Role of creatine supplementation on exercise-induced cardiovascular function and oxidative stress

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Many degenerative diseases are associated with increased oxidative stress. Creatine has the potential to act as an indirect and direct antioxidant; however, limited data exist to evaluate the antioxidant capabilities of creatine supplementation within in vivo human systems. This study aimed to investigate the effects of oral creatine supplementation on markers of oxidative stress and antioxidant defenses following exhaustive cycling exercise. Following preliminary testing and two additional familiarization sessions, 18 active males repeated two exhaustive incremental cycling trials (T1 and T2) separated by exactly 7 days. The subjects were assigned, in a double-blind manner, to receive either 20 g of creatine (Cr) or a placebo (P) for the 5 days preceding T2. Breath-by-breath respiratory data and heart rate were continually recorded throughout the exercise protocol and blood samples were obtained at rest (preexercise), at the end of exercise (postexercise), and the day following exercise (post24 h). Serum hydroperoxide concentrations were elevated at postexercise by 17 ± 5% above preexercise values (p = 0.030). However, supplementation did not influence lipid peroxidation (serum hydroperoxide concentrations), resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations, resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations, resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations, resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations, resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations, resistance of low density lipoprotein to oxidative stress (t1/2max LDL oxidation) and plasma concentrations. Creatine supplementation did not influence the mRNA expression of genes involved in oxidative stress. Creatine supplementation has been demonstrated to increase plasma creatine concentrations and enhance total creatine concentrations, which is mostly stored as creatine phosphate (CrP) within skeletal muscles. Consequently, creatine monohydrate supplementation has the potential to improve performance during high-intensity exercise.

In addition to its ergogenic properties, creatine might augment cellular energy supply and attenuate the loss of cellular Ca2+ homeostasis that has been associated with a range of disease populations. As the accumulation of intracellular Ca2+ has been widely implicated in the formation of reactive oxygen species (ROS) and oxidative damage, creatine supplementation has the potential to improve Ca2+ homeostasis, reduce ROS production and lessen oxidative damage. Lawler et al. were the first to report direct antioxidant properties of creatine. In a series of highly controlled in-vitro studies, these authors demonstrated that creatine was effective in scavenging a range of radicals, including superoxide anion. More recently, exogenous creatine has been demonstrated to scavenge ROS (including hydroxyl radical) as well as reactive nitrogen species in cultured human cells.

There is a growing body of evidence to suggest that creatine supplementation might be beneficial in the prevention and/or treatment of a number of neuromuscular and cardiovascular diseases, for which oxidative stress has been consistently associated. For example, creatine has been demonstrated to improve a variety of outcome measures in animal models of neurodegenerative diseases, such as Huntington’s disease, Parkinson’s disease, amyotrophic lateral sclerosis and other metabolic disorders. Also, pretreatment with creatine has been shown to protect against ischemic injury in rodent models of stroke and creatine supplementation is an effective adjunct therapy in the protection of cardiac function and antioxidant defences in ovariectomized hamsters.

Oxidative stress within the mitochondrial has been associated with aging, where oxidized nucleotides have been demonstrated to accumulate in the brain and muscle with age. Additionally, several studies have linked aging with reductions in the mRNA of genes
associated with mitochondrial structure and function in skeletal muscle and the accumulation of mitochondrial DNA mutations. Consequently, mitochondrial dysfunction has been implicated in a downward spiral, whereby elevated ROS production causes further mitochondrial damage, mitochondrial dysfunction and increased oxidative stress.

Although a small number of reports, mainly anecdotal in nature, suggest that creatine supplementation has the potential to reduce oxidative stress in diseased populations, the individualized nature of these degenerative diseases makes long-term randomized controlled clinical trials difficult to perform. Consequently, the efficacy of creatine in reducing oxidative stress in humans is unclear. It has been estimated that approximately 2–5% of the oxygen flux through the mitochondria is not completely reduced by antioxidative defences are able to prevent oxidative damage to biological molecules including lipids, amino acids and nucleotides. Exercise provoking large increases in whole-body oxygen uptake has the potential to generate ROS (particularly superoxide anions) through the reperfusion of hypoxic tissues in addition to ROS produced through the elevated mitochondrial oxygen flux. Therefore, strenuous concentric muscle activity could increase ROS production, overwhelm normal antioxidant defences, and induce oxidative stress; indeed, exhaustive cycling has been demonstrated to elevate measurements of ROS and lipid peroxidation in apparently healthy humans.

