



## UWS Academic Portal

### **Hemoglobin, hematocrit and plasma volume variations following combined sprint and strength**

Sellami, M.; Ben Abderrahmen, A.; Dhahbi, W.; Hayes, L.D.; Zouhal, H.

*Published in:*  
Science and Sports

*DOI:*  
[10.1016/j.scispo.2019.10.012](https://doi.org/10.1016/j.scispo.2019.10.012)

E-pub ahead of print: 25/01/2020

*Document Version*  
Peer reviewed version

[Link to publication on the UWS Academic Portal](#)

*Citation for published version (APA):*

Sellami, M., Ben Abderrahmen, A., Dhahbi, W., Hayes, L. D., & Zouhal, H. (2020). Hemoglobin, hematocrit and plasma volume variations following combined sprint and strength: effect of advanced age. *Science and Sports*, 36(1), e13-e21. <https://doi.org/10.1016/j.scispo.2019.10.012>

**General rights**

Copyright and moral rights for the publications made accessible in the UWS Academic Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

**Take down policy**

If you believe that this document breaches copyright please contact [pure@uws.ac.uk](mailto:pure@uws.ac.uk) providing details, and we will remove access to the work immediately and investigate your claim.

1 **Summary and Keywords Page**

2 **Hemoglobin, Hematocrit and plasma volume variations following combined sprint**  
3 **and strength training: Effect of advanced age**

4 **Summary**

5 **Objectives:** The study investigated the effect of combined sprint and resistance training  
6 (CSRT) on red blood cell (RBC) count, hemoglobin (Hb), hematocrit (Hct), plasma  
7 volume (PV) variation at rest and during exercise.

8 **Equipment and methods:** Twenty-eight moderately trained were randomly assigned  
9 into a young trained (YT), young control (YC), middle-aged trained (MAT), and middle-  
10 aged control (MAC) group. Before (P1), and after (P2) CSRT, blood samples were  
11 collected at rest and after exercise.

12 **Results:** At P1, Hct was significantly ( $p < .05$ ) greater in young compared to middle-aged  
13 groups. At P1, PV decrease during exercise was significantly ( $p < .05$ ) higher in middle-  
14 aged compared to young groups. Following CSRT, resting RBC count and Hb increased  
15 significantly ( $p < .05$ ) in MAT. At P2, Following CSRT, Hct decreased significantly  
16 ( $p < .05$ ) in trained groups. At P2, no significant ( $p > .05$ ) age- effect between MAT and  
17 YT was observed for Hct. In conclusion, CSRT increases RBC count and Hb in middle-  
18 aged men, and ameliorates the effect of age in Hct. Such adaptations may improve  
19 cardiovascular fitness of middle-aged individuals, and may be preventative of subsequent  
20 declines with age.

21 **Keywords**

22 Blood viscosity; muscle blood flow; red blood cells; sports anemia; training effect.

1 **1. Introduction**

2 It is well known that older adults experience decreased total body water,  
3 baroreceptor sensitivity, cell-producing marrow, and blood viscosity compared to  
4 younger counterparts [1, 2]. Homeostasis observed in older adults is paralleled by  
5 increases in heart rate and blood pressure and underpins a lowered exercise-induced  
6 muscle blood flow [2]. During continuous exercise, oxygen delivery to the mitochondria  
7 of muscle cells represents a main determinant of performance [3]. As such, age-associated  
8 reductions in muscle blood flow contribute to reduced functional capacity in older adults  
9 [4].

10 Acute exercise induces a variation in plasma volume (PV) because of transient  
11 fluid shifts into (hemodilution), and out of (hemoconcentration), the intravascular space  
12 [5]. This change in PV is dependent upon exercise intensity and type [6], posture [7],  
13 ambient temperature [8], and fluid consumption [9]. A reduction in PV may limit  
14 endurance capacity, via reduced blood supply to the working muscle. Moreover, PV  
15 variation (PVV) has the potential to hamper interpretation of plasma biomarkers [5] as  
16 specific quantities of plasma are required for common assays. In addition, PV influences  
17 end diastolic volume, stroke volume, and therefore cardiac output [10].

18 Whilst the PV changes during acute exercise are well defined, the chronic  
19 adaptation to exercise training is currently poorly understood. Some authors suggest that  
20 magnitude of change of PV and hematocrit is dependent to exercise intensity and training  
21 type (strength, aerobic, sprint) [11]. An increase in blood volume, red cell mass, and PV  
22 and a decrease in hematocrit (Hct) following continuous aerobic and intermittent interval  
23 exercise training in young athletes has been observed [11]. However, others found no

24 changes in PV, Hct, hemoglobin (Hb), and red blood cell (RBC) count in young and older  
25 (>50 years) men after prolonged aerobic or resistance training (RT) [1].

26 Numerous studies have found greater PV variation following chronic sprint  
27 training [12], and in young sprinters compared with endurance athletes [11]. In addition,  
28 Vechin and colleagues [13] found greater improvements in muscle blood flow following  
29 high-intensity RT when compared with low and moderate intensity training in elderly  
30 participants. Ahmadizad and colleagues [14] observed slight differences in Hct, RBC  
31 count, and blood viscosity between young, middle-aged, and old males following  
32 endurance exercise. However, Bongers and colleagues [15] recently observed no  
33 differences in PV changes between octogenarians and sexagenarians following a 30 km  
34 march.

35 High intensity training (HIT) involves repeated bouts of high-intensity exercise,  
36 interspersed with recover periods, proclaimed as a time-efficient “healthogenic” strategy  
37 [16] despite falling short of the recommended exercise volume to improve and maintain  
38 cardiovascular health [16]. Given recent interest in HIT, it is imperative to determine  
39 PVV following this training type to allow appropriate interpretation of serum biomarkers,  
40 changes in fluids balances, and cardiac output following this exercise modality.

41 Whilst PVV has been quantified in younger cohorts following HIT [17], the effect  
42 of HIT and age on PVV is yet to be examined. Therefore, the main aim of the present  
43 study was to investigate the effect of HIT on PVV, Hct, RBC count, and Hb in young and  
44 middle-aged participants. It was hypothesized *a priori* that an age effect would exist pre-  
45 training, and this effect would be ameliorated post-training.

46

47 **2. Materials and methods**

48 **2.1. Participants**

49 Twenty-eight moderately trained men were recruited for participation in the  
50 present study. Eligible participants were subsequently randomized to receive 13-weeks'  
51 combined sprint and resistance training (CSRT), or control. Thus, four groups existed: a  
52 young trained group (YT, age:21.2±1.2 years, height:179.5±4.2 cm; n=7), a young control  
53 group (YC, age: 21.5±2.5 years, height:179.8±6.8 cm; n=7), middle-aged trained group  
54 (MAT, age:40.8±1.8 years, height: 176.3±6.7 cm; n=7) and middle-aged control group  
55 (MAC, age:40.9±2.1years, height:175.2±5.2 cm; n=7). Participants gave their written  
56 informed consent to participate in the study after receiving a thorough explanation of the  
57 study's protocol. The protocol conformed to internationally-accepted policy statements  
58 regarding the use of human participants in accordance with the Declaration of Helsinki  
59 and was approved by the University's Ethics Committee. Training status was assessed  
60 using an adapted version of the Baecke questionnaire, to identify those with a medical  
61 contraindication to performing specific assessments, participants completed medical  
62 history, and 3-day-food record. Inclusion criteria included no contraindications to  
63 maximal exercise testing such as cardiovascular or pulmonary risk factors, no history of  
64 chronic disease, illness, surgeries, hospitalizations, and musculoskeletal or joint injuries.

65 During design of the study, statistical power analysis was performed to determine  
66 sample size. This procedure showed that seven participants for each group was needed to  
67 achieve a statistical power of 80% and detect a small effect ( $d=0.2$ ) when assessed by  
68 four-factor mixed analysis of variance (ANOVA) with a level of significance of 5%.

69

## 70 **2.2. Study design**

71 A randomized controlled trial study design was used. This study investigated the  
72 effects of HIT on PVV, Hct, RBC count and Hb in young and middle-aged participants.  
73 Trained participants participated in 13-weeks of CSRT. Briefly, CSRT consisted of one  
74 sprint running, one sprint cycling, and one RT session per week, separated by a minimum  
75 of 48 h (13 sessions of each training unit). Each age group (young and middle-aged) was  
76 randomly divided between control (n = 7) and trained (n = 7) groups. Data were collected  
77 before starting training, and immediately after the 13<sup>th</sup> week. On both occasions, data  
78 were collected in the same conditions, at the same time of day. The protocol included the  
79 Astrand-Ryhming test, a repeated sprint cycling test, the Wingate Anaerobic Test with  
80 concomitant heart rate measurement, a lactate threshold test, and systolic and diastolic  
81 blood pressure and hematological markers levels (more details below).

