



Evidence-Based Supplementation Strategies for Wrestlers: A Systematic Review

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Accepted: 7 June 2025 / Published online: 25 June 2025
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Abstract

Background Wrestling is a popular combat sport that requires muscular strength, power, agility, and endurance. Weight classes have motivated wrestlers to compete at a lower weight to optimise power-to-weight ratio and performance. To achieve these characteristics, athletes may use dietary supplements, however, their efficacy in wrestlers has not been systematically evaluated.

Objective The purpose was to systematically review the literature to determine the efficacy of dietary supplements to improve body composition, physiological status, and performance in wrestlers.

Methods A systematic search was conducted in PubMed, ProQuest Medline, Web of Science, Cochrane Library, and Scopus on the 21st of January 2024 and updated on the 6th of January 2025. Studies were included if the participants were healthy wrestlers ingesting any type of dietary supplement in comparison to a control. Data associated with intervention type and characteristics, target populations, outcomes, and analysis methods were extracted.

Results A total of 24 eligible original articles were included that assessed various supplementation strategies on body composition, exercise performance, and metabolic markers in wrestlers. Individual studies revealed significant effects of sodium citrate, creatine monohydrate, spirulina, green tea and oolong tea extracts, and branched-chain amino acids on body mass or composition. β -Hydroxy- β -methylbutyrate (HMB-FA), creatine monohydrate, and iron supplementation improved recovery and may improve exercise performance. Beet-root juice supplementation enhanced muscular strength and balance. BCAA supplementation produced mixed results on muscle damage biomarkers and performance, while sodium citrate, creatine, and spirulina can act as buffering agents. Thyme tea appears to improve antioxidant capacity.

Key points

- **Supplementation Benefits for Body Composition:** Certain supplements, like sodium citrate, creatine monohydrate, and spirulina improved body mass recovery, fat loss, and muscle composition in wrestlers. However, results were mixed for some other supplements such as chromium picolinate and Spatone®.
- **Performance and Muscle Recovery:** Supplements such as creatine monohydrate and HMB-FA were shown to enhance exercise performance and reduce muscle damage biomarkers after intense activities like simulated wrestling matches. Beetroot juice also improved balance and strength, while BCAA supplementation had varied effects on muscle damage and performance.
- **Metabolic and Hormonal Impacts:** Supplementation with sodium citrate, spirulina, and iron influenced important metabolic markers, including pH and lactate levels, which may help improve recovery and reduce fatigue after intense exercise. Interestingly, sex differences were observed in some of these responses, particularly with iron supplementation, which benefited male participants more.
- These findings suggest that certain supplements can help optimize body composition, improve performance, and support recovery in wrestlers, though the effectiveness varies based on the supplement and individual responses.

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Conclusions Overall, individual studies show some promise for several dietary supplements to alter body mass and body composition, improve exercise recovery and performance, delay fatigue, and modify serum biomarkers; nevertheless, effect sizes were often small, and results were often mixed.

Keyword Wrestling · Sports nutrition · Exercise performance · Ergogenic aids

Introduction

Wrestling is one of the oldest sports, dating back to ancient Egypt and Greece [1]. Currently, wrestling is one of the most popular sports in the world, having more than 1.9 million individuals engaged in wrestling in the U.S. alone [2]. Since competitors are divided into weight classes, this has motivated wrestlers to compete at a lower weight to optimise power-to-weight ratio, body composition, and performance [3]. However, the methods used to alter body mass (BM) and composition are often non-evidence based such as severely restricting food intake, dehydration, or excessive workouts [4, 5]. Furthermore, unlike other combat sports, wrestlers typically compete much more frequently [6] thereby inducing repeated changes in weight that have been linked to a reduction in anaerobic exercise performance, depleted muscle glycogen, reduction in lean body mass, depression, and fatigue [7]. In addition, rapid weight loss achieved through the reduction of energy and fluid intake, as well as increased exercise, increases fatigue and decreases peak power [8].

Multiple factors such as energy production, hormones, inflammation, and oxidative stress could influence a wrestlers' physical performance. As a high-intensity sport, wrestlers depend on glycolysis to provide the necessary energy to the muscles [9]. In anaerobic conditions, this leads to muscle acidification through the accumulation of hydrogen ions (H^+), which is associated with muscle fatigue [10]. The physiological process of acidification and muscle fatigue can be explained by: 1) the competition of H^+ ions with calcium ions (Ca^{2+}) for the troponin binding site that prevents contraction from occurring properly; 2) the inhibition of phosphocreatine resynthesis; 3) the inhibition of key enzymes (e.g., phosphofructokinase) of the glycolytic pathway [11]; 4) and a decreased production of energy in muscle cells due to a reduced proton gradient between the mitochondrial matrix and the cellular cytoplasm [12]. Therefore, maintaining a pH in physiological ranges is critical for sustained muscle contractions to occur [13]. During high-intensity exercise, intramuscular acidity is regulated both intra and extracellularly with bicarbonate being one of the major contributors [14, 15]. Oral supplementation with sodium bicarbonate leads to

alkalemia, which creates a greater efflux of H^+ and lactate out of the active muscles and into the circulation [16]; further metabolic alkalosis results in the acceleration of glycogenesis [17] and may potentially reduce membrane depolarization that could lead to increased performance [18]. As such, supplementation with sodium bicarbonate could improve performance due to buffering with H^+ and the efflux of lactate from the muscle [16, 19].

The physiological stress that wrestlers undergo can also affect the endocrine system, which has been described for testosterone and cortisol levels. In summary, testosterone is an anabolic hormone which promotes muscle and bone mass; however, endogenous production plays a small role in muscle adaptation in comparison to the muscle's androgen receptor content [20]. Cortisol is a catabolic hormone that increases during physiological stress and stimulates glucose production, decreases amino acid uptake by the muscle, and reduces muscle and bone formation [21]. Elevations in total testosterone and cortisol have been observed in competitive wrestling [22–24], with the testosterone response reducing with subsequent tournaments. This is important given that the response of testosterone has been observed to be greater in the winners [22–24].

Wrestling has also been associated with abnormal levels of inflammatory and oxidative stress markers. Studies have reported that interleukin 1 beta (IL-1B), interleukin 6, tumour necrosis factor-alpha (TNF- α), glutathione peroxidase, superoxide dismutase, lipid hydroperoxides, and total glutathione levels, are altered through training in wrestlers [25]. Some dietary supplements have been purported to help athletes with oxidative stress and inflammatory-related damage, enhance performance and improve recovery. For instance, blueberry supplementation might reduce inflammatory and oxidative stress markers, as well as enhance recovery after exhaustive exercise [26], vitamin D supplementation might be useful to increase strength, lower inflammatory markers, and decrease the risk of injury [27], and zinc supplementation might improve appetite, increase BM, reduce fatigue, and increase endurance [28]. Interestingly, some studies have reported that a proper diet might be just as effective (or more) than supplementation with creatine or glutamine in improving exercise performance [29].

Other commonly used supplements include vitamins, minerals, botanicals or herbs, botanical compounds, amino acids and their derivatives, amongst others [30]. Despite safety and efficacy concerns for some, dietary supplements are generally considered to be part of a well-rounded approach to BM management, and using dietary supplements to modify an individual's body composition has become a common strategy when attempting to alter BM or body composition [31, 32]. As such, while supplementation promises great benefits, there is a need to investigate the efficacy in specific populations. Therefore, in this systematic review, we evaluated studies examining the effects of dietary supplements on markers of inflammation, oxidative stress, body composition, exercise performance, and recovery in wrestlers.

Methodology

This systematic review was pre-registered in the PROSPERO database (ID:CRD42023458266) and was performed according to the recommendations established by the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) [33].

Literature Search

An extensive search in PubMed, ProQuest Medline, Web of Science, Cochrane Library, and Scopus was conducted on the 21 st of January 2024 and then updated on the 6th of January 2025 to find studies that assessed the effects of supplementation on body composition, markers of oxidative stress or inflammation, endocrine responses, exercise performance, muscle damage, and recovery in wrestlers. The search expression consisted of: (“wrestlers” OR “wrestling”) AND (“supplement” OR “supplementation” OR “oral”) AND (“weight” OR “body” OR “composition” OR “antioxidant” OR “oxidative stress” OR “inflammation” OR “anti-inflammation*” OR “hormone” OR “muscle” OR “strength” OR “recovery” OR “performance” OR “aerobic” OR “anaerobic” OR “power” OR “exhaustion”). Articles relevant to our investigation that were referenced in any of the included studies were also taken into consideration. No limitation was made on the publication date or time length of the studies. Only studies in English were included. The search strategy and inclusion/exclusion criteria based on population, intervention, comparison, outcomes and study design (PICOS) have been summarised in Table 1.

Study Selection

Included studies were randomized and non-randomized controlled trials in humans that had a control group as a

Table 1 Search strategy and inclusion/exclusion criteria based on population, intervention, comparison, outcomes and study design (PICOS)

Databases	Search Terms	PICOS	Inclusion criteria	Exclusion criteria
PubMed, ProQuest Medline, Web of Science, Cochrane, Scopus	(“wrestlers” OR “wrestling”) AND (“supplement” OR “supplementation” OR “oral”) AND (“weight” OR “body” OR “composition” OR “antioxidant” OR “oxidative stress” OR “inflammation” OR “anti-inflammation*” OR “hormone” OR “muscle” OR “strength” OR “recovery” OR “performance” OR “aerobic” OR “anaerobic” OR “power” OR “exhaustion”)	Population	Wrestlers between 18 to 45 years old	Unhealthy individuals; individuals below or above the established age range; other types of athletes or non-sport practitioners
		Intervention	Any dietary supplement	Multiple ingredients used as a single intervention; non-dietary supplementation
		Comparison	Supplementation vs. no supplementation/ placebo	
		Outcome	Changes in markers of physiological status, body composition, muscle damage, exercise performance and/or recovery	Alterations in any of the markers before the intervention
		Study design	Randomized, non-randomized, controlled, crossover, and quasi-experimental studies	Meta-analysis, systematic review, cross-sectional, case-control, case reports, animal, and in vitro research studies

comparator to assess the beneficial effects of dietary supplements on markers of physiological status, exercise performance, and recovery. The participants of all the studies were wrestlers. The excluded studies were observational, animal, and in vitro studies. Two reviewers independently assessed the titles and abstracts against the inclusion and exclusion criteria. The eligible full-text articles were retrieved. The full-text screening was completed independently by the two reviewers. Any disagreements were resolved by establishing a consensus.

Data Extraction

The following data was retrieved from each study: type of intervention, target population characteristics, outcomes, and analysis of the outcomes. All data was summarised and described as qualitative and quantitative variables. A narrative synthesis was performed for the demographic characteristics of the participants such as age, sex, health status, and exercise performance activity, the characteristics of the interventions such as dose, frequency, and intervention time of the supplementation, as well as the characteristics of the placebo and the assessment tools used to determine the antioxidant, anti-inflammatory, and muscle damage markers, as well as recovery time, quantitative and qualitative exercise performance assessments.

Risk of Bias Assessment

The scientific quality of the studies was assessed independently by two reviewers using the Risk of Bias 2 tool (RoB2) for randomised and crossover trials [34], and the ROBINS-I tool for non-randomised trials [35]. The assessment of randomised trials was based on the following domains: randomisation process, assignment and adherence to intervention, missing data, measurement of outcome, and selection of the reported results. For crossover studies, the risk of bias arising from period and carryover effects was also considered. For non-randomised studies, the assessment also evaluated the bias due to confounding in addition to the domains stated for randomised studies. The studies were then categorised as having a low, some concerns, or high risk of bias. If assessment outcomes were conflicting, reviewers discussed and came to a consensus. Visualisation of the risk of bias assessments was performed using the robvis online tool [36].

Results

Study Selection

The review identified 267 records by searching the five databases. After removing duplicates ($n = 84$), 183 articles

remained, from which 120 non-clinical trial articles were identified and removed before the screening. A total of 63 articles were then screened by title, abstract, and keywords by the reviewers independently. A total of 39 articles were excluded for different reasons: study included underage participants ($n = 20$), study included multiple sports ($n = 9$), the complete article was not found ($n = 4$), the article was not in English ($n = 2$), the study was a thesis/poster ($n = 2$), the study used non-diet supplementation ($n = 1$), or study was of low quality ($n = 1$). A total of 24 articles were assessed for eligibility. The details of the study selection process are shown in Fig. 1.