Creatine has potential antioxidant properties (direct and indirect) and has been shown to reduce free radical production and oxidative stress in various animal models of disease; however, limited in-vivo data exists to evaluate the effects of oral creatine monohydrate supplementation on oxidative stress in humans. Therefore, the purpose of the current study was to investigate the proposed antioxidant effects of short-term creatine monohydrate supplementation on antioxidant defences and oxidative stress following exhaustive cycling exercise.

**Results**

The environmental conditions were similar during all trials, ambient temperature being 21.8 ± 0.1 and 21.9 ± 0.2°C, barometric pressure 766 ± 1 and 766 ± 2 mmHg, and humidity 45.2 ± 1.8 and 44.3 ± 1.9%, for T1 and T2 respectively. The calculated daily diet comprised of 8.9 ± 0.3 MJ·d⁻¹, of which 45 ± 2, 32 ± 2, 17 ± 1 and 6 ± 3% of energy intake was obtained from carbohydrates, fat, protein and alcohol, respectively. No differences were identified between groups or trials for calculated antioxidant vitamin intakes (Vitamin A: 528 ± 2.6 μg·d⁻¹; Vitamin C: 47 ± 6 mg·d⁻¹; Vitamin E: 6.1 ± 0.5 mg·d⁻¹).

Peak exercising heart rate and whole-body oxygen uptake were similar during all trials (Table 1), being 194 ± 1 beats·min⁻¹ and 3.97 ± 0.07 L·min⁻¹, respectively. Net whole-body oxygen uptake during exercise (55.5 ± 2.1 L) was similar between supplementation groups and did not change following supplementation (Table 1). Supplementation did not influence changes in plasma volume (trial x timing x group effect, p = 0.359) and these values were similar between trials (trial effect, p = 0.146), being -9.7 ± 0.8% from preexercise values at postexercise and returning to preexercise values at post24 h (1.6 ± 1.3%).

Blood lactate concentrations were similarly elevated from preexercise (1.12 ± 0.06 mmol·L⁻¹) at postexhaustion (8.31 ± 0.21 mmol·L⁻¹) and postexercise (4.98 ± 0.20 mmol·L⁻¹) during all trials (p < 0.001) (Table 2). Blood glucose concentrations were not significantly different for preexercise, postexercise and post24 h, being 4.17 ± 0.09, 4.35 ± 0.07 and 4.32 ± 0.07 mmol·L⁻¹, respectively, for all trials (Table 2).

Exercise did not change t₁/₂max LDL oxidation (time of sample effect, p = 0.502) and supplementation did not influence the pattern of these values (trial x timing of sample x group effect, p = 0.331), with preexercise, postexercise and post24 h values being 97.5 ± 3.9, 96.6 ± 3.8, and 96.1 ± 4.3 min, respectively (Table 2).

Serum hydroperoxide concentrations were elevated at postexercise by 17 ± 5% above preexercise values (time of sample effect, p = 0.030) in all trials (Fig. 1). The three-way interaction effect was not significant (trial x timing of sample x group effect, p = 0.205) and no difference existed between trials (trial effect, p = 0.959).

Supplementation did not influence the pattern of response in any of the measured antioxidant vitamins (trial x timing of sample x group effect, p ≥ 0.176). Exercise elevated plasma concentrations of retinol by 7.2 ± 3.0% (time of sample effect, p = 0.015). Plasma concentrations of α-carotene, β-carotene, α-tocopherol, γ-tocopherol, lycopene and vitamin C were similar throughout the duration of each trial and were not significantly different between trials (time of sample effect, p ≥ 0.078) (Table 3).