82

## 83 **2.3. Evaluation and Procedures**

84 Sessions were performed during the morning and lasted no longer than 70 min,  
85 inclusive of 15 min warm-up (jogging and stretching) and 15 min cool-down (jogging  
86 and stretching). Sprint running sessions entailed 3-5 sets of 3-5 short bouts at maximum  
87 velocity. A recovery of 2-3 min was permitted between each set. Sprint cycling sessions  
88 comprised 3-5 repetitions of 10-30 s. The 10-30 s trials were performed maximally.  
89 Participants recovered actively (50%  $\text{VO}_{2\text{max}}$ ) for 3-5 min between each sprint. RT  
90 sessions entailed 5-6 exercises targeting all major muscle groups. The load used during  
91 exercise was progressively increased from 40% to 65% of one-repetition maximum (1-  
92 RM) [18]. To produce maximal power output (in other words; velocity  $\times$  load), the

93 concentric phase of each exercise was performed as fast as possible [19]. Repetitions were  
94 maintained at 10-15 per sets and the number of sets increased from 3 to 4 during the  
95 training period. Hence, training volume increased progressively during the CSRT  
96 program. Rest periods between sets were 3-5 min for upper body muscles [19] and a  
97 minimum of 1 min for lower limbs. To adjust load during RT session and monitor  
98 adaptation, we determined strength using a 1-RM for the six resistance exercises, pre-  
99 training (P1), during the sixth week, and post-training (P2).

100

### 101 2.3.1. Testing Schedule

102 During experimental period, participants completed anthropometric  
103 measurements (pre-, mid-, and post training) and a dietary assessment using a 3-day food  
104 record by a sports nutritionist. One week before training-cycle, participants were  
105 familiarized with testing procedures to minimize learning effect. Participants avoided  
106 physical activity for 48 h preceding each test. The testing period was divided into two  
107 phases: before (P1), and after (P2) training and included three consecutive laboratory  
108 visits separated by 48h. P2 commenced 48 h after training cessation and finished 7-days  
109 later.

110 On day 1, participants performed the Astrand-Ryhming test on a cycle ergometer  
111 to estimate maximal oxygen uptake ( $VO_{2max}$ ). On day 2, participants performed a repeated  
112 sprint cycling test on a cycle ergometer. It consisted of five short trials (6 s) against  
113 increasing resistance (2 kg per sprint) until exhaustion and when the velocity began to  
114 decrease during the 6 s trials. Recovery time between each trial was 5 min. On day 3,  
115 participants performed the WAnT on a mechanically braked Monark cycle ergometer.

116

117 *2.3.2. Physiological parameters*

118 Systolic (SBP) and diastolic (DBP) blood pressure were measured in a sitting  
119 position. Heart rate variability during WAnT was also measured continuously using Heart  
120 rate monitor.

121 During day 3, blood samples were collected to determine hematological markers.  
122 Upon arriving, a heparinized catheter (Insyte-W, 1.1 mm o.d. × 30 mm) was inserted into  
123 an antecubital vein, following 20-min sitting. Blood was drawn 8:00-9:00 h following  
124 overnight fasting. Venous blood samples were drawn at four times: rest (<sub>0</sub> [after 20 min  
125 sitting on the bike]), after warm-up, immediately post-WAnT (<sub>end</sub>) and 10 min post-  
126 WAnT (<sub>10</sub>). Hct and [Hb] were determined directly in quadruplicate, automatically by  
127 using standard laboratory procedures. PVV was calculated using Dill and Costill [20]  
128 method.

129

130 *2.4. Statistical Analysis*

131 Data analyses were performed using SPSS version 23.0 for Windows (SPSS, Inc.  
132 Chicago, IL, USA). Means and SD were calculated after verifying the normality of  
133 distributions using the Kolmogorov-Smirnov procedure. For anthropometric,  
134 physiological, and physical performances indices, data were analyzed using a  
135 multifactorial three-way (time [P1, P2] × age [young, middle-aged] × group [trained,  
136 control]) ANOVA and Fisher “F” value was given. Blood variables changes were  
137 analyzed using a four-factor ANOVA (time [P1, P2] × Wingate time [warm-up,  
138 immediately post-WAnT and 10 min post-WAnT] × age [young, middle-aged] × group  
139 [trained, control]). To help protect against type II errors, an estimate of power ( $\hat{\omega}$ ) and



140 effect size ( $\eta^2_p$ ) were calculated. Bonferroni-adjusted pairwise post hoc comparisons were  
141 performed where appropriate. Pearson's product-moment correlation coefficients were  
142 calculated to assess relationships between variables. Significance level was fixed to  
143  $p < .05$ .

144

### 145 3. Results

#### 146 3.1. Morphological Data and Physical Performances

147 For body mass ((BM) kg), there was no significant age-effect ( $F=1.61$ ,  $p=0.26$ ,  
148  $\eta^2_p=0.11$ ) at P1 or P2. Following CSRT, both training groups experienced a decrease in  
149 BM ( $72.8 \pm 6.3$  to  $70.9 \pm 6.7$  kg for YT and  $73.0 \pm 12.5$  to  $72.3 \pm 10.6$  kg for MAT respectively  
150 with  $F= 8.79$ ,  $p < 0.001$ ,  $\eta^2_p=0.27$ ).

151 At P1, there was no age-effect for body fat percentage (BF %) ( $11.6 \pm 3.1\%$ ,  
152  $10.4 \pm 2.4\%$ ,  $12.3 \pm 1.6\%$  and  $12.5 \pm 1.4\%$  for YT, YC, MAT and MAC respectively with  
153  $F=2.33$ ,  $p=0.16$  and  $\eta^2_p=0.09$ ). Following CSRT, both training groups experienced a  
154 decrease in body fat from P1 ( $10.3 \pm 5.5\%$  and  $10.4 \pm 1.1\%$  for YT and MAT respectively  
155 with  $F= 10.32$ ,  $p < 0.001$ ,  $\eta^2_p=0.28$ ), while the control groups' body fat percentages were  
156 not significantly different from P1 ( $p > .05$ ).

157 At P1, there was an age-effect for fat-free mass ( $F=??$ ,  $p < 0.001$ ,  $\eta^2_p=??$ ).  
158 Following CSRT, fat-free mass increased significantly ( $F= 8.21$ ,  $p=0.03$ ) only in MAT  
159 and was  $63.9 \pm 5.3$  kg.

Commented [LH1]: Please change as according.

160 For estimated  $VO_{2max}$ , there was no significant age-effect ( $F= 2.64$ ,  $p=0.15$ ,  
161  $\eta^2_p=0.32$ ), but we observed a significant effect of time ( $F=17.35$ ,  $p < 0.001$ ,  $\eta^2_p=0.30$ ). In  
162 fact, estimated  $VO_{2max}$  increased significantly ( $p < .001$ ) after CSRT in both trained  
163 groups, but not in control groups ( $p > .05$ ).

Commented [LH2]: From what at P1?

164  $W_{\text{peak}}$  during the WAnT exhibited a significant effect of age for at P1 ( $F= 8.32$ ,  
165  $p<.001, \eta^2_{\text{P}}= 0.99$ ), which was ameliorated at P2 ( $p>.05$ ).  $W_{\text{peak}}$  was significantly ( $F=5.88$ ,  
166  $p=0.02, \eta^2_{\text{P}}=0.25$ ) higher after training in both YT ( $1025\pm 187$  to  $1187\pm 165$  W) and MAT  
167 ( $934\pm 178$  to  $1096\pm 145$  W).

168  $W_{\text{mean}}$  increased significantly only in MAT after CSRT ( $422\pm 56$  to  $560\pm 67$  W).

169 Only at P2,  $[\text{La}]_{\text{peak}}$  increased significantly ( $F=20.12, p<0.001, \eta^2_{\text{P}}=0.89$ ) in both  
170 trained participants (YT and MAT), while remained stable in their control matched groups  
171 ( $16.7\pm 2.1, 16.3\pm 3.6, 14.8\pm 2.8$ , and  $13.1\pm 3.1 \text{mmol}\cdot\text{l}^{-1}$  respectively for YT, YC, MAT and  
172 MAC.

173

### 174 **3.2. Blood Pressure, Heart Rate Characteristics and Hematological Markers**

175 At P1, a significant age effect ( $F=5.43, p=0.02, \eta^2_{\text{P}}=0.64$ ) was observed in systolic  
176 blood pressure (SBP). After CSRT, there was a training effect ( $F=9.43, p=0.03, \eta^2_{\text{P}}=0.98$ )  
177 decrease in SBP in MAT at rest and at the end of exercise. Post-hoc and pairwise  
178 comparisons were represented in Table 1.

179 *\*\*\*Insert Table 1 here\*\*\**

180 RBC ( $10^{12}\cdot\text{L}^{-1}$ ) levels are described in Table 2. There was no significant age-  
181 effect ( $F=2.09, p=0.16, \eta^2_{\text{P}}=0.06$ ) at P1 and P2. In addition, a significant ( $F=14.50$ ,  
182  $p=0.04, \eta^2_{\text{P}}=0.86$ ) effect of training was observed in MAT but not ( $p>.05$ ) in the other  
183 three groups. Post-hoc and pairwise comparisons were represented in Table 2.

184 *\*\*\*Insert Table 2 here\*\*\**

185 Hemoglobin (Hb) concentration (g/100ml) levels are represented in Table 3.  
186 There was no significant age-effect ( $F=1.87, p>.05, \eta^2_{\text{P}}=0.15$ ). However, a slight increase  
187 ( $F=9.10, p<.001, \eta^2_{\text{P}}=0.34$ ) in basal Hb concentration were observed in MAT after CSRT.

188 \*\*\*Insert Table 3 here\*\*\*

189 Hematocrit (Hct) changes (%) are represented in Table 4. A significant effect of  
190 time ( $F=13.50$ ,  $p<0.001$ ,  $\eta^2_p=0.92$ ), Wingate time ( $F=14.12$ ,  $p<0.001$ ,  $\eta^2_p=0.10$ ), age  
191 ( $F=8.55$ ,  $p<0.001$ ,  $\eta^2_p=0.06$ ) and also group ( $F=9.21$ ,  $p<0.001$ ,  $\eta^2_p=0.85$ ) was present.  
192 For YT, Hct<sub>0</sub>, Hct<sub>w</sub>, and Hct<sub>end</sub> were significantly lower at P2 as compared to P1  
193 ( $p<0.001$ ). For MAT, Ht<sub>w</sub>, and Hct<sub>end</sub> were significantly lower at P2 as compared to P1  
194 ( $p<0.001$ ). Moreover, Hct<sub>0</sub> were significantly higher ( $p<0.001$ ) in YT as compared to YC  
195 at P2 (see Table 4).

196 \*\*\*Insert Table 4 here\*\*\*

197 During WAnT, plasma volume decreased significantly ( $p<0.001$ ) from warm-up  
198 (PVV<sub>w</sub>) to the end of the WAnT (PVV<sub>end</sub>) in all groups, then increased from the WAnT  
199 to recovery time (PVV<sub>10</sub>) at P1 and P2 ( $p<.05$ ) (Table 5). This decrease of PVV during  
200 exercise was significantly ( $p<.05$ ) greater in middle-aged groups compared to younger  
201 groups at P1.

202 During warm-up and WAnT, the PVV decrease was significantly ( $p=0.04$ ) higher  
203 in MAT as compared to young groups at P1 (in other words; PVV<sub>w</sub>:  $-8.19\pm 2.88\%$  for YT  
204 vs.  $-12.75\pm 7.41\%$  for MAT,  $F=10.31$ ,  $p=0.02$ ,  $\eta^2_p=0.14$ ). Significant increases in PVV  
205 were observed in YT and MAT following CSRT ( $p<0.001$ ). The age effect was not  
206 present at P2 ( $F=1.25$ ,  $p=0.35$ ,  $\eta^2_p=0.03$ ) between YT and MAT, whilst, for the control  
207 groups, the age-effect remained statistically significant ( $F=8.46$ ,  $p<0.001$ ,  $\eta^2_p=0.12$ ).

208 \*\*\*Insert Table 5 here\*\*\*

209

#### 210 4. Discussion

211           The main finding of the current study was the increased resting RBC count and  
212 Hb in MAT after CSRT. A decrease in Hct in response to WAnT was observed in trained  
213 groups following CSRT with a reduction in age-related difference between age-groups.  
214 PVV changes suggest a moderate to high increase in PV after warm-up, at the end of the  
215 WAnT, and during recovery in the trained groups. Furthermore, the age-related effect on  
216 PVV during exercise was not seen between groups after training. In addition, we observed  
217 decreased SBP in MAT with diminution in age-related difference between trained groups  
218 after CSRT. Although the increase in PV following exercise training is well known [21,  
219 22], we believe we are the first to describe a change to acute PVV pre- and post-training  
220 in middle-aged men after training exercise. Ben Abderrahman and colleagues [22],  
221 described increased PV in 15 young males following interval training, which is consistent  
222 with the present investigation. Moreover, they observed greater PVV following training,  
223 which is consistent with the present study in observing increases in PVV following CSRT.  
224 Our finding that the age-effect was not present post-CSRT suggests that CSRT can  
225 improve haematological regulation in middle-aged individuals, to the point where it is  
226 similar to young adults.

227           An aim of the present study was to provide insight into the effect of acute and  
228 chronic intense exercise on hematological profiles in young and middle-aged men. Sosner  
229 and colleagues [23] reported that training in general resulted in an increase of vagal  
230 parasympathetic activity in the myocardium, an improvement in endothelial function,  
231 resulting in decreased arterial resistances, and improved aortic compliance. As we age,  
232 this aortic compliance declines and arterial resistance increases, leading to higher blood  
233 pressure [24]. Hence, to mediate blood pressure disturbance, the American College of  
234 Sports Medicine (ACSM) Position Stand [25], suggested that aerobic activities and

235 resistance exercises performed 3 times per week are the best alternative to counteract age-  
236 related blood disorders. Rezk and colleagues [26] measured a decrease in SBP in  
237 normotensive young participants following 90-minutes of low intensity RT. RT reduced  
238 BP by 3.2-3.5 mmHg in young trained men [27]. Interestingly, results of the present study  
239 reported greater decreased in SBP (~10 mmHg) in trained groups after CSRT. Typically,  
240 attenuated effects of aerobic exercise on blood pressure are observed in trials lasting 3-6  
241 months, because of poor adherence. As such, Weston and colleagues [28], suggested that  
242 short term HIT (2-3 months) allow greater decline in SBP in individuals with  
243 hypertension.

244 Age-related differences in body fat and fat free mass increase risk of hypertension  
245 and aortic stiffness in older individuals [29]. In the current study, BP improvements  
246 following CSRT were associated with decreased BM and increased FFM in MAT. These  
247 improvements occurred alongside improved anaerobic ( $W_{peak}$  and  $W_{mean}$ ) and aerobic  
248 ( $VO_{2max}$ ) performance following CSRT.

249 Greater acidosis is usually detected in patients with severe depletion of body fluids  
250 [30]. At the end of WAnT, the higher blood lactate was associated with higher PV  
251 decrease in young and middle-aged men. Before intervention, the PV decrease was higher  
252 in middle-aged compared to young groups suggesting a greater fluid depletion during the  
253 WAnT. Interestingly, PV increased after CSRT in both YT and MAT with a reduction in  
254 the age effect on PV. Hence, the PV increase in trained groups after CSRT intervention  
255 suggests 1) improvements in water balance in the extracellular compartment driven  
256 indirectly by the lower blood pressure detected in MAT and 2) better nutrient exchanges  
257 through the compartment leading to low blood viscosity.

258 A decrease in post-training Hct has been detected in trained (endurance or  
259 resistance) participants when compared to untrained ones [11]. Hct decreases are usually  
260 associated with higher red cell mass as well as plasma volume in young endurance-  
261 trained individuals [11] but not following strength training in young and middle-aged men  
262 [31]. In our study, we found that the combination of strength and sprint training improved  
263 resting RBC count and Hb in middle-aged trained group. However, further research is  
264 required to determine underlying mechanisms that decrease Hct and increase RBC count  
265 and Hb following the HIT.

266 The present study is not without limitations. For example, evaluation of water and  
267 sodium status, antidiuretic hormone (ADH) and aldosterone, would have furthered our  
268 understanding of the fluid movements during acute and chronic exercise. However, this  
269 was outside the scope of the present study. Moreover, although changes in the present  
270 study reached statistical significance, they may not be considered clinically meaningful.  
271 However, in the present investigation where RBC count at rest increased by ~25% in  
272 MAT from P1 to P2, this exceeds the critical difference of ~9% determined using flow  
273 cytometry [32]. When resting RBC count decreased by ~20% in YC from P1 to P2 it  
274 exceeded the critical difference, but did not reach statistical significance, suggesting that  
275 an increased sample size should be used in future investigations. Moreover, the difference  
276 between biological and statistical significance should be considered.

277

## 278 **5. Conclusion**

279 In summary, 13 weeks' sprint and resistance training appears to reduce the age-  
280 related decline in substrate metabolism (in other words; lactate) with increased  
281 performance levels during strenuous exercise in middle-aged men. In addition, this

282 training intervention reduced systolic blood pressure in middle-aged trained men at rest  
283 and in response to exercise. These results occurred alongside increased resting RBC  
284 count, Hb, and PV in MAT. Moreover, the age-related differences among groups RBC  
285 count and PV changes following short-term exercise, were reduced at P2. Hence, short-  
286 term intense training with mixed exercises (sprints and resistances) prescription would  
287 allow lower blood pressure for a short period. Typically, individuals with hypertension  
288 have been dissuaded from engaging in long duration interventions and a poor adherence  
289 is usually registered. However, from this study, it appears 13-weeks' exercise training  
290 may be recommended as part of a program that reduces cardiovascular disease risk.

291

#### 292 **Disclosure of interest**

293 The authors declare that they have no competing interest.

#### 294 **Funding**

295 This research did not receive any specific grant from funding agencies in the public,  
296 commercial, or not-for-profit sectors.

#### 297 **Acknowledgements**

298 The authors are grateful to all the participants for their enthusiasm and commitment to  
299 the completion of this study.

#### 300 **References**

- 301 1. Bobeuf F, Labonté M, Khalil A, Dionne IJ. Effect of resistance training on  
302 hematological blood markers in older men and women: a pilot study. *Current gerontology  
303 and geriatrics research*. 2009;2009.
- 304 2. Lionakis N, Mendrinou D, Sanidas E, Favatas G, Georgopoulou M. Hypertension in  
305 the elderly. *World journal of cardiology*. 2012;4(5):135.

- 306 3. Bassett Jr DR, Howley ET. Limiting factors for maximum oxygen uptake and  
307 determinants of endurance performance. *Medicine & Science in Sports & Exercise*.  
308 2000;32(1):70.
- 309 4. Joyner MJ, Casey DP. Regulation of increased blood flow (hyperemia) to muscles  
310 during exercise: a hierarchy of competing physiological needs. *Physiological reviews*.  
311 2015;95(2):549-601.
- 312 5. Kargotich S, Goodman C, Keast D, Morton AR. The influence of exercise-induced  
313 plasma volume changes on the interpretation of biochemical parameters used for  
314 monitoring exercise, training and sport. *Sports medicine*. 1998;26(2):101-17.
- 315 6. Miles DS, Sawka MN, Glaser RM, Petrofsky JS. Plasma volume shifts during  
316 progressive arm and leg exercise. *Journal of applied physiology*. 1983;54(2):491-5.
- 317 7. Tsampoukos A, Stokes K, Nevill M. Separate and combined influence of posture and  
318 sprint running on plasma volume changes. *European journal of sport science*.  
319 2014;14(sup1):S267-S74.
- 320 8. Edwards R, Harrison M, Cochrane LA, Mills F. Blood volume and protein responses  
321 to skin cooling and warming during cycling exercise. *European journal of applied*  
322 *physiology and occupational physiology*. 1983;50(2):195-206.
- 323 9. Zappe DH, Tankersley CG, Meister TG, Kenney WL. Fluid restriction prior to cycle  
324 exercise: effects on plasma volume and plasma proteins. *Medicine and science in sports*  
325 *and exercise*. 1993;25(11):1225-30.
- 326 10. Kanstrup I, Ekblom B. Acute hypervolemia, cardiac performance, and aerobic power  
327 during exercise. *Journal of applied physiology*. 1982;52(5):1186-91.
- 328 11. Mairbäurl H. Red blood cells in sports: effects of exercise and training on oxygen  
329 supply by red blood cells. *Frontiers in physiology*. 2013;4.



- 330 12. Hoffman J. Physiological aspects of sport training and performance. *Human Kinetics*;  
331 2014.
- 332 13. Vechin FC, Libardi CA, Conceição MS, Damas FR, Lixandrão ME, Berton RP et al.  
333 Comparisons between low-intensity resistance training with blood flow restriction and  
334 high-intensity resistance training on quadriceps muscle mass and strength in elderly. *The*  
335 *Journal of Strength & Conditioning Research*. 2015;29(4):1071-6.
- 336 14. Ahmadizad S, Moradi A, Nikookheslat S, Ebrahimi H, Rahbaran A, Connes P. Effects  
337 of age on hemorheological responses to acute endurance exercise. *Clinical hemorheology*  
338 *and microcirculation*. 2011;49(1-4):165-74.
- 339 15. Bongers CC, Eijssvogels TM, Nyakayiru J, Veltmeijer MT, Thijssen DH, Hopman  
340 MT. Thermoregulation and fluid balance during a 30-km march in 60-versus 80-year-old  
341 subjects. *Age*. 2014;36(6):9725.
- 342 16. Grace F, Herbert P, Elliott AD, Richards J, Beaumont A, Sculthorpe NF. High  
343 intensity interval training (HIIT) improves resting blood pressure, metabolic (MET)  
344 capacity and heart rate reserve without compromising cardiac function in sedentary aging  
345 men. *Experimental Gerontology*. 2017.
- 346 17. Bloomer RJ, Farney TM. Acute plasma volume change with high-intensity sprint  
347 exercise. *The Journal of Strength & Conditioning Research*. 2013;27(10):2874-8.
- 348 18. Padulo J, Laffaye G, Chaouachi A, Chamari K. Bench press exercise: the key points.  
349 *The Journal of sports medicine and physical fitness*. 2015;55(6):604-8.
- 350 19. Kawamori N, Haff GG. The optimal training load for the development of muscular  
351 power. *The Journal of Strength & Conditioning Research*. 2004;18(3):675-84.
- 352 20. Dill DB, Costill DL. Calculation of percentage changes in volumes of blood, plasma,  
353 and red cells in dehydration. *Journal of applied physiology*. 1974;37(2):247-8.

- 354 21. Convertino VA, Brock PJ, Keil LC, Bernauer EM, Greenleaf JE. Exercise training-  
355 induced hypervolemia: role of plasma albumin, renin, and vasopressin. *J Appl Physiol*  
356 *Respir Environ Exerc Physiol.* 1980;48(4):665-9.
- 357 22. Ben Abderrahman A, Prioux J, Chamari K, Ben Ounis O, Tabka Z, Zouhal H.  
358 Running interval training and estimated plasma-volume variation. *Int J Sports Physiol*  
359 *Perform.* 2013 Jul;8(4):358-65.
- 360 23. Sosner P, Gremeaux V, Bosquet L, Herpin D, editors. High blood pressure and  
361 physical exercise. *Annales de cardiologie et d'angiologie*; 2014.
- 362 24. Pickering TG, Hall JE, Appel LJ, Falkner BE, Graves J, Hill MN et al.  
363 Recommendations for blood pressure measurement in humans and experimental animals.  
364 *Circulation.* 2005;111(5):697-716.
- 365 25. Donnelly JE, Blair SN, Jakicic JM, Manore MM, Rankin JW, Smith BK. Appropriate  
366 physical activity intervention strategies for weight loss and prevention of weight regain  
367 for adults. *Medicine & Science in Sports & Exercise.* 2009;41(2):459-71.
- 368 26. Rezk C, Marrache R, Tinucci T, Mion D, Forjaz C. Post-resistance exercise  
369 hypotension, hemodynamics, and heart rate variability: influence of exercise intensity.  
370 *European journal of applied physiology.* 2006;98(1):105-12.
- 371 27. Cornelissen VA, Fagard RH. Effect of resistance training on resting blood pressure:  
372 a meta-analysis of randomized controlled trials. *LWW*; 2005.
- 373 28. Weston KS, Wisløff U, Coombes JS. High-intensity interval training in patients with  
374 lifestyle-induced cardiometabolic disease: a systematic review and meta-analysis. *British*  
375 *journal of sports medicine.* 2014;48(16):1227-34.

- 376 29. Corden B, Keenan NG, de Marvao AS, Dawes TJ, DeCesare A, Diamond T et al.  
377 Body Fat Is Associated With Reduced Aortic Stiffness Until Middle Age Novelty and  
378 Significance. *Hypertension*. 2013;61(6):1322-7.
- 379 30. Mero A, Hulmi J, Salmijärvi H, Katajavuori M, Haverinen M, Holviala J et al.  
380 Resistance training induced increase in muscle fiber size in young and older men.  
381 *European journal of applied physiology*. 2013;113(3):641-50.
- 382 31. McCarthy JP, Bamman MM, Yelle JM, LeBlanc AD, Rowe RM, Greenisen MC et al.  
383 Resistance exercise training and the orthostatic response. *European journal of applied*  
384 *physiology and occupational physiology*. 1997;76(1):32-40.
- 385 32. Buoro S, Mecca T, Seghezzi M, Manenti B, Cerutti L, Dominoni 2, Napolitano G,  
386 Resmini S, Crippa A, Ottomano C3 Lippi G. Assessment of blood sample stability for  
387 complete blood count using the Sysmex XN-9000 and Mindray BC-6800 analyzers.  
388 *Revista brasileira de hematologia e hemoterapia*. 2016;38:225-39.

389

390

391

392

393

394

395

396 **Illustrations**

397 **Tables**

398 **Table 1.** Blood pressure and heart rate variation determined before (P1) and after (P2)  
399 training

400 **Table 2.** Red blood cell count ( $10^{12}/L$ ) determined before (P1) and after (P2) training

401 **Table 3.** Hemoglobin concentration (g/100 ml) determined before (P1) and after (P2)

402 training

403 **Table 4.** Hematocrit variation (%) determined before (P1) and after (P2) training

404 **Table 5.** Plasma volume variation (%) determined before (P1) and after (P2) training