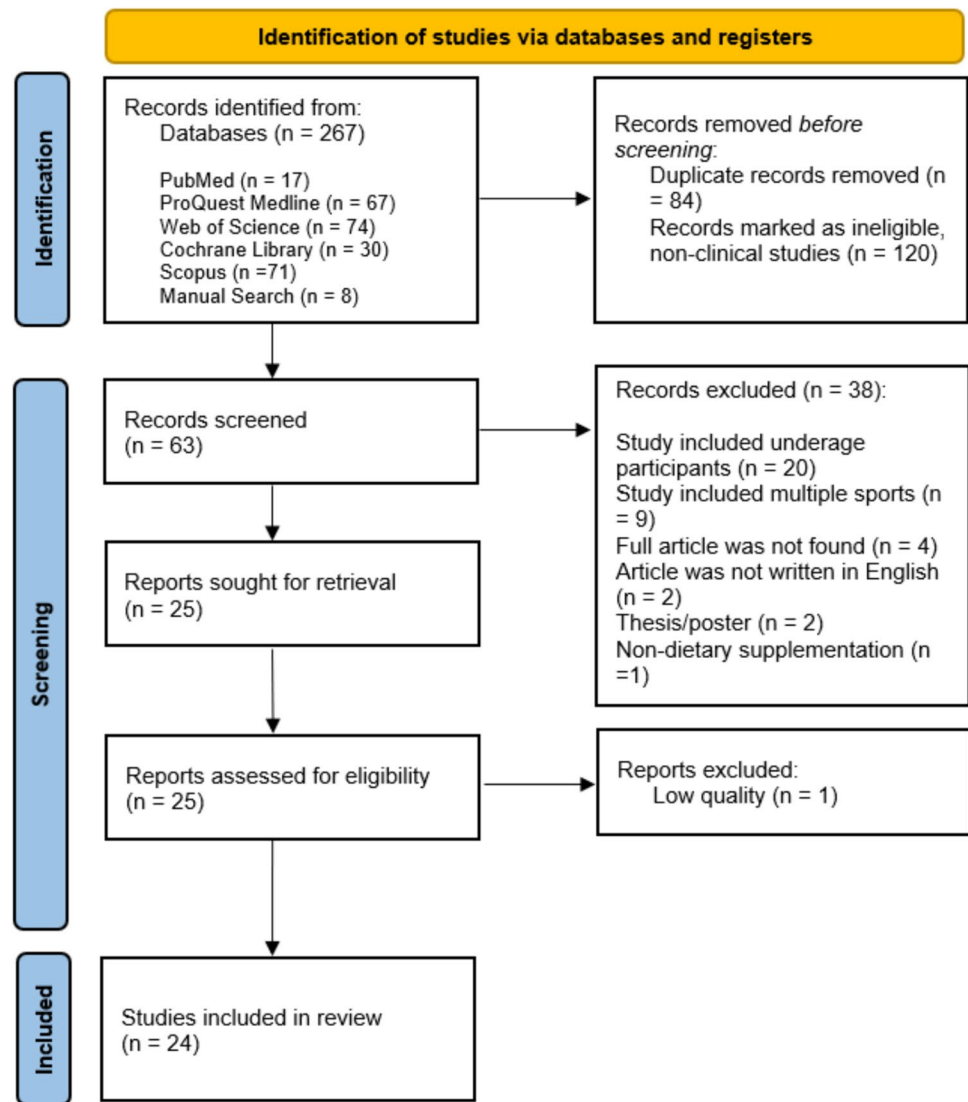
Characteristics of the Included Studies

Among the included studies, one study was a non-randomised, double-blind, placebo-controlled trial [37]; five studies were randomised, double-blind, placebo-controlled trials [38–42]; eight studies were randomised, single-blind, placebo-controlled trials [43–48]; and ten studies were placebo controlled, crossover trials [49–56]. The total supplementation duration ranged from an acute dose 30 min before the test to 14 weeks. Supplementation was given as a single dose or up to four times a day [29, 37–43, 45–53, 56–58]. Nine articles reviewed the effects on body composition [37, 38, 40, 42, 43, 47, 50, 56, 57]; 17 studies evaluated the effects on exercise performance or muscle damage [29, 37–39, 41, 43, 45, 48–52, 54–58], 11 studies analysed the effects on hormonal and metabolic markers [37, 38, 40, 42, 46, 50, 52–55, 57], and six articles reported other outcomes [40, 46–48, 55, 56]. The details of supplementation used in each study are described in Table 2.

General Findings

This systematic review included 24 studies (23 randomised and 1 non-randomised trial). The studies included between five to forty wrestlers who were considered healthy and young (18–29 years old). Sixteen studies evaluated men, only one study evaluated women, another one evaluated both men and women, and six did not specify sex. A total of 415 participants were studied, 28 of which were women and 274 were men; the sex of 113 participants was not disclosed. The studies analysed the effects of supplementation after various types of trials such as running, cycling, callisthenics, using exercise machines, wrestling match simulations, and habitual training. The different types of supplementations included sodium citrate, chromium picolinate, creatine monohydrate, arginine, branched-chain amino acids (BCAA), beetroot juice, β -Hydroxy- β -methylbutyrate (HMB-FA), spirulina, Spatone®, carbohydrates, and glutamine. The relevant changes observed with each type of supplementation are

Fig. 1 Preferred reporting items for systematic reviews and meta-analyses (PRISMA) flow diagram



discussed throughout Sects. 3.3.1 to 3.3.4 and summarized in Table 2 and Fig. 2 [29, 37–43, 45–54, 56–58].

Body Composition

A study evaluating the effects of sodium citrate supplementation reported a higher BM recovery after rapid body mass loss in the experimental group [37]. Neither group achieved the same BM as before the rapid body mass loss, but the deficit was significantly higher in the placebo group. No significant differences between groups were found in underwater weighing, skinfold thickness, body circumference, BM, lean BM, body fat %, or fat mass with Chromium picolinate (CrPic) supplementation [38]. Three studies evaluated the effects of creatine monohydrate (CM) on body composition [59]. Kocak & Karli [43] found a significant increase in the mean BM gained during 5 days of supplementation.

Zahabi and colleagues (2024) found similar results in female wrestlers, where 25 days of supplementation significantly increased BM and BMI [59]. Oopik and colleagues (1998) examined acute creatine monohydrate + glucose supplementation but revealed no effect on BM regain during a 17-h rapid body mass loss recovery period [50]. A study investigating the effects of spirulina supplementation found significantly lower body fat %, skeletal muscle mass (SMM), and fat mass in the experimental group [42]. Sung and colleagues (2018) examined Spatone®, which is a natural iron-rich water supplementation, on SMM, body fat, and body mass index (BMI), but found no significant differences between groups [56]. Oolong (OTG) tea extracts were able to significantly influence BM reduction in the oolong group at weeks 4 and 6. The BM reduction was accompanied by a significant decrease in adipose tissue. Additionally, there was a significant difference in the percentage of fat reduction

Table 2 Summary of clinical-based trials evaluating the effect of dietary supplementation on physiological status and/or performance-related outcomes of wrestlers

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Sodium citrate	Timpmann, S., et al., 2012	Non-randomised double-blind placebo controlled parallel trial	n = 16 n = 8 Experience: 5–13 years of training Sex: N/S Age: 18–26 years	Controlled diet with a single dose of 600 mg/kg	n = 8 Controlled diet with a single dose of wheat flour	Acute dose 16 h before testing	BM USG MP FI Blood pH Blood HCO ₃ BE Blood lactate	UBISP test	BC: The average gain in BM was significantly greater in the CIT group ($p = 0.008$) with a significant interaction effect between test and treatment ($p = 0.006$). No difference between groups in mean USG MDEP: No significant difference in MP, PP, or FI between groups HMM: Significantly higher HCO ₃ concentrations ($p = 0.018$), higher pH levels ($p = 0.001$), and significantly higher base excess in the CIT group ($p < 0.0001$)
Chromium picolinate	Walker, S., et al., 1998	Randomised double-blind placebo-controlled parallel trial	n = 20 n = 7 Experience: NCAA division I Sex: male Age: 18–23 years	Gelcap with 200 µg once a day	n = 7 Gelcap with sodium diphosphate n = 6 No supplementation	14 weeks	Insulin Glucose BM LBM FM Fat % Leg power Upper body power endurance Peak AnP Rel AnC Rel VO ₂ max	Progressive resistance training program and metabolic conditioning regimen Bruce protocol Wingate 30-s cycle ergometer test Upper body endurance (maximal repetitions of seated low-pulls) Absolute lower body endurance (maximal repetitions of leg press performed) Global muscular power (Olympic power clean) Maximal upper body strength (IRM bench press)	BC: No significant differences in BM, LMB, Fat %, or FM between groups MDEP: No significant differences in leg power, upper body power, endurance, Peak AnP, Rel AnC, and Rel VO ₂ max between groups HMM: No significant differences in fasting serum glucose levels or insulin concentrations between groups
Creatine monohydrate	Koçak, S., et al., 2003	Randomised placebo-controlled parallel trial	n = 20 n = 10 Experience: Turkish National Team Sex: male Age: 22–27 years	5 g four times a day dissolved in 250 ml of water 1 h prior to eating	n = 10 5 g of milk powder four times a day dissolved in 250 ml of water 1 h prior to eating	5 days	BM AP PP	Wingate 30 s anaerobic test	BC: Significantly higher weight gain in CM group ($p < 0.01$) MDEP: Significant difference between pre-and post-test scores of AP ($p < 0.01$), PP ($p < 0.01$)

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Creatine monohydrate	Mohamed, E., and Tammam, A., 2020	Randomised, double-blind placebo controlled trial	Experience: Shooting wrestling club Sex: N/S Age: 21–23 years	n = 16 n = 8 1 st week: 20 g of CM dissolved in 250 ml of mil juice for four days Week 2–8: 5 g of CM dissolved in 250 ml of mil juice for 33 days (245 g in 8 weeks)	n = 8 250 ml of mil juice	8 weeks	CPK Muscular endurance Power Agility	Back-throw dummy test Bridge skill test Performance of the skill of the bridge test	MDEP : Statistically significant effects of the time (i.e. pre- to post-training) for all measured variables ($p \leq 0.05$) in the CM group. Statistically significant differences for CPK ($p = 0.000$), muscular endurance ($p = 0.002$), power ($p = 0.003$), and agility ($p = 0.036$) in the CM group at the POST evaluation
Creatine monohydrate + glucose	Özpik, V., et al., 2002	Randomised double-blind placebo controlled crossover trial	Experience: 7 years average Sex: male Age: 19–21 years	n = 5 n = 5 80 g of glucose + 7.5 g of CM four times a day dissolved in 300–350 ml of natural fruit juice	n = 5 320 g of glucose four times a day dissolved in 300–350 ml of natural fruit juice	Acute dose immediately after the first test (17 h before the final test) The next two doses, three and six hours after the first test, respectively The last dose, at least two hours before the final test	BM Submaximal work Wtot Wmax	Isokinetic performance of the knee extensors for 5 min	BC : No effect of treatment on the extent of BM regained during 17 h recovery MDEP : Significant increase in Wtot from test 2 to test 3 in CM + GLC trial. A 13.8%–44.5% increase in Wmax in several time points in glc + cr trial ($p = 0.02$) HMM : No significant difference in ammonia, lactate, glucose, or urea levels between groups
Carbohydrates/ Creatine monohydrate/Glutamine	Abbasalipour, M., et al., 2012	Randomised, single blind parallel group controlled trial	Experience: elite wrestlers Sex: male Age: 18 to 25 years	n = 28 n = 7 Carbohydrate solution made of 5% honey n = 7 0.3 g/kg of CM per day n = 7 Glutamine supplement 0.3 g/kg/day	n = 7 Control group	15 days	Grip strength Agility	Bicycle ergometer till the point of complete exhaustion 9 × 4 agility test Hand grip	MDEP : Significant increase in grip strength and agility performance in CM group ($p < 0.05$)
Creatine Monohydrate	Zahabi, G., et al., 2024	Randomised, single blind parallel group controlled trial	Experience: International/professional wrestlers Sex: female Age: 18 to 19 years	n = 18 n = 6 Loading phase (5 days): 5 g of CM four times per day Maintenance phase (20 days): 5 g of CM 30 min before and immediately after training	n = 6 training without supplementation n = 6 no training, no supplementation	25 days	BM BMI BF% FFM VO ₂ max RHR IRM Agility Muscular power	Resistance training Chest and leg press Modified Bruce protocol Chest and leg press IAT VJT	BC : Significant increase in BM and BMI in CM group MDEP : Significant increase in VO ₂ max, IRM, agility, and muscular power in CM group