**Discussion**

The main findings of this study were that creatine supplementation (20 g·d⁻¹ of creatine administered for 5 d) was not effective in attenuating oxidative stress, did not influence the concentrations of non-enzymatic antioxidant vitamins, and did not alter the susceptibility of LDL particles to oxidative stress in previously familiarized healthy individuals.
Creatine supplementation and oxidative stress

Table 2
Conjugated diene lag times and blood concentrations of lactate and glucose before (trial 1) and after (trial 2) supplementation with creatine (Cr) and placebo (P)

<table>
<thead>
<tr>
<th></th>
<th>Trial 1 Preexercise</th>
<th>Trial 1 Postexhaustion</th>
<th>Trial 1 Postexercise</th>
<th>Trial 1 Post24 h</th>
<th>Trial 2 Preexercise</th>
<th>Trial 2 Postexhaustion</th>
<th>Trial 2 Postexercise</th>
<th>Trial 2 Post24 h</th>
<th>Effect: P interaction, timing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lactate (mmol·L⁻¹)</td>
<td>Cr 1.02 ± 0.08</td>
<td>8.21 ± 0.52</td>
<td>5.00 ± 0.38</td>
<td>0.89 ± 0.08</td>
<td>1.04 ± 0.10</td>
<td>8.51 ± 0.49</td>
<td>5.23 ± 0.48</td>
<td>1.09 ± 0.20</td>
<td>0.585</td>
</tr>
<tr>
<td>Glucose (mmol·L⁻¹)</td>
<td>P 1.08 ± 0.14</td>
<td>8.25 ± 0.42</td>
<td>4.90 ± 0.41</td>
<td>1.29 ± 0.28</td>
<td>1.33 ± 0.16</td>
<td>8.26 ± 0.34</td>
<td>4.80 ± 0.35</td>
<td>1.34 ± 0.20</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>t₁/₂max LDL (min)</td>
<td>Cr 89.0 ± 7.1</td>
<td>-</td>
<td>87.3 ± 6.3</td>
<td>88.1 ± 6.3</td>
<td>93.0 ± 8.4</td>
<td>-</td>
<td>91.1 ± 7.7</td>
<td>88.0 ± 6.1</td>
<td>0.502</td>
</tr>
<tr>
<td></td>
<td>P 102.5 ± 8.7</td>
<td>-</td>
<td>105.0 ± 7.9</td>
<td>100.9 ± 7.5</td>
<td>104.4 ± 7.3</td>
<td>-</td>
<td>103.1 ± 7.6</td>
<td>107.4 ± 12.4</td>
<td>0.393</td>
</tr>
</tbody>
</table>

Values are presented as mean ± SEM (N = 9). Definitions: Cr, Creatine group; P, placebo group. Effects present p-values for supplementation group x trial x timing of sample interaction effect and timing effect.

Figure 1. Exercise-induced lipid peroxidation as measured by serum hydroperoxide concentrations. Exhaustive cycling caused increases in lipid peroxidation (time of sample effect, p = 0.030), with serum hydroperoxide concentrations being elevated after exercise (post-exercise) when compared with values before (Pre-exercise) and 24 hours after exercise (Post-24 h). The pattern of response was similar following supplementation with creatine (Cr) and placebo (P) (trial x timing of sample x group effect, p = 0.205). Values represent mean ± SEM (N = 9).

The cycling protocol increased lipid peroxidation above preexercise values, as measured by increased postexercise serum hydroperoxide concentrations. The relative change in lipid peroxidation (17 ± 5%) was less than we previously reported after intermittent running, where increases of 24 ± 5% were measured. Exhaustive cycling does not involve significant amounts of eccentric muscle activity and structural muscle damage; therefore, unlike intermittent running, cycling does not lead to inflammation-mediated free radical production that follows structural muscle damage. Consequently, during the current study, it is likely that increased mitochondrial oxygen flux led to elevated ROS production that overwhelmed the antioxidant defences and caused oxidative stress. In support of this premise, incremental cycling has previously been demonstrated to increase the intensity of the electron spin resonance (ESR) signal in the plasma of healthy young subjects. These authors suggested that the enhanced postexercise ESR signal reflected an increase in the concentration of alkoxyl radicals formed as a consequence of primary ROS (such as superoxide anion and hydroxyl radicals) attack on membrane phospholipids.

