - Changes in velocity and HR

<u>Reviewer</u>: I also think that the Results text needs better subheadings to clarify its organization e.g. change in HR and running velocity in response to..; change in muscle deoxygenation in response to, etc.

<u>Authors' response</u>: We appreciate the reviewer suggesting better subheadings within the results section. These have subsequently been renamed as below.

P12 L196 - Changes in SpO2 and muscle oxygenation

- P12 L201 Changes in exercise-related sensations
- P13 L222 <u>Changes in [La⁺] and attention and executive function</u>

Reviewer: Line 186 needs a clearer subheading e.g. change in recovery and .. in response to..

Authors' response: Please see response above regarding this comment.

<u>Reviewer</u>: Line 244, Discussion; so what is the importance of these findings to the athlete, coach, clinician, etc.? Please cite this here.

<u>Authors' response</u>: We acknowledge the reviewers comment that the importance of our findings should be stated here for those in an applied setting. We have subsequently added this information in to the manuscript, which can also be found below.

P17 L261-266 – <u>A matched internal workload for a decreased external workload during</u> perceptually-regulated HIIT in hypoxia *versus* normoxia may assist athletes to reach intended session goals with minimal over-induced physiological stress. However, perceptually-regulated HIIT exacerbates exercise-related sensations and blood lactate concentrations in hypoxia compared to normoxia. This may then have negative carry-over effects on training responsiveness in the following days.

Reviewer: Line 269; so what is the importance or application of this particular finding?

<u>Authors' response</u>: We understand the reviewers comment that this take home message can be further developed for importance and application. We have subsequently added this information in to the manuscript, which can also be found below.

P18 L291-294 – <u>This finding may be of benefit to athletes who are unable or advised by</u> their coach not to be training at a full intensity. Completing perceptually-regulated HIIT in hypoxia that requires slower running velocities compared to normoxia may in turn minimise mechanical constraints and eventually injury risk.

<u>Reviewer</u>: Line 302: what do you mean by 'metabolic by product?' this is blood lactate so just represent this as is to be most clear. And why does this excess accumulation of BLa matter to the scientist or athlete/practitioner?

<u>Authors' response</u>: We thank the reviewer for suggesting more clarity regarding the take home message of this paragraph in the discussion, and have replaced '*metabolic by-product*' with ' $[La^+]$ '. We have also added in a sentence regarding the implications of this finding in an applied setting.

P19 L327 – ... to increased $[La^+]$ at slower...

P19 L328-P20 L332 – <u>Practitioners should be aware that perceptually-regulated HIIT in</u> hypoxia is a viable method for matching indices of physiological stress to normoxia. <u>However, the blood lactate concentration increases after exercise were larger in hypoxia</u> <u>compared to normoxia. This may have negative implications on the muscle fatigue</u> <u>recovery process.</u>

<u>Reviewer</u>: Lines 307-15: are there any data showing that hypoxia reduces O2 delivery to the brain which then may alter perceptions of exercise which are regulated by various brain centers? Please comment on this.

<u>Authors' response</u>: We thank the reviewer for raising this interesting point. Accordingly, we have added some information from the work of Subudhi et al. (2007 & 2009) relating to the negative implication hypoxia has on oxygen delivery to the brain, potentially altering perceptions of exercise. We also believe that this point should be of consideration for further research investigations, attempting to capture further measurements in response to HIIT in hypoxia at a perceptually-regulated intensity compared to normoxia.

P20 L342-348 – <u>Further, it could be postulated that cerebral deoxygenation was greater</u> <u>during HIIT in hypoxia versus normoxia, as demonstrated by Subudhi et al. during</u>

422	
423	incremental cycling (40, 41). Accordingly, cerebral deoxygenation during HIIT may
424	
425	contribute to an integrative decision regarding negative perceptions, in which hypoxia
426	hastens this effect (41). Given that the perceptually-regulated exercise model is governed
427	centrally, this may provide a potential explanation as to why exercise-related sensations
428	were more elevated in the hypoxic trial.
429	
430	P28 L522-524 – Subudhi, A. W., Dimmen, A. C., & Roach, R. C. Effects of acute hypoxia
431	on cerebral and muscle oxygenation during incremental exercise. J App Phys.
432 433	
433	<u>2007;103(1):177-183.</u>
434	
436	P28 L525-527 – Subudhi, A. W., Miramon, B. R., Granger, M. E., & Roach, R. C. Frontal
430	and motor cortex oxygenation during maximal exercise in normoxia and hypoxia. J App
438	Phys. 2009;106(4):1153-1158.
439	
440	
441	
442	
443	<u>Reviewer</u> : This text seems to be way too much of a 'stretch' from your particular study and
444	should be removed, as it is not relevant; thank you. Athletes and clinical patients awaiting or
445	shortly proceeding surgery may exercise in hypoxia to increase the internal workload similar
446	to that achieved in normoxia for a lower external workload. This could decrease joint pain of
447	the lower extremities during ambulation.
448	the lower extremities during amoutation.
449	
450	Authors' response: We understand the reviewers concern regarding some of the statements
451	presented in the limitations and perspectives section. We have since removed the points raised
452	by the reviewer.
453	
454	
455	
456	
457	
458	
459	
460	
461	
462	
463 464	
465	
466	
400	
468	
469	
470	
471	
472	
473	
474	
475	
476	
477	
478	8
479	
480	

Reviewer 2

 <u>Reviewer</u>: This study was primarily aimed at investigating potential differences in running velocities between perceptually-regulated high-intensity intervals in hypoxia and normoxia. It has been demonstrated that running velocity progressively decreased from interval 1 to 4, more pronounced in hypoxic conditions. Negative exercise-related sensations increased over time, again more pronounced in hypoxia.

The authors deal with an interesting topic especially from a training-practical point of view however, findings are not really unexpected and novelty of this study should be more highlighted. Methods are well and reproducible described and results are nicely presented. Nevertheless, the authors may respond to the following comments:

<u>Authors' response</u>: We thank the reviewer for their careful consideration of our work. We have responded to the comments individually raised below.

<u>Reviewer</u>: In the introduction session, you may refer to studies demonstrating differences between RPE and cardiorespiratory responses in hypoxia/altitude compared to normoxia.

<u>Authors' response</u>: We thank the reviewer for raising our attention to this point and have added in this information to the introduction section as outlined below.

P5 L36-38 – <u>Although HR was similar between conditions</u>, <u>RPE has been reported to be higher in hypoxia compared to normoxia during fixed-intensity interval runs (5) and repeated-sprint cycling (12).</u>

P26 L443-445 – <u>Brocherie, F., Millet, G. P., & Girard, O. Psychophysiological responses</u> to repeated-sprint training in normobaric hypoxia and normoxia. International J of <u>Sports Phys Perform. 2017;12(1):115-123.</u>

<u>Reviewer</u>: Nineteen experienced runners have been recruited. As you might expect, I am not happy with the inclusion of only 3 females. Such a sex distribution may reduce the conclusiveness of the findings and does not allow to analyse potential sex differences.

<u>Authors' response</u>: We understand the reviewers concern regarding the small population of females within our participant cohort. We would like to highlight that within the literature, it is inconclusive as to whether gender distributions impact on acute responses to hypoxia – with some confirming (Lombardi et al., 2013; Mortola & Saiki, 1996) and others rejecting (Loeppky et al., 2001; Sandoval & Matt, 2002) this hypothesis.

In response to the reviewers comment, our aim was not to investigate the effect of gender in response to perceptually-regulated interval running in hypoxia and normoxia. Our aim was to investigate the effect of perceptually-regulated interval running in hypoxia and normoxia on trained runners. During our recruitment period, three eligible females volunteered for the study who matched the inclusion criteria (P6 L67-69). We have calculated groups means \pm SD and re-analysed the main findings of our study (velocity, heart rate, SpO₂ and exercise-related sensations) without the presence of the three females (located at the bottom of this response document) which does not change the overall message of the study. Therefore, we believe that keeping the three females within the current participant total (n = 19) is warranted since our power calculation indicated 21 participants are requied to yield suffificent power in the statistical tests carried out (P6 L73-P7 L76), which would not be achieved if the data from the three females were removed (n = 16).

 We do believe that the reviewer raises an important comment, and have since highlighted this in the limitations and perspectives section of the paper for future investigations.

P22 L384-387 – <u>In addition, whether there are gender differences in response to hypoxic</u> <u>exposure during perceptually-regulated HIIT should be investigated, given that our final</u> <u>sample size (n = 19) included only three females.</u>

Loeppky, J. A., Scotto, P., Charlton, G. C., Gates, L., Icenogle, M., & Roach, R. C. (2001). Ventilation is greater in women than men, but the increase during acute altitude hypoxia is the same. Respiration physiology, 125(3), 225-237.

Lombardi, C., Meriggi, P., Agostoni, P., Faini, A., Bilo, G., Revera, M., ... & Gregorini, F. (2013). High-altitude hypoxia and periodic breathing during sleep: gender-related differences. Journal of sleep research, 22(3), 322-330.

Mortola, J. P., & Saiki, C. (1996). Ventilatory response to hypoxia in rats: gender differences. Respiration physiology, 106(1), 21-34.

Sandoval, D. A., & Matt, K. S. (2002). Gender differences in the endocrine and metabolic responses to hypoxic exercise. Journal of Applied Physiology, 92(2), 504-512.

Reviewer: Did you perform some type of power calculation? (at least a-posteriori).

<u>Authors' response</u>: We thank the reviewer for raising this comment. We did indeed carry out a power analysis to determine the number of participants required for sufficient power in our results. Information regarding this has been added to the manuscript, as found below.

P6 L72-P7 L78 – <u>A-priori sample size was calculated using G*Power (Version 3.1.9.3)</u>. This was determined using published power output data by Jeffries et al. (19), whereby healthy individuals cycled at a perceptually-regulated intensity (RPE = 16) in hypoxic ($FiO_2 = 15.0\%$) and normoxic conditions. Twenty-one participants were deemed sufficient to yield a power of 0.8 at an α probability of 0.05. Two individuals dropped out due to injuries sustained during their time enrolled onto the study, not associated with the HIIT protocols we employed.

P26 L467-469 – <u>Jeffries, O., Patterson, S. D., & Waldron, M. The effect of severe and</u> moderate hypoxia on exercise at a fixed level of perceived exertion. Euro J App Phys.

<u>2019;1-12.</u>

<u>Reviewer</u>: What means experienced runners? Can you report race times, VO2max, etc.?

<u>Authors' response</u>: We appreciate the reviewers comment regarding our use of the term 'experienced runners'. As we did not collect information regarding race times and race history nor did we assess VO_{2MAX} , we have replaced this term with 'trained runners' throughout the manuscript. Further, the runners we recruited included those with middle and long distance backgrounds. As such, it would be difficult to estimate or predict one particular race distance (i.e., 10 km) for all participants if they had not competed in this distance previously. We believe the term 'trained' better reflects the demographic of individuals recruited for the study, whom may have a knowledge of different race times and race history, but we have no quantifiable data to present.

<u>Reviewer</u>: Please, provide a table depicting more characteristics of study participants, beside anthropometric data, e.g. also performance parameters, regular physical sports/exercise activity, medical history, coffee and/or alcohol drinking, smoking, medications, etc.

<u>Authors' response</u>: We appreciate the reviewer suggesting to add further characteristics of the participants who completed the study protocol. Prior to enrolling onto the study, we screened volunteers (using a health questionnaire) for their medical history. As described in the methods section (P6 L66-69), we only included individuals who were free of clinical signs of disease, orthopedic, neurological, cardiovascular or respiratory problems. Further, runners were recruited who trained for ≥ 6 h/wk. Therefore, we are unable to provide any further information regarding the characteristics of participants as this was not collected, but re-assure the reviewer that they were of a healthy status.

<u>Reviewer</u>: The limitation section should be extended and the statements that "athletes and clinical patients awaiting or shortly proceeding surgery may exercise in hypoxia to increase the internal workload similar to that achieved in normoxia for a lower external workload" and "this could decrease joint pain of the lower extremities during ambulation" are in my opinion not justified based on the presented findings and should rather be deleted.

<u>Authors' response</u>: We understand the reviewers concern regarding some of the statements presented in the limitations and perspectives section. We have since removed the points raised by the reviewer, and added in sentences regarding the need to investigate gender differences in response to perceptually-regulated HIIT in hypoxia.

P22 L384-387 – In addition, whether there are gender differences in response to hypoxic

exposure during perceptually-regulated HIIT should be investigated, given that our final

sample size (n = 19) included only three females.

<u>Reviewer</u>: The conclusion might a bit more focus on the training-practical importance of the findings.

<u>Authors' response</u>: We thank the reviewer for raising this important point, and have subsequently added a sentence within the conclusion section of our manuscript highlighting the practical considerations of our findings.

P22 L396-401 – <u>Our results suggest that athletes under the influence of hypoxia require</u> lower external workloads to reach a perceptually-regulated target during HIIT than normoxia. If employed in a practical setting, coaches should consider the potential of negatively implicated exercise-related sensations and blood lactate concentrations which may have further negative carry-over effects on training responsiveness in the following days.

Primary outcome measures processed and analysed with males and females (n = 19) and males only (n = 16).

Statistical analysis presented here used the methods presented in the manuscript. A two-way ANOVA investigated the main effect of condition, time and interaction. Group means \pm SD, *p* values and effects sizes are presented for velocity, heart rate, SpO₂ and exercise-related sensations.