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Citrulline Malate	Jafari, R., et al., 2024	Randomised, crossover trial	Experience: Varsity Junior team 5+ years Sex: N/S Age: 18 to 21 years	n = 12 n = 12 8 g of citrulline malate	n = 12 N/S	Acute dose one hour before testing	Hypoxanthine HGPRT Grip strength BAT test BLC strength test RPE	JSWPT protocol Hand grip Burpees Isometric force with a dynamometer	HMM: No significant differences in hypoxanthine or HGPRT levels MDEP: No significant differences in grip strength, BAT test, or BLC strength test
Arginine	Ulas, H., et al., 2012	Randomised placebo-controlled crossover trial	Experience: National and international level Sex: male Age: 20–28 years	n = 10 n = 10 1.5 g/10 kg capsules after a 12 h fasting once a day	n = 10 Capsules with starch	Acute dose 60 min before the test and after an overnight fast	Amino acid levels	Cycle ergometer Warm up at 60 rpm without any load for 3 min, followed by 90 watts load increasing 30 watts every 3 min till the exhaustion	HMM: Significantly higher pre- and post-exercise arginine, ornithine and citrulline concentrations in arginine trial ($p < 0.05$) Higher post-exercise TAA, BCAA, glutamine, tyrosine, methionine, phenylalanine, isoleucine and leucine concentrations in arginine trial ($p < 0.05$)
Arginine	Yavuz, H. U., et al., 2014	Randomised crossover trial	Experience: national and international level wrestlers Sex: male Age: 24.7 ± 3.8 years	n = 9 n = 9 Single dose of 1.5 g/10 kg body weight arginine capsules	n = 9 Equal number of capsules containing starch	Two weeks One week wash-out period	Lactate HR Time to exhaustion Maximum oxygen consumption	Incremental bicycle ergometer test to exhaustion	HMM: No significant difference in mean lactate levels MDEP: No significant difference in maximum oxygen consumption or in maximum heart rate Time to exhaustion was longer with arginine supplementation compared to placebo ($p < 0.05$)
Arginine	Zembron, A., et al., 2020	Randomised, double-blind, parallel group, placebo-controlled trial	Experience: Members of the Polish national team Sex: male Age: 20 to 29 years	n = 32 n = 7 Mem-Arginine 2 × 6 g per day n = 9 Hypoxia and arginine 2 × 6 g per day n = 6 Hypoxia	n = 10 Placebo 2 × 6 g per day	12 days	BM FFM FM CK NO H2O2 CRP HGF IGF-1 PDGF VEGF BDNF TC HDL LDL TG Hb RBC RET HTC MCV MCH MCHC RDW	14-day training camp	BC: No significant difference between groups HMM: No significant differences between groups OO: NO levels significantly increased in arginine group ($p < 0.05$)

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Carbohydrate, BCAAs and arginine	Jang, T. R., et al., 2022	Randomised, double-blind, cross-over trial	Experience: At least 4 years and experience in national or international competitions Sex: male Age: 19.2 ± 0.4 years	n = 9 1.2 g/kg glucose (CH trial) 1 g/kg glucose + 0.1 g/kg Arg + 0.1 g/kg BCAA (leucine: isoleucine: valine = 2:1:1, CH + AA trial)	n = 9 600 ml lemon flavored water	One day Wash-out period of at least 2 weeks	PP MP Glucose Insulin Glycerol NEFA Lactate	3 wrestling matches	MDEP: No significant differences between groups HMM: Significantly higher glucose and insulin levels in CH + AA trial and lower glycerol and non-esterified fatty acid concentrations
BCAA	Armisan, R., et al., 2014	Randomised placebo-controlled semi-experimental trial	Experience: trained Mahabadi City wrestlers Sex: N/S Age: 22 years	n = 29 Low Dose n = 10 68 mg/kg × 3 times/day before meals for six days before the test and two days after 210 mg/kg 30 min before and after the test High Dose n = 10 68 mg/kg × 3 times/day before meals for six days 450 mg/kg 30 min before and after the test	n = 9 Dextrin 68 mg/kg three times a day before meals for six days before the test and two days after 210 mg/kg 30 min before and after the test	8 days Acute dose 30 min before and after the test	CK CKMB LDH	80% IRM leg presses, chest presses, lat pull downs, leg extensions, arm curls, leg curls, and abdominal crunches	MDEP: No significant difference in CK, CKMB, or LDH levels between groups
Caloric restriction/BCAA	Mourier, A., et al., 1996	Randomised, single-blind, parallel-group, controlled trial	Experience: wrestlers of the French National Institute of Sports Sex: male Age: N/S	n = 31 n = 7 Hypocaloric high-protein n = 6 Hypocaloric high-branched-chain amino acid n = 6 Hypocaloric low-protein	n = 6 Normocaloric control n = 6 Hypocaloric control	19 days	BM BMI BF SAT VAT AT MT VO2max MVC Glycerol T3	Treadmill Right knee isometric extensions Arm ergometer Wingate Anaerobic Capacity Test	BC: Significantly higher BM loss on hBCAA, with a significant loss on SAT ($p < 0.05$). Significant effect on time and diet on thigh muscle adipose tissue with a greater loss in hBCAA group ($p < 0.05$) MDEP: No significant differences in VOMax, peak power, or endurance time of the extensor muscles HMM: No significant differences in glucose, lactate, or insulin plasma levels

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Beetroot Juice	Tatlici, A., et al., July 2021	Randomised double-blind crossover trial	Experience: trained wrestlers Sex: male Age: 19–24	n = 8 140 ml of BRJ	n = 8 140 ml of cherry juice with lemon juice	Acute dose 150 min before the test	OSI APSI MLSI	Dynamic and static balance in a biodex balance system Maximal contraction knee extension and flexion	MDEP: At rest- static MLSI, dynamic OSI, and dynamic APSI significantly improved in BRJ. ($p=0.00$, 0.03 , 0.01 , respectively) At fatigue- static OSI, static APSI, dynamic OSI, dynamic APSI, and dynamic MLSI significantly improved in BRJ. ($p=0.00$, 0.01 , 0.01 , 0.02 , 0.02 , respectively)
Beetroot Juice	Tatlici, A., June 2021	Randomised double-blind crossover, placebo-controlled trial	Experience: trained wrestlers Sex: male Age: 19–24	n = 8 140 ml of BRJ	n = 8 140 ml of cherry juice with lemon juice	Acute dose 150 min before the test	PE PF PIR PER AE AF AIR AER	Extension and flexion strength of the knee Internal and external rotation strength of the shoulder	MDEP: No statistically significant difference in ExP and FlexP. Significant increase in IntP and ExP in BRJ. ($p=0.048$ and $p=0.024$, respectively) Significant increase in ExtAvg, FlexAvg, IntAvg, and ExtAvg in BRJ ($p=0.023$, 0.027 , 0.023 , and 0.021 , respectively)
HMB-FA	B. Tartibian, B. Rezaei, 2021	Randomised double-blind placebo-controlled parallel trial	Experience: elite wrestlers Sex: N/S Age: 19–26	n = 20 n = 10 3 g/d of HMB-FA	N/S	Single dose	CK LDH PRS	Five simulated wrestling protocols	MDEP: Significantly lower levels of LDH after the first, third, and fifth tests. ($p<0.05$) Significantly lower level of CK after the fifth test. ($p<0.05$)
Spirulina	Bagheri, R., et al., 2021	Randomised, placebo-controlled double-blind trial	Experience: wrestling training at least three times a week for at least 3 years before the study Sex: N/S Age: 22 years	n = 40 Designed diet for weight loss Two tablets containing 500 mg of spirulina	n = 20 Designed diet for weight loss Two tablets containing 500 mg of placebo	12 days	BM BFP FM SMM FST MST IGF-1 AST ALT	Daily moderate physical activity lasting 20–40 min, which consisted of technical training	BC: Significantly lower BFP, SMM, and FM in SP group ($p<0.001$) HMM: Significantly lower MST, AST, and ALT concentrations in SP group ($p=0.005$). Significantly lower FST and IGF-1 concentrations in PL group ($p<0.05$)

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Spatone®	Sung, Y., et al., 2018	Randomised, single-blind, crossover, group controlled trial	Experience: amateur wrestlers Sex: male Age: N/S	n = 9 n = 7 Spatone water containing 5 mg of iron in orange juice, two times a day for 1 week of weight-loss period	n = 6 Weight loss of 7% over 7 days with placebo	7 days 4 weeks wash-out period	BM SMM FM BMI VO2max Lactate Calcium Magnesium Iron RBC Hb Hct MCV Platelet WCV MCH MCHC IL-10 TNF-alpha IL-6	Morning session, consisted of running and dashing, on Monday, Wednesday, and Thursday. The interval training and fartlek training were on Tuesday and Friday. For the afternoon session, mat training For the evening session, the subjects performed weight training on Tuesday and Thursday. Training protocol twice or thrice per day and the training session that consisted 30 h a week of exercise, 5 h each day	BC: No significant differences between groups MDEP: Significantly higher endurance capacity, VO2max, and lactate accumulation in Spatone group ($p < 0.05$) OO: No significant differences between groups
Thyme tea	Berkan, C., et al., 2013	Randomised, single-blind, parallel-group, placebo controlled trial	Experience: wrestler students of Nigde University Physical Education and Sports School Sex: male Age: 18 to 28 years	n = 18 n = 9 Thyme tea three times/day and a loading dose of 150 cm ³	n = 9 Control group	35 days	MDA TAC RSH	5 wrestling matches	HMM: Significant increase in TAC ($p < 0.05$) OO: Significantly lower MDA levels in thyme group ($p < 0.01$)

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Green tea extract/ Oolong tea extract	Bajerska, J., et al., 2010	Randomised, single-blind, parallel group, placebo controlled trial	n = 35 Experience: Solesky wrestling team Sex: male Age: 18 to 24 years	n = 10 Two capsules three times/day of 400 mg, containing 60% green tea extracts n = 10 Two capsules three times/day of 400 mg, containing 40% oolong tea extracts	n = 10 Two capsules three times/day of 400 mg, containing 100% cellulose	6 weeks	BM FM FFM MEB		BC: Significantly lower BM in GTE ($p < 0.05$) and OTE ($p < 0.01$) groups. Significantly lower FM in OTE group. ($p < 0.05$) OO: No significant differences between groups
Iron	Sung, J. Y., 2021	Randomised, single-blind, crossover trial, placebo-controlled trial	n = 23 Experience: N/S Sex: 13 males and 10 females Age: 21.6 ± 0.8 years for males and 20 ± 1 for females	n = 23 Water containing 5 mg of iron and orange juice two pouches per day	n = 23 Orange juice	7 days 3 week washout period	BM RBC Hb Hct WCW MCH MCHC TIBC Fe Transferrin EPO VO2max Lactate	Training protocol two or three times each day, totaling around 5 h of exercise each day, and 30 h over the week Morning session, running and dashing on Monday, Wednesday and Thursday. Interval day. Interval training on Tuesday and Friday Afternoon session, mat training Evening session, weight training on Tuesday and Thursday	OO: Significantly lower Fe, ($p < 0.041$); transferrin, ($p < 0.004$); TIBC ($p < 0.031$) in males Both groups experienced decreases in erythropoietin (males, $p < 0.021$; females, $p < 0.027$) MDEP: Significant decrease in VO2max ($p < 0.001$) after weight loss HMM: Blood lactate in the intake group decreased after maximal exercise immediately after the test for the male group ($p < 0.031$) and during resting time and immediately after exercise in the female group ($p < 0.001$ and $p < 0.050$, respectively)
Whey protein	Shawwy, A., 2013	Randomised, single-blind, parallel group, controlled trial	n = 18 Experience: trained wrestlers Sex: male Age: N/S	n = 18 Supplement pre-exercise 1.4 g/kg of BM/day n = 8 Post-exercise supplement 1.4 g/kg of BM/day	n = 18 Control group	Twelve weeks	Total protein Albumin Urea Creatinine Leg extension Barbell Bench Press Barbell Front Raise	Squat, chest and arm exercises	MDEP: Significant increase in strength on the barbell bench press ($p < 0.05$) OO: Significant higher total protein and albumin levels ($p < 0.05$)

Table 2 (continued)

Supplement	Study	Study design	Participants	Supplemented group	Placebo/control group	Duration	Measured outcomes	Training Load	Key Findings
Caffeine	Negaraesh, et al., 2018	Randomised double-blind, crossover trial, placebo-controlled trial	<p>n = 12 n = 12</p> <p>Experience: professional male freestyle, wrestling experience if at least 10 years</p> <p>Sex: male</p> <p>Age: 24 ± 3 years</p>	<p>High-dose of caffeine 10 mg/kg 5 times a day</p> <p>Moderate-dose of caffeine 4 mg/kg 5 times a day</p> <p>n = 12</p> <p>Repeated dose caffeine 5 × 2 mg/kg 5 times a day</p> <p>n = 12</p> <p>Selective caffeine consumption 6.16 ± 1.58 mg/kg 5 times a day</p>	<p>n = 12</p> <p>Placebo</p>	1 day	<p>PWPT time</p> <p>HR</p> <p>Fatigue rating</p> <p>Lactate</p> <p>Urine osmolality</p> <p>USG</p>	<p>Simulated wrestling tournament PWPT</p>	<p>MDEP: Significantly lower PWPT time before the first match with high dose of caffeine ($p < 0.05$)</p> <p>Significantly lower PWPT time during the third and fourth matches with repeated doses and selective administration ($p < 0.05$)</p> <p>Significant time effect for hip/back strength and vertical jump height ($p < 0.05$)</p> <p>Significantly lower fatigue rating before the fourth match with selective and repeated doses ($p < 0.05$)</p>

at weeks 2, 4, and 6, as well as in the green tea extract group (GTG) at week 6 [47]. A study found that high doses of caffeine caused a higher urine volume output and dehydration index [49]. Caloric restriction and branched-chain amino acid (BCAA) supplementation were found to have a significant main effect of time on BM reduction; interestingly, the hypocaloric branched-chain amino acid group (hBCAA) was the one with the greatest BM loss, where a significant loss in subcutaneous adipose tissue was found. A significant effect on time and diet was also present on the thigh muscles adipose tissue, where the hBCAA group had a significantly higher loss of adipose tissue than the other hypocaloric groups [57].