The amount of external work undertaken during the individualised exhaustive protocol was matched during T1 and T2; furthermore, heart rate, oxygen uptake and blood lactate concentrations reached similar peak values at the end of exercise during all trials suggesting that the protocol challenged all subjects to an equal extent. Net whole-body oxygen uptake was similar between groups and trials (Table 2). Assuming that the rate of mitochondrial ROS production remained approximately constant.
for all individuals, the exercise protocol was effective in producing similar amounts of ROS during T1 and T2. Consequently, this repeated measures design was successful in challenging the antioxidant defences of all subjects to a comparable extent during the main exercise trials.

The finding that creatine supplementation did not influence oxygen uptake during exhaustive exercise is in agreement with previous literature. Further evidence to suggest that cardiopulmonary function during exhaustive exercise was unaffected by creatine supplementation was provided by the similarity in heart rate and net oxygen uptake responses before and after supplementation. In support of this conclusion, most previous studies confirm previous in-vitro data showing that creatine acts a direct antioxidant by scavenging ROS (particularly hydroxyl radicals) in cultured cells. Later, intracellular creatine was also shown to act as a direct antioxidant by scavenging ROS (particularly hydroxyl radicals) in cultured cells. Mitochondrial ROS production (including superoxide anions and hydroxyl radicals) is likely to be the primary source of free radicals species in highly controlled cell-free environments. Later, intracellu lar creatine was also shown to act as a direct antioxidant by scavenging ROS (particularly hydroxyl radicals) in cultured cells.

It has been repeatedly demonstrated that the current creatine monohydrate supplementation regime is effective in increasing circulatory creatine concentrations and pharmacokinetic analysis of plasma creatine concentrations following a 5-g oral bolus of creatine monohydrate showed that plasma creatine decayed with a half-life of approximately 2 h. Therefore, it is probable that the current supplementation protocol elevated plasma and intramuscular creatine concentrations.

Calculated intakes of the main dietary antioxidant vitamins were similar during all trials and supplementation had no effect on plasma concentrations of the antioxidant vitamins (retinol, α-carotene, β-carotene, α-tocopherol, γ-tocopherol, and lycopene, and vitamin C). In addition, the two main exercise trials were completed within 7 d; therefore, it is likely that the enzymatic antioxidant defences remained similar for each individual during both main exercise trials. As the enzymatic and non-enzymatic vitamin antioxidant defences were comparable between exercise trials, this repeated measures design was suitable to assess the influence of short-term oral creatine supplementation as an in-vivo antioxidant.

Supplementation did not influence the pattern of response in serum hydroperoxide concentrations (trial x timing of sample x group effect, p = 0.205). Although there was a tendency for mean serum hydroperoxide concentrations to be reduced following supplementation with creatine (Fig. 1), creatine supplementation did not significantly influence lipid peroxidation following exhaustive cycling. Therefore, the current finding does not confirm previous in-vitro data showing that creatine acts a direct antioxidant. Lawler et al. were the first to report that creatine effectively quenches superoxide anions and other aqueous radical species in highly controlled cell-free environments. Later, intracellular creatine was also shown to act as a direct antioxidant by scavenging ROS (particularly hydroxyl radicals) in cultured cells.

Differences between in-vivo and in-vitro systems might partially explain the lack of statistical evidence to support an antioxidant effect in the current study. Although the current supplementation regime has been shown to “load” muscle cells to threshold levels, considerable heterogeneity exists in intramuscular creatine uptake following oral supplementation. It is, therefore, possible that the

### Table 3: Plasma antioxidant vitamin concentrations before (trial 1) and after (trial 2) supplementation with creatine (Cr) and placebo (P)

<table>
<thead>
<tr>
<th>Vitamin</th>
<th>Trial 1 Preexercise</th>
<th>Trial 1 Postexercise</th>
<th>Trial 1 Post24 h</th>
<th>Trial 2 Preexercise</th>
<th>Trial 2 Postexercise</th>
<th>Trial 2 Post24 h</th>
<th>Effect P (interaction, timing)</th>
</tr>
</thead>
<tbody>
<tr>
<td>retinol</td>
<td>Cr</td>
<td>2.1 ± 0.2</td>
<td>2.2 ± 0.2</td>
<td>2.3 ± 0.2</td>
<td>2.1 ± 0.2</td>
<td>2.3 ± 0.2</td>
<td>2.3 ± 0.2</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>2.3 ± 0.3</td>
<td>2.3 ± 0.3</td>
<td>2.3 ± 0.4</td>
<td>2.2 ± 0.3</td>
<td>2.5 ± 0.3</td>
<td>2.2 ± 0.2</td>
</tr>
<tr>
<td>α-carotene</td>
<td>Cr</td>
<td>0.20 ± 0.05</td>
<td>0.22 ± 0.04</td>
<td>0.21 ± 0.04</td>
<td>0.18 ± 0.04</td>
<td>0.16 ± 0.04</td>
<td>0.17 ± 0.03</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>0.15 ± 0.02</td>
<td>0.17 ± 0.01</td>
<td>0.16 ± 0.02</td>
<td>0.15 ± 0.02</td>
<td>0.16 ± 0.02</td>
<td>0.15 ± 0.02</td>
</tr>
<tr>
<td>β-carotene</td>
<td>Cr</td>
<td>0.61 ± 0.15</td>
<td>0.70 ± 0.13</td>
<td>0.70 ± 0.18</td>
<td>0.55 ± 0.12</td>
<td>0.49 ± 0.12</td>
<td>0.47 ± 0.10</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>0.42 ± 0.07</td>
<td>0.46 ± 0.07</td>
<td>0.47 ± 0.12</td>
<td>0.43 ± 0.07</td>
<td>0.53 ± 0.14</td>
<td>0.49 ± 0.11</td>
</tr>
<tr>
<td>α-tocopherol</td>
<td>Cr</td>
<td>28.3 ± 1.8</td>
<td>29.1 ± 2.2</td>
<td>29.9 ± 2.5</td>
<td>28.7 ± 1.6</td>
<td>28.2 ± 1.8</td>
<td>28.2 ± 1.6</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>29.5 ± 2.7</td>
<td>31.0 ± 2.6</td>
<td>31.2 ± 2.6</td>
<td>29.5 ± 2.6</td>
<td>31.0 ± 2.6</td>
<td>31.2 ± 2.6</td>
</tr>
<tr>
<td>γ-tocopherol</td>
<td>Cr</td>
<td>1.4 ± 0.1</td>
<td>1.3 ± 0.1</td>
<td>1.4 ± 0.2</td>
<td>1.2 ± 0.1</td>
<td>1.1 ± 0.1</td>
<td>1.2 ± 0.1</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>1.5 ± 0.2</td>
<td>1.5 ± 0.1</td>
<td>1.5 ± 0.1</td>
<td>1.5 ± 0.2</td>
<td>1.7 ± 0.2</td>
<td>1.7 ± 0.2</td>
</tr>
<tr>
<td>lycopene</td>
<td>Cr</td>
<td>1.4 ± 0.3</td>
<td>2.0 ± 0.4</td>
<td>1.9 ± 0.5</td>
<td>1.5 ± 0.5</td>
<td>1.3 ± 0.3</td>
<td>1.2 ± 0.3</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>1.3 ± 0.5</td>
<td>1.5 ± 0.3</td>
<td>1.2 ± 0.1</td>
<td>1.0 ± 0.1</td>
<td>1.8 ± 0.4</td>
<td>1.0 ± 0.2</td>
</tr>
<tr>
<td>vitamin C</td>
<td>Cr</td>
<td>45.0 ± 3.9</td>
<td>48.1 ± 4.5</td>
<td>42.7 ± 6.0</td>
<td>49.7 ± 4.9</td>
<td>44.5 ± 4.1</td>
<td>47.1 ± 5.6</td>
</tr>
<tr>
<td></td>
<td>P</td>
<td>48.1 ± 2.9</td>
<td>43.0 ± 4.3</td>
<td>47.4 ± 4.6</td>
<td>41.7 ± 3.5</td>
<td>42.3 ± 5.1</td>
<td>44.1 ± 5.6</td>
</tr>
</tbody>
</table>

Values are presented as mean ± SEM (N = 9). Definitions: Cr, Creatine group; P, Placebo group. Effects present p-values for supplementation group x trial x timing of sample interaction effect and timing effect.
change in intracellular creatine concentrations from T1 to T2 in 
Cr was insufficient to cause significant differences in lipid 
peroxidation. This finding suggests that increasing intramuscular 
creatine in healthy young males does not afford additional protec-
tion against oxidative damage derived from strenuous concentric 
muscle activity.

Supplementation did not influence $t_{1/2\text{max}}$ LDL oxidation 
(Table 3). As $t_{1/2\text{max}}$ LDL oxidation measures the resistance of 
LDL particles to oxidation, these data suggest that creatine did not 
alter the antioxidant defences of LDL particles in healthy young 
males. Unlike the current data, some recent studies have previ-
ously demonstrated that prolonged exercise involving consider-
able eccentric muscular activity causes significant increases in LDL 
resistance to peroxidation in healthy males.23,37 These equivocal 
results probably reflect increased mobilisation of antioxidant 
defences following prolonged exercise that leads to muscle damage 
and acute inflammation.

Similar doses of creatine have been repeatedly demonstrated to 
enhance short-term exercise performance in healthy individuals, 
which has generally been ascribed to enhanced temporal energy 
buffering through increased intramuscular CrP.1 Supplementation 
did not influence peak values for heart rate, oxygen uptake, and 
blood glucose and lactate concentrations (Table 3); therefore the 
metabolic responses to matched exhaustive exercise were not 
changed by creatine supplementation. Nevertheless, Andrews 
et al.38 showed that creatine supplementation attenuated blood 
concentrations of lactate and ammonia responses to exhaustive 
handgrip dynamometry exercise in patients with chronic heart 
failure. Consequently, creatine supplementation might be more 
effective in maintaining cellular energy supply in patient popula-
tions; thereby, reducing the accumulation of intracellular Ca²⁺38, 
limiting the formation of reactive oxygen species (ROS) and 
attenuate oxidative damage.5

Although the current data does not support in-vivo antioxidant 
benefits, mediated by direct or indirect mechanisms, from short-
term creatine supplementation in healthy young males, it remains 
possible that individuals with diseases that are associated with 
reduced antioxidant defences and elevated radical production, 
probably through an impairment in cellular energy metabolism, 
might benefit from creatine supplementation. Creatine is unique, 
within the currently available antioxidants, in that it has the 
potential to reduce ROS production through better maintenance of 
cellular energeticas as well as quench ROS and other radical 
species. Indeed, oral creatine supplementation has been demon-
strated to reduce oxidative stress and prevent paracrystalline 
inclusions in a patient with mitochondrial dysfunction.19 A small 
number of randomised clinical trials have evaluated the effects of 
short- to moderate-term creatine supplementation in patients 
with diseases that have been traditionally associated with oxidative 
stress, such as Huntington’s disease,39 Parkinson’s disease,40-42 
amyotrophic lateral sclerosis,43-45 rheumatoid arthritis,46 chronic 
obstructive pulmonary disease47 and congestive heart failure.58 
Although equivocal, these studies have generally reported small 
positive effects in a range of outcome measures. Nevertheless, 
no data from randomised clinical trials are currently available to 
evaluate the influence of creatine supplementation on oxidative 
stress in patient populations.

In summary, this is the first study to evaluate the effects of 
creatine supplementation on acute exercise-derived oxidative stress 
in healthy young males. Short-term creatine supplementation was 
in ineffective in attenuating oxidative stress (as measured by serum 
hydroperoxide concentrations) induced by exhaustive cycling; in 
addition, supplementation did not influence LDL susceptibility to 
oxidative stress exercise or change plasma concentrations of anti-
oxidant vitamins. Nevertheless, it remains possible that creatine 
supplementation could benefit patient populations that have 
increased susceptibility to oxidative stress due to impaired cellular 
energy metabolism and decreased antioxidant defences.

Methods

Subjects. Eighteen healthy male volunteers (age: 21.5 ± 0.4 
years; stature: 1.77 ± 0.01 m; body mass: 81.1 ± 2.0 kg; body 
composition: 16.8 ± 0.7%BF) were informed about the potential 
risks of the study and gave written informed consent for their 
participation in the study, which was approved by a local research 
ethics committee. No subject had prior history of cardiovascular 
or respiratory disease and all subjects were non-smokers. Potential 
subjects attended an interview prior to undertaking the study and 
were subsequently excluded if they engaged in a structured weight 
training programme, or had taken creatine or other nutritional 
supplements in the 8 weeks prior to the study.

Experimental design. Prior to the main exercise trials, all subjects 
completed three incremental exercise tests (separated by exactly 7 
d) in order to ensure that the subjects were fully familiarized 
and able to repeat the exhaustive exercise protocol. Subjects then 
performed two main exercise trials, which consisted of incremental 
cycling that was continued until the individualized predetermined 
point of exhaustion (determined during the preliminary trials), 
and separated by exactly 7 d. Two days after completing the first 
main exercise trial (T1) the subjects were assigned, in a randomised 
double-blind fashion, to either a creatine (Cr) group or a placebo (P) 
group and instructed to take supplements for 5 d, which included 4 
d prior to and the day of trial two (T2). The subject characteristics 
of each group are presented in Table 4. The Cr group ingested 22.8 
g·d⁻¹ Cr·H₂O (equivalent to 5 g Cr x 4 daily) for 5 d. Each supple-
ment dose consisted of 5.7 g Cr·H₂O and 5 g of glucose polymer
dissolved in 500 ml of warm water. This regimen was adopted in light of the work by Harris et al.2 who found that this protocol resulted in significant increases in total creatine content of resting quadriceps femoris muscle; in addition, the majority of studies that have shown a direct or indirect increase in total creatine concentration have used similar supplementation regimes. Dissolving creatine in warm water prevented any detectable formation of creatinine, with no parts of the supplement remaining undissolved. The placebo group consumed 40 g·d⁻¹ of glucose polymer. Both supplements had similar taste, texture and appearance, and were administered in generic packaging. Subjects were instructed to maintain their normal diet and activity patterns throughout the study. Food was weighed and recorded by the subjects for 2 d prior to each exercise trial and for 1 d afterwards. These food records were subsequently analyzed using commercial software (CompEat version v5.8.0; Nutrition Systems, UK). In addition, the subjects were instructed to abstain from strenuous exercise for 3 d before and 1 d following each trial. At the completion of the study all subjects gave their verbal assurance that they had complied with all instructions. The experimental design is illustrated in Figure 2.

**Preliminary procedures.** Subjects were habituated with the exercise protocol on three separate occasions prior to the main exercise trials. After a 5-min warm-up at 60 W and a 3-min rest period the subjects completed an incremental exercise test on an electromagnetically braked cycle ergometer (Lode Excalibur Sport; Lode, Holland). The work rate began at 60 W and thereafter increased in 10-W increments every 1 min until volitional exhaustion, which was defined as an inability to maintain a cadence of 60 rev min⁻¹ despite verbal encouragement. The time to exhaustion during the preliminary tests were used to determine the exercise time for the main exercise trials; therefore, the individualised exercise protocol was recorded and repeated during both main exercise trials. Heart rate (S810; Polar Electro Oy, Finland) and breath-by-breath respiratory parameters (Oxycon Pro; Jaeger, Germany) were simultaneously recorded throughout exercise. Breath-by-breath oxygen uptake during exercise was used to calculate net oxygen uptake.

During the initial preliminary session, mass (model 712; Seca, Germany) and stature (Portable Stadiometer; Holtain, UK) were measured; additionally, percentage body fat (BF) was determined using bioelectrical impedance analyses (Quadscan; Bodystat, Isle of Man), as previously described.⁴⁸

**Main trial procedures.** On the day of each main exercise trial the subjects reported to the laboratory at approximately the same time of day (±1 h) and following an overnight fast. After 5-min of cycling at 60 W and a 3-min rest period the subjects completed the individualised incremental cycling protocol. Subjects were instructed to pedal at a constant cadence between 65 and 75 rev min⁻¹. Heart rate (S810; Polar Electro Oy, Finland), and breath-by-breath respiratory parameters (Oxycon Pro; Jaeger, Germany) were simultaneously recorded throughout exercise. Blood samples were taken by venepuncture (Vacutainer system; Becton-Dickinson Ltd., UK) in a seated position from an antecubital vein before exercise (preexercise), 20 min after the completion of exercise (postexercise), and 24 h after the trial (Post24 h). Additional capillary blood samples were taken immediately after exhaustion (postexhaustion) for the analysis of blood lactate and glucose concentrations. Ambient temperature and humidity were recorded at the beginning and end of exercise (ETHG-912; Oregan Scientific, USA).

**Blood sampling and analysis.** Venous blood was collected in a 5 mL container (Becton-Dickinson Ltd., UK) containing the anticoagulant ethylenediamine tetracetic acid (EDTA). Several small aliquots were removed for the triplicate determination of blood lactate concentration, glucose concentration (YSI 2300, Yellow Springs Instruments, US), hemoglobin concentration (Hemocue Ltd., UK), hematocrit (Micro hematocrit MK IV, Hawksley, UK) and changes in plasma volume were estimated as previously described.⁴⁹ The remaining blood was centrifuged at 3,000 g for 15 min to obtain plasma. An aliquot of plasma (100 μL) was added to 900 μL of freshly prepared 10% metaphosphoric acid, mixed and frozen at -70°C for subsequent vitamin C analysis. The remaining plasma was frozen in aliquots and stored at -70°C prior to subsequent analysis. Two additional 7 mL blood samples were collected in serum separation tubes (Becton-Dickinson Ltd., UK), left to stand for 15 min then centrifuged at 3,000 g for 15 min to obtain serum. The serum was transferred to appropriate containers and subsequently stored at -70°C prior to analysis.

Serum hydroperoxide concentrations were measured as described in McEneny et al.⁵⁰ Low density lipoprotein was isolated from plasma and oxidised according to the method of McDowell et al.⁵¹ Subsequently, the production of conjugated dienes was followed in triplicate at 234 nm (SpectraMax 190; Molecular Devices Corp., US) using the computer software SoftMax Pro Version 3.0 (Molecular Devices Corp., US); the change in absorbance (from baseline to the maximum absorbance following the propagation

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![Figure 2. Schematic diagram representing the experimental design that was used to evaluate the role of creatine supplementation on exercise-induced cardiovascular function and oxidative stress. After repeated familiarization, the subjects performed two main exercise trials that consisted of exhaustive incremental cycling. Two days after completing the first main exercise trial (T1) the subjects were assigned, in a randomised double-blind fashion, to either a creatine (C) group or a placebo (P) group and instructed to take supplements for 5 d, which included 4 d prior to and the day of trial two (T2).](image-url)
phase) was used to quantify conjugated diene concentration, and the time taken to reach half of the maximum oxidation (t_{1/2max} of LDL oxidation), which directly correlates with conjugated diene lag time, was taken as a measure of the resistance of the particle to oxidation. The duration of the conjugated diene lag time has been shown to correlate with endogenous antioxidants in LDL. Vitamin C concentrations were determined using a fluorimetric assay using a centrifugal analyser with fluorescence detection according to the method of Vuilleumier and Keck. Plasma concentrations of retinol, α-carotene, β-carotene, α-tocopherol, γ-tocopherol and lycopene were measured by high-performance lipid chromatography with electrochemical detection according to the methods of Craft. The intra-assay coefficient of variance (%CV) for these assays ranged from 1.4 for α-tocopherol to 12.0 for lycopene.

**Statistical analysis.** Statistical analysis was carried out using SPSS software (version 13.0; SPSS Inc., IL, US). Group data were expressed as mean ± SEM and statistical significance was set at the p < 0.05 level. All data were assessed for normality (Shapiro-Wilk's t) and data that were not normally distributed (plasma retinol, α-tocopherol and lycopene concentrations) were log transformed prior to analyses. Subject characteristics were compared under supplementation groups using independent samples t-tests (Table 1). Environmental conditions were compared using paired samples t-tests. Body composition, heart rate and whole-body oxygen uptake measurements were assessed using mixed-model repeated measures ANOVA (within-subject factors: trials; between-subject factor: supplementation groups). The remaining data, which contained multiple time points during each trial, were analysed using mixed-model repeated measures ANOVA (within subject factors: trial x time of sample; between subject factor: group). Mauchly's test was consulted and Greenhouse-Geisser correction was applied if the assumption of sphericity was violated. If a significant p value was identified for the three-way interaction (group x trial x time of sample), the subject groups reacted differently and supplementation was deemed to have had a significant effect. Where a significant p value was identified for the main effect of time (time of sample), multiple pairwise comparisons were made using Bonferroni confidence interval adjustment.

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**References**


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