The Effect of Curcumin Supplementation on Selected Markers of Delayed Onset Muscle Soreness (DOMS)

Babak Nakhostin-Roohi, Arash Nasirvand Moradlou, Sahar Mahmoodi Hamidabad, Babak Ghanivand

ABSTRACT

Inflammation and pain induced by delayed onset muscle soreness (DOMS) can be induced by eccentric exercise or an unaccustomed activity. The condition can cause problems in exercising and for athletes. The purpose of this study was to assess the effect of 150mg curcumin supplementation immediately after intensive eccentric exercise. Evaluations were made for total antioxidant capacity (TAC), muscle damage markers, and DOMS induced pain. Ten healthy young males (age, 25.0 ± 1.6 years; height, 178.9 ± 4.1 cm; body mass, 81.1 ± 6.8 kg; fat%, 14.2 ± 2.1) completed a double blind randomized-controlled crossover trial to estimate the effects of oral curcumin supplementation (150mg) and a placebo on squat performance and DOMS following unaccustomed heavy eccentric exercise. Curcumin (CU) or placebo (P) was taken at the prescribed dose immediately after eccentric squat exercises; administrations were separated by a 14-day washout period. Measurements were made at the baseline, immediately, 24, 48, and 72h after exercise comprising: (a) limb pain (1–10 cm visual analogue scale; VAS), (b) total antioxidant capacity (TAC) (c) serum markers of muscle damage and inflammation. Measurements taken after exercise showed significantly reduced levels of pain, creatine kinase (CK), alanine aminotransferase (ALT), and aspartate aminotransferase (AST) in C group compared with group P group (P<0.05). TAC remained significantly high in group C after exercise compared with levels in group P (P<0.05). The findings of this study suggest that a 150mg dose of curcumin may have antioxidant, anti-inflammatory and analgesic effects on DOMS.

INTRODUCTION

Delayed onset muscle soreness (DOMS) from training or exercise produces unpleasant and undesirable feelings of pain and muscle stiffness. That condition can discourage a beginner or an experienced athlete from continuing with the exercise (1-4). DOMS results in discomfort at the site of the injury, inflammation, oxidative stress, edema and loss of muscle function and strength (5, 6). Extensive research has been done to investigate procedures to reduce DOMS and many studies have reported that nutrition interventions can reduce experiences of DOMS (7). Commonly known nutritional interventions

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Curcumin Supplementation and DOMS

include caffeine, omega-3 fatty acids, and taurine (8-10).

Recent research attention has focused on the effects of nutraceutical bioactive compounds such as polyphenols (11). Polyphenols are a class of organic chemical compound, mainly found in plants that are characterized by the presence of multiples of phenol structural units (12). Curcumin is a hydroxycinnamic acid derivative and its structure contains two hydrophobic polyphenolic rings with two carbonyl groups. It is the main curcuminoid found in the spice turmeric, a plant alkaloid obtained from ground rhizome of the perennial herb Curcuma longa (13). According to investigations, curcumin has multiple bio-functional activities including anti-inflammatory, antioxidant, anti-cancer, anti-diabetes and chemo preventative activities (14-18).

Evidence from tests suggest that in some conditions, curcumin possess similar anti-inflammatory activity as some of the common non-steroidal anti-inflammatory drugs (NSAIDs) such as indomethacin, Vioxx, Celebrex, and ibuprofen, but without many of the side effects, such as gastrointestinal distress and cardiovascular complications (19, 20). Accordingly, some investigations have suggested that curcumin could be effective in controlling inflammation and oxidative stress induced by DOMS.

Some investigations have evaluated for the effect of curcumin on DOMS (21-24). However, to the best of our knowledge, no study to date has tested administration of 150mg curcumin supplement immediately after intensive eccentric exercise. Therefore, the aim of this study was to evaluate the effect of acute curcumin supplementation on the DOMS markers immediately after intensive eccentric exercise.

MATERIALS AND METHODS

Participants. 10 healthy young males gave written informed consent and took part in this study. This study was conducted according to the guidelines laid down in the Declaration of Helsinki and approved by the ethics committee of Ardabil Branch, Islamic Azad University. All volunteers completed a questionnaire on physical activity, exercise, dietary intake and health history prior to participation in the study.

Study Design. The study was designed as a double-blind randomized-controlled crossover trial to compare the effects of taking oral curcumin to those placebo on against evaluation markers of DOMS immediately after a bout of unaccustomed squat exercises. All subjects completed two test trials at an interval of at least 2 weeks. Randomization was applied to both groups (curcumin and placebo treatments). Exclusion criteria comprised regular leg weight training in the 3 months prior to the tests; current lower limb musculoskeletal injury; current use of non-steroidal anti-inflammatory drugs and antioxidant supplement, as well as neurological disease involving the lower limb.

Dietary. Participants were instructed verbally and in writing to follow an average polyphenol-contain diet throughout the duration of participation in the study starting 3 weeks prior to the start of the study period until the last blood sampling. A list of high polyphenol-contain foods was provided to participants to help them avoid major sources of dietary polyphenols.