Muscle Damage and Exercise Performance

Sodium citrate supplementation resulted in no significant differences between groups performing upper body intermittent sprint performance [37]. CrPic supplementation showed no significant differences between groups on peak aerobic power, peak anaerobic power, maximal anaerobic capacity, upper and lower body endurance, upper body power, bench press power, leg press power, or global muscular power [38]. Two studies analysed the effects of BCAA supplementation on muscle damage biomarkers and exercise performance. Amirsasan and colleagues (2011) found no significant difference in creatine kinase (CK), creatine kinase-MB (CKMB), or lactate dehydrogenase (LDH) between groups [45]. Mourier and colleagues (1997) found no significant differences in $VO_2\text{max}$, peak power output, or endurance time of the extensor muscles with caloric restriction and BCAA supplementation [57]. Jafari and colleagues (2024) found that citrulline malate supplementation had no significant effects on hand grip strength, the back-chest-leg strength test, nor the burpee agility test [60]. Supplementation with β -Hydroxy- β -methylbutyrate (HMB-FA) resulted in significantly lower CK and LDH after simulated wrestling protocols [41]. Four studies evaluated the effects of CM supplementation on exercise performance, all of which reported favourable results. Zahabi and colleagues (2024) reported a significant increase in $VO_2\text{max}$, one repetition max on chest and leg press, force applied during a vertical jump test, as well as a faster time during the Illinois agility test [59]. Kocak & Karli (2003) reported a significantly higher average power and peak power after supplementation [43]; Oopik and colleagues (1998) reported a significant increase in maximal work (W_{max}) and total work (W_{tot}); additionally, a strong correlation was established between whole-body creatine retention and the extent of change in W_{max} [50]. Sabry & Tammam (2020) reported significantly better effects of the time for creatine phosphokinase (CPK), muscular endurance, power, and agility [39].

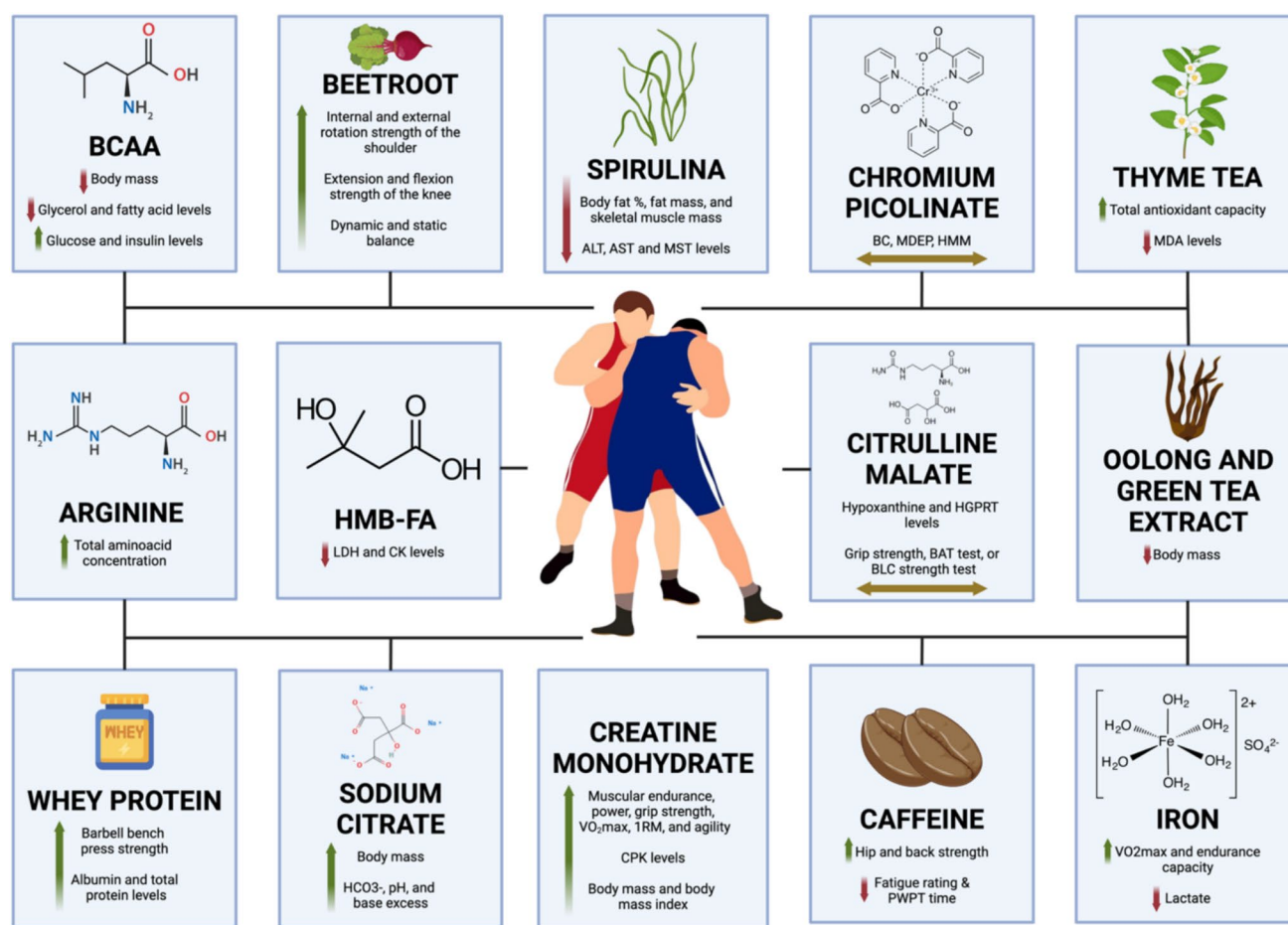


Fig. 2 Summary of the effects of different dietary supplements on body composition, exercise performance, and hormonal and metabolic markers

Two studies evaluated beetroot juice supplementation (BRJ). Tatlici and colleagues (2021) found significantly improved balance, resulting in a better at-rest static medial–lateral stability index (MLSI), dynamic overall stability index (OSI), and anterior–posterior stability index (APSI). At fatigue, significant improvements in static OSI, APSI, and dynamic OSI, APSI, and MLSI were also found. The second study reported no statistically significant difference in peak extension and flexion knee strength, however, a significant increase in peak strength of internal and external rotation of the shoulder was found after supplementation. Additionally, all average strength values for extension and flexion of the knee, as well as strength values for internal and external rotation of the shoulder, significantly increased [58].

Two studies analysed the effects of arginine supplementation on muscle damage biomarkers and exercise performance. Yavuz and colleagues (2014) found that supplementation significantly increased time to exhaustion by 5.8% compared to placebo [53]. The second study evaluated the

effect of intermittent hypoxic exposure and supplementation with high doses (12 g) of arginine and found no significant differences in CK levels between groups [40]. Iron supplementation was found to either diminish the reduction of VO₂max after exercise [55] or augment VO₂max [56]. A study that analysed caffeine intake found that the Pittsburgh Wrestling Performance Test (PWPT) time was lower in the high-dose (10 mg/kg) caffeine group. Additionally, a repeated dose of caffeine as well as selective supplementation, reduced the PWPT times before the third and fourth match. A significant time effect for hip/back strength and vertical jump height was found as well, however, there were no differences in performance between caffeine-intake protocols. Further, the fatigue rating was lower before the fourth match in the selective and repeated dose administration group, while the placebo group reported higher fatigue ratings before the third and final matches [49]. With regards to whey protein supplementation, exercise performance was not improved when taken either immediately after a training session or 40 min prior on leg extensions or barbell front

raises; nevertheless, it did increase performance on the barbell bench press [48].

Hormonal and Metabolic Markers

Sodium citrate supplementation reported higher pH, HCO₃⁻ levels, and base excess in the experimental group [37]. CrPic supplementation reported no significant differences in insulin or glucose concentrations [38]. One study evaluated the effects of CM supplementation on metabolic markers and reported no significant differences in ammonia, lactate, glucose, or urea [50]. Two studies evaluated arginine supplementation on metabolic parameters. The first one reported that the concentrations of glutamine, tyrosine, methionine, phenylalanine, leucine, isoleucine, and total BCAA (branched chain amino acids –valine, leucine, and isoleucine) levels were significantly higher following exercise in the experimental group. Additionally, pre- and post-exercise concentrations of arginine, ornithine, and citrulline were higher in the supplementation group [52]. The second one reported no significant effect on lactate concentrations [53]. Jafari and colleagues (2024) reported that an acute dose of citrulline malate had no effect on hypoxanthine or hypoxanthine–guanine phosphoribosyltransferase (HGPRT) levels [60]. The study evaluating the effects of spirulina supplementation described a significant main effect of time for follistatin (FST) concentrations, a significant group for time effect for myostatin (MST) and FST:MST, as well as significantly lower aspartate aminotransferase (AST) and alanine aminotransferase (ALT) levels. Additionally, the MST concentrations significantly decreased in the supplementation group, the FST:MST ratio was significantly lower in the placebo group, and the FST and insulin-like growth factor 1 (IGF-1) levels significantly decreased in this group as well [42].

Two studies analysed the effects of BCAA supplementation on metabolic markers. Jang and colleagues (2011) investigated the effects of carbohydrate, BCAA, and arginine supplementation and reported significantly higher concentrations of glucose at 30 min and significantly higher insulin concentrations after 30, 60, and 90 min [54]. Mourier et al. (1997) found that the concentrations of glucose, lactate, and insulin in plasma had no significant changes with caloric restriction and BCAA supplementation. Nonetheless, the nitrogen-enriched (N-enriched) diets lowered the concentrations of free fatty acids (FFA) and raised the concentrations of glycerol. Additionally, triiodothyronine (T3) concentrations were significantly lower in the N-enriched diets [57].

Thyme tea supplementation resulted in a significant increase in total antioxidant capacity (TAC) [46]. Iron supplementation significantly lowered lactate levels after 10 min of recovery in the supplement group; interestingly, not only did male participants show a significant decrease in lactate

after rapid body mass loss, but they also had significantly higher levels during the resting period and immediately after testing, than women [55]; another trial observed a decrease in lactate accumulation in the early phase after exhaustive exercise but resulted in no significant changes in calcium and magnesium concentrations [56]. Repeated and selective (administration of caffeine based on performance decrement before the wrestling) caffeine supplementation increased lactate levels after the third match. With selective administration, lactate levels were lower before the fourth match, but higher with repeated administration [49].

Effects on Other Outcomes

Other outcomes include anti-inflammatory and antioxidant markers, as well as growth factors, protein and albumin levels, and blood parameters. Thyme tea significantly decreased malondialdehyde levels [46]. No significant differences in mean energy balance values were observed after supplementation with GTE or OTE [47]. Iron supplementation resulted in a significant increase in haemoglobin (Hb) and hematocrit (Hct) levels. Interestingly, iron concentrations, transferrin, and total iron bound capacity (TIBC) increased significantly, but only in the male supplement group [55]; further, the proinflammatory cytokines interleukin 10 and 6 (IL-10 and IL-6), and tumour necrosis factor-alpha (TNF-alpha) had no significant changes between groups [56]. Intermittent hypoxic exposure (IHE) and a high dose of arginine significantly increased nitric oxide (NO) and hydrogen peroxide (H₂O₂) concentrations, nevertheless, arginine alone was not as effective. A similar pattern was observed with C reactive protein (CRP), multiple haematological markers (i.e., haemoglobin, hematocrit, amongst others), as well as with tissue regeneration mediators such as hepatocyte growth factor (HGF), insulin-like growth factor 1 (IGF-1), platelet-derived growth factor (PDGF), brain-derived neurotrophic factor (BDNF), and vascular endothelial growth factor (VEGF) [40]. A 12-week whey protein supplementation resulted in significantly higher total protein and albumin levels [48].

