

Post-exercise hypotension after exercising in hypoxia with and without tart cherry supplementation

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1	Post-exercise hypotension after exercising in hypoxia with and without tart cherry
2	supplementation
3	
4	Short running title: Hypoxia and tart cherry effects on post-exercise hypotension
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20 Abstract

21 Background: This study investigated the effects of hypoxic exercise with and without tart cherry supplementation on post-exercise hypotension (PEH). Method: In a randomized order, 22 12 healthy young adults (9 men and 3 women) completed cycle exercise to exhaustion i) in 23 normoxia without any supplementation (Norm), ii) in hypoxia (13% O₂) with placebo (Hypo), 24and iii) in hypoxia with tart cherry supplementation (Hypo+TC). Supplements were supplied 25 for 5 days pre-trial (TC was 200 mg anthocyanin per day for 4 days and 100 mg on day 5). 26 27 Results: Cycle exercise total energy expenditure was greater in Norm than Hypo and Hypo+TC (P<0.001) with no difference between Hypo and Hypo+TC (P=0.41). Mean arterial pressure 28 29 (MAP) decreased during recovery in all trials (main effect of time, P<0.001), with no difference in PEH between the trials (P>0.05, change (Δ) in MAP from pre-exercise at 60 min recovery, 30 mean difference, Norm Δ -4.4 mmHg, Hypo Δ -6.1 mmHg, and Hypo+TC Δ -5.2 mmHg). 31 Cardiac baroreflex sensitivity decreased during recovery in all trials (P<0.001) and was lower 32 in Hypo than Norm and Hypo+TC (main effect of trial, P=0.02). Conclusion: Post-exercise 33 hypotension was not increased after exercise in hypoxia, with or without tart cherry 34 supplementation, compared to exercise in normoxia. 35

Keywords: baroreflex sensitivity, hypoxic vasodilation, mean arterial pressure, polyphenol
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39 Introduction

40 Arterial blood pressure (BP) is reduced for up to 24 h following a single session of physical exercise; a phenomenon called "Post Exercise Hypotension (PEH)" (Halliwill et al., 2014). It 41 42 is clinically important to investigate factors that enhance PEH as the magnitude of PEH after acute exercise relates to the beneficial BP-lowering effects of exercise training (Kleinnibbelink 43 et al., 2020). While various factors such as exercise mode, intensity, and duration, and 44 environmental temperature may influence PEH, few studies have investigated the effect of 45 46 hypoxia on PEH (Halliwill et al., 2014; Horiuchi and Oliver, 2023). PEH follows a decrease in peripheral vascular resistance (Brito et al., 2014), and as hypoxia enhances vasodilation (Joyner 47 48 and Casey, 2014), greater PEH may be anticipated after exercising in hypoxia than normoxia, which has been confirmed in some (Horiuchi et al., 2016a; 2018; Saito et al., 2019), but not all 49 previous studies (Fornasiero et al., 2021; Horiuchi et al., 2022; Kleinnibbelink et al., 2020). BP 50 may not be reduced after exercise in hypoxia due to an attenuation of baroreflex sensitivity 51 52 (BRS) and a shift in cardiac autonomic function to sympathetic activity (Bourdillon et al., 2023; Halliwill et al., 2014). 53

Tart cherries, and other dark-coloured berries, are rich in antioxidants and polyphenols including anthocyanins (Keane et al., 2016). In normoxic conditions, anthocyanin-rich supplements have been shown to increase peripheral artery diameter and blood flow (Barnes et al., 2020; Cook et al., 2023; Matsumoto et al., 2005), and reduce peripheral vascular resistance

58	(Barnes, 2020), which precedes PEH (Halliwill et al., 2014). Moreover, a recent study in
59	normoxia reported a larger decrease in post-exercise systolic blood pressure, but not diastolic
60	or mean arterial pressure (MAP), following 7 days of an anthocyanin-rich supplement
61	compared to a placebo (Shan and Cook, 2023). These vascular effects may be mediated by
62	polyphenols and circulating metabolites' improving nitric oxide bioavailability (Bell and
63	Gochenaur, 2006; Xu et al., 2004) and reducing oxidative stress, which is elevated post-exercise
64	and in hypoxic environments. PEH may also be expected to be greater after hypoxic exercise
65	and anthocyanin-rich supplementation compared to normoxic or hypoxic exercise alone, as
66	antioxidant supplementation has previously been shown to restore the imbalance of cardiac
67	autonomic nervous activity, as assessed by heart rate variability (HRV) in humans (Weggen et
68	al., 2021), and improve BRS in rats (Alves et al., 2015; Garcia et al., 2017).
69	Accordingly, this study investigated the effects of tart cherry (TC) supplementation on
70	PEH after exercising in hypoxia. We hypothesized that the magnitude of PEH would be greater
71	in hypoxia compared to normoxia, and PEH would be further accentuated with TC
72	supplementation.

74 Methods

75 Participants

76 The present report presents additional recovery and normoxia data from previously published

77	investigations that examined tart cherry supplementation effects on hypoxic exercise
78	performance (Horiuchi et al., 2023). This study was approved by the Ethical Committee of
79	Mount Fuji Research Institute in Japan and was performed following Declaration of Helsinki
80	guidelines (No. 202001). Of the 13 participants in the previous study, 12 (9 men and 3 women)
81	performed an additional normoxic exercise and recovery experimental trial. The participants'
82	age, height, and body mass were 21 \pm 1 years, 169 \pm 7 cm, and 62.1 \pm 8.9 kg, respectively
83	(values are mean \pm standard deviation [SD]). All participants were non-smokers, had no history
84	of cardiovascular disease, and had not been exposed to an altitude higher than 1,500 m in the 6
85	months before the study.

87 Study design

88 This study consisted of three trials (Figure 1): (1) normobaric normoxic exercise without any 89 supplementation (Norm); (2) normobaric hypoxic exercise (13% O₂) with a placebo (Hypo), 90 and (3) normobaric hypoxic exercise (13% O₂) with TC supplementation (Hypo+TC). In a 91 double-blinded and randomized manner, each participant ingested a placebo or TC capsule (Tart 92 cherry 1200 mg containing 100 mg of anthocyanin, Nature's Life, Orem, UT, USA) twice per 93 day for 4 days before the experimental trial, and once on the day of the experimental trial 2 h before beginning excise, which is consistent with studies reporting hemodynamic changes after 94 95 single doses and 4-7 days of anthocyanin-rich supplementation (Matsumoto et al., 2005).

96 Participants were provided a list of antioxidant-rich foods and instructed to avoid these while
97 in the study.

98

99 Experimental procedur

100 The exercise was performed on a cycle ergometer (COMBI232-C, COMBI, Japan) in an 101 environmental chamber (24 C°, 50% relative humidity, TBR-4, 5SA2GX, Tabai Espec Co. Ltd., Tokyo, Japan). After a 15-minute semi-recumbent rest, participants performed incremental leg 102 cycling exercise to exhaustion, consisting of three 4 min incremental stages (40-80-120 Watts 103 104 [W] for men, and 30-60-90 W for women, with each stage lasting 3 min), followed by an 105 increase in workload of 20 W (men) or 10 W (women) per min until exhaustion. The pedal cadence was set at 60 rpm using a metronome. After exhaustion, the participants sat semi-106 107 recumbent for 60 minutes in normoxia in all trials. 108

109 Measurements

At rest and during exercise, pulmonary oxygen uptake (VO₂) and carbon dioxide output (VCO₂) were measured by a metabolic cart (AE-310S, Minato Medical Science, Osaka, Japan) and beatby-beat BP was measured using finger photoplethysmography at the middle or index finger (MUB-101; Medisens Inc., Tokyo, Japan) as the time-averaged from the beat-by-beat pressure wave (Horiuchi et al., 2016b). Beat-by-beat BP data were stored with a sampling frequency of

115	200 Hz by a field data recorder (es8; TEAC, Tokyo, Japan), and transferred to a laptop computer
116	for further analysis. Based on a previous study (Horiuchi and Thijssen, 2020), heart rate (HR)
117	was measured using a portable HR monitor (Check-My-Heart, TRYTECH Co., Ltd., Tokyo,
118	Japan), and HRV was calculated by accompanying HRV analysis software. Participants were
119	instructed to breathe normally throughout testing. Fingertip blood samples (0.3 μ L) were taken
120	to measure blood lactate concentration (Lactate Pro 2LT-1730; Arkray, Tokyo, Japan) pre-
121	exercise, 5, 20, and 60 min of recovery. Total urine samples were collected pre-exercise and 1
122	h post-exercise and analyzed for urinary 8-hydro-2' deoxyguanosine (8-OHdG), an index of
123	oxidative DNA damage, as described previously (Horiuchi et al., 2023).

125 Data Analysis

To calculate spontaneous cardiac BRS (cBRS), the beat-to-beat systolic BP (SBP) time series 126 127 and RR interval were analyzed for more than 3 consecutive beats, with increasing or falling 128 direction from a 5-min steady-state data segment at rest and during recovery (Carrington and 129 White, 2001; Horiuchi and Oliver, 2023; Ogoh et al., 2005). Linear regression was applied to 130 each baroreflex sequence, with only sequences with an $R^2 > 0.85$ accepted (Horiuchi and Oliver, 131 2023; Iellamo et al., 1994). The overall average slope of the SBP-RR interval was calculated as spontaneous cBRS. Time domain HRV was calculated by the standard deviation of the 132 133 normal-to-normal intervals (SDNN) and the root-mean-square of successive differences in R- 134 R interval (RMSSD). In the frequency domain, the extent of very low-frequency oscillations (0.0033-0.04 Hz), low-frequency oscillations (LF: 0.04-0.15 Hz), and high-frequency 135 oscillations (HF: 0.15–0.4 Hz) was quantified using a fast Fourier transformation (Horiuchi and 136 Thijssen, 2020). Total exercise energy expenditure (EE) was calculated using \dot{VO}_2 and \dot{VCO}_2 137 as follows: Total EE (J s⁻¹) = $(3.869 \times \dot{VO}_2) + (1.195 \times \dot{VCO}_2) \times 4.168 / 60 \times 1000$ 138 where, the unit of VO_2 and VCO_2 were liter per minute (Horiuchi et al., 2017). 139 140 141 **Statistics** 142 Data are presented mean \pm SD. Statistical analyses were performed using commercial software

143 (Jamovi, 3.2.3). One-way repeated measures analysis of variance (ANOVA) compared the total EE across the three trials, and changes in urinary 80HdG excretion. A two-way (time × trials) 144repeated ANOVA compared time course changes in all physiological variables (BPs, HR, HRV, 145 and blood lactate). For further comparisons, Tukey's post hoc test was used. Effect size was 146 calculated as η^2 , defined as small ($\eta^2 = 0.01$), medium ($\eta^2 = 0.06$), and large ($\eta^2 = 0.14$) (Lakens, 147 2013). Statistical significance was set at P < 0.05. The normality of the data was examined 148 using the Bartlett and Levene test. If equal variance failed, logarithmic transformation data were 149 used for further analysis (HF and LF/HF). 150

151

152 Results

153 Cycle exercise total EE was detected to be different between the trials (F=34.5, P<0.001, 154 η^2 =0.21), where total exercise EE in Norm (846±189 J s⁻¹) was greater than Hypo (672±125 J 155 s⁻¹) and Hypo+TC (692±153 J s⁻¹) (P<0.001, respectively), with no differences detected 156 between Hypo and Hypo+TC (P=0.41).

157

During the 60 min recovery, an interaction effect was found for MAP (F=1.86, *P*=0.045, η^2 =0.013), but not for SBP and DBP (**Figure 2**). Mean arterial pressure decreased in all trials (main effect of time, F=14.51, *P*<0.001, η^2 =0.15), with no difference detected in PEH between trials (*P*>0.05, change (Δ) in MAP from pre-exercise at 60 min recovery, mean difference [95% confidence interval], Norm Δ -4.4 [-6.0, -2.8] mmHg, Hypo Δ -6.0 [-8.5, -3.7] mmHg, and Hypo+TC Δ -5.2 [-8.8, -1.6] mmHg, **Figure 2A**).

164

165 Cardiac BRS was reduced during recovery compared to pre-exercise (main effect of time, 166 F=59.55, P<0.001, $\eta^2=0.62$). Moreover, a main effect of trial was detected (F=4.45, P=0.02, 167 $\eta^2=0.02$), where overall cBRS was lower in Hypo than Norm (P=0.03) and Hypo+TC (P=0.06), 168 with no difference between Norm and Hypo+TC (P=0.74, **Figure 3A**). No trial or time effects 169 were detected for HR. An interaction was detected for HR due to higher resting HR on Hypo 170 and Hypo+TC than Norm (F=2.29, P=0.01, $\eta^2=0.01$) (**Figure 3B**). There was no interaction or 171 and trial effects in blood lactate (**Figure 3C**). For HRV metrics, no interactions or main effects

172	of time were detected. However, regardless of the trial, cardiac parasympathetic activity indices
173	(SDNN, RMSDD, log [HF]) were lower, and cardiac sympathetic activity index (log [LF/HF])
174	was higher during recovery compared with pre-exercise (Table 1). At 1 h post-exercise, changes
175	in urinary 8-OHdG excretion from pre-exercise were 5.2±4.4 in Norm, 5.3±3.1 in Hypo, and
176	3.4±2.7 ng kg ⁻¹ h ⁻¹ in Hypo+TC, with a trend for a smaller increase in 8-OHdG excretion on
177	Hypo+TC than Hypo (<i>P</i> =0.08).

- 178
- 179 **Discussion**

180 Our study showed that incremental leg cycling until exhaustion leads to reductions in 181 MAP of 4-6 mmHg after exercise in untrained men, supporting the presence of PEH. These findings confirm the results of previous studies showing PEH after various exercise intensities, 182 durations, and types (Jones et al., 2021; Marcal et al., 2021; Pimenta et al., 2019). In contrast 183 to our hypothesis, PEH was not increased after exercise in hypoxia, with or without tart cherry 184 supplementation, compared to exercise in normoxia. One possible explanation is the exercise 185 186 was performed until exhaustion, which resulted in greater exercise energy expenditure and absolute work in Norm than Hypo or Hypo+TC. This is consistent with a recent study that 187 revealed the magnitude of PEH was not different between normoxia and hypoxia when the 188 absolute work of exercise was matched (Fornasiero et al., 2021). These findings have good 189 190 ecological validity as those exercising in hypoxic conditions normally reduce workload due to 191 increased perception of effort (Rossetti et al., 2017).

192 Tart cherry supplementation before exercise in hypoxia did not further accentuate PEH compared to exercise in hypoxia alone. These unique findings build upon the limited research 193 194 in normoxia to examine the effect of anthocyanin-rich supplementation on PEH (Shan and Cook, 2023). Consistent with this previous study we reported no difference in MAP or DBP post-195 exercise after placebo and anthocyanin-rich supplementation. In contrast, we did not observe a 196 larger decrease in post-exercise SBP, which may be explained by the different types (tart cherry 197 vs New Zealand blackcurrant) and dose of anthocyanin-rich supplementation (7 vs 4 days, and 198 199 210 vs 100 mg anthocyanin on the final day).

200 In the present study, HRV indices during recovery indicated a shift in cardiac 201 autonomic balance compared to pre-exercise, i.e., increased cardiac sympathetic activity and decreased cardiac parasympathetic activity; however, these indices were not influenced by 202hypoxia or tart cherry supplementation. cBRS was lowest during recovery after exercise in 203 Hypo, which is consistent with previous research indicating hypoxia lowers cBRS (Bourdillon 204 et al., 2023). cBRS was similar during recovery in Hypo+TC to Norm, suggesting tart cherry 205supplementation restored cBRS, lowered by exercise in hypoxia. One possible explanation is 206 oxidative stress tended to be lower after hypoxic exercise with tart cherry supplementation 207 compared to a placebo. This explanation is supported by animal research reporting 208 improvements in baroreflex sensitivity after antioxidant supplementation (Alves et al., 2015; 209

210	Garcia et al., 2017). Improvements in oxidative stress and cBRS sensitivity with tart cherry
211	supplementation at the same time as similar magnitude of PEH in all trials suggests a limited
212	regulatory role of oral antioxidants and cBRS in PEH. Previous research has also shown the
213	intravenous infusion of antioxidants did not influence PEH (Romero et al., 2015). Therefore,
214	non-antioxidant mechanisms, like increased NO bioavailability, may explain the greater
215	reductions in post-exercise BP observed after consuming anthocyanin-rich supplements (Shan
216	and Cook, 2023).
217	
218	Conclusion
219	Post-exercise hypotension was not increased after exercise in hypoxia, with or without
220	tart cherry supplementation, compared to exercise in normoxia.
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222	

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228	manuscript (MH); revising the manuscript (MH, SO); approval of the final manuscript (MH,
229	SO).
230	
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387 Figure legends

Figure 1. Experimental procedure. BP, blood pressure; cBRS, cardiac baroreflex sensitivity;
HRV, heart rate variability; Suppl., supplementation; VO₂, oxygen uptake; VCO₂, carbon
dioxide output



Figure 2. Mean arterial blood pressure (MAP; panel A), systolic blood pressure (SBP: panel
B), and diastolic blood pressure (DBP; panel C) during a 1 h recovery period after exercising
in normoxia (Norm; white circles), hypoxia with placebo (Hypo; black squares), and hypoxia
with antioxidants (Hypo+TC; gray triangles) trials. Values are mean ± standard deviation (SD).
*†‡ indicates a difference compared with the pre-exercise value in Norm, Hypo, and Hypo+
TC trials, respectively.



Figure 3. Cardiac baroreflex sensitivity (cBRS; panel A), heart rate (HR: panel B), and blood
lactate (panel C) during a 1 h recovery period after exercising in Norm (white circles), Hypo
(black squares), and Hypo+TC (gray triangles) trials. Values are mean ± SD. # and \$ indicate
differences compared with Norm trial.