Induction of delayed-onset muscle soreness. DOMS was induced in all subjects by standardized repetitive quadriceps muscle exercise using a squat machine. The researchers selected one repeated maximum (1RM) on the squat machine to start with a low but reasonable weight that then increased with each repetition until subjects reached their 1RM. 1RM is defined as the highest weight under which a subject is able to perform a squat position just for one repetition.

For the main trials, subjects performed exercises (7 sets of 20 squats) using a squat machine at under 50% 1RM. Each repetition was performed through the full pain-free range of motion in a slow controlled manner. Subjects performed the concentric portion of the repetition for 2 seconds, paused at full contraction for 1 second and then completed the eccentric portion over a 4-second period (for a total of 7 seconds per repetition). Subjects performed each set according to their level of tolerance and were given a rest period of 1 to 3 minutes between sets (25).

Intervention. Each participant underwent 2 laboratory-based trials in randomized order: (1) placebo (control), (2) curcumin supplementation. There was an interval of at least 2 weeks between tests.
between trials. Curcumin was provided by the Thera values Corporation design (Tokyo, Japan) (26). The product was developed to result in much higher plasma concentration and bioavailability after intake compared to conventional curcumin powder. The curcumin capsule in this study consisted of 10% curcumin, 2% curcuminoids without curcumin, 3.2% gum ghatti, 0.27% citric acid, 54.53% dextrin and 30% maltose. The placebo capsule consisted of 5% tartrazine, 3.5% gum ghatti, 0.3% citric acid, 59.2% dextrin and 32% maltose. Participants received oral administration of 150mg curcumin or the same capsules of placebo from the researchers after exercise (just after second blood sampling)(27).

Measurement of muscle damage and oxidative stress markers. Muscle damage and oxidative stress markers were measured immediately before, after, 24, 48, and 72h after eccentric strength exercise. Muscle soreness was assessed using a visual analogue pain scale (VAS), for which participants placed a mark on a 10cm line to indicate a degree of soreness. Soreness was measured by distance in centimeters from the left end of the scale to the mark. Establishment of validity and reliability of VAS as a measure for subjective soreness was cited in another report (28). Markers for muscle damage were as follows: creatine kinase (CK), alanine aminotransferase (ALT), and aspartate aminotransferase (AST); measured using spectrophotometer commercial kits (Pars-Azmoon, Iran). Total antioxidant capacity (TAC) was used to mark oxidative stress and measured by the method cited in Varga et al. (29).

Statistical Analysis. All data are presented as mean ± SEM, and statistical significance was set at p<0.05. Data with multiple time points during the main trial were analyzed using the mixed-model repeated-measures ANOVA. Mauchly’s test was consulted and Greenhouse-Geisser correction was applied if the sphericity assumption was violated. If a significant p value was identified for the main effect of time (time of sample), multiple pair wise comparison was made using Bonferroni confidence interval adjustment. Moreover, comparison was made of dependent variable data in multiple time points between the two groups using the independent sample t-test.

RESULTS

Physical characteristics of the participants were as follows (expressed as mean ± standard deviation [SD]): age, 25.0 ± 1.6 years; height, 178.9 ± 4.1 cm; body mass, 81.1 ± 6.8 kg; fat%, 14.2 ± 2.1.

Total Antioxidant Capacity (TAC). Comparison of baseline resting plasma TAC showed no difference between the groups (P>0.05). TAC was significantly higher in C group than in group P at 24 and 48h after exercise (P=0.001 and P=0.01, respectively) (Figure1).

Figure 1. TAC plasma level. Values represent means ± SEM. † demonstrate a significant increase in group C compared with group P (p<0.05).

CK. Baseline resting serum CK showed no difference different between groups C and P (P>0.05). CK showed a significant increase immediately, and at 72h after exercise in both groups compared with pre-exercise (P<0.05). CK was significantly lower in C group compared with group P 24h after exercise (P=0.019) (Figure 2).

Figure 2. CK before and after exercise. Values represent means ± SEM. * show significant increase compared with pre-exercise in both groups (p<0.05). † demonstrates a significant increase compared with pre-exercise only in group P and a significant increase in group P compared with group C (p<0.05).
Visual Analogue Scale (VAS). VAS showed significant increase at the following times; immediately, 24, 48, and 72h after exercise in both groups compared with pre-exercise (P<0.05). VAS was significantly lower in group C compared with group P at the times; 48, and 72h after exercise (P<0.05) (Figure 3).

ALT and AST. Baseline resting serum ALT and AST were not different between groups (P>0.05). ALT significantly increased immediately, and 24h after exercise in both groups (P<0.05). ALT was significantly lower in C group compared with the P group 24h after exercise (P=0.019) (Table 2). AST was significantly lower in C group compared with the P group just 24h after exercise (P=0.034) (Table 1).

DISCUSSION

To the best of our knowledge, the present study was the first to examine the effects of 150mg curcumin supplement on antioxidant capacity and muscle damage markers after intensive eccentric exercise in humans. The main findings in this study were as follows: (1) curcumin supplement maintained TAC high, 24 and 48h after exercise (2) It enabled attenuation of CK, LDH, ALT, AST, and experience of pain related to DOMS.

Table 1. ALT and AST before and after exercise. Values represent means ± SEM.

<table>
<thead>
<tr>
<th>Groups</th>
<th>ALT (U/L)</th>
<th>Baseline</th>
<th>Immediately after Exercise</th>
<th>24h after Exercise</th>
<th>48h after Exercise</th>
<th>72h after Exercise</th>
</tr>
</thead>
<tbody>
<tr>
<td>Placebo</td>
<td>17.4 ± 0.62</td>
<td>21.0 ± 1.16*</td>
<td>25.5 ± 1.73†</td>
<td>23.0 ± 1.76</td>
<td>20.1 ± 1.97</td>
<td></td>
</tr>
<tr>
<td>Curcumin</td>
<td>17.7 ± 0.61</td>
<td>22.9 ± 1.48*</td>
<td>20.4 ± 0.90†</td>
<td>19.8 ± 0.91</td>
<td>18.2 ± 1.34</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Groups</th>
<th>AST (U/L)</th>
<th>Baseline</th>
<th>Immediately after Exercise</th>
<th>24h after Exercise</th>
<th>48h after Exercise</th>
<th>72h after Exercise</th>
</tr>
</thead>
<tbody>
<tr>
<td>Placebo</td>
<td>21.2 ± 2.32</td>
<td>24.4 ± 1.53</td>
<td>31.1 ± 2.21†</td>
<td>23.0 ± 1.30</td>
<td>24.4 ± 1.51</td>
<td></td>
</tr>
<tr>
<td>Curcumin</td>
<td>20.9 ± 0.81</td>
<td>21.3 ± 1.11</td>
<td>24.0 ± 2.23</td>
<td>26.6 ± 1.93</td>
<td>26.6 ± 1.89</td>
<td></td>
</tr>
</tbody>
</table>

* shows significant increase compared with pre-exercise (p<0.05).
† demonstrates a significant increase compared with pre-exercise in both groups and significant increase in group P compared with group C (p<0.05).

It is known that Curcumin has antioxidant quality (18). According to the findings of this study, TAC remained significantly higher in group C than in group P at 24 and 48h after exercise; this can possibly be attributed to antioxidant properties of curcumin (Figure 1). In parallel, markers of muscle damage enzyme showed a reduction in group C group compared to group P. Observations shown in figures 2, 3, and Table 1, alteration patterns of CK, ALT, and AST were almost the same as TAC. CK, and ALT were significantly enhanced after exercise demonstrating occurrence of DOMS, but there was a significant reduction at some points of the time series in group C. One explanation is that the antioxidant properties of curcumin caused a decline in some enzymes.

Elevations of CK and ALT serum levels show increased leakage of these enzymes through cell membranes after exercise. Oxidative stress induced by lipid peroxidation may lead to membrane permeability that would allow muscle constituents such as CK and aminotransferases to escape (30). The phenolic compound of curcumin acts as a scavenger of reactive oxygen species and a quencher of the lipid peroxidative side chain (31). Thus, the phenolic OH groups of curcumin may have reduced lipid hydroperoxides. Finally, the inhibitory effects of curcumin on lipid peroxidation may have
Curcumin Supplementation and DOMS


prevented CK leakage and other related enzymes from cell membranes and consequently less increase in CK serum level. Previous studies have confirmed that supplements containing antioxidants are able to attenuate these enzymes (32, 33).

Attenuation of these enzymes could also be attributed to the anti-inflammatory effects of curcumin. Several studies have confirmed that curcumin blocks the activity of transcription factor NF-kappaB, reduces AP-1 binding to DNA and causes decreased production of COX-2, all of which play a role in inflammation (22, 34, 35). Curcumin appears to target NF-kappaB as opposed to COX-2, indicating the potential for less serious side effects than NSAIDs (22). This mechanism could also be responsible for reduced inflammatory response to exercise observed in this study.

Some investigations have evaluated the analgesic efficacy of curcumin. Di Pierro et al. reports that 400mg curcumin had a well-defined pain-relieving effect, even greater than that of acetaminophen 500 mg, and was better tolerated than nimesulide (36). Drobnic et al. also shows that curcumin supplement caused a decrease in DOMS-induced pain (23). The findings of this study support those results of earlier research. According to our data, 150mg curcumin supplement resulted in pain reduction of 48 and 72h after exercise (Figure 4). This acute effect was probably related to desensitization or inhibition of a series of transient receptor potential ion channels involved in the generation of painful stimuli such as TRPV1 and TRPA1 (37, 38).

CONCLUSION
The results of this study demonstrate that acute curcumin supplement after intensive eccentric exercise can not only keep antioxidant capacity responses high, but also serve to decrease muscle damage and pain in humans.

APPLICABLE REMARKS
- Athletes wanting to prevent development of DOMS may benefit from the anti-inflammatory and antioxidant properties of curcumin supplement taken after an intensive bout of eccentric exercise.

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REFERENCES