Risk of Bias Assessment

A total of 23 studies were randomised trials and only one was a non-randomised trial [29, 38–43, 45–48, 50–54, 56–58] (Figs. 3 and 4). Out of the 23 studies, one presented a high risk of bias, and three raised some bias concerns [41, 43, 48]. The first one posed a high risk of bias because of missing data. This study evaluated CK, LDH, and perceived recovery status (PRS) but only presented a graph for LDH values over time. No data was presented for the other two variables [41]. The second study raised some bias concerns regarding the randomisation process and deviations from the intended interventions since the

participants were not randomly allocated and because there was insufficient information to determine if this deviation arose because of the trial context [43]. The last study raised some bias concerns since it is not clear on how the allocation sequences were concealed, however, baseline imbalances do not suggest a problem. It is also not clear if the outcome assessors were blinded when evaluating results [48]. Other than the lack of randomisation, the non-randomised study was deemed to have a low risk of bias [37].

Discussion

This systematic review evaluated the effects of dietary supplements on body composition, muscle damage, exercise performance, and hormonal and metabolic markers. Out of multiple ergogenic aids, we found that some of them had significant effects on exercise performance, such as creatine monohydrate [29, 39, 43, 50]; while others had little or no effects, such as BCAA and arginine [45, 54, 57].

Body Composition

The ingestion of a single 600 mg/kg dose of sodium citrate was useful for regaining BM. Supplementation favours rehydration and restoration of BM through water retention, as well as through a reduced volume of urine excretion within a couple of hours after consumption. This in turn leads to a higher plasma volume (PV), which might be beneficial for the cardiovascular system and exercise performance [61, 62]. In addition, previous reports suggest that lower doses (200–300 mg) are better tolerated and produce less gastrointestinal effects than higher doses (500–600 mg), however this also diminishes the overall effect of supplementation [63, 64].

CrPic supplementation, with a dose of 200 µg/day for 14 days was not able to change body composition in wrestlers. CrPic has been implicated in inducing the effects of insulin, which should in theory promote muscle anabolism via a reduction in muscle protein breakdown [65–67]. Nevertheless, only a few studies have reported desirable effects in body composition with supplementation [68–70]. This might be because of estimation errors associated with the use of tools other than hydrodensitometry to determine body composition or because of the lack of control groups to assess the possibility of a placebo effect. This premise is supported by similar results obtained in other studies where CrPic supplementation coupled with strength training did not result in significant alterations in body fat or lean BM [71, 72].

CM supplementation for five days was able to significantly increase BM. These findings were consistent with other studies where weight gain was reported with dosages

ranging from 1.5 to 25 g/d for at least three days in untrained and trained participants [73]. Even high doses (20–25 g/d) for short periods of time have been reported to increase BM between 0.7 to 1.6 kg [74–76]. These effects are due to water retention and/or protein synthesis. However, an acute supplementation of CM + glucose in wrestlers did not accelerate the restoration of BM over 17 h. Potentially, the supplementation duration was too short, however, previous studies have demonstrated that there is a higher accumulation of BM and muscle creatine when combining creatine monohydrate with carbohydrates, nonetheless, it is likely that these effects occur within days, not hours, even when ingested with glucose [77]. These finding might not come as a surprise when other studies analysing a similar dosage for longer periods of time reported only a 0.1 kg/day increase in BM [78, 79]. Another explanation might be that the effect of creatine on BM restoration was masked by the marked (~ 2.4 kg) and fast (17 h) weight increase. In short, the absence of a change in BM does not strictly determine that the creatine supplementation has failed to increase the muscle's creatine content [29].

Spirulina supplementation, with 500 mg/day for 12 days, significantly decreased body fat percentage and fat mass. In theory, these effects are caused by activating an AMP protein kinase signalling pathway and sirtuin 1 in adipose tissue and skeletal muscle. Sirtuin 1 deacetylates nuclear factor-κB (NF-κB), c-Myc, forkhead transcription factor 1 (FKHR), and peroxisome proliferator-activated receptors (PPAR-γ), increasing the levels of adiponectin which in turn decreases fat mass [80–85]. Not only does it raise the expression of PPAR-γ coactivator 1-α and uncoupling protein 2 in skeletal muscle, which results in the up-regulation of adiponectin, but it also promotes mitochondrial biogenesis and fat oxidation [82–84]. Supplementation with oolong tea extracts, with a dose of 2400 mg containing 40% oolong tea for 6 weeks, had a positive effect on BM reduction. While these effects have only been observed in other studies in obese people, these results might be attributable to the effect of catechins present in the tea, which cause an increment in lipid catabolism [86, 87]. Another effect that may contribute is the inhibition of gastric and pancreatic lipases, the modulation of appetite, the stimulation of thermogenesis by the inhibition of catechol-O-methyl-transferase, and the suppression of fatty acid synthesis [88].

Caloric restriction and branched-chain amino acid (BCAA) supplementation were found to have a positive effect on BM reduction, with a high-branched-chain amino acid diet producing the highest losses in body fat. The mechanism by which BCAA improves weight reduction alongside caloric restriction is not fully understood, however, it has been theorised that N-enriched hypocaloric diets further increase growth hormone (GH) release, which consequently mobilises fat storage for energy production [89, 90].

(a)

		Risk of bias						
		D1	D2	D3	D4	D5	D6	Overall
Study	Bagheri, R., et al., 2021.							
	Sung, Y., et al., 2018							
	Yavuz, H., et al., 2014.							
	Walker, S., et al., 1998.							
	Koçak, S., et al., 2003.							
	Oopik, V, et al., 2002.							
	Tatlici, A., et al., 2021a.							
	Mourier, A., et al, 1996.							
	Armisan, R., et al., 2011.							
	Shawary, A., 2013.							
	Negaresh, R., et al., 2018.							
	B. Tartibian, B. Rezaei, 2021.							
	Zembrom, et al., 2020.							
	Jang, T. et al., 2022.							
	Berkan, C., el at., 2013.							
	Bajerska, J., et al., 2010.							
	Sung, J., et al., 2022.							
	Abbasalipour, M., et al., 2012.							
	Ulas, H., et al., 2012.							
	Tatlici, A., et al., 2021b.							
	Mohamed, E., & Tammam, A., 2020.							
	Jafari, R., et al., 2024.							
	Zahabi, G., et al., 2024.							

D1: Bias arising from the randomization process

D2: Bias arising from period and carryover effects

D3: Bias due to deviations from intended intervention

D4: Bias due to missing outcome data

D5: Bias in measurement of the outcome

D6: Bias in selection of the reported result

Judgement

High

Unclear

Low

Not applicable

Fig. 3 Assessment of bias of the randomized studies according to RoB 2 tool – (a) traffic light plot and (b) summary plot

(b)

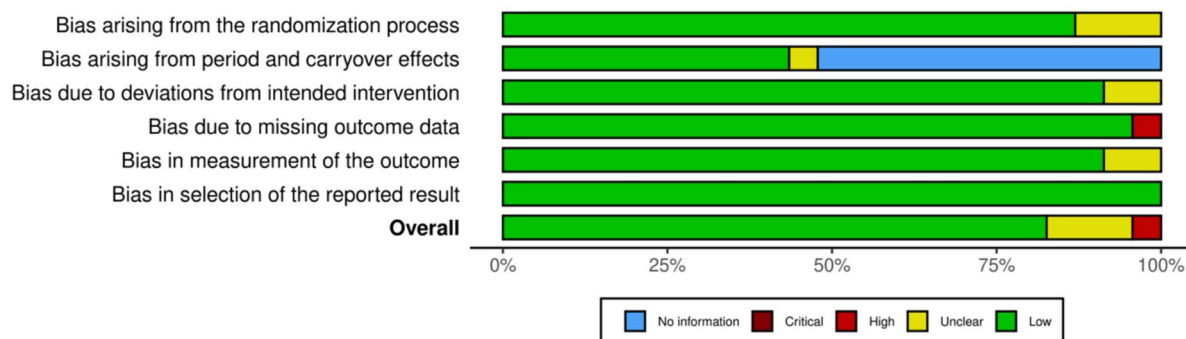
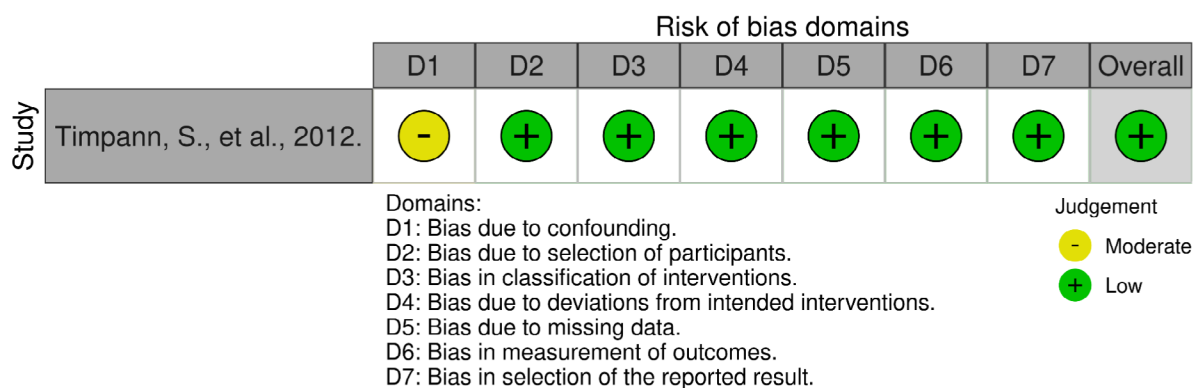


Fig. 3 (continued)

(a)



(b)

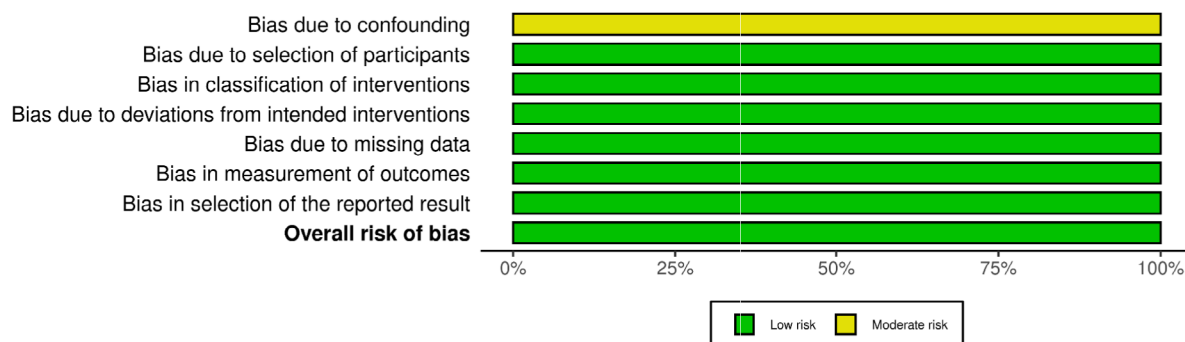


Fig. 4 Assessment of bias of the non-randomised studies according to the ROBINS-I tool – (a) traffic light plot and (b) summary plot

Muscle Damage and Exercise Performance

Sodium citrate supplementation had no effect on exercise performance, however, rapid body mass loss resulted in a significant decrease in mean power. Considering that rapid body mass loss in the range of 5–8% may be accompanied

by a significant (36–54%) decrease in muscle glycogen concentration, a reduction in muscle glycogen could have contributed to the decline in performance [91, 92]. While induced metabolic alkalosis would, in theory, be very effective with respect to performance during activities causing extensive perturbations in acid–base balance by reducing the

rate of developing muscular fatigue by delaying the decrease in intracellular pH and enhancing muscle energy supply through glycolysis [93], the metabolic alkalosis induced through sodium citrate ingestion did not improve UBISP in the experimental group, which may suggest that the degree of alkalosis was insufficient for enhancing glycolytic ATP production. This could also mean that the disturbance in acid–base balance is not the cause for the decreased performance in UBISP; it may appear that, if sufficient carbohydrate consumption is maintained during rapid body mass loss, blood buffering capacity is not a factor in anaerobic performance. Another point to consider is that the buffering capacity of sodium citrate might be sex-dependent like with sodium bicarbonate, showing greater benefits in men [94]. A few theories have been proposed to explain this: 1) type II muscle fibers mainly rely on glycolysis and females have smaller type II fibers than men [95, 96]; 2) males have a greater glycolytic capacity [97, 98]; and 3) the pH drops to a lesser extent in females than in males during the same type of exercise [97]. Sodium citrate supplementation is also able to induce a significant PV increase. An acute increase in intravascular volume after consuming alkalizing substances has been shown to improve performance in sprinters through better muscle perfusion [99]. While this was not the case in this study, the possibility that exercise performance in athletes could be improved if a sufficient degree of alkalosis is achieved along with an increase in PV makes further research worthwhile.