		<u>Males &</u>	<u>x females</u>				<u>Male</u>	<u>sonly</u>		
Measure		Con	dition		p value (effect size)		Con	dition		p value (effect siz
	Н	YP	Ν	OR	Condition	Н	YP	Ν	Or	Condition:
	13.9	± 0.6	14.7	± 0.3	p < 0.01 (0.91)	14.1	± 0.7	14.9	± 0.3	p < 0.01 (0.92
Velocity		inte	erva		Time		Int	erval		Time:
(km/h-1)	1	2	3	4	p < 0.01 (0.65)	1	2	3	4	p < 0.01 (0.66
	14.8 ± 0.2	14.4 ± 0.5	14.1 ± 0.7	13.8 ± 0.8	Interaction:	15.1 ± 0.2	14.6 ± 0.4	14.2 ± 0.6	14.0 ± 0.8	Interaction:
					p < 0.01 (0.42)					p < 0.01 (0.45
	н	ΥP	Ν	OR	Condition	н	YP	Ν	OR	Condition:
	170 ± 3		169 ± 4		p = 0.65 (0.01)	169 ± 3		168 ±4		p = 0.53 (0.03
Heart rate			erval		Time			nterval		Time:
(bpm)	1	2	3	4	p < 0.01 (0.62)	1	2	3	4	p < 0.01 (0.66
	165 ± 1	169 ± 0	171 ± 0	173 ±1	Interaction:	164 ± 1	168 ± 1	170 ± 0	173 ± 1	Interaction:
					p = 0.28 (0.07)					p = 0.23 (0.09
	н	YP	Ν	OR	Condition:	н	YP	Ν	OR	Condition:
	86 ± 0		95 ± 0		p < 0.01 (0.99)	86	86 ± 0		± 0	p < 0.01 (0.99
SpO2		inte	erval		Time:		int	erval		Time:
(%)	1	2	3	4	p = 0.37 (0.06)	1	2	3	4	p = 0.51 (0.50
	90 ± 6	91 ± 6	90 ± 6	90 ± 7	Interaction:	90 ± 5	91 ± 6	90 ± 6	90 ± 7	Interaction:
					p = 0.17 (0.09)					p = 0.14 (0.12
	Н	YP	Ν	OR	Condition:	Н	YP	Ν	OR	Condition:
	6 :	± 2	7	± 1	p < 0.01 (0.61)	6 :	± 2	7	± 1	p < 0.01 (0.61
Recovery		Inte	erval		Time:		Int	erval		Time:
(au)	1	2	3	4	p < 0.01 (0.81)	1	2	3	4	p < 0.01 (0.80
	9 ± 0	7 ± 0	6 ± 1	5 ± 1	Interaction:	9 ± 0	7 ± 0	6 ± 1	5 ± 1	Interaction:
					p < 0.01 (0.24)					p < 0.01 (0.35
	н	YP	Ν	OR	Condition:	н	YP	Ν	or	Condition:
	13 ± 2		14 ± 1		p < 0.01 (0.65)	12 ± 3		14 ± 1		p < 0.01 (0.71
Motivation		Inte	erval		Time:		Int	erval		Time:
(au)	1	2	3	4	p < 0.01 (0.50)	1	2	3	4	p < 0.01 (0.52
	15 ± 0	14 ± 1	12 ± 2	12 ± 2	Interaction	15 ± 0	14 ± 1	12 ± 2	12 ± 2	Interaction:
					p < 0.01 (0.25)					p < 0.01 (0.28
	HYP		NOR		Condition:	HYP		NOR		Condition:
	9 ± 1		7 ± 1		p < 0.01 (0.56)	9 ± 1		7 ± 1		p < 0.01 (0.61
Breathlessness		Inb	erval		Time:		Int	erval		Time:
(au)	1	2	3	4	p < 0.01 (0.55)	1	2	3	4	p < 0.01 (0.61
	7 ± 1	8 ± 1	8 ± 1	8 ± 1	Interaction:	7 ± 1	8 ± 1	8 ± 1	8 ± 1	Interaction:
					p = 0.80 (0.02)					p = 0.58 (0.04
	н	YP	Ν	OR	Condition:	н	YP	Ν	or	Condition:
Lingh	6 :	± 1	5	± 1	p = 0.02 (0.30)	6 :	± 1	5	± 1	p = 0.04 (0.26
Limb discomfort		int	erval		Time:		int	erval		Time:
	1	2	3	4	p < 0.01 (0.59)	1	2	3	4	p < 0.01 (0.59
(au)	5 ± 1	6 ± 0	6 ± 1	7 ± 0	Interaction:	4 ± 0	6 ± 0	6 ± 1	7 ± 1	Interaction:
					p = 0.78 (0.02)					p = 0.66 (0.59
	HYP		NOR		Condition	HYP		NOR		Condition:
	7 :	± 2	10	± 1	p < 0.01 (0.59)	7 :	± 2	10	± 1	p < 0.01 (0.63
Pleasure		Inte	erval		Time:		int	erval		Time:
(au)	1	2	3	4	p < 0.01 (0.60)	1	2	3	4	p < 0.01 (0.68
	10 ± 1	9 ± 2	8 ± 3	7 ± 3	Interaction:	10 ± 2	9 ± 2	8 ± 3	7 ± 3	Interaction:
					p = 0.07 (0.12)					p = 0.02 (0.19)

1		
2 3 4 5 6	1 2 3	Title Psycho-physiological responses to perceptually-regulated interval runs in hypoxia and normoxia
7 8	4	
9 10 11	5 6	Author names and affiliations Hobbins L ^a , Gaoua N ^a , Hunter S ^a , Girard O ^b
12	7	
13 14 15 16 17 18 19	8 9 10 11 12	 ^aSport and Exercise Science Research Centre (SESRC), London South Bank University, London, United Kingdom ^bMurdoch Applied Sports Science (MASS) Laboratory, Murdoch University, Perth, Australia <u>cAthlete Health and Performance Research Center, ASPETAR, Qatar Orthopedic and</u> <u>Sports Medicine Hospital, Doha, QATAR</u>
20 21	13	
	13 14 15 16 17 18 19 20 21	Corresponding author Mr. Liam Hobbins hobbinsl@lsbu.ac.uk Sport and Exercise Science Research Centre London South Bank University 103 Borough Road London SE1 0AA
54 55 56 57 58		1
59		

1 Abstract

We investigated whether perceptually-regulated high-intensity intervals in hypoxia are associated with slower running velocities versus normoxia, when physiological responses and exercise-related sensations remain the same. Nineteen trained runners $(33.4 \pm 9.1 \text{ years})$ completed a high-intensity interval running protocol (4 \times 4-min intervals at a clamped perceived rating exertion of 16 on the 6–20 Borg scale, 3-min passive recoveries) in either hypoxic (HYP; FiO₂ 15.0%) or normoxic (NOR; FiO₂ 20.9%) conditions. Participants adjusted to a progressively slower running velocity from interval 1-4 (-7.0%), and more so in HYP vs. NOR for intervals 2, 3 and 4 (-4.6%, -6.4% and -7.9%, respectively; p < 0.01). Heart rate increased from interval 1–4 (+4.8%; p < 0.01), independent of condition. Arterial oxygen saturation was lower in HYP vs. NOR (86.0% vs. 94.8%; p < 0.01). Oxyhemoglobin (-23.7%) and total hemoglobin (-77.0%) decreased, whilst deoxyhemoglobin increased (+44.9%) from interval 1–4 (p < 0.01), independent of condition. Perceived recovery (-41.6%) and motivation (-21.8%) were progressively lower from interval 1-4, and more so in HYP vs. NOR for intervals 2, 3 and 4 (recovery: -8.8%, -24.2% and -29.3%; motivation: -5.3%, -20.3% and -22.4%, respectively; p < 0.01). Perceived breathlessness (+18.6%), limb discomfort (+44.0%) and pleasure (-32.2%) changed from interval 1-4, with significant differences (+21.8%, +11.3% and -31.3%, respectively) between HYP and NOR (p < 0.01). Slower interval running velocities in hypoxia achieve similar heart rate and muscle oxygenation responses to those observed in normoxia when perceptually-regulated, yet at the expense of less favourable exercise-related sensations.

23	Key words
24	High-intensity intermittent running; normobaric hypoxia; perceptually-regulated exercise;
25	ratings of perceived exertion; near-infrared spectroscopy; effort perception.
	3
	-
	24

Introduction

High-intensity interval training (HIIT) is a popular exercise format in athletic and clinical populations (1,2). HIIT includes repeated short-to-long (2–5 min) intense exercise bouts (80– 90% of the velocity associated with maximal oxygen uptake or vVO_{2Max}) interspersed with shorter (1–3 min) recoveries (3). Compared to moderate-intensity continuous running, HIIT leads to similar improvements in cardiorespiratory fitness that is achieved with a shorter effective exercise duration per session (2). Due to the reduced time-commitment and exercise training volume, investigations surrounding the potential physiological and performance benefits of **HIIT** have surged (4).

HIIT in normobaric hypoxia (a lower inspired oxygen fraction or FiO₂) is receiving attention for its potential in further advancing athletic performance compared to HIIT in normoxia. Buchheit et al. (5) employed a HIIT protocol (3×5 -min, 90-s recovery) carried out in hypoxia (vVO_{2Max} = 84%; FiO₂ = 15.4%) and normoxia (vVO_{2MAX} = 90%) at a fixed-intensity (determined in normoxia) in highly-trained runners. A reduced physiological stress (i.e., lower heart rate or HR) was observed during hypoxia compared to normoxia, likely due to a lower vVO_{2Max} in hypoxia versus normoxia. However, fixed exercise intensities, regardless of environmental conditions, do not permit adjustments (i.e., increases or decreases of workload) during exercise to match the intensity target (i.e., vVO_{2MAX}). In turn, over-induced physiological stress may be counter-productive (i.e., greater deoxygenated muscle heamoglobin, lower oxygenated haemoglobin) for intended session goals (6). Furthermore, matched absolute fixed exercise intensities (i.e., a similar percentage of vVO_{2MAX}) lead to greater physiological stress (i.e., compensatory increase in HR) in hypoxia compared to normoxia due to reduced FiO₂ (7). Perceptually-regulated exercise intensities, that allow velocity adjustments based upon exercise-related sensations in order to maintain a target effort

25 level, may offer a viable solution, and is perhaps more reflective of how exercisers modify 26 <u>intensity during acute exercise</u>.

Perceptually-regulated exercise permits the individual exercising to self-regulate external workload (i.e., running velocity/cycling power production) based upon Borg's rating of perceived exertion (RPE) scale (8). The validity and usefulness of using RPE for perceptually-regulating exercise has been described (9). The reduced oxygen availability in hypoxia makes the expectation tenable that there would be a slower self-selected running velocity in hypoxia for a given RPE, while velocity in normoxia would be more preserved, as evidenced previously (10). Chacaroun et al. (11) demonstrated for a lower power output (-15%), vastus lateralis muscle deoxyhemoglobin was higher and oxyhemoglobin lower in hypoxia (FiO₂ = 13.5%) compared to normoxia during a single interval session (15×1 -min at 75% of maximal HR, 1-min recoveries). Although HR was similar between conditions, RPE has been reported to be higher in hypoxia compared to normoxia during fixed-intensity interval runs (5) and repeated-sprint cycling (12). Employing self-paced exercise, in replace of fixed-intensity exercise, may assist in overcoming the over-excessive physiological stress observed when exercising in hypoxia versus normoxia, due to the likelihood of greater velocity preservations in the latter than the former.

In normoxia at pre-determined fixed intensities, HIIT is perceived as more enjoyable compared to moderate-intensity continuous running (13). However, during HIIT at fixed-intensities, exercise-related sensations decrease when the exercise intensity rises above threshold preference (14). Further, **HIIT** in hypoxia at fixed-intensities typically surpasses the preferred threshold in normoxia (15). Implementing a self-paced exercise model may permit modifications required (i.e., slower running velocities) to maintain exercise-related sensations contributing to RPE (16) in hypoxia and normoxia. Cyling continuously for 10 min at a fixed-intensity (corresponding to 50% VO_{2Max}) in hypoxia versus normoxia negatively

50 <u>impacts cognitive function (17).</u> Slower self-selected running velocities may assist <u>with</u>
 51 <u>mitigating</u> hypoxic-induced negative cognitive function compared to normoxia (18). These
 52 potential findings may benefit athletes exercising intensely in hypoxia, shortly followed by
 53 skills requiring attention and accuracy.

Therefore, the aim of this study was to investigate the effect of HIIT at a clamped RPE of 16 (typically used by athletes during HIIT) (19) in hypoxia and normoxia on adjustments in running velocity and associated exercise-related sensations of trained runners. We hypothesized that running velocity would be progressively slower in hypoxia compared to normoxia across intervals, whilst physiological and cognitive responses, and exercise-related sensations would not differ between conditions. Decreasing external load with matched internal load during perceptually-regulated HIIT in hypoxia compared to normoxia may benefit athletes during heavy training blocks prior to competition.

63 Methods

64 Participants

Nineteen trained runners (3 females, 16 males; age: 33.4 ± 9.1 years; height: 176 ± 88 cm; weight: 76.3 ± 10.9 kg) provided written informed consent to participate. Participants had no musculoskeletal injuries and met the following eligibility criteria: a training volume >6 h/wk. free of clinical signs of disease, orthopedic, neurological, cardiovascular or respiratory problems, and no hypoxic exposure >2000 m for >48 h 6 months before the study. The study was carried out in accordance with the Declaration of Helsinki and was approved by the Ethics Committee of the Anti-Doping Lab Qatar institutional review board (Agreement SCH-ADL-170). A-priori sample size was calculated using G*Power (Version 3.1.9.3). This was

73 determined using published power output data by Jeffries et al. (20), whereby healthy 74 individuals cycled at a perceptually-regulated intensity (RPE = 16) in hypoxic (FiO₂ = 75 15.0%) and normoxic conditions. Twenty-one participants were deemed sufficient to 76 yield a power of 0.8 at an α probability of 0.05. Two individuals dropped out due to 77 injuries sustained during their time enrolled onto the study, not associated with the HIIT 78 protocols we employed.

79 Experimental design

Participants reported to the laboratory on three occasions, each separated by >48 h. The first session included study familiarisation. The second and third visits included completing a HIIT protocol in either hypoxia or normoxia in a randomized, conterbalanced order. Physiological, perceptual and cognitive responses were assessed continuously, immediately before and after each interval, and before and after the **HIIT** protocol, respectively. Participants were instructed to refrain from any intense exercise 48 h prior to each visit and consume their last meal at least 2 h prior to the HIIT sessions. To minimise the impact of social desirability bias, participants were made aware of the purpose of the study but were naïve to experimental hypotheses. Laboratory conditions were similar throughout all sessions (mean temperature \sim 22°C, relative humidity \sim 50%) and time of day was standardized for each participant.

Familiarization session

At the preliminary visit to the laboratory, participants were familiarised with the perceptual scales and cognitive test. Preferred running velocity (PRV) was determined for each participant in normoxia using a modified version of identifying preferred walking speed (21). After a 5min warm up at 10 km/h⁻¹, participants completed four ramped treadmill runs (increasing and decreasing velocities) on an instrumented treadmill (ADAL3D-WR, Medical Development– HEF Tecmachine, France). After every 20 s per ramp, participants rated their RPE of the current velocity (controlled by the investigator and out of sight of the participant) in accordance with Borg's (20) 6 ("no exertion at all") – 20 ("maximal exertion") numeric scale. Ramp one started at 10 km/h⁻¹, increasing by 0.8 km/h⁻¹ every 20 s until the velocity was considered as RPE \geq 18; ramp two started at +1.5 km/h⁻¹ the previous end velocity, decreasing by 0.8 km/h⁻¹ until the velocity was considered as RPE ≤ 12 ; ramp three started at the velocity considered as an RPE of 14 in ramp two, increasing by 0.5 km/h⁻¹ until the velocity was considerd as RPE \geq 18; and ramp four started at +1.0 km/h⁻¹ the previous end velocity, decreasing by 0.5 km/h⁻¹ until the velocity was considered as RPE ≤ 12 . Ramps two, three and four began once the participants declared their perceived recovery level as a 7 out of 10 following the previous ramp (23). HR was recorded every 20-s through each ramp. PRV corresponded to the velocity participants considered as a RPE of 16 (between "hard" and "very hard") or closest to a HR of 160 bpm. After 10 min of rest, participants completed one 4-min interval composing the HIIT protocol (see below) for habituation.

110 *Experimental trials*446

Participants completed two experimental trials in normoxia (NOR; $FiO_2 = 20.9\%$) and hypoxia (HYP; FiO₂ = 15.0%, equivalent to \sim 2700 m above sea level). After a standardised warm up (5-min at 10 km/h⁻¹), a facemask connected to a portable hypoxic generator (See *Hypoxic* simulation section) was attached. Participants rested for 1-min (quiet standing) before a 1-min run at their PRV (RPE = 16). Participants then rested for 3 min before completing the HIIT protocol. The **HIIT** protocol was based upon aerobic interval-training (2). Participants completed four, 4-min intervals, interspersed with 3-min recoveries (quiet standing). The first 30 s of each 4-min interval began at participants' PRV; participants were then free to decide if or how treadmill velocity needed to be adjusted (manually by one experimenter) to ensure maintenance of a RPE of 16 every 30 s. Participants hand-signalled in response to the current

velocity (finger up to increase, finger down to decrease, and circle using index finger and thumb to maintain); and signalled again to inform how much of an increase/decrease in velocity is required [1, 2 or 3 fingers up (faster) or down (slower) for 0.5, 1.0 or 1.5 km/h⁻¹ changes, respectively]. Signals were trialled during familiarisation. Mild verbal encouragement to keep running at an RPE of 16 was used throughout HIIT. Total hypoxic exposure corresponded to exactly 28 min.

127 *Hypoxic simulation*

Participants were fitted with a facemask fastened with a Velcro headset connected via plastic tubing to a hypoxic generator (Altitrainer, SMTec SA, Nyon, Switzerland) to simulate hypoxia. The gas mixing system enriches inspired air by adding a fixed quantity of nitrogen via a 30-L mixing chamber, with the dilution being constantly controlled by a PO_2 probe (precision = T0.82 torr, safety $FiO_2 = 9.7\%$). The hypoxic generator was hidden from participant viewing to ensure condition blinding. When breathing 'normal air' during normoxia, the hypoxic generator was on (for background noise) and set at a simulated altitude of 100 m to increase the strength of blinding.

- **136** *Measures*
- **137** *Exercise intervals*

HR was monitored telemetrically with a Polar transmitter-receiver (Polar S810, Kempele, Finland) and recorded 20 s before and every 30 s during each interval. Arterial oxygen saturation (SpO₂) was assessed via finger pulse oximetery (Palmsat 2500, NONIN Medical Inc., Plymouth, MI, USA) at the same time intervals. HR and SpO₂ were obtained before (i.e., after a 2-min seated period) and at the end of the warm-up procedure (i.e., prior to HIIT). Both

the HR watch (RS400, Polar) and oximeter receiver were attached on the handrails of the
 treadmill outside of the participants' view.

Muscle oxygenation trends of the right vastus lateralis muscle were recorded using near-infrared spectroscopy (NIRS; Portalite, Artinis, Netherlands) in real-time. A wireless bi-polar optode sensor was attached (~10 cm above the proximal patella border) and secured to skin via adhesive tape. Sampling frequency was set at 10 Hz (11) following a 'zero set' of all signals. Bandages were fastened around the lower limb and optode to prevent external light distorting readings. Oxy- (Δ ; [O₂Hb]), deoxy- (Δ ; [HHb]) and total haemoglobin (µmol; [tHb]) were exported (1 Hz). For analysis, each interval was averaged and normalized to a 10 s sample prior to interval one (reference value) for each respective condition and presented as percentage change.

During recovery

Perceived recovery and motivation to exercise were assessed 30 s before each interval. Perceived recovery was assessed by answering 'how recovered do you feel currently?' via a numeric scale, ranging from 0 ("very poorly recovered") to 10 ("very well recovered") (23). Recovery was assessed before interval one to determine perceptions following the warm up. Perceived motivation to exercise was assessed via a 20-cm visual analog scale (24). Participants were asked 'how motivated do you feel to exercise right now?' and answered by adjusting the level on the scale between 0 ("*extremely low*"; white colored) and 20 ("*extremely* high"; black colored). Immediately after each interval, ratings of perceived breathlessness, limb discomfort and pleasure were assessed. Perceived breathlessness was assessed by answering 'how does your breathing feel currently?' via a numeric scale, ranging from 0 ("nothing at all") to 10 ("very, very severe") (25). Using the same scale, perceived limb discomfort was assessed by answering 'how do your legs feel currently?'. A 20-cm visual analog scale (same

as motivation above) was used to assess 'how pleasant was that run?' ranging from 0 ("extremely unpleasant") and 20 ("extremely pleasant").

Pre- and post-exercise

A capillary blood sample taken from the fingertip was analyzed for blood lactate concentration ([La⁺]) with the Lactate Pro (LT-1710, Arkray, Japan) portable analyzer before the warm-up and 2 min after HIIT. An offline Stroop colour-word test (26) assessed attention and executive function. Using one hand and as quickly as possible, participants selected the colored key on the keyboard representing the color of the text appearing on the screen (red, vellow, green or blue). The cognitive test lasted for 3 min, and took place in a silent environment before the warm up and 3 min after **HIIT**. Reaction time (ms; time taken to select a color) and accuracy (%; correct color selected) were averaged over each test for analysis.

Statistical analysis

Data distribution was assessed via a Shapiro-Wilk test. A parametric within-subject two-way analysis of variance was used to investigate the main effect of condition (NOR vs. HYP), time (interval 1, 2, 3 vs. 4 or pre vs. post) and the condition \times time interaction for normally distributed data. Partial eta-squared (n^2) was calculated as a measure of effect size. Values of 0.01, 0.06 and above 0.14 were considered as small, medium and large, respectively (27). A related samples Friedman's non-parametric test was used for data not normally distributed. Bonferroni post-hoc pairwise comparisons were used to identify locations of significant effects. Statistical testing was carried out in SPSS (v21; CED, Cambridge, United Kingdom). Data was considered significant if $p \le 0.05$. All data are presented as group means \pm SD.

- **Results**

<u>Changes in velocity and HR</u>

191 Compared to interval 1, participants adjusted to a progressively slower running velocity during

192 intervals 2, 3 and 4 (-2.8%, -5.2% and -7.0%, respectively; p < 0.01), and more so in HYP vs.

193 NOR for intervals 2, 3 and 4 (-4.6%, -6.4% and -7.9%, respectively; p < 0.01; Figure 1A).

194 Compared to interval 1, HR increased during intervals 2, 3 and 4 (+2.3%, +3.6% and 4.8%,

195 respectively; p < 0.01; Figure 1B), independently of condition (p = 0.65).

196 <u>Changes in SpO₂ and muscle oxygenation</u>

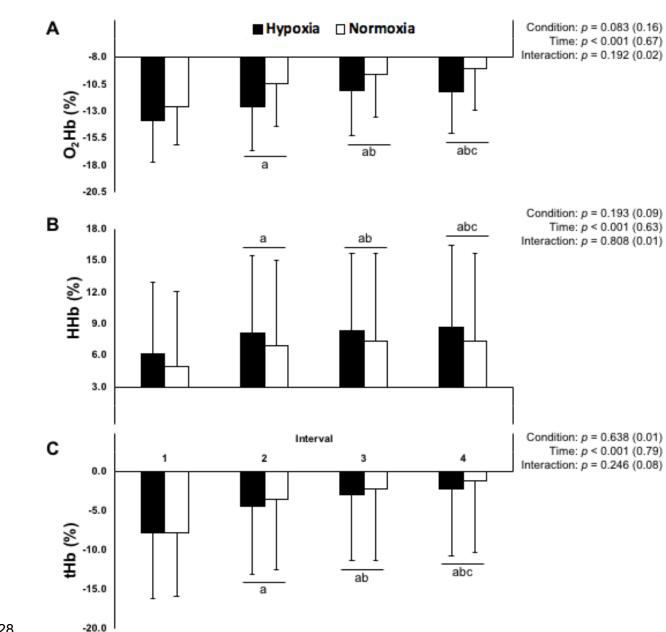
197 SpO₂ was globally lower in HYP *vs.* NOR (-9.3% average across intervals; p < 0.01; Figure 198 1C), independently of time (p = 0.37). From interval 1 to 4, [O₂Hb] and [tHb] decreased (-199 23.7% and -77.0%, respectively) whilst [HHb] increased (+44.9%; p < 0.01; Figures 2A–C), 200 independently of condition (p > 0.08).

9 201 <u>Changes in exercise-related sensations</u>

Perceived recovery decreased progressively from interval 1 to 4 (-41.6%; p < 0.01), and more so in HYP vs. NOR before intervals 2, 3 and 4 (-8.8%, -24.2% and -29.3%, respectively; p =0.02; Figure 3A). Perceived motivation decreased progressively from interval 1 to 4 (-21.8%; p < 0.01), and more so in HYP vs. NOR before intervals 3 and 4 (-20.3% and -22.4%), respectively; p < 0.01; Figure 3B). Compared to interval 1, perceived breathlessness increased following intervals 2, 3 and 4 (+14.0%, +13.6% and +18.6%, respectively; p < 0.01; Figure 3C), independently of condition. Breathlessness was rated globally higher in HYP vs. NOR (+21.8%; p < 0.05), irrespective of time. Compared to interval 1, perceived limb discomfort increased following intervals 2, 3 and 4 (+23.3%, +35.3% and +44.0%, respectively; p < 0.01; Figure 3D), independently of condition. Limb discomfort was rated globally higher in HYP vs. NOR (+11.3%; p = 0.01), irrespective of time. The time-dependent decreases in perceived

pleasure across intervals (-14.7%, -25.4% and -32.3%, intervals 2, 3 and 4 vs. 1, respectively; p < 0.01; Figure 3E) tended to be larger in HYP vs. NOR (-31.3%, p = 0.06). Condition: p < 0.001 (0.91) А Hypoxia Normoxia Time: p < 0.001 (0.65) Interaction: p < 0.001 (0.41) *abc а *ab Velocity (km/h⁻¹) в Condition: p = 0.652 (0.01) Time: p < 0.001 (0.62) abc а Interaction: p = 0.280 (0.07) а Heart rate (bpm) Condition: p < 0.001 (0.99) С Time: p = 0.370 (0.06) Arterial oxygen Interaction: p = 0.170 (0.10) saturation (%) Interval Figure 1 Changes in velocity (A), heart rate (B) and arterial oxygen saturation (C) during the high-intensity intermittent running protocol. Data are presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. * denotes a statistically significant difference between conditions for a given interval (p < 0.05), a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05). Changes in [La⁺] and attention and executive function The pre- to post-exercise increase in [La⁺] was larger (p = 0.001) in HYP ($1.7 \pm 0.8 \text{ vs.} 13.1 \pm$ 3.8 mmol/l⁻¹) vs. NOR (2.1 ± 0.9 vs. 10.1 ± 3.9 mmol/l⁻¹). During the Stropp test, accuracy was

unaffected by condition and time (Figure 4A). Participants' reaction time was faster (+11%) post *vs.* pre **HIIT** (p < 0.01; Figure 4B), independently of condition.



813 228

Figure 2 Changes in Oxygenated (A; O_2Hb), deoxygenated (B; HHb) and total hemoglobin (C; tHb) during the high-intensity intermittent running protocol. Data are calculated as a percentage difference from baseline (%) and presented as mean ± SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05).

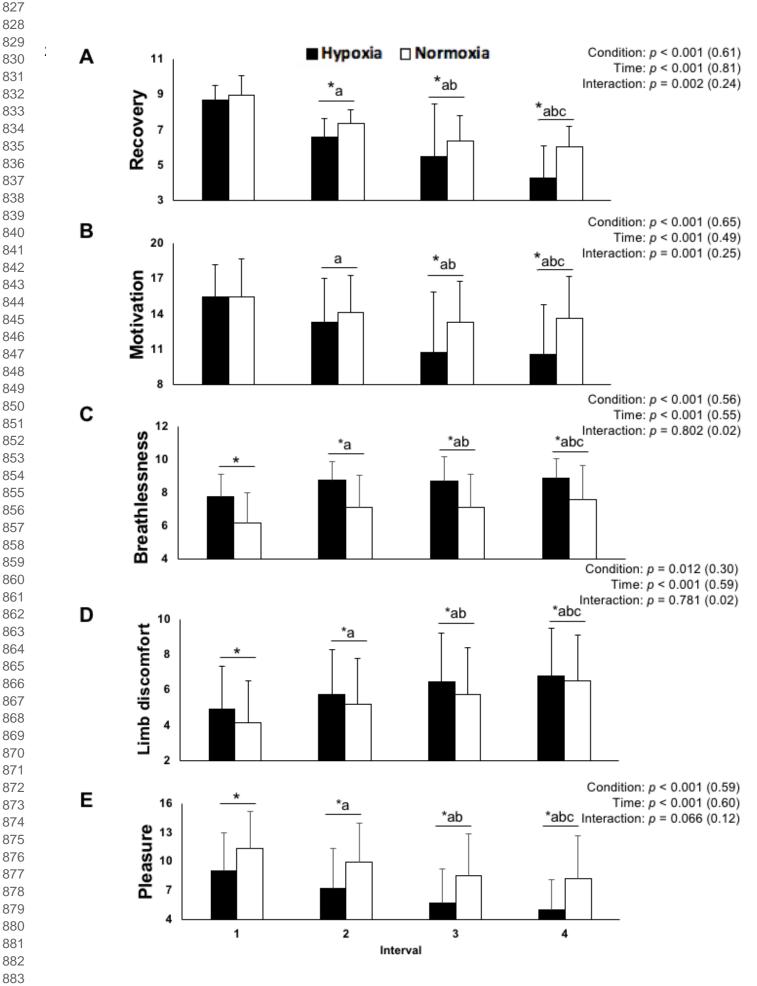


Figure 3 Changes in perceived recovery (A), motivation (B), breathlessness (C), limb discomfort (D) and pleasure (E) during the high-intensity intermittent running protocol. Data are presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. * denotes a statistically significant difference between conditions for a given interval (p < 0.05), a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05).

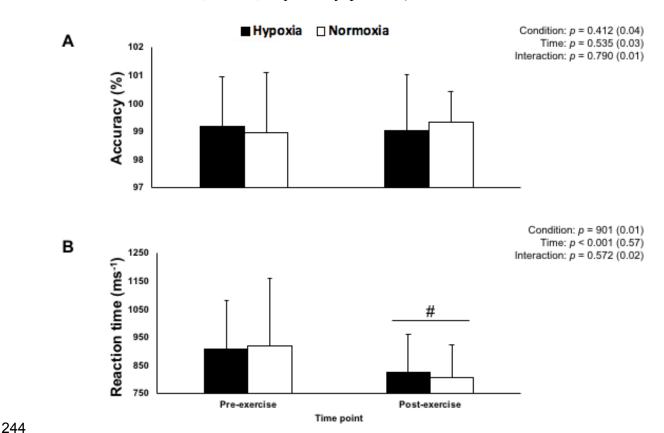


Figure 4 Changes in accuracy (A) and reaction time (B) pre and post high-intensity intermittent running protocol. Data are averaged over 3 mins and presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. # denotes a statistically significant difference vs. pre-exercise (p < 0.01).

Discussion

Using a perceptually-regulated (RPE = 16) exercise model, we observed: 1) participants ran progressively slower during HIIT with larger decreases in HYP versus NOR, 2) HR and muscle oxygenation trends (during intervals) and cognitive responses (pre vs. post HIIT) were similar between conditions, 3) greater breathlessness and limb discomfort, and lower recovery,

motivation and pleasure scores were stated during recovery between HYP vs. NOR, and 4) blood lactate concentration was larger after HYP vs. NOR. Overall, using a manipulation of oxygen availability, reduced external workload (i.e., running velocity) during perceptuallyregulated interval running is associated with a similar internal load (i.e., physiological responses). Although no cognitive function differences were found between conditions, this is achieved with less favourable exercise-related sensations. A matched internal workload for a decreased external workload during perceptually-regulated HIIT in hypoxia versus normoxia may assist athletes to reach intended session goals with minimal over-induced physiological stress. However, perceptually-regulated HIIT exacerbates exercise-related sensations and blood lactate concentrations in hypoxia compared to normoxia. This may then have negative carry-over effects on training responsiveness in the following days.

Exercise intervals

The velocity deemed equal to RPE 16 (PRV) was as expected for trained runners (~15 km/h-¹) (28). Interestingly, running velocity did not differ between conditions during the first HIIT interval, despite lower SpO₂ in hypoxia versus normoxia. Smith & Billaut (29) found maintained SpO₂ during repeated-sprinting in normoxia (20×5 -s all out, 25-s recovery) until after the fifth sprint in national-level soccer players, whereby peak power significantly decreased compared to sprint one. Overall, it seems that initial decreases in SpO₂ (within interval one) do not necessarily impact on HIIT compared to sprint intervals.

We found that participants selected a progressively slower running velocity during **HIIT** in both conditions. In highly-trained middle to long-distance runners, a 6% reduction in vVO_{2MAX} when running in hypoxia versus normoxia is acceptable to match the acute physiological stress induced (5). It can be suggested that self-selected velocity adjustments found in the current study to maintain RPE 16 are matched with modifications in hypoxic versus normoxic traning sessions employed by coaches and sport scientists for athletes (30). Decreased external workloads have been reported by Pramsohler et al. (31) during continuous cycling (seven 30-min sessions over 3-wk), whereby participants cycled at -28% lower power output in hypoxia $(FiO_2 = 15.3\%)$ versus normoxia for a similar HR. Differences in these findings and ours may be due to the inclusion of geriatric patients completing pre-set (in normoxia) fixed-intensity cycling compared to trained runners self-regulating HIIT in the current study. However, Fernández-Menéndez et al. (10) reported preferred walking velocity (RPE of 10) in hypoxia $(FiO_2 = 15.3\%)$ was 7% slower than normoxia in obese adults over 3 weeks. Using a self-paced model, irrespective of RPE target, population demographics and training block duration, lower external workloads are selected in hypoxia compared to normoxia. Overall, decreases in self-paced running velocity occured to a greater extent in hypoxia than normoxia to maintain RPE 16, suggesting of a lower external workload. This finding may be of benefit to athletes who are unable or advised by their coach not to be training at a full intensity. Completing perceptually-regulated HIIT in hypoxia that requires slower running velocities compared to normoxia may in turn minimise mechanical constraints and eventually injury risk.

Our data show HR increased progressively during HIIT, irrespective of condition. This matches our hypothesis that HR will be comparable between hypoxia and normoxia, even though running velocity was lower in hypoxia. Other studies employing moderate continuous-intensity exercise have also found matched HR responses between hypoxic and normoxic training interventions (~4 weeks) when cycling at a -21.0% power output in healthy males (32) and walking/running at a -17.5% velocity in obese adults (33) in hypoxia verus normoxia. Although exercise intensities in these studies were fixed, we believe similar increases in HR between conditions occur due to the environmental stressor (hypoxia) augmenting autonomic cardiac regulation (34). Overall, it seems self-paced exercise in hypoxia provides an added

 304 environmental stressor that is able to mimick HR responses in normoxia for a lower external305 load.

Lower [O₂Hb] and [tHb], and greater [HHb] of the *vastus lateralis* were recorded across HIIT, irrespective of condition. Decreases in [O₂Hb] and increases in [HHb] were expected during **HIIT** as oxygen delivery is outweighed by utilisation, whilst decreases in [tHb] reflect a lower localised blood flow (35). Active musculature oxygenation is negatively impacted during fixed-intensity exercise in hypoxia compared to normoxia due to a lower FiO_2 (7). In support of this, Chacaroun et al. (11) reported lower [O₂Hb] and greater [HHb] with maintained [tHb] of the vastus lateralis during fixed, relative high-intensity cycling in hypoxia (85% maximal power output in normoxia; $FiO_2 = 13.5\%$) versus normoxia. Where we employed a self-paced exercise model, similar [O₂Hb] and [HHb] responses are achieved between conditions. This is likely explained through the decreased workload (i.e., slower running velocity) in hypoxia compared to normoxia, subsequently lowering oxygen utilisation. Discrepants findings in [tHb] may be due to different exercise modalities (cycling versus running) modifying blood flow regulation (36). Similar to HR responses (central) previously discussed, it can be suggested here that local (tissue oxygenation) physiological stress is matched between conditions during **HIIT** in hypoxia at a slower velocity compared with normoxia.

Elevations in [La⁺] following **HIIT** were higher in HYP than NOR. Values in the current study $(10-13 \text{ mmol/l}^{-1})$ are somewhat higher than those $(5-6 \text{ mmol/l}^{-1})$ reported elsewhere following a single HIIT session (6×4 -min intervals at a RPE ~17, 4-min recoveries) (19). This maybe due to a 1:0.75 work:rest ratio implemented during our protocol compared to 1:1 employed by Seiler & Sjursen (19). [La⁺] normalization during shorter recovery periods may not occur to the extent following longer recovery periods due to excess pyruvate accumulation (37). This suggests that HIIT in hypoxia *per se* leads to increased [La⁺] at slower running velocities compared to normoxia for similar physiological stress amounts. Practitioners should be

329 aware that perceptually-regulated HIIT in hypoxia is a viable method for matching
 330 indices of physiological stress to normoxia. However, the blood lactate concentration
 331 increases after exercise were larger in hypoxia compared to normoxia. This may have
 332 negative implications on the muscle fatigue recovery process.

333 During recovery

Perceptual responses to **HIIT** were negatively impacted (i.e., lower recovery, and motivation) when assessed before intervals, with further exacerbations in hypoxia. Participants were instructed to maintain a RPE of 16 throughout HIIT by adjusting their velocity where necessary. It might be surprising at first that perceptual responses were worse in hypoxia compared to normoxia. However, perceived recovery and motivation are important affects associated with exercise intensity regulation (38). Our results indicate that hypoxia negatively impacts these affects during **HIIT** compared with normoxia. This may be explained through lower perceived capabilities of hypoxic HIIT completion over normoxia (39), lowering perceived recovery and motivation. Further, it could be postulated that cerebral deoxygenation was greater during HIIT in hypoxia versus normoxia, as demonstrated by Subudhi et al. during incremental cycling (40, 41). Accordingly, cerebral deoxygenation during HIIT may contribute to an integrative decision regarding negative perceptions, in which hypoxia hastens this effect (41). Given that the perceptually-regulated exercise model is governed centrally, this may provide a potential explanation as to why exerciserelated sensations were more elevated in the hypoxic trial. Overall, our data poses a disconnection between RPE and exercise-related sensations (i.e., recovery and motivation). Further research should look to optimise **HIIT** in hypoxia for positive perceptual responses. Perceptual responses after intervals were negatively impacted (i.e., higher breathlessness and

³ 352 limb discomfort, lower pleasure), and to a further extent in hypoxia than normoxia. Buchheit

et al. (5) reported that 3-min absolute-intenxity running intervals (84% vVO_{2MAX}) in hypoxia $(FiO_2 = 15.4\%)$ led to larger perceived limb discomfort compared to a lower absolute intensity in normoxia (90% vVO_{2MAX}). We expected exercise-related sensations to be similar between conditions as parcipants could adjust their velocity where necessary. However, this was not the case. Similar responses have been shown elsewhere (42), with greater perceived overall discomfort, breathlessness and limb discomfort following progressive, sub-maximal, self-paced cycling intervals (RPE = 3; modified CR10 Borg scale) in hypoxia (FiO₂ = 13.0%) compared to normoxia at a similar power output. Perceived breathlessness, limb discomfort and pleasure are exercise-related sensations contributing to overall RPE during exercise (16). However, there is a detachment between these when immediately assessed after HIIT intervals. We suggest that self-paced **HIIT** in hypoxia leads to unfavourable exercise-related sensations before and after running intervals, compared to normoxia.

Pre- and post-exercise

During the Stroop test, alertness increased (i.e., faster reaction time) whilst accuracy was maintained following **HIIT**, irrespective of condition. It is well known that **HIIT** in normoxia generally increases cognitive performance versus rest (i.e., faster reaction time, better accuracy) (43). However, during fixed-intensity exercise in hypoxia, cognitive performance (i.e., attention and executive function) is worsened compared to normoxia (17,18). We report that even though exercise-related sensations were worsened during HIIT, cognitive performance (assessed post-HIIT) was not negatively affected. Ochi et al. (18) reported decreased Stroop performance 15 mins after 10 mins of moderate-continuous intensity exercise (50% peak oxygen uptake) in hypoxia (FiO₂ = 13.5%) versus normoxia. Our results likely differ to the aforementioned study due to cognitive testing performed in normoxia and following different exercise modalities. Our data show that alertness is increased following **HIIT**, and not negatively impacted by hypoxia.

Limitations and perspectives

During self-paced exercise at a perceptually-regulated intensity in hypoxia, HR and muscle oxygenation responses are similar to normoxia for a lower running velocity. However, we used a single "hypoxic dose" (i.e., hypoxic severity and duration), target RPE and exercise duration during **HIIT**. Further investigations should refine self-selected protocols in hypoxia, such as the "hypoxic dose", target RPE and exercise duration to minimise the negative side effects of worsened exercise-related sensations found under the present circumstances. In addition, whether there are gender differences in response to hypoxic exposure during perceptually-regulated HIIT should be investigated, given that our final sample size (n = 19) included only three females.

389 Conclusion

When carrying out **HIIT** at a perceptually-regulated intensity (RPE equal to 16), larger running velocity decreases are needed in hypoxia than normoxia. This is accompanied by similar physiological stress (i.e., HR and muscle oxygenation) during **HIIT**, and cognitive function adjustments after. In hypoxia, exercise-related sensations and blood lactate concentrations were higher-than-normal with larger peripheral oxygen desaturation. Overall, perceptually-regulated running velocity in hypoxia compared to normoxia may be an effective alternative, at the expense of less favourable exercise-related sensations. Our results suggest that athletes under the influence of hypoxia require lower external workloads to reach a perceptually-regulated target during HIIT than normoxia. If employed in a practical setting, coaches should consider the potential of negatively implicated exercise-related sensations and blood lactate concentrations which may have further negative carry-over effects on training responsiveness in the following days.

1299		
1300		
1301 1302 1303	402	Acknowledgements
1304 1305	403	The authors would like to thank the Sport and Exercise Science Research Centre of London
1306 1307	404	South Bank University for a travel bursary and Dr. Sebastien Racinais of ASPETAR for
1308 1309 1310	405	accommodating Liam Hobbins. This work was supported by Qatar National Research Fund
1310 1311 1312	406	(grant number: NPRP 4–760–3–217).
1313 1314	407	
1315 1316 1317	408	Conflict of interest
1318 1319	409	The authors have no conflicts of interest or financial ties to disclose and no current or past
1320 1321 1322	410	relationship with companies or manufacturers who could benefit from the results of the present
1323 1324	411	study.
1325 1326		
1327 1328		
1320		
1330		
1331		
1332 1333		
1334		
1335		
1336		
1337 1338		
1339		
1340		
1341		
1342		
1343 1344		
1345		
1346		
1347		
1348		
1349 1350		
1351		
1352		
1353		
1354		
1355		22
1356 1357		23

1359			
1360	412	Refere	ences
1361 1362			
1363	413	1	Zais I Dishan D and Australia D High intensity warm was offered during
1364	413	1.	Zois, J., Bishop, D. and Aughey, R., High-intensity warm-ups: effects during
1365	414		subsequent intermittent exercise. Int J Sports Phys Perf. 2015;10(4):498-503.
1366			subsequent interimitent excreme. Into sports rings ren. 2015,10(1). 190 505.
1367 1368	415	2.	Gibala, M. J., Gillen, J. B. and Percival., M. E. Physiological and health-related
1369			
1370	416		adaptations to low-volume interval training: influences of nutrition and sex. Sports
1371	447		Mad 2014.44 2: 127 127
1372 1373	417		Med. 2014;44.2: 127-137.
1374	418	3.	Buchheit, M. and Laursen, P. B. High-intensity interval training, solutions to the
1375			
1376	419		programming puzzle. Sports Med. 2013;43(10):927-954.
1377 1378	400	4	Ciller I.D. and Cihele M. I. Is high internativinterval training a time officient eventing
1379	420	4.	Gillen, J. B. and Gibala, M. J. Is high-intensity interval training a time-efficient exercise
1380	421		strategy to improve health and fitness? App Phys Nutr Met. 2013;39(3):409-412.
1381 1382			
1383	422	5.	Buchheit, M., Kuitunen, S., Voss, S. C., Williams, B.K., Mendez-Villanueva, A. and
1384	423		Bourdon, P.C. Physiological strain associated with high-intensity hypoxic intervals in
1385	423		bourdon, F.C. Flyslological strain associated with high-intensity hypoxic intervals in
1386 1387	424		highly trained young runners. J Strength Cond Res. 2012;26(1):94-105.
1388			
1389	425	6.	Laurent, C. M., Vervaecke, L. S., Kutz, M. R. and Green, J. M., Sex-specific responses
1390 1391	426		to self-paced, high-intensity interval training with variable recovery periods. J Strength
1392	120		to som paeed, men mensity mer var daming with variable recovery periods. I Strengar
1393	427		Cond Res. 2014;28(4):920-927.
1394 1395		_	
1395	428	7.	Heinonen, I. H., Boushel, R. and Kalliokoski, K. K. The circulatory and metabolic
1397	429		responses to hypoxia in humans-with special reference to adipose tissue physiology
1398			
1399 1400	430		and obesity. Front Endo. 2016;7:116.
1401	40.4	0	
1402	431	8.	Parfitt, G., Blisset, A., Rose, E. A. and Eston, R. Physiological and perceptual responses
1403 1404	432		to affect-regulated exercise in healthy young women. Psychophysiology.
1405	-		
1406	433		2012;49(1):104-110.
1407	40.4	0	
1408 1409	434	9.	Hardy, C. J. and Rejeski, W. J. Not what, but how one feels: The measurement of affect
1410	435		during exercise. J Sport Exerc Psych. 1989;11(3):304-317.
1411			
1412 1413			
1414			
1415			24
1416			

1417		
1418		
1419 1420	436	10. Fernández Menéndez, A., Saudan, G., Sperisen, L., Hans, D., Saubade, M., Millet, G.P.
1421 1422 1423	437	and Malatesta, D. Effects of Short-Term Normobaric Hypoxic Walking Training on
1423 1424 1425	438	Energetics and Mechanics of Gait in Adults with Obesity. Obesity. 2018;26(5):819-
1426 1427	439	827.
1428 1429	440	11. Chacaroun, S., Gonzalez, I. V. E., Flore, P., Doutreleau, S. and Verges, S. Physiological
1430 1431	441	responses to hypoxic constant-load and high-intensity interval exercise sessions in
1432 1433 1434	442	healthy subjects. Euro J App Phys. 2018;1-12.
1434 1435 1436	443	12. Brocherie, F., Millet, G. P., & Girard, O. Psychophysiological responses to
1437 1438	444	repeated-sprint training in normobaric hypoxia and normoxia. International J of
1439 1440	445	Sports Phys Perform. 2017;12(1):115-123.
1441 1442 1443	446 447	13. <u>Thum, J., Parsons, G., Whittle, T., & Astorino, T. (2017). High-intensity interval</u> training elicits higher enjoyment than moderate intensity continuous exercise.
1443 1444 1445	448	PloS one. 2017;12(1):e0166299.
1446 1447	449	14. Lind, E., Joehns-Matre, R. R., and Ekkekakis, P. What intensity of physical activity do
1448 1449	450	previously sedentary middle-aged women select? Evidence of a coherent pattern from
1450 1451 1452	451	physiological, perceptual and affective markers. Prev Med. 2005;40:407-419.
1453 1454	452	15. Friedmann, B., Bauer, T., Menold, E. and Bärtsch, P. Exercise with the intensity of the
1455 1456	453	individual anaerobic threshold in acute hypoxia. Med Scie Sports Exerc.
1457 1458 1459	454	2004;36(10):1737-1742.
1460 1461	455	16. Abbiss, C. R., Peiffer, J. J., Meeusen, R. and Skorski, S. Role of ratings of perceived
1462 1463	456	exertion during self-paced exercise: what are we actually measuring? Sports Med.
1464 1465	457	2015;45(9):1235-1243.
1466 1467 1468	458	17. Ochi, G., Yamada, Y., Hyodo, K., Suwabe, K., Fukuie, T., Byun, K., Dan, I. and Soya,
1469 1470	459	H. Neural basis for reduced executive performance with hypoxic exercise. Neuroimage.
1471 1472	460	2018;171:75-83.
1473 1474 1475		25

1476		
1477		
1478 1479	461	18. McMorris, T., Hale, B. J., Barwood, M., Costello, J. and Corbett, J. Effect of acute
1480 1481	462	hypoxia on cognition: A systematic review and meta-regression analysis. Neuro
1482 1483	463	Biobehavioral Revs. 2017;74:225-232.
1484 1485 1486	464	19. Seiler, S. and Sjursen, J. E. Effect of work duration on physiological and rating scale
1487 1488	465	of perceived exertion responses during self-paced interval training. Scand J Med Sci
1489 1490	466	Sports. 2004;14(5):318-325.
1491 1492	467	20. Jeffries, O., Patterson, S. D., & Waldron, M. The effect of severe and moderate
1493 1494	468	<u>hypoxia on exercise at a fixed level of perceived exertion. Euro J App Phys. 2019;1-</u>
1495 1496 1497	469	<u>12.</u>
1497 1498 1499	470	21. Martin, P. E., Rothstein, D. E., Larish, D. D. Effects of age and physical activity status
1500 1501	471	on the speed-aerobic demand relationship of walking. J Appl Phys 1992;73:200–206.
1502 1503	472	22. Borg, G. A. V. Perceived exertion as an indicator of somatic stress. Scand J Rehab Med.
1504 1505	473	1970;2:92-98.
1506 1507 1508	474	23. Laurent, C.M., Green, J. M., Bishop, P. A., Sjökvist, J., Schumacker, R. E., Richardson,
1508 1509 1510	475	M. T. and Curtner-Smith, M. A practical approach to monitoring recovery:
1511 1512	476	development of a perceived recovery status scale. J Strength Cond Res.
1513 1514	477 478	2011;25(3):620-628. 24. Crewther, B. T., Carruthers, J., Kilduff, L. P., Sanctuary, C. E. and Cook, C. J.
1515 1516	479	Temporal associations between individual changes in hormones, training motivation
1517 1518 1519	480	and physical performance in elite and non-elite trained men. Bio Sport. 2016;33(3):215.
1520 1521	481	25. Ward, S. A. and Whipp, B. J. Effects of peripheral and central chemoreflex activation
1522 1523	482	on the isopnoeic rating of breathing in exercising humans. J Phys. 1989;411(1):27-43.
1524 1525	483	26. Stroop, J. R. Studies of interference in serial verbal reactions. J Exp Psychol.
1526 1527 1528	484	1935;18:643-662.
1520 1529 1530		
1531		
1532		
1533		26
1534		

1536
1537
1538
485
27. Cohen, J. Statistical power analysis for the behavioral sciences, 2nd ed. 1988. New
1539
1540
486
Jersey: Lawrence Erlbaum Associates.
1541

- 1542 487
 1543 28. da Silva, D. F., Simões, H. G. and Machado, F. A., vVO2max versus Vpeak, what is
 1543 the best predictor of running performances in middle-aged recreationally-trained
 1545 1546 489 runners?. Science & Sports. 2015;30(4):e85-e92.
- 1548
154949029. Smith, K. J. and Billaut F. Influence of cerebral and muscle oxygenation on repeated-1550
1551491sprint ability. Eur J Appl Physiol. 2010;109(5):989-99
- 30. Dufour, S. P., Ponsot, E., Zoll, J., Doutreleau, S., Lonsdorfer-Wolf, E., Geny, B., Lampert, E., Fluck, M., Hoppeler, H., Billat, V. and Mettauer, B. Exercise training in normobaric hypoxia in endurance runners. I. Improvement in aerobic performance capacity. J App Phys. 2006;100(4):1238-1248.
- 496
 496
 31. Pramsohler, S., Burtscher, M., Faulhaber, M., Gatterer, H., Rausch, L., Eliasson, A. and
 497
 497
 497
 497
 497
 497
 497
 498
 498
 498
 498
 498
 498
 498
 498
 498
- 1567
156849932. Haufe, S., Wiesner, S., Engeli, S. et al. Influences of normobaric hypoxia training on1569
1570500metabolic risk markers in human subjects. Med Sci Sports Exerc. 2008;40(11):1939–1571
15725011944.
- 157450233. Wiesner, S., Haufe, S., Engeli S, et al. Influences of normobaric hypoxia training on15751576503157750315785041579504
- 1580
158150534. Krejčí, J., Botek, M. and McKune, A.J., 2018. Dynamics of the heart rate variability1582
1583and oxygen saturation response to acute normobaric hypoxia within the first 10 min of1584
1585exposure. Clin Phys Funct Imag. 2018;38(1):56-62.

1595		
1596 1597	508	35. Van Beekvelt, M. C., Colier, W. N., Wevers, R. A., Van Engelen, B. G. Performance
1598 1599	509	of near-infrared spectroscopy in measuring local O2 consumption and blood flow in
1600 1601	510	skeletal muscle. J Appl Physiol. 2001;90:511–519.
1602 1603	511	36. Joyner, M. J. and Casey, D. P. Regulation of increased blood flow (hyperemia) to
1604 1605	512	muscles during exercise: a hierarchy of competing physiological needs. Phys Revs.
1606 1607	513	2015;95(2):549-601.
1608 1609 1610	514	37. Howlett, R. A., Heigenhauser, G. J., Hultman, E., Hollidge-Horvat, M. G. and Spriet,
1610 1611 1612	515	L. L. Effects of dichloroacetate infusion on human skeletal muscle metabolism at the
1612 1613 1614	516	onset of exercise. Am J Phys Endo Met. 1999;277(1):E18-E25.
1615 1616	517	38. Renfree, A., Martin, L., Micklewright, D., and Gibson, A. S. C. Application of decision-
1617 1618	518	making theory to the regulation of muscular work rate during self-paced competitive
1619 1620	519	endurance activity. Sports Med. 2014;44:147-158.
1621 1622	520	39. Stork, M. J., Gibala, M. J. and Martin, K. G. Psychological and Behavioral Responses
1623 1624	521	to Interval and Continuous Exercise. Med Scie Sports Exerc. 2018;50(10):2110-2121.
1625 1626	522	40. Subudhi, A. W., Dimmen, A. C., & Roach, R. C. Effects of acute hypoxia on
1627 1628 1629	523	<u>cerebral and muscle oxygenation during incremental exercise. J App Phys.</u>
1630 1631	524	<u>2007;103(1):177-183.</u>
1632 1633	525	41. Subudhi, A. W., Miramon, B. R., Granger, M. E., & Roach, R. C. Frontal and
1634 1635	526	motor cortex oxygenation during maximal exercise in normoxia and hypoxia. J
1636 1637	527	<u>App Phys. 2009;106(4):1153-1158.</u>
1638 1639	528	42. Christian, R. J., Bishop, D., Girard, O., & Billaut, F. The role of sense of effort on self-
1640 1641	529	selected cycling power output. Front in Phys. 2014;5:115.
1642 1643	530	43. Lambourne, K. and Tomporowski, P. The effect of exercise-induced arousal on
1644 1645	531	cognitive task performance: a meta-regression analysis. Brain Res. 2010;1341:12-24.
1646 1647		
1648 1649		
1650 1651		28
1652		

Psycho-physiological responses to perceptually-regulated interval runs in hypoxia and normoxia

- Our primary aim was to investigate whether perceptually-regulated high-intensity intervals in hypoxia are associated with slower running velocities *versus* normoxia, whilst physiological responses and exercise-related sensations do not differ.
- Our findings show that participants adjusted to a progressively slower running velocity over the course of the protocol, and more so in hypoxic compared to normoxic conditions.
- Whilst SpO₂ was intuitively lower in hypoxia *versus* normoxia, heart rate and muscle oxygenation haemodynamics values changed over time but were matched between environmental conditions.
- Further, exercise-related sensations (*i.e.*, perceived recovery, motivation, breathlessness, limb discomfort and pleasure) were negatively impacted over time, and more so in hypoxic compared with normoxic conditions.
- Overall, slower interval running velocities in hypoxia achieve similar heart rate and muscle oxygenation responses to those observed in normoxia when perceptually-regulated, yet at the expense of less favourable exercise-related sensations.

Title

- Psycho-physiological responses to perceptually-regulated interval runs in hypoxia and
- normoxia

Author names and affiliations

- Hobbins La, Gaoua Na, Hunter Sa, Girard Obc

- ^aSport and Exercise Science Research Centre (SESRC), London South Bank University,
- London, United Kingdom
- ^bMurdoch Applied Sports Science (MASS) Laboratory, Murdoch University, Perth, Australia
- ^cAthlete Health and Performance Research Center, ASPETAR, Qatar Orthopedic and Sports
- Medicine Hospital, Doha, QATAR

Corresponding author

- Mr. Liam Hobbins
- hobbinsl@lsbu.ac.uk
- Sport and Exercise Science Research Centre
 - London South Bank University
 - 103 Borough Road
 - London
- SE1 0AA

Abstract

We investigated whether perceptually-regulated high-intensity intervals in hypoxia are associated with slower running velocities versus normoxia, when physiological responses and exercise-related sensations remain the same. Nineteen trained runners $(33.4 \pm 9.1 \text{ years})$ completed a high-intensity interval running protocol (4 \times 4-min intervals at a clamped perceived rating exertion of 16 on the 6–20 Borg scale, 3-min passive recoveries) in either hypoxic (HYP; FiO₂ 15.0%) or normoxic (NOR; FiO₂ 20.9%) conditions. Participants adjusted to a progressively slower running velocity from interval 1-4 (-7.0%), and more so in HYP vs. NOR for intervals 2, 3 and 4 (-4.6%, -6.4% and -7.9%, respectively; p < 0.01). Heart rate increased from interval 1–4 (+4.8%; p < 0.01), independent of condition. Arterial oxygen saturation was lower in HYP vs. NOR (86.0% vs. 94.8%; p < 0.01). Oxyhemoglobin (-23.7%) and total hemoglobin (-77.0%) decreased, whilst deoxyhemoglobin increased (+44.9%) from interval 1–4 (p < 0.01), independent of condition. Perceived recovery (-41.6%) and motivation (-21.8%) were progressively lower from interval 1-4, and more so in HYP vs. NOR for intervals 2, 3 and 4 (recovery: -8.8%, -24.2% and -29.3%; motivation: -5.3%, -20.3% and -22.4%, respectively; p < 0.01). Perceived breathlessness (+18.6%), limb discomfort (+44.0%) and pleasure (-32.2%) changed from interval 1-4, with significant differences (+21.8%, +11.3% and -31.3%, respectively) between HYP and NOR (p < 0.01). Slower interval running velocities in hypoxia achieve similar heart rate and muscle oxygenation responses to those observed in normoxia when perceptually-regulated, yet at the expense of less favourable exercise-related sensations.

119		
120		
121	23	Key words
122 123		·
123	24	High-intensity intermittent running; normobaric hypoxia; perceptually-regulated exercise;
125		
126	25	ratings of perceived exertion; near-infrared spectroscopy; effort perception.
127		
128		
129		
130		
131		
132		
133		
134		
135		
136		
137 138		
139		
140		
141		
142		
143		
144		
145		
146		
147		
148 149		
149 150		
151		
152		
153		
154		
155		
156		
157		
158		
159		
160		
161 162		
163		
164		
165		
166		
167		
168		
169		
170		
171		
172		
173		
174 175		
175 176		3
177		5
-		

Introduction

High-intensity interval training (HIIT) is a popular exercise format in athletic and clinical populations (1,2). HIIT includes repeated short-to-long (2-5 min) intense exercise bouts (80-90% of the velocity associated with maximal oxygen uptake or vVO_{2Max}) interspersed with shorter (1–3 min) recoveries (3). Compared to moderate-intensity continuous running, HIIT leads to similar improvements in cardiorespiratory fitness that is achieved with a shorter effective exercise duration per session (2). Due to the reduced time-commitment and exercise training volume, investigations surrounding the potential physiological and performance benefits of HIIT have surged (4).

HIIT in normobaric hypoxia (a lower inspired oxygen fraction or FiO₂) is receiving attention for its potential in further advancing athletic performance compared to HIIT in normoxia. Buchheit et al. (5) employed a HIIT protocol $(3 \times 5\text{-min}, 90\text{-s recovery})$ carried out in hypoxia $(vVO_{2Max} = 84\%; FiO_2 = 15.4\%)$ and normoxia $(vVO_{2MAX} = 90\%)$ at a fixed-intensity (determined in normoxia) in highly-trained runners. A reduced physiological stress (i.e., lower heart rate or HR) was observed during hypoxia compared to normoxia, likely due to a lower vVO_{2Max} in hypoxia versus normoxia. However, fixed exercise intensities, regardless of environmental conditions, do not permit adjustments (i.e., increases or decreases of workload) during exercise to match the intensity target (i.e., vVO_{2MAX}). In turn, over-induced physiological stress may be counter-productive (i.e., greater deoxygenated muscle heamoglobin, lower oxygenated haemoglobin) for intended session goals (6). Furthermore, matched absolute fixed exercise intensities (i.e., a similar percentage of vVO_{2MAX}) lead to greater physiological stress (i.e., compensatory increase in HR) in hypoxia compared to normoxia due to reduced FiO₂ (7). Perceptually-regulated exercise intensities, that allow velocity adjustments based upon exercise-related sensations in order to maintain a target effort

level, may offer a viable solution, and is perhaps more reflective of how exercisers modifyintensity during acute exercise.

Perceptually-regulated exercise permits the individual exercising to self-regulate external workload (i.e., running velocity/cycling power production) based upon Borg's rating of perceived exertion (RPE) scale (8). The validity and usefulness of using RPE for perceptually-regulating exercise has been described (9). The reduced oxygen availability in hypoxia makes the expectation tenable that there would be a slower self-selected running velocity in hypoxia for a given RPE, while velocity in normoxia would be more preserved, as evidenced previously (10). Chacaroun et al. (11) demonstrated for a lower power output (-15%), vastus lateralis muscle deoxyhemoglobin was higher and oxyhemoglobin lower in hypoxia (FiO₂ = 13.5%) compared to normoxia during a single interval session (15×1 -min at 75% of maximal HR, 1-min recoveries). Although HR was similar between conditions, RPE has been reported to be higher in hypoxia compared to normoxia during fixed-intensity interval runs (5) and repeatedsprint cycling (12). Employing self-paced exercise, in replace of fixed-intensity exercise, may assist in overcoming the over-excessive physiological stress observed when exercising in hypoxia versus normoxia, due to the likelihood of greater velocity preservations in the latter than the former.

In normoxia at pre-determined fixed intensities, HIIT is perceived as more enjoyable compared to moderate-intensity continuous running (13). However, during HIIT at fixed-intensities, exercise-related sensations decrease when the exercise intensity rises above threshold preference (14). Further, HIIT in hypoxia at fixed-intensities typically surpasses the preferred threshold in normoxia (15). Implementing a self-paced exercise model may permit modifications required (i.e., slower running velocities) to maintain exercise-related sensations contributing to RPE (16) in hypoxia and normoxia. Cyling continuously for 10 min at a fixed-intensity (corresponding to 50% VO_{2Max}) in hypoxia versus normoxia negatively impacts

cognitive function (17). Slower self-selected running velocities may assist with mitigating
hypoxic-induced negative cognitive function compared to normoxia (18). These potential
findings may benefit athletes exercising intensely in hypoxia, shortly followed by skills
requiring attention and accuracy.

Therefore, the aim of this study was to investigate the effect of HIIT at a clamped RPE of 16 (typically used by athletes during HIIT) (19) in hypoxia and normoxia on adjustments in running velocity and associated exercise-related sensations of trained runners. We hypothesized that running velocity would be progressively slower in hypoxia compared to normoxia across intervals, whilst physiological and cognitive responses, and exercise-related sensations would not differ between conditions. Decreasing external load with matched internal load during perceptually-regulated HIIT in hypoxia compared to normoxia may benefit athletes during heavy training blocks prior to competition.

63 Methods

Participants

Nineteen trained runners (3 females, 16 males; age: 33.4 ± 9.1 years; height: 176 ± 88 cm; weight: 76.3 ± 10.9 kg) provided written informed consent to participate. Participants had no musculoskeletal injuries and met the following eligibility criteria: a training volume >6 h/wk, free of clinical signs of disease, orthopedic, neurological, cardiovascular or respiratory problems, and no hypoxic exposure >2000 m for >48 h 6 months before the study. The study was carried out in accordance with the Declaration of Helsinki and was approved by the Ethics Committee of the Anti-Doping Lab Qatar institutional review board (Agreement SCH-ADL-170). A-priori sample size was calculated using G*Power (Version 3.1.9.3). This was

determined using published power output data by Jeffries et al. (20), whereby healthy individuals cycled at a perceptually-regulated intensity (RPE = 16) in hypoxic (FiO₂ = 15.0%) and normoxic conditions. Twenty-one participants were deemed sufficient to yield a power of 0.8 at an α probability of 0.05. Two individuals dropped out due to injuries sustained during their time enrolled onto the study, not associated with the HIIT protocols we employed.

78 Experimental design

Participants reported to the laboratory on three occasions, each separated by ≥ 48 h. The first session included study familiarisation. The second and third visits included completing a HIIT protocol in either hypoxia or normoxia in a randomized, conterbalanced order. Physiological, perceptual and cognitive responses were assessed continuously, immediately before and after each interval, and before and after the HIIT protocol, respectively. Participants were instructed to refrain from any intense exercise 48 h prior to each visit and consume their last meal at least 2 h prior to the HIIT sessions. To minimise the impact of social desirability bias, participants were made aware of the purpose of the study but were naïve to experimental hypotheses. Laboratory conditions were similar throughout all sessions (mean temperature ~22°C, relative humidity \sim 50%) and time of day was standardized for each participant.

89 Familiarization session

At the preliminary visit to the laboratory, participants were familiarised with the perceptual scales and cognitive test. Preferred running velocity (PRV) was determined for each participant in normoxia using a modified version of identifying preferred walking speed (21). After a 5min warm up at 10 km/h⁻¹, participants completed four ramped treadmill runs (increasing and decreasing velocities) on an instrumented treadmill (ADAL3D-WR, Medical Development– HEF Tecmachine, France). After every 20 s per ramp, participants rated their RPE of the current velocity (controlled by the investigator and out of sight of the participant) in accordance with Borg's (20) 6 ("no exertion at all") – 20 ("maximal exertion") numeric scale. Ramp one started at 10 km/h⁻¹, increasing by 0.8 km/h⁻¹ every 20 s until the velocity was considered as RPE \geq 18; ramp two started at +1.5 km/h⁻¹ the previous end velocity, decreasing by 0.8 km/h⁻¹ until the velocity was considered as RPE ≤ 12 ; ramp three started at the velocity considered as an RPE of 14 in ramp two, increasing by 0.5 km/h⁻¹ until the velocity was considerd as RPE \geq 18; and ramp four started at +1.0 km/h⁻¹ the previous end velocity, decreasing by 0.5 km/h⁻¹ until the velocity was considered as RPE ≤ 12 . Ramps two, three and four began once the participants declared their perceived recovery level as a 7 out of 10 following the previous ramp (23). HR was recorded every 20-s through each ramp. PRV corresponded to the velocity participants considered as a RPE of 16 (between "hard" and "very hard") or closest to a HR of 160 bpm. After 10 min of rest, participants completed one 4-min interval composing the HIIT protocol (see below) for habituation.

109 *Experimental trials*

Participants completed two experimental trials in normoxia (NOR; $FiO_2 = 20.9\%$) and hypoxia (HYP; FiO₂ = 15.0%, equivalent to \sim 2700 m above sea level). After a standardised warm up (5-min at 10 km/h⁻¹), a facemask connected to a portable hypoxic generator (See Hypoxic simulation section) was attached. Participants rested for 1-min (quiet standing) before a 1-min run at their PRV (RPE = 16). Participants then rested for 3 min before completing the HIIT protocol. The HIIT protocol was based upon aerobic interval-training (2). Participants completed four, 4-min intervals, interspersed with 3-min recoveries (quiet standing). The first 30 s of each 4-min interval began at participants' PRV; participants were then free to decide if or how treadmill velocity needed to be adjusted (manually by one experimenter) to ensure maintenance of a RPE of 16 every 30 s. Participants hand-signalled in response to the current velocity (finger up to increase, finger down to decrease, and circle using index finger and thumb

to maintain); and signalled again to inform how much of an increase/decrease in velocity is required [1, 2 or 3 fingers up (faster) or down (slower) for 0.5, 1.0 or 1.5 km/h⁻¹ changes, respectively]. Signals were trialled during familiarisation. Mild verbal encouragement to keep running at an RPE of 16 was used throughout HIIT. Total hypoxic exposure corresponded to exactly 28 min.

Hypoxic simulation

Participants were fitted with a facemask fastened with a Velcro headset connected via plastic tubing to a hypoxic generator (Altitrainer, SMTec SA, Nyon, Switzerland) to simulate hypoxia. The gas mixing system enriches inspired air by adding a fixed quantity of nitrogen via a 30-L mixing chamber, with the dilution being constantly controlled by a PO_2 probe (precision = T0.82 torr, safety $FiO_2 = 9.7\%$). The hypoxic generator was hidden from participant viewing to ensure condition blinding. When breathing 'normal air' during normoxia, the hypoxic generator was on (for background noise) and set at a simulated altitude of 100 m to increase the strength of blinding.

Measures

Exercise intervals

HR was monitored telemetrically with a Polar transmitter-receiver (Polar S810, Kempele, Finland) and recorded 20 s before and every 30 s during each interval. Arterial oxygen saturation (SpO₂) was assessed via finger pulse oximetery (Palmsat 2500, NONIN Medical Inc., Plymouth, MI, USA) at the same time intervals. HR and SpO₂ were obtained before (i.e., after a 2-min seated period) and at the end of the warm-up procedure (i.e., prior to HIIT). Both the HR watch (RS400, Polar) and oximeter receiver were attached on the handrails of the treadmill outside of the participants' view.

Muscle oxygenation trends of the right vastus lateralis muscle were recorded using near-infrared spectroscopy (NIRS; Portalite, Artinis, Netherlands) in real-time. A wireless bi-polar optode sensor was attached (~10 cm above the proximal patella border) and secured to skin via adhesive tape. Sampling frequency was set at 10 Hz (11) following a 'zero set' of all signals. Bandages were fastened around the lower limb and optode to prevent external light distorting readings. Oxy- (Δ ; [O₂Hb]), deoxy- (Δ ; [HHb]) and total haemoglobin (µmol; [tHb]) were exported (1 Hz). For analysis, each interval was averaged and normalized to a 10 s sample prior to interval one (reference value) for each respective condition and presented as percentage change.

153 During recovery

Perceived recovery and motivation to exercise were assessed 30 s before each interval. Perceived recovery was assessed by answering 'how recovered do you feel currently?' via a numeric scale, ranging from 0 ("very poorly recovered") to 10 ("very well recovered") (23). Recovery was assessed before interval one to determine perceptions following the warm up. Perceived motivation to exercise was assessed via a 20-cm visual analog scale (24). Participants were asked 'how motivated do you feel to exercise right now?' and answered by adjusting the level on the scale between 0 ("*extremely low*"; white colored) and 20 ("*extremely high*"; black colored). Immediately after each interval, ratings of perceived breathlessness, limb discomfort and pleasure were assessed. Perceived breathlessness was assessed by answering 'how does your breathing feel currently?' via a numeric scale, ranging from 0 ("nothing at all") to 10 ("very, very severe") (25). Using the same scale, perceived limb discomfort was assessed by answering 'how do your legs feel currently?'. A 20-cm visual analog scale (same as motivation above) was used to assess 'how pleasant was that run?' ranging from 0 ("extremely unpleasant") and 20 ("extremely pleasant").

Pre- and post-exercise

A capillary blood sample taken from the fingertip was analyzed for blood lactate concentration ([La⁺]) with the Lactate Pro (LT-1710, Arkray, Japan) portable analyzer before the warm-up and 2 min after HIIT. An offline Stroop colour-word test (26) assessed attention and executive function. Using one hand and as quickly as possible, participants selected the colored key on the keyboard representing the color of the text appearing on the screen (red, vellow, green or blue). The cognitive test lasted for 3 min, and took place in a silent environment before the warm up and 3 min after HIIT. Reaction time (ms; time taken to select a color) and accuracy (%; correct color selected) were averaged over each test for analysis.

177 Statistical analysis

Data distribution was assessed via a Shapiro-Wilk test. A parametric within-subject two-way analysis of variance was used to investigate the main effect of condition (NOR vs. HYP), time (interval 1, 2, 3 vs. 4 or pre vs. post) and the condition \times time interaction for normally distributed data. Partial eta-squared (n^2) was calculated as a measure of effect size. Values of 0.01, 0.06 and above 0.14 were considered as small, medium and large, respectively (27). A related samples Friedman's non-parametric test was used for data not normally distributed. Bonferroni post-hoc pairwise comparisons were used to identify locations of significant effects. Statistical testing was carried out in SPSS (v21; CED, Cambridge, United Kingdom). Data was considered significant if $p \le 0.05$. All data are presented as group means \pm SD.

637 187

188

643 189 *Changes in velocity and HR*

Results

190 Compared to interval 1, participants adjusted to a progressively slower running velocity during 191 intervals 2, 3 and 4 (-2.8%, -5.2% and -7.0%, respectively; p < 0.01), and more so in HYP vs. 192 NOR for intervals 2, 3 and 4 (-4.6%, -6.4% and -7.9%, respectively; p < 0.01; Figure 1A). 193 Compared to interval 1, HR increased during intervals 2, 3 and 4 (+2.3%, +3.6% and 4.8%, 194 respectively; p < 0.01; Figure 1B), independently of condition (p = 0.65).

*Changes in SpO*₂ and muscle oxygenation

196 SpO₂ was globally lower in HYP *vs.* NOR (-9.3% average across intervals; p < 0.01; Figure 197 1C), independently of time (p = 0.37). From interval 1 to 4, [O₂Hb] and [tHb] decreased (-198 23.7% and -77.0%, respectively) whilst [HHb] increased (+44.9%; p < 0.01; Figures 2A–C), 199 independently of condition (p > 0.08).

Changes in exercise-related sensations

Perceived recovery decreased progressively from interval 1 to 4 (-41.6%; p < 0.01), and more so in HYP vs. NOR before intervals 2, 3 and 4 (-8.8%, -24.2% and -29.3%, respectively; p =0.02; Figure 3A). Perceived motivation decreased progressively from interval 1 to 4 (-21.8%; p < 0.01), and more so in HYP vs. NOR before intervals 3 and 4 (-20.3% and -22.4%), respectively; p < 0.01; Figure 3B). Compared to interval 1, perceived breathlessness increased following intervals 2, 3 and 4 (+14.0%, +13.6% and +18.6%, respectively; p < 0.01; Figure 3C), independently of condition. Breathlessness was rated globally higher in HYP vs. NOR (+21.8%; p < 0.05), irrespective of time. Compared to interval 1, perceived limb discomfort increased following intervals 2, 3 and 4 (+23.3%, +35.3% and +44.0%, respectively; p < 0.01; Figure 3D), independently of condition. Limb discomfort was rated globally higher in HYP vs. NOR (+11.3%; p = 0.01), irrespective of time. The time-dependent decreases in perceived pleasure across intervals (-14.7%, -25.4% and -32.3%, intervals 2, 3 and 4 vs. 1, respectively; p < 0.01; Figure 3E) tended to be larger in HYP vs. NOR (-31.3%, p = 0.06).

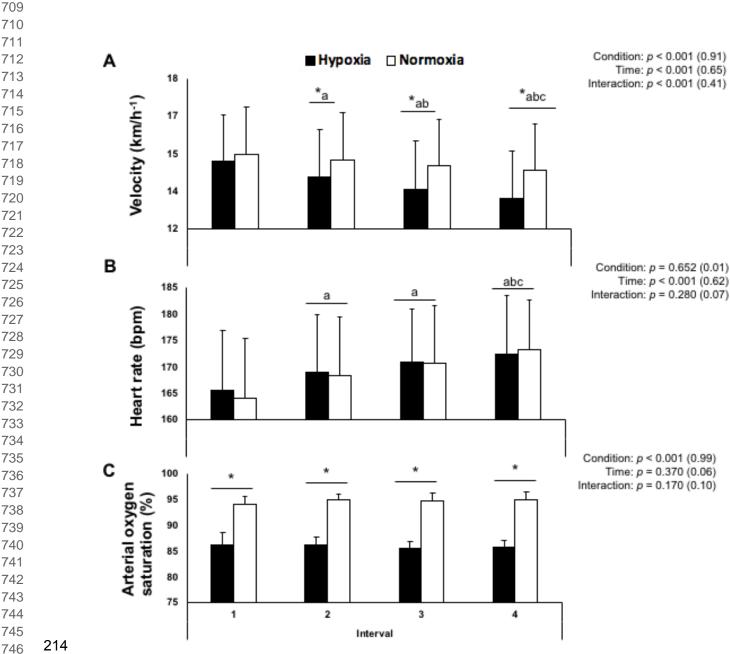


Figure 1 Changes in velocity (A), heart rate (B) and arterial oxygen saturation (C) during the high-intensity intermittent running protocol. Data are presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition: white bars = normoxic condition. * denotes a statistically significant difference between conditions for a given interval (p < 0.05), a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05).

221 Changes in $[La^+]$ and attention and executive function

The pre- to post-exercise increase in [La⁺] was larger (p = 0.001) in HYP ($1.7 \pm 0.8 \text{ vs. } 13.1 \pm$ 3.8 mmol/l⁻¹) vs. NOR (2.1 ± 0.9 vs. 10.1 ± 3.9 mmol/l⁻¹). During the Stropp test, accuracy was unaffected by condition and time (Figure 4A). Participants' reaction time was faster (+11%)post vs. pre HIIT (p < 0.01; Figure 4B), independently of condition.

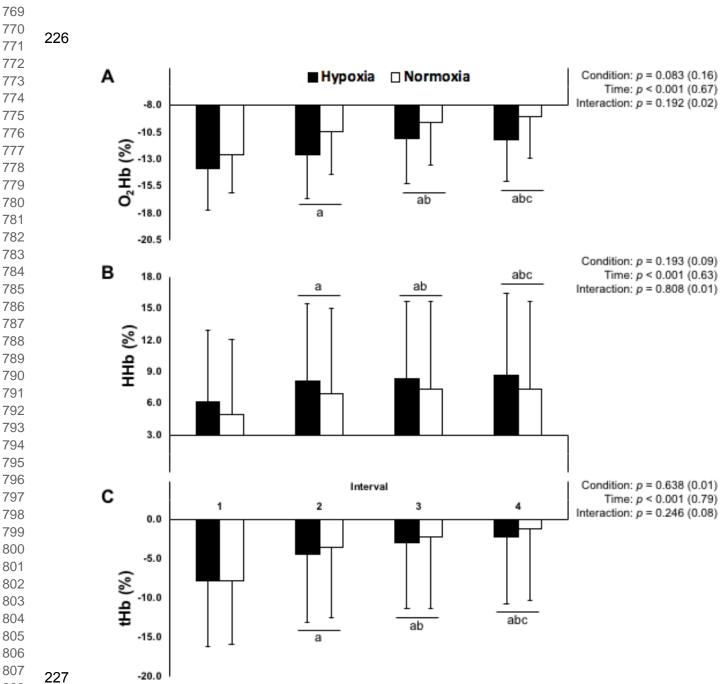


Figure 2 Changes in Oxygenated (A; O_2Hb), deoxygenated (B; HHb) and total hemoglobin (C; tHb) during the high-intensity intermittent running protocol. Data are calculated as a percentage difference from baseline (%) and presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05).

- ⁸¹⁵ 234

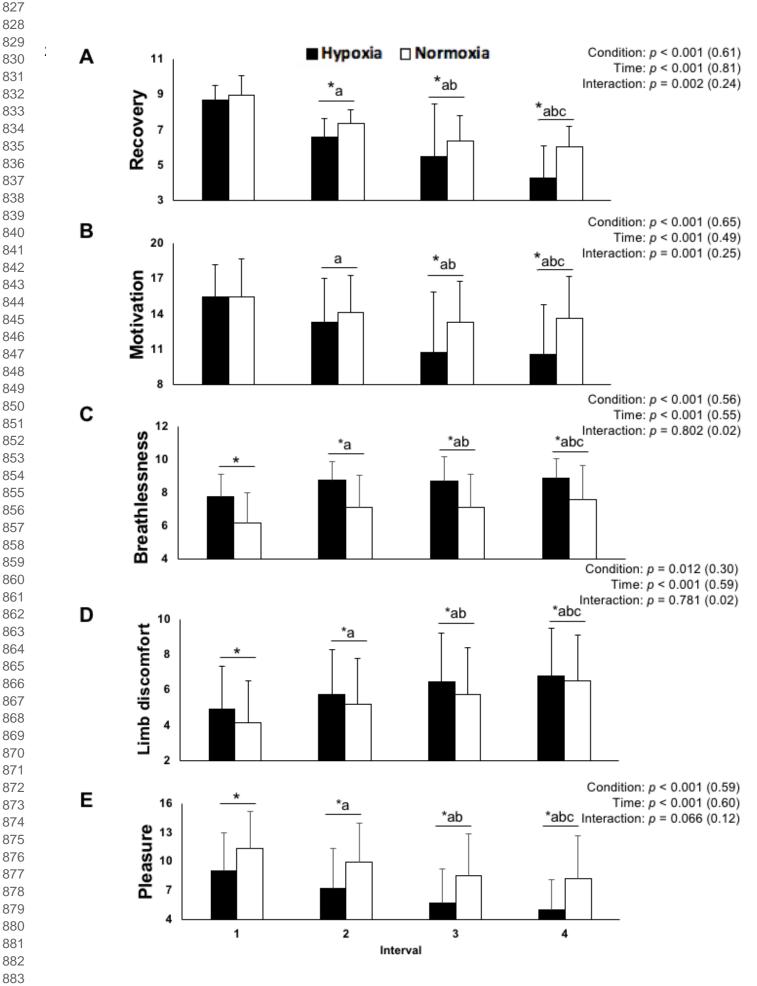
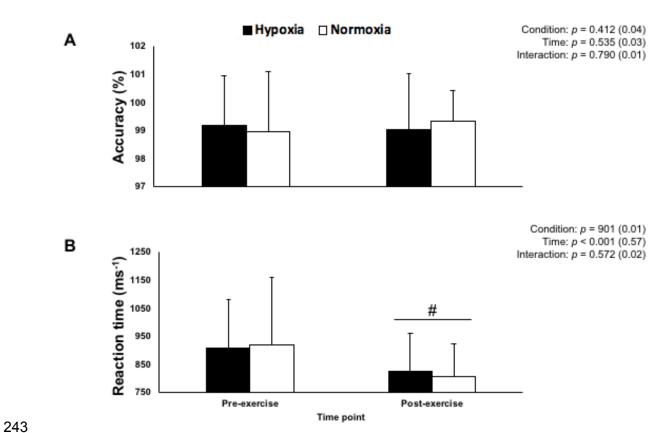


Figure 3 Changes in perceived recovery (A), motivation (B), breathlessness (C), limb discomfort (D) and pleasure (E) during the high-intensity intermittent running protocol. Data are presented as mean \pm SD. ANOVA main effects of time, condition and interaction are presented along with partial-eta squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic condition. * denotes a statistically significant difference between conditions for a given interval (p < 0.05), a, b and c denotes a statistically significant difference vs. interval 1, 2 and 3, respectively (p < 0.05).



922244Figure 4 Changes in accuracy (A) and reaction time (B) pre and post high-intensity923245intermittent running protocol. Data are averaged over 3 mins and presented as mean \pm SD.924246ANOVA main effects of time, condition and interaction are presented along with partial-eta925247squared for effect size into brackets. Black bars = hypoxic condition; white bars = normoxic926248

250 Discussion

Using a perceptually-regulated (RPE = 16) exercise model, we observed: 1) participants ran progressively slower during HIIT with larger decreases in HYP versus NOR, 2) HR and muscle oxygenation trends (during intervals) and cognitive responses (pre vs. post HIIT) were similar between conditions, 3) greater breathlessness and limb discomfort, and lower recovery,

motivation and pleasure scores were stated during recovery between HYP vs. NOR, and 4) blood lactate concentration was larger after HYP vs. NOR. Overall, using a manipulation of oxygen availability, reduced external workload (i.e., running velocity) during perceptually-regulated interval running is associated with a similar internal load (i.e., physiological responses). Although no cognitive function differences were found between conditions, this is achieved with less favourable exercise-related sensations. A matched internal workload for a decreased external workload during perceptually-regulated HIIT in hypoxia versus normoxia may assist athletes to reach intended session goals with minimal over-induced physiological stress. However, perceptually-regulated HIIT exacerbates exercise-related sensations and blood lactate concentrations in hypoxia compared to normoxia. This may then have negative carry-over effects on training responsiveness in the following days.

Exercise intervals

The velocity deemed equal to RPE 16 (PRV) was as expected for trained runners (~15 km/h⁻¹) (28). Interestingly, running velocity did not differ between conditions during the first HIIT interval, despite lower SpO2 in hypoxia versus normoxia. Smith & Billaut (29) found maintained SpO₂ during repeated-sprinting in normoxia (20×5 -s all out, 25-s recovery) until after the fifth sprint in national-level soccer players, whereby peak power significantly decreased compared to sprint one. Overall, it seems that initial decreases in SpO₂ (within interval one) do not necessarily impact on HIIT compared to sprint intervals.

We found that participants selected a progressively slower running velocity during HIIT in both conditions. In highly-trained middle to long-distance runners, a 6% reduction in vVO_{2MAX} when running in hypoxia versus normoxia is acceptable to match the acute physiological stress induced (5). It can be suggested that self-selected velocity adjustments found in the current study to maintain RPE 16 are matched with modifications in hypoxic versus normoxic traning

sessions employed by coaches and sport scientists for athletes (30). Decreased external workloads have been reported by Pramsohler et al. (31) during continuous cycling (seven 30-min sessions over 3-wk), whereby participants cycled at -28% lower power output in hypoxia $(FiO_2 = 15.3\%)$ versus normoxia for a similar HR. Differences in these findings and ours may be due to the inclusion of geriatric patients completing pre-set (in normoxia) fixed-intensity cycling compared to trained runners self-regulating HIIT in the current study. However, Fernández-Menéndez et al. (10) reported preferred walking velocity (RPE of 10) in hypoxia $(FiO_2 = 15.3\%)$ was 7% slower than normoxia in obese adults over 3 weeks. Using a self-paced model, irrespective of RPE target, population demographics and training block duration, lower external workloads are selected in hypoxia compared to normoxia. Overall, decreases in self-paced running velocity occured to a greater extent in hypoxia than normoxia to maintain RPE 16, suggesting of a lower external workload. This finding may be of benefit to athletes who are unable or advised by their coach not to be training at a full intensity. Completing perceptually-regulated HIIT in hypoxia that requires slower running velocities compared to normoxia may in turn minimise mechanical constraints and eventually injury risk.

Our data show HR increased progressively during HIIT, irrespective of condition. This matches our hypothesis that HR will be comparable between hypoxia and normoxia, even though running velocity was lower in hypoxia. Other studies employing moderate continuous-intensity exercise have also found matched HR responses between hypoxic and normoxic training interventions (~4 weeks) when cycling at a -21.0% power output in healthy males (32) and walking/running at a -17.5% velocity in obese adults (33) in hypoxia verus normoxia. Although exercise intensities in these studies were fixed, we believe similar increases in HR between conditions occur due to the environmental stressor (hypoxia) augmenting autonomic cardiac regulation (34). Overall, it seems self-paced exercise in hypoxia provides an added

1065
1066303environmental stressor that is able to mimick HR responses in normoxia for a lower external1067
1068304load.

Lower [O₂Hb] and [tHb], and greater [HHb] of the *vastus lateralis* were recorded across HIIT, irrespective of condition. Decreases in [O₂Hb] and increases in [HHb] were expected during HIIT as oxygen delivery is outweighed by utilisation, whilst decreases in [tHb] reflect a lower localised blood flow (35). Active musculature oxygenation is negatively impacted during fixed-intensity exercise in hypoxia compared to normoxia due to a lower FiO_2 (7). In support of this, Chacaroun et al. (11) reported lower [O₂Hb] and greater [HHb] with maintained [tHb] of the vastus lateralis during fixed, relative high-intensity cycling in hypoxia (85% maximal power output in normoxia; $FiO_2 = 13.5\%$) versus normoxia. Where we employed a self-paced exercise model, similar [O₂Hb] and [HHb] responses are achieved between conditions. This is likely explained through the decreased workload (i.e., slower running velocity) in hypoxia compared to normoxia, subsequently lowering oxygen utilisation. Discrepants findings in [tHb] may be due to different exercise modalities (cycling versus running) modifying blood flow regulation (36). Similar to HR responses (central) previously discussed, it can be suggested here that local (tissue oxygenation) physiological stress is matched between conditions during HIIT in hypoxia at a slower velocity compared with normoxia.

Elevations in [La⁺] following HIIT were higher in HYP than NOR. Values in the current study $(10-13 \text{ mmol/l}^{-1})$ are somewhat higher than those $(5-6 \text{ mmol/l}^{-1})$ reported elsewhere following a single HIIT session (6×4 -min intervals at a RPE ~17, 4-min recoveries) (19). This maybe due to a 1:0.75 work:rest ratio implemented during our protocol compared to 1:1 employed by Seiler & Sjursen (19). [La⁺] normalization during shorter recovery periods may not occur to the extent following longer recovery periods due to excess pyruvate accumulation (37). This suggests that HIIT in hypoxia per se leads to increased [La⁺] at slower running velocities compared to normoxia for similar physiological stress amounts. Practitioners should be aware

328 that perceptually-regulated HIIT in hypoxia is a viable method for matching indices of 329 physiological stress to normoxia. However, the blood lactate concentration increases after 330 exercise were larger in hypoxia compared to normoxia. This may have negative implications 331 on the muscle fatigue recovery process.

During recovery

Perceptual responses to HIIT were negatively impacted (i.e., lower recovery, and motivation) when assessed before intervals, with further exacerbations in hypoxia. Participants were instructed to maintain a RPE of 16 throughout HIIT by adjusting their velocity where necessary. It might be surprising at first that perceptual responses were worse in hypoxia compared to normoxia. However, perceived recovery and motivation are important affects associated with exercise intensity regulation (38). Our results indicate that hypoxia negatively impacts these affects during HIIT compared with normoxia. This may be explained through lower perceived capabilities of hypoxic HIIT completion over normoxia (39), lowering perceived recovery and motivation. Further, it could be postulated that cerebral deoxygenation was greater during HIIT in hypoxia versus normoxia, as demonstrated by Subudhi et al. during incremental cycling (40, 41). Accordingly, cerebral deoxygenation during HIIT may contribute to an integrative decision regarding negative perceptions, in which hypoxia hastens this effect (41). Given that the perceptually-regulated exercise model is governed centrally, this may provide a potential explanation as to why exercise-related sensations were more elevated in the hypoxic trial. Overall, our data poses a disconnection between RPE and exercise-related sensations (i.e., recovery and motivation). Further research should look to optimise HIIT in hypoxia for positive perceptual responses.

Perceptual responses after intervals were negatively impacted (i.e., higher breathlessness and
 limb discomfort, lower pleasure), and to a further extent in hypoxia than normoxia. Buchheit

et al. (5) reported that 3-min absolute-intenxity running intervals (84% vVO_{2MAX}) in hypoxia $(FiO_2 = 15.4\%)$ led to larger perceived limb discomfort compared to a lower absolute intensity in normoxia (90% vVO_{2MAX}). We expected exercise-related sensations to be similar between conditions as parcipants could adjust their velocity where necessary. However, this was not the case. Similar responses have been shown elsewhere (42), with greater perceived overall discomfort, breathlessness and limb discomfort following progressive, sub-maximal, self-paced cycling intervals (RPE = 3; modified CR10 Borg scale) in hypoxia (FiO₂ = 13.0%) compared to normoxia at a similar power output. Perceived breathlessness, limb discomfort and pleasure are exercise-related sensations contributing to overall RPE during exercise (16). However, there is a detachment between these when immediately assessed after HIIT intervals. We suggest that self-paced HIIT in hypoxia leads to unfavourable exercise-related sensations before and after running intervals, compared to normoxia.

Pre- and post-exercise

During the Stroop test, alertness increased (i.e., faster reaction time) whilst accuracy was maintained following HIIT, irrespective of condition. It is well known that HIIT in normoxia generally increases cognitive performance versus rest (i.e., faster reaction time, better accuracy) (43). However, during fixed-intensity exercise in hypoxia, cognitive performance (i.e., attention and executive function) is worsened compared to normoxia (17,18). We report that even though exercise-related sensations were worsened during HIIT, cognitive performance (assessed post-HIIT) was not negatively affected. Ochi et al. (18) reported decreased Stroop performance 15 mins after 10 mins of moderate-continuous intensity exercise (50% peak oxygen uptake) in hypoxia (FiO₂ = 13.5%) versus normoxia. Our results likely differ to the aforementioned study due to cognitive testing performed in normoxia and following different exercise modalities. Our data show that alertness is increased following HIIT, and not negatively impacted by hypoxia.

Limitations and perspectives

During self-paced exercise at a perceptually-regulated intensity in hypoxia, HR and muscle oxygenation responses are similar to normoxia for a lower running velocity. However, we used a single "hypoxic dose" (i.e., hypoxic severity and duration), target RPE and exercise duration during HIIT. Further investigations should refine self-selected protocols in hypoxia, such as the "hypoxic dose", target RPE and exercise duration to minimise the negative side effects of worsened exercise-related sensations found under the present circumstances. In addition, whether there are gender differences in response to hypoxic exposure during perceptually-regulated HIIT should be investigated, given that our final sample size (n = 19) included only three females.

388 Conclusion

When carrying out HIIT at a perceptually-regulated intensity (RPE equal to 16), larger running velocity decreases are needed in hypoxia than normoxia. This is accompanied by similar physiological stress (i.e., HR and muscle oxygenation) during HIIT, and cognitive function adjustments after. In hypoxia, exercise-related sensations and blood lactate concentrations were higher-than-normal with larger peripheral oxygen desaturation. Overall, perceptually-regulated running velocity in hypoxia compared to normoxia may be an effective alternative, at the expense of less favourable exercise-related sensations. Our results suggest that athletes under the influence of hypoxia require lower external workloads to reach a perceptually-regulated target during HIIT than normoxia. If employed in a practical setting, coaches should consider the potential of negatively implicated exercise-related sensations and blood lactate concentrations which may have further negative carry-over effects on training responsiveness in the following days.

1299		
1300		
1301 1302 1303	401	Acknowledgements
1304 1305	402	The authors would like to thank the Sport and Exercise Science Research Centre of London
1306 1307	403	South Bank University for a travel bursary and Dr. Sebastien Racinais of ASPETAR for
1308 1309 1310	404	accommodating Liam Hobbins. This work was supported by Qatar National Research Fund
1311 1312	405	(grant number: NPRP 4–760–3–217).
1313 1314	406	
1315 1316 1317	407	Conflict of interest
1318 1319	408	The authors have no conflicts of interest or financial ties to disclose and no current or past
1320 1321 1322	409	relationship with companies or manufacturers who could benefit from the results of the present
1323 1324	410	study.
1325 1326		
1327		
1328		
1329		
1330 1331		
1332		
1333		
1334		
1335		
1336 1337		
1338		
1339		
1340		
1341		
1342 1343		
1343		
1345		
1346		
1347		
1348		
1349 1350		
1351		
1352		
1353		
1354		
1355		22
1356 1357		23

1359			
1360	411	Refere	nces
1361 1362			
1363		1	
1364	412	1.	Zois, J., Bishop, D. and Aughey, R., High-intensity warm-ups: effects during
1365	413		subsequent intermittant eversion. Int I Sports Phys. Parf. 2015:10(4):408-502
1366	413		subsequent intermittent exercise. Int J Sports Phys Perf. 2015;10(4):498-503.
1367	414	2	Gibala, M. J., Gillen, J. B. and Percival., M. E. Physiological and health-related
1368 1369		2.	Giodia, M. J., Ginen, J. D. and Fereival., M. D. Thysiological and neural related
1309	415		adaptations to low-volume interval training: influences of nutrition and sex. Sports
1371			
1372	416		Med. 2014;44.2: 127-137.
1373		2	
1374 1375	417	3.	Buchheit, M. and Laursen, P. B. High-intensity interval training, solutions to the
1376	418		programming puzzle. Sports Med. 2013;43(10):927-954.
1377	10		programming puzzle. Sports Med. 2013,45(10).727 734.
1378	419	4.	Gillen, J. B. and Gibala, M. J. Is high-intensity interval training a time-efficient exercise
1379			
1380 1381	420		strategy to improve health and fitness? App Phys Nutr Met. 2013;39(3):409-412.
1382	404	-	
1383	421	5.	Buchheit, M., Kuitunen, S., Voss, S. C., Williams, B.K., Mendez-Villanueva, A. and
1384	422		Bourdon, P.C. Physiological strain associated with high-intensity hypoxic intervals in
1385 1386			bourdon, r.e. r hystological strain associated with high intensity hypoxic intervals in
1387	423		highly trained young runners. J Strength Cond Res. 2012;26(1):94-105.
1388			
1389	424	6.	Laurent, C. M., Vervaecke, L. S., Kutz, M. R. and Green, J. M., Sex-specific responses
1390 1391	425		to self-paced, high-intensity interval training with variable recovery periods. J Strength
1392	725		to sen-paced, ingh-intensity interval training with variable recovery periods. J Strength
1393	426		Cond Res. 2014;28(4):920-927.
1394			
1395 1396	427	7.	Heinonen, I. H., Boushel, R. and Kalliokoski, K. K. The circulatory and metabolic
1390	400		
1398	428		responses to hypoxia in humans-with special reference to adipose tissue physiology
1399	429		and obesity. Front Endo. 2016;7:116.
1400	720		
1401 1402	430	8.	Parfitt, G., Blisset, A., Rose, E. A. and Eston, R. Physiological and perceptual responses
1403			
1404	431		to affect-regulated exercise in healthy young women. Psychophysiology.
1405	400		2012 40(1) 104 110
1406 1407	432		2012;49(1):104-110.
1407	433	9.	Hardy, C. J. and Rejeski, W. J. Not what, but how one feels: The measurement of affect
1409	100	2.	
1410	434		during exercise. J Sport Exerc Psych. 1989;11(3):304-317.
1411			
1412 1413			
1414			
1415			24
1416			

1417		
1418		
1419 1420 1421	435	10. Fernández Menéndez, A., Saudan, G., Sperisen, L., Hans, D., Saubade, M., Millet, G.P.
1421 1422 1423	436	and Malatesta, D. Effects of Short-Term Normobaric Hypoxic Walking Training on
1424 1425	437	Energetics and Mechanics of Gait in Adults with Obesity. Obesity. 2018;26(5):819-
1426 1427	438	827.
1428 1429	439	11. Chacaroun, S., Gonzalez, I. V. E., Flore, P., Doutreleau, S. and Verges, S. Physiological
1430 1431	440	responses to hypoxic constant-load and high-intensity interval exercise sessions in
1432 1433	441	healthy subjects. Euro J App Phys. 2018;1-12.
1434 1435 1436	442	12. Brocherie, F., Millet, G. P., & Girard, O. Psychophysiological responses to repeated-
1430 1437 1438	443	sprint training in normobaric hypoxia and normoxia. International J of Sports Phys
1439 1440	444	Perform. 2017;12(1):115-123.
1441 1442	445	13. Thum, J., Parsons, G., Whittle, T., & Astorino, T. (2017). High-intensity interval
1443 1444	446	training elicits higher enjoyment than moderate intensity continuous exercise. PloS one.
1445 1446	447	2017;12(1):e0166299.
1447 1448	448	14. Lind, E., Joehns-Matre, R. R., and Ekkekakis, P. What intensity of physical activity do
1449 1450 1451	449	previously sedentary middle-aged women select? Evidence of a coherent pattern from
1452 1453	450	physiological, perceptual and affective markers. Prev Med. 2005;40:407-419.
1454 1455	451	15. Friedmann, B., Bauer, T., Menold, E. and Bärtsch, P. Exercise with the intensity of the
1456 1457	452	individual anaerobic threshold in acute hypoxia. Med Scie Sports Exerc.
1458 1459	453	2004;36(10):1737-1742.
1460 1461	454	16. Abbiss, C. R., Peiffer, J. J., Meeusen, R. and Skorski, S. Role of ratings of perceived
1462 1463 1464	455	exertion during self-paced exercise: what are we actually measuring? Sports Med.
1464 1465 1466	456	2015;45(9):1235-1243.
1467 1468	457	17. Ochi, G., Yamada, Y., Hyodo, K., Suwabe, K., Fukuie, T., Byun, K., Dan, I. and Soya,
1469 1470	458	H. Neural basis for reduced executive performance with hypoxic exercise. Neuroimage.
1471 1472	459	2018;171:75-83.
1473 1474 1475		25

1476		
1477		
1478 1479	460	18. McMorris, T., Hale, B. J., Barwood, M., Costello, J. and Corbett, J. Effect of acute
1480 1481	461	hypoxia on cognition: A systematic review and meta-regression analysis. Neuro
1482 1483 1484	462	Biobehavioral Revs. 2017;74:225-232.
1485 1486	463	19. Seiler, S. and Sjursen, J. E. Effect of work duration on physiological and rating scale
1487 1488	464	of perceived exertion responses during self-paced interval training. Scand J Med Sci
1489 1490	465	Sports. 2004;14(5):318-325.
1491 1492	466	20. Jeffries, O., Patterson, S. D., & Waldron, M. The effect of severe and moderate hypoxia
1493 1494	467	on exercise at a fixed level of perceived exertion. Euro J App Phys. 2019;1-12.
1495 1496 1497	468	21. Martin, P. E., Rothstein, D. E., Larish, D. D. Effects of age and physical activity status
1497 1498 1499	469	on the speed-aerobic demand relationship of walking. J Appl Phys 1992;73:200–206.
1500 1501	470	22. Borg, G. A. V. Perceived exertion as an indicator of somatic stress. Scand J Rehab Med.
1502 1503	471	1970;2:92-98.
1504 1505	472	23. Laurent, C.M., Green, J. M., Bishop, P. A., Sjökvist, J., Schumacker, R. E., Richardson,
1506 1507	473	M. T. and Curtner-Smith, M. A practical approach to monitoring recovery:
1508 1509 1510	474	development of a perceived recovery status scale. J Strength Cond Res.
1510 1511 1512	475	2011;25(3):620-628.
1513 1514	476	24. Crewther, B. T., Carruthers, J., Kilduff, L. P., Sanctuary, C. E. and Cook, C. J.
1515 1516	477	Temporal associations between individual changes in hormones, training motivation
1517 1518	478	and physical performance in elite and non-elite trained men. Bio Sport. 2016;33(3):215.
1519 1520	479	25. Ward, S. A. and Whipp, B. J. Effects of peripheral and central chemoreflex activation
1521 1522	480	on the isopnoeic rating of breathing in exercising humans. J Phys. 1989;411(1):27-43.
1523 1524 1525	481	26. Stroop, J. R. Studies of interference in serial verbal reactions. J Exp Psychol.
1525 1526 1527	482	1935;18:643-662.
1528 1529	483	27. Cohen, J. Statistical power analysis for the behavioral sciences, 2nd ed. 1988. New
1530 1531	484	Jersey: Lawrence Erlbaum Associates.
1532 1533 1534		26

1535 1536		
1537 1538	485	28. da Silva, D. F., Simões, H. G. and Machado, F. A., vVO2max versus Vpeak, what is
1539 1540	486	the best predictor of running performances in middle-aged recreationally-trained
1541 1542	487	runners?. Science & Sports. 2015;30(4):e85-e92.
1543 1544	488	29. Smith, K. J. and Billaut F. Influence of cerebral and muscle oxygenation on repeated-
1545 1546 1547	489	sprint ability. Eur J Appl Physiol. 2010;109(5):989-99
1548 1549	490	30. Dufour, S. P., Ponsot, E., Zoll, J., Doutreleau, S., Lonsdorfer-Wolf, E., Geny, B.,
1550 1551	491	Lampert, E., Fluck, M., Hoppeler, H., Billat, V. and Mettauer, B. Exercise training in
1552 1553	492	normobaric hypoxia in endurance runners. I. Improvement in aerobic performance
1554 1555 1556	493	capacity. J App Phys. 2006;100(4):1238-1248.
1557 1558	494	31. Pramsohler, S., Burtscher, M., Faulhaber, M., Gatterer, H., Rausch, L., Eliasson, A. and
1559 1560	495	Netzer, N. C. Endurance Training in Normobaric Hypoxia Imposes Less Physical Stress
1561 1562	496	for Geriatric Rehabilitation. Front Phys, 2017;8:514.
1563 1564	497	32. Haufe, S., Wiesner, S., Engeli, S. et al. Influences of normobaric hypoxia training on
1565 1566	498	metabolic risk markers in human subjects. Med Sci Sports Exerc. 2008;40(11):1939-
1567 1568 1569	499	1944.
1570 1571	500	33. Wiesner, S., Haufe, S., Engeli S, et al. Influences of normobaric hypoxia training on
1572 1573	501	physical fitness and metabolic risk markers in overweight to obese subjects. Obesity
1574 1575	502	(Silver Spring). 2010;18(1):116–120.
1576 1577	503	34. Krejčí, J., Botek, M. and McKune, A.J., 2018. Dynamics of the heart rate variability
1578 1579	504	and oxygen saturation response to acute normobaric hypoxia within the first 10 min of
1580 1581 1582	505	exposure. Clin Phys Funct Imag. 2018;38(1):56-62.
1583 1584	506	35. Van Beekvelt, M. C., Colier, W. N., Wevers, R. A., Van Engelen, B. G. Performance
1585 1586	507	of near-infrared spectroscopy in measuring local O ₂ consumption and blood flow in
1587 1588	508	skeletal muscle. J Appl Physiol. 2001;90:511–519.
1589 1590 1591 1592 1593		27

1594		
1595		
1596 1597	509	36. Joyner, M. J. and Casey, D. P. Regulation of increased blood flow (hyperemia) to
1598 1599	510	muscles during exercise: a hierarchy of competing physiological needs. Phys Revs.
1600 1601 1602	511	2015;95(2):549-601.
1602 1603 1604	512	37. Howlett, R. A., Heigenhauser, G. J., Hultman, E., Hollidge-Horvat, M. G. and Spriet,
1605 1606	513	L. L. Effects of dichloroacetate infusion on human skeletal muscle metabolism at the
1607 1608	514	onset of exercise. Am J Phys Endo Met. 1999;277(1):E18-E25.
1609 1610	515	38. Renfree, A., Martin, L., Micklewright, D., and Gibson, A. S. C. Application of decision-
1611 1612	516	making theory to the regulation of muscular work rate during self-paced competitive
1613 1614	517	endurance activity. Sports Med. 2014;44:147-158.
1615 1616 1617	518	39. Stork, M. J., Gibala, M. J. and Martin, K. G. Psychological and Behavioral Responses
1618 1619	519	to Interval and Continuous Exercise. Med Scie Sports Exerc. 2018;50(10):2110-2121.
1620 1621	520	40. Subudhi, A. W., Dimmen, A. C., & Roach, R. C. Effects of acute hypoxia on cerebral
1622 1623	521	and muscle oxygenation during incremental exercise. J App Phys. 2007;103(1):177-
1624 1625	522	183.
1626 1627	523	41. Subudhi, A. W., Miramon, B. R., Granger, M. E., & Roach, R. C. Frontal and motor
1628 1629	524	cortex oxygenation during maximal exercise in normoxia and hypoxia. J App Phys.
1630 1631 1632	525	2009;106(4):1153-1158.
1633 1634	526	42. Christian, R. J., Bishop, D., Girard, O., & Billaut, F. The role of sense of effort on self-
1635 1636	527	selected cycling power output. Front in Phys. 2014;5:115.
1637 1638	528	43. Lambourne, K. and Tomporowski, P. The effect of exercise-induced arousal on
1639 1640	529	cognitive task performance: a meta-regression analysis. Brain Res. 2010;1341:12-24.
1641		
1642 1643		
1644		
1645		
1646		
1647		
1648		
1649 1650		
1651		28
1652		20