CrPic supplementation did not improve exercise performance in wrestlers. It was believed that improvements in the cellular uptake and storage of glucose could have a positive effect on metabolic performance through glycogen utilisation [100]. Nonetheless, previous studies have also not reported changes in aerobic or anaerobic performance as a result of enhanced insulin activity and no correlation was able to be made between the increased BM with strength [69, 72]. BCAA supplementation had no effect on CK, CKMB, or LDH levels. These results are inconsistent with previous studies where CK and LDH levels decreased after a dose of 10 gr [101–103]. These inconsistencies might be explained by the type of population studied which were long-distance runners versus trained wrestlers. Other factors that have been shown to affect these enzyme levels include oestrogen, which has a protective effect on the muscle cell membrane that might diminish their increase in blood serum [104]. Citrulline malate supplementation had no effects neither on grip strength nor on strength or agility tests. Citrulline malate supplementation improves ATP production by enhancing ammonia, arginine, and lactic acid buffering mechanisms, which might be useful in reducing fatigue and increasing endurance; however, a longer supplementation period might be needed to produce the desired effects [105, 106].

The study evaluating the effects of HMB-FA supplementation reported significantly lower LDH and CK levels, as well as an increased index of perceived recovery status after the tests; however, the latter two findings are only mentioned, and no data is shown to support these claims. While HMB-FA has been shown to increase intramuscular anabolic signalling, stimulate muscle protein synthesis, and attenuate muscle protein breakdown in humans [107], the effects on markers of muscle damage and perceived recovery following resistance exercise have yielded mixed results in the past [108–111]. Last but not least, the article raises high bias concerns regarding the poor attention to detail in which the article was described, as well as when presenting its results, as previously detailed in Sect. 3.4.

Three studies reported similar effects of CM supplementation on exercise performance. The first one reported enhanced short-term high-intensity exercise performance. Phosphocreatine (PCr) concentrations are higher in fast-twitch muscle fibres compared to slow-twitch muscle fibres, which explains why creatine uptake is helpful in these types of exercises [112]. This effect could also be explained by: 1) an increased intramuscular PCr storage, which increases the amount of exercise to be done before it is depleted and decreases the amount of anaerobic glycolysis required; 2) the suppression of pH reduction in skeletal muscle cells by decreasing the amount of ATP resynthesis made through anaerobic glycolysis [113, 114]. The second study reported similar results, where the muscle endurance test was designed to simulate wrestling conditions by implementing no recovery periods between maximal contractions, but a lower level of intensity between each one. As such, the rate of restoration of physical performance capacity was the intended evaluation. Even though all subjects improved their W_{tot} and W_{max} results in the experimental group compared to the two in the control group, this study only involved 5 subjects in total, which increases the bias of finding significant differences between groups. The third one reported a significant increase in CPK levels, muscular endurance, power, and agility. These results suggest once again that CM supplementation increases intramuscular creatine storage and the ability to reconstitute ATP, which helps create energy reserves that contribute to improved physical performance [115].

Although the two studies evaluating beetroot juice supplementation, acute dose of 140 ml before the test, showed improved balance performance at rest and at fatigue, as well as increased average strength in knee and shoulder exercises, it is worth noting that both studies were written by the same author. While it is true that fatigue has a negative effect on proprioception [116, 117], it can cause poor motor coordination by decreasing muscle stimulation and a gradual decrease in muscle force [118], that nitrate-rich beetroot juice may reduce muscle metabolic perturbation by expanding the

antioxidant pool, delay the depletion of ATP reserves, and facilitate muscle glucose and creatine uptake [119–121], there are a couple of limitations with these studies: 1) the first study only induced fatigue in the thigh muscles, but not on other muscles such as hip or leg muscles that also contribute to balance, and 2) both had a small population size.

Iron supplementation, 10 mg/day for 7 days, helped prevent a reduction in endurance capacity during the recovery phase of the rapid weight loss control program. Previous studies have reported that iron deficiency is related to inflammatory and oxidative processes [122, 123], and it also promotes an increased rate of lactate production in muscle [124, 125], which could be prevented with iron supplementation.

Arginine supplementation, ranging from 1.2 to 2 g/kg with different durations, reduces the O_2 cost of moderate-intensity cycle exercise, and the VO_2 slow component amplitude, and increases the time to task failure in severe intensity exercise. L-arginine is a main element for the synthesis of NO, NO synthase converts L-arginine into NO and L-citrulline in the presence of some cofactor [126]. NO is a potent vasodilator which acts by increasing cyclic guanosine monophosphate (cGMP), which in turn causes the relaxation of smooth muscle that consequently increases perfusion causing better performance and post-exercise recovery [126, 127]. While caffeine supplementation was able to reduce the PWPT time in this article, other articles show contradictory evidence for supplementation with caffeine, with not only neutral but also negative effects. On the one hand, caffeine has an ergogenic effect, in which there is a calcium-induced calcium release by acting on the ryanodine receptor that enhances calcium signalling, as well as the extracellular secretion of proteins such as myokines [128]. On the other hand, the ergogenic effects of caffeine can be variable between individuals because of a polymorphism in the CYP1A2 gene that codifies a protein that metabolises caffeine into paraxanthine and methylxanthines, which might prove detrimental for some individuals [129, 130].

Hormonal and Metabolic Markers

Even though the addition of sodium citrate to a high carbohydrate diet increased blood pH, blood buffering capacity, and PV during the 16 h recovery period after rapid body mass loss, it did not affect exercise performance. CrPic supplementation did not have any effect on fasting serum blood glucose and insulin concentrations. It was believed that improvements in the cellular uptake and storage of glucose through insulin potentiation or increased insulin sensitivity could improve glycogen mobilisation and breakdown, as well as reduce the concentrations of fasting insulin and glucose in the blood [100]. This absence of change might be explained because of the heightened insulin sensitivity in highly trained groups of people such as professional

wrestlers [131]. With CM supplementation, plasma ammonia concentrations did not significantly differ between groups, although they did decrease significantly after exercise, a known effect after short-term high-intensity exercise [132]. Lactate and glucose levels did not vary between groups, which suggests that the mobilisation of carbohydrates is not dependent on this supplement during the recovery period of rapid body mass loss. There was no effect of the exercise test on plasma urea concentrations, even though it increased significantly after rapid body mass loss, which reflects an increased rate of protein degradation [133]. On the other hand, the whole-body creatine retention levels were not found to be as expected in the experimental group. It has been reported that a single 5 g dose of CM considerably raises the concentration of creatine in plasma for at least 2 h [134]. Therefore, it can be assumed that the short consumption period of the last dose (around two hours before performing the last test) probably resulted in the last dose not reaching the muscle.

Supplementation with a single dose of arginine increased total amino acid (TAA) concentrations following exercise in both trials, which could be explained by haemoconcentration, however, pre- and post-exercise TAA concentrations were still significantly higher in the supplementation group. These changes could be helpful with fatigue. The causes of fatigue are complex, and influenced by events occurring in both the periphery and the central nervous system [135]. A theory of fatigue developing from the central nervous system is based on the observation that exercise promotes an increase in plasma-free fatty acids and a decrease in large neutral amino acids such as leucine, methionine, valine, phenylalanine and tyrosine due to uptake by skeletal muscle [136]. As both conditions favour more tryptophan entering the CNS, increased production of brain serotonin might be expected, and this could account for the decreased motor drive and increased sensation of fatigue [137]. A significant increase in plasma concentrations of all these large neutral amino acids was observed, except for valine (probably because of the small study population), which might delay the effects of central fatigue. It has also been reported that the plasma BCAA increase produced by arginine administration may attenuate fatigue because BCAAs and free tryptophan compete for transport through the blood–brain barrier [138]. Citrulline supplementation was not able to reduce hypoxanthine or HGPRT levels. Hypoxanthine is derived from the degradation of purines and can indicate metabolic stress in the muscles; while it is not frequently used, it can directly correlate with the amount of ATP consumed inside the cell, making it a theoretically good marker of muscle fatigue [139]. On the other hand, HGPRT levels are also not a commonly used biomarker for muscle damage, but have been used in muscle dystrophies research [140]. Similar to the results obtained in exercise performance, a high but acute

dose might not be sufficient to bring down their levels, which opens the possibility of different doses and longer durations of supplementation.

Spirulina supplementation caused a significant decrease in MST concentrations that can mediate FM reduction. MST inhibition results in adipose tissue loss in high-fat diet-induced mice [141, 142]. Also, the catabolic state decreases with lower levels of FST and increases with higher MST concentrations, resulting in a diminished FST:MST ratio (which is present in the study) [143]. The lower levels of liver enzymes might be caused by the presence of beta carotene, superoxide dismutase, and phyco-cyanin, all of which reduce cell damage, induce the regeneration of damaged hepatocytes, and also reduce oxidative stress and inflammation [144, 145].

Thyme tea three times a day (with a loading dose of 150 cm³ for 35 days), might help improve exercise performance by increasing antioxidant capacity and increasing the resistance to oxidative stress [146, 147]. The study investigating the effects of carbohydrate, BCAA, and arginine supplementation reported significantly higher concentrations of glucose and insulin due to the impaired insulin-dependent glucose disposal and glycogen synthesis in skeletal muscle caused by the increase in the inhibitory insulin receptor substrate-1 phosphorylation and decreasing PI3K activity [148]. Iron supplementation significantly decreased lactate levels, suggesting that it increased the efficacy of muscle contractions by improving the maximum oxygen uptake [56]. Supplementation might prevent iron deficiency, which causes the iron-containing enzymes in skeletal muscle and liver to be altered and promote an increased rate of lactate production [124, 125].

Other Outcomes

The study that investigated thyme tea supplementation reported a significant decrease in malondialdehyde (MDA) levels. MDA is one of the indicators of oxidative stress and one of the main products of lipid peroxidation [147]. As such, thyme tea might be able to reduce oxidative stress in the muscles and improve exercise performance. While IHE significantly increased NO, H₂O₂, tissue regeneration mediators, and multiple haematological parameters, it appears as if hypoxic exposure is the key factor for these changes. When compared to the other groups, arginine alone was not sufficient to elicit significant results in any of the measured outcomes. Nonetheless, the combination of IHE and arginine supplementation was superior to IHE alone in most cases [40]. Whey protein supplementation resulted in significantly higher total protein and albumin levels. Whey protein has one of the highest amounts of BCAA, which promotes the signalling pathways of muscle

protein synthesis and also serves as a donor of nitrogen to alanine and glutamine during protein modulation [149].

Limitations and Future Perspectives

One limitation of this review was the heterogeneity between studies since some involved either multiple sports or underage participants. In addition, the type of supplementation, but also in dose, frequency, or duration of administration, as well as in the measured outcomes, considerably differed. It is also worth mentioning that some authors appear on multiple studies, for example, Tlatici, A. was the main author for both BRJ articles [51, 58]; Ööpik, V. and Timpman, S. were involved in two CM articles as well as in the sodium citrate article [37, 50]; Jun-Young Sung was the main author for both Spatone® and iron supplementation articles [55, 56]; and Ulas, H. was the first author for two arginine articles [52, 53].

As such, future research in this area can be improved: 1) studies should ensure they are sufficiently powered to detect statistically significant group effects by performing a priori sample size calculations; 2) more studies focusing on wrestling are needed. Each type of sport demands specific abilities, which makes the measurement of exercise performance outcomes uneven since the results are usually mixed between participants; 3) underage participants should not be included in the same studies as adults since the metabolism and excretion of certain substances, as well as their performance outcomes, are more likely to differ, providing confusing results.

Conclusions

Based on the findings from the current systematic review, there is some evidence that:

- I) Sodium citrate supplementation has a positive effect on BM re-gain [37].
- II) Supplementation with CrPic has no significant effects in BC, MDEP, HMM or OO [38].
- III) CM supplementation generally improved exercise performance but had mixed outcomes regarding weight gain [29, 39, 43, 50].
- IV) Arginine supplementation significantly increased NO and HGF levels, but had mixed outcomes regarding exercise performance [40, 52–54].
- V) BCAA supplementation showed mixed results regarding glucose and insulin levels, but had significant effects on reducing body and fat mass; lastly, it generally had no significant effect on exercise performance [45, 57].
- VI) Beetroot juice generally improves exercise performance [51, 58].

- VII) HMB-FA supplementation requires further investigation since the results of the included article possess a high risk of bias [41].
- VIII) Spirulina supplementation helped decrease BM, BFP, FM and SMM [42].
- IX) Supplementation with iron had contradictory outcomes: one study reported that VO_2max increased while the other reported it decreased; interestingly, similar to sodium bicarbonate supplementation, it appears iron supplementation might be sex dependent [55, 56].
- X) Thyme tea supplementation exhibited a meaningful increase in TAC and lower MDA levels [46].
- XI) Green and oolong tea extracts were useful for body mass loss [47].
- XII) Whey protein supplementation had generally no significant effect on exercise performance [48].
- XIII) Supplementation with caffeine enhanced exercise performance [49].

Overall, this review might be useful for the creation of safer, evidence-based weight-cutting protocols compared to the standard practices used today, however, more studies are needed to carefully determine whether each type of supplementation is helpful for modifying body composition, physiological status, or exercise performance in wrestlers.

Key References

- Jafari RA, Hosseini S, Rashidlamir A, Nobari H. Evaluating the Impact of Active and Passive Recovery Strategies and Citrulline-Malate Supplementation in Wrestling: Do the Results Add Up? *Acta kinesiologicala*. 2024 08/09;18.

This study examined the effects of active versus passive recovery and citrulline malate supplementation on performance and biomarkers in trained wrestlers during a simulated tournament. While no overall significant effects were found, differences in HGPRT levels, agility, and perceived exertion suggest that recovery strategies may influence specific aspects of performance.

- Bagheri R, Negaresh R, Motevalli MS, Wong A, Ashtary-Larky D, Kargarfard M, et al. Spirulina supplementation during gradual weight loss in competitive wrestlers. *Br J Nutr*. 2022 Jan 28;127(2):248–56.

This study highlights that spirulina supplementation during gradual weight loss enhances fat loss, reduces myostatin and liver enzyme levels, and helps maintain IGF-1 and follistatin concentrations in competitive

wrestlers. These findings suggest that spirulina may be beneficial for optimizing body composition and metabolic markers during weight reduction.

- Zahabi G, García Ramos A, Ilic V, Nedeljkovic A, Štajer V, Žugaj N, et al. Effects of Short-Term Creatine Monohydrate Supplementation Combined with Strength Training on the Physical Fitness Characteristics and Muscle Hypertrophy in Junior Women Wrestlers. *Journal of Health and Allied Sciences NU*. 2024 07/29.

This study demonstrates that short-term creatine supplementation, combined with strength training, significantly enhances muscle hypertrophy and physical fitness in junior female wrestlers. The findings suggest that creatine could be a valuable addition to strength training programs for improving athletic performance in this population.

Abbreviations AA: Amino acid; AE: Average extension; AER: Average external rotation; AF: Average flexion; AIR: Average internal rotation; ALT: Alanine aminotransferase; AP: Average power; APSI: Anterior-posterior stability index; AST: Aspartate aminotransferase; AT: Adipose Tissue; BAT: Burpee agility test; BC: Body composition; BCAA : Branched-chain amino acids; BDNF: Brain derived neurotrophic factor; BE: Base excess; BF: Body fat; BFP: Body fat percentage; BLC: Back-leg-chest; BM: Body mass; BMI: Body mass index; BRJ: Beetroot juice; BW: Body weight; CH: Carbohydrate; CK: Creatine kinase; CKMB: Creatine kinase- MB; CM: Creatine monohydrate; CPK: Creatine phosphokinase; CRP: C-reactive protein; EPO: Eritropoietin; Fe: Iron; FFM: Fat free mass; FI: Fatigue index; FlexAvg: Average flexion; FlexP: Peak flexion; FM: Fat mass; FST: Follistatin; GLC: Glucose; GTE: Green tea extract; H2O2: Hydrogen peroxide; Hb: Haemoglobin; hBCAA: Hypocaloric BCAA group; HCO_3^- : Bicarbonate; HDL: High density lipoprotein; HGF: Hepatocyte growth factor; HGPRT: Hypoxanthine-guanine phosphoribosyltransferase test; HMB-FA: β -Hydroxy- β -methylbutyrate; HMM: Hormonal and metabolic markers; HR: Heart rate; HTC: Haematocrit; IAT: Illinois agility test; IGF-1: Insulin-like growth factor 1; IL: Interleukin; JSWPT: Jafari's simulated wrestling performance test; LBM: Lean body mass; LDH: Lactate dehydrogenase; LDL: Low density lipoprotein; MEB: Mean energy balance; MCH: Mean corpuscular haemoglobin; MCHC: Mean corpuscular haemoglobin concentration; MCV: Mean corpuscular volume; MDA: Malondialdehyde; MDEP: Muscle damage and exercise performance; MLSI: Medial-lateral stability index; MP: Mean power; MST: Myostatin; MT: Muscle tissue; MVC: Maximal isometric volume contraction; N/S: Not specified; NEFA: Non-esterified fatty acid; NO: Nitric oxide; OO: Other outcomes; OSI: Overall stability index; OTE: Oolong tea extract; PDGF: Platelet derived growth factor; Peak AnP: Peak anaerobic power; PE: Peak extension; PER: Peak external rotation; PF: Peak flexion; PIR: Peak internal rotation; PP: Peak power; PRS: Perceived recovery status; PWPT: Pittsburgh wrestling performance test; RBC: Red blood cells; RDW: Red cell distribution width; REL Anc: Relative anaerobic capacity; REL VOMax: Relative peak aerobic power; RET: Reticulocytes; RHR: Reserve heart rate; RM: Repetition maximum; RPE: Rating of perceived exertion; RPF: Rating of perceived fatigue; RSH: Total sulphhydryl group; SAT: Subcutaneous adipose tissue; SMM: Skeletal muscle mass; T3: Triiodothyronine; TAA: Total aminoacid concentration; TAC: Total antioxidant capacity; TC: Total cholesterol; TG: Triglycerides; TIBC: Total iron binding capacity; TNF-alpha: Tumor necrosis factor alpha; UBISP: Upper body intermittent sprint performance test; USG: Urine specific gravity; VAT: Visceral adipose tissue;

VEGF: Vascular endothelial growth factor; VJT: Vertical jump test; WCV: White cell volume; Wmax: Maximum work; Wtot: Total work

Author Contributions Conceptualisation: RZ and AAR. Methodology: AAR, RZ, APM, and MCD. Investigation, data curation, formal analysis, writing – original draft: APM and MCD. Writing—Review & Editing: RZ, AAR, and SCF. Project administration and supervision: AAR.

Funding Not applicable.

Data Availability No datasets were generated or analysed during the current study.

Code Availability Not applicable.

Declarations

Ethical Approval Not applicable.

Consent to Participate Not applicable.

Consent for Publication Not applicable.

Conflicts of interest All authors declare that they have no conflict of interest relevant to the content of this review. SCF serves as a scientific advisor for Bear Balanced, a company that sells creatine products. SCF has received creatine donations for scientific studies. SCF is a sport nutrition advisor for the International Society of Sports Nutrition (ISSN).

Human and Animal Rights and Informed Consent This article does not contain any studies with human or animal subjects performed by any of the authors

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References

- Molnár S, Mensch K, Bacskai K, Körösi É, Sántha Á, Gáspár K. Wrestling. Epidemiology of injuries in sports. Berlin, Heidelberg: Springer; 2022. p. 155–162.
- Department SR. Number of participants in wrestling in the United States from 2006 to 2017. 2020 Available from: <https://www.statista.com/statistics/191989/participants-in-wrestling-in-the-us-since-2006/>. Accessed 22 Jan 2024.
- Buford TW, Rossi SJ, Smith DB, O'Brien MS, Pickering C. The effect of a competitive wrestling season on body weight, hydration, and muscular performance in collegiate wrestlers. *J Strength Cond Res*. 2006;20(3):689–92.
- Oppliger RA, Case HS, Horswill CA, Landry GL, Shelter AC. American College of Sports Medicine position stand. Weight loss in wrestlers. *Med Sci Sports Exerc*. 1996;28(10):135–8.
- Marttinen RH, Judelson DA, Wiersma LD, Coburn JW. Effects of self-selected mass loss on performance and mood in collegiate wrestlers. *J Strength Cond Res*. 2011;25(4):1010–5.
- Allen TW, DO MPH. Prevention of Heat Exhaustion: Focus on Wrestling. *Int J Wrestling Sci*. 2020;10:54–6.
- Finn KJ, Dolgener FA, Williams RB. Effects of carbohydrate refeeding on physiological responses and psychological and physical performance following acute weight reduction in collegiate wrestlers. *J Strength Cond Res*. 2004;18(2):328–33.
- Cengiz A. Effects of self-selected dehydration and meaningful rehydration on anaerobic power and heart rate recovery of elite wrestlers. *J Phys Ther Sci*. 2015;27(5):1441–4.
- Karnincic H, Tocilj Z, Uljevic O, Erceg M. Lactate profile during greco-roman wrestling matchx. *J Sports Sci Med*. 2009;8(CSSI3):17–9.
- Del Coso J, Hamouti N, Aguado-Jimenez R, Mora-Rodriguez R. Restoration of blood pH between repeated bouts of high-intensity exercise: effects of various active-recovery protocols. *Eur J Appl Physiol*. 2010;108(3):523–32.
- Cardozo LF, Pedruzzi LM, Stenvinkel P, Stockler-Pinto MB, Daleprane JB, Leite M Jr, et al. Nutritional strategies to modulate inflammation and oxidative stress pathways via activation of the master antioxidant switch Nrf2. *Biochimie*. 2013;95(8):1525–33.
- Ciechanowski K. To dialyze or to alkalyze? (Dializować czy alkalizować?). *Forum Nefrol*. 2012;5:347–50.
- Heisler N. Buffering and H⁺ ion dynamics in muscle tissues. *Respir Physiol Neurobiol*. 2004;144(2–3):161–72.
- Juel C. Regulation of pH in human skeletal muscle: adaptations to physical activity. *Acta Physiol (Oxf)*. 2008;193(1):17–24.
- Lancha Junior AH, Painelli Vde S, Saunders B, Artioli GG. Nutritional Strategies to Modulate Intracellular and Extracellular Buffering Capacity During High-Intensity Exercise. *Sports Med*. 2015;45(Suppl 1):S71–81.
- Carr BM, Webster MJ, Boyd JC, Hudson GM, Scheett TP. Sodium bicarbonate supplementation improves hypertrophy-type resistance exercise performance. *Eur J Appl Physiol*. 2013;113(3):743–52.
- Hollidge-Horvat MG, Parolin ML, Wong D, Jones NL, Heigenhauser GJ. Effect of induced metabolic alkalosis on human skeletal muscle metabolism during exercise. *Am J Physiol Endocrinol Metab*. 2000;278(2):E316–29.
- Sostaric SM, Skinner SL, Brown MJ, Sangkabutra T, Medved I, Medley T, et al. Alkalosis increases muscle K⁺ release, but lowers plasma [K⁺] and delays fatigue during dynamic forearm exercise. *J Physiol*. 2006;570(Pt 1):185–205.
- McNaughton LR, Gough L, Deb S, Bentley D, Sparks SA. Recent Developments in the Use of Sodium Bicarbonate as an Ergogenic Aid. *Curr Sports Med Rep*. 2016;15(4):233–44.
- Morton RW, Sato K, Gallagher MPB, Oikawa SY, McNicholas PD, Fujita S, et al. Muscle androgen receptor content but not systemic hormones is associated with resistance training-induced skeletal muscle hypertrophy in healthy. *Young Men Front Physiol*. 2018;9:1373.
- Ratamess NA, Hoffman JR, Kraemer WJ, Ross RE, Tranchina CP, Rashti SL, et al. Effects of a competitive wrestling season on body composition, endocrine markers, and anaerobic exercise performance in NCAA collegiate wrestlers. *Eur J Appl Physiol*. 2013;113(5):1157–68.
- Barbas I, Fatouros IG, Douroudos II, Chatzinikolaou A, Michailidis Y, Draganidis D, et al. Physiological and performance adaptations of elite Greco-Roman wrestlers during a one-day tournament. *Eur J Appl Physiol*. 2011;111(7):1421–36.

23. Fry AC, Schilling BK, Fleck SJ, Kraemer WJ. Relationships between competitive wrestling success and neuroendocrine responses. *J Strength Cond Res.* 2011;25(1):40–5.
24. Kraemer WJ, Fry AC, Rubin MR, Triplett-McBride T, Gordon SE, Koziris LP, et al. Physiological and performance responses to tournament wrestling. *Med Sci Sports Exerc.* 2001;33(8):1367–78.
25. Jowko E, Gierczuk D, Cieslinski I, Kotowska J. SOD2 gene polymorphism and response of oxidative stress parameters in young wrestlers to a three-month training. *Free Radic Res.* 2017;51(5):506–16.
26. Prieto Martinez A, Coutino Diaz M, Anaya Romero L, Ali Redha A, Zare R, Ventura Hernandez S, et al. Effects of Vaccinium berries (blueberries, cranberries and bilberries) on oxidative stress, inflammation, exercise performance, and recovery - a systematic review. *Food Funct.* 2024;15(2):444–59.
27. Barcal JN, Thomas JT, Hollis BW, Austin KJ, Alexander BM, Larson-Meyer DE. Vitamin D and weight cycling: Impact on injury, illness, and inflammation in collegiate wrestlers. *Nutrients.* 2016;8(12):775.
28. Micheletti A, Rossi R, Rufini S. Zinc status in athletes: relation to diet and exercise. *Sports Med.* 2001;31(8):577–82.
29. Abbasalipour M, Parsay S, Melkumyan K, Minasyan S. Effects of creatine and glutamine supplements in comparison with proper nutrition on performance factors of wrestlers. *Advances in Environmental Biology.* 2012;6(10):2726–30.
30. FDA. FDA 101: Dietary Supplements. 2022 [cited 2024; Available from: <https://www.fda.gov/consumers/consumer-updates/fda-101-dietary-supplements>
31. Blanck HM, Serdula MK, Gillespie C, Galuska DA, Sharpe PA, Conway JM, et al. Use of nonprescription dietary supplements for weight loss is common among Americans. *J Am Diet Assoc.* 2007;107(3):441–7.
32. Sharpe PA, Granner ML, Conway JM, Ainsworth BE, Dobre M. Availability of weight-loss supplements: Results of an audit of retail outlets in a southeastern city. *J Am Diet Assoc.* 2006;106(12):2045–51.
33. Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, Moher D. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *Bmj.* 2021;372:n71.
34. Sterne JA, Savović J, Page MJ, Elbers RG, Blencowe NS, Boutron I, Cates CJ, Cheng HY, Corbett MS, Eldridge SM, Emberson JR. RoB 2: a revised tool for assessing risk of bias in randomised trials. *BMJ.* 2019;66:14898.
35. Sterne J, Hernán MA, Reeves BC, Savović J, Berkman ND, Viswanathan M, Henry D, Altman DG, Ansari MT, Boutron I, Carpenter JR. ROBINS-I: a tool for assessing risk of bias in non-randomized studies of interventions. *British Med J.* 2016;355:i4919.
36. McGuinness LA, Higgins JP. Risk-of-bias VISualization (robvis): An R package and Shiny web app for visualizing risk-of-bias assessments. *Res Synthesis methods.* 2021;12(1):55–61.
37. Timpmann S, Burk A, Medijainen L, Tamm M, Kreegipuu K, Vahi M, et al. Dietary sodium citrate supplementation enhances rehydration and recovery from rapid body mass loss in trained wrestlers. *Appl Physiol Nutr Metab.* 2012;37(6):1028–37.
38. Walker LS, Bemben MG, Bemben DA, Knehans AW. Chromium picolinate effects on body composition and muscular performance in wrestlers. *Med Sci Sports Exerc.* 1998;30(12):1730–7.
39. Mohamed ES, Tammam AH. The Effect of Creatine Supplementation on the CPK Enzyme in Blood Serum and Some Physical Abilities for Wrestlers during the Competition Period. *Amazonia Investiga.* 2020;9(34):76–84.
40. Zembron-Lacny A, Gramacki A, Wawrzyniak-Gramacka E, Tylutka A, Hertmanowska N, Kasperska A, et al. Intermittent hypoxic exposure with high dose of arginine impact on circulating mediators of tissue regeneration. *Nutrients.* 2020;12(7):1933.
41. Tartibian B, Rezaei B. Effect of HMB-FA Supplementation on Muscle damage indices in a Simulated Wrestling Protocols in elite wrestlers. *Sport Physiology.* 2021;13(50):137–62.
42. Bagheri R, Negaresh R, Motevalli MS, Wong A, Ashtary-Larky D, Kargarfard M, et al. Spirulina supplementation during gradual weight loss in competitive wrestlers. *Br J Nutr.* 2022;127(2):248–56.
43. Kocak S, Karli U. Effects of high dose oral creatine supplementation on anaerobic capacity of elite wrestlers. *J Sports Med Phys Fitness.* 2003;43(4):488–92.
44. Abbasalipour M, Parsay S, Melkumyan KV, Minasyan S. Effects of creatine and glutamine supplements in comparison with proper nutrition on performance factors of wrestlers. *Adv Environ Biol.* 2012;6:2726–30.
45. Amirsasan R, Nikookheslat S, Sari-Sarraf V, Kaveh B, Letafatkar A. The effects of two different dosages of BCAA supplementation on A serum indicators of muscle damage in wrestlers. *Int J Wrestl Sci.* 2014;1(2):32–6.
46. Alpay C, Hazar S, Gökdemir K, Atalay Guzel N, Gonenc A, Simsek B. The effects of thyme tea supplement on free radicals formation and antioxidant system of elite wrestlers. *Pakistan J Nutr.* 2013;12:433–40.
47. Bajerska J, Jan J, Tarnowska A, Czlapka-Matyasik M. The Effect of Green and Oolong Tea Extracts Supplementation on Body Composition in Wrestlers. *Pakistan Journal of Nutrition.* 2010;9(7):696–702.
48. Sharawy A. The effects of a pre- and post-exercise whey protein supplement on protein metabolism and muscular strength among elite wrestlers. *Ovidius University Annals, Series Physical Education and Sport/Science, Movement and Health.* 2013;13:5.
49. Negaresh R, Del Coso J, Mokhtarzade M, Lima-Silva AE, Baker JS, Willems MET, et al. Effects of different dosages of caffeine administration on wrestling performance during a simulated tournament. *Eur J Sport Sci.* 2019;19(4):499–507.
50. Oopik V, Paasuke M, Timpmann S, Medijainen L, Ereline J, Smirnova T. Effect of creatine supplementation during rapid body mass reduction on metabolism and isokinetic muscle performance capacity. *Eur J Appl Physiol Occup Physiol.* 1998;78(1):83–92.
51. Tathici A, Lima Y, Yilmaz S, Ekin A, Okut S, Ceviz E. The Effects of Beetroot Juice Supplementation on Balance Performance of Wrestlers. *Pakistan J Med Health Sci.* 2021;15:2234–40.
52. Yavuz H. Effect of pre-exercise single dose arginine ingestion on plasma amino acid profile during exhaustive exercise in elite male wrestlers. *Turkish J Biochem.* 2012;37:139–45.
53. Yavuz HU, Turnagol H, Demirel AH. Pre-exercise arginine supplementation increases time to exhaustion in elite male wrestlers. *Biol Sport.* 2014;31(3):187–91.
54. Jang TR, Wu CL, Chang CM, Hung W, Fang SH, Chang CK. Effects of carbohydrate, branched-chain amino acids, and arginine in recovery period on the subsequent performance in wrestlers. *J Int Soc Sports Nutr.* 2011;22(8):21.
55. Sung J-Y, Oh T-W, Lim S-G. Effects of natural iron supplement intake on EPO, Hp, and iron metabolism during rapid weight loss in university wrestlers. *J Men's Health.* 2022;18:114.
56. Sung JY, Park S, Lim SG, Lee SK, Kang DM, Lee M, et al. Effect of Spatone Supplement on Endurance Capacity and Inflammatory Cytokines in a Rapid Weight Control Program in University Wrestlers: A Pilot Study. *J Med Food.* 2018;21(8):832–9.
57. Mourier A, Bigard AX, de Kerviler E, Roger B, Legrand H, Guezennec CY. Combined effects of caloric restriction and branched-chain amino acid supplementation on body composition and exercise performance in elite wrestlers. *Int J Sports Med.* 1997;18(1):47–55.
58. Tathici A. The effects of acute beetroot juice supplementation on lower and upper body isokinetic strength of the wrestlers. *J Men s Health.* 2021;17:249–54.

59. Zahabi G, García Ramos A, Ilic V, Nedeljkovic A, Štajer V, Žugaj N, et al. Effects of Short-Term Creatine Monohydrate Supplementation Combined with Strength Training on the Physical Fitness Characteristics and Muscle Hypertrophy in Junior Women Wrestlers. *J Health Allied Sci NU*. 2024;14:34–43.
60. Jafari RA, Hosseini S, Rashidlamir A, Nobari H. Evaluating the Impact of Active and Passive Recovery Strategies and Citrulline-Malate Supplementation in Wrestling: Do the Results Add Up? *Acta kinesiologica*. 2024;18(2):58–69.
61. Oopik V, Saaremets I, Timpmann S, Medijainen L, Karelson K. Effects of acute ingestion of sodium citrate on metabolism and 5-km running performance: a field study. *Can J Appl Physiol*. 2004;29(6):691–703.
62. Oopik V, Timpmann S, Hackney AC, Kadak K, Medijainen L, Karelson K. Ingestion of sodium citrate suppresses aldosterone level in blood at rest and during exercise. *Appl Physiol Nutr Metab*. 2010;35(3):278–85.
63. Oopik V, Saaremets I, Medijainen L, Karelson K, Janson T, Timpmann S. Effects of sodium citrate ingestion before exercise on endurance performance in well trained college runners. *Br J Sports Med*. 2003;37(6):485–9.
64. Schabert EJ, Wilson G, Noakes TD. Dose-related elevations in venous pH with citrate ingestion do not alter 40-km cycling time-trial performance. *Eur J Appl Physiol*. 2000;83(4–5):320–7.
65. Murray RK, Mayers PK, Granner DK, Rodwell VW. Harper's Biochemistry (22nd). Chemical constituents of blood and body fluids. Appleton & Lange: Connecticut; 1990. p. 577.
66. Trent LK, Thieding-Cancel D. Effects of chromium picolinate on body composition. *J Sports Med Phys Fitness*. 1995;35(4):273–80.
67. Wagner JC. Use of chromium and cobamide by athletes. *Clin Pharm*. 1989;8:832–4.
68. Evans GW. The effect of chromium picolinate on insulin controlled parameters in humans. *Int J Bios Med Re*. 1989;11:163–80.
69. Hasten DL, Rome EP, Franks BD, Hegsted M. effects of chromium picolinate on beginning weight training students. *Int J Sport Nutr Exercise Metab*. 1992;2(4):343–50.
70. Hasten DL, Rome EP, Franks BD, Hegsted M. Effects of chromium picolinate on beginning weight training students. *Int J Sport Nutr*. 1992;2(4):343–50.
71. Clancy SP, Clarkson PM, DeCheke ME, Nosaka K, Freedson PS, Cunningham JJ, et al. Effects of chromium picolinate supplementation on body composition, strength, and urinary chromium loss in football players. *Int J Sport Nutr*. 1994;4(2):142–53.
72. Hallmark MA, Reynolds TH, DeSouza CA, Dotson CO, Anderson RA, Rogers MA. Effects of chromium and resistive training on muscle strength and body composition. *Med Sci Sports Exerc*. 1996;28(1):139–44.
73. Balsom PD, Soderlund K, Ekblom B. Creatine in humans with special reference to creatine supplementation. *Sports Med*. 1994;18(4):268–80.
74. Balsom PD, Harridge SD, Soderlund K, Sjodin B, Ekblom B. Creatine supplementation per se does not enhance endurance exercise performance. *Acta Physiol Scand*. 1993;149(4):521–3.
75. Lemon PWR, Boska MD, Bredle DL, Rogers MD, Ziegenfuss TN, Newcomer BR. Effect of Oral Creatine Supplementation on Energetics During Repeated Maximal Muscle Contraction. *Med Sci Sports Exercise*. 1995;27:S204.
76. Vandenberghe K, Gillis N, Van Leemputte M, Van Hecke P, Vanstapel F, Hespel P. Caffeine counteracts the ergogenic action of muscle creatine loading. *J Appl Physiol* (1985). 1996;80(2):452–7.
77. Green AL, Hultman E, Macdonald IA, Sewell DA, Greenhaff PL. Carbohydrate ingestion augments skeletal muscle creatine accumulation during creatine supplementation in humans. *Am J Physiol*. 1996;271(5 Pt 1):E821–6.
78. Jacobs I, Bleue S, Goodman J. Creatine ingestion increases anaerobic capacity and maximum accumulated oxygen deficit. *Can J Appl Physiol*. 1997;22(3):231–43.
79. Mujika I, Chatard JC, Lacoste L, Barale F, Geyssant A. Creatine supplementation does not improve sprint performance in competitive swimmers. *Med Sci Sports Exerc*. 1996;28(11):1435–41.
80. Bagheri R, Rashidlamir A, Ashtary-Larky D, Wong A, Alipour M, Motevalli MS, et al. Does green tea extract enhance the anti-inflammatory effects of exercise on fat loss? *Br J Clin Pharmacol*. 2020;86(4):753–62.
81. Galic S, Loh K, Murray-Segal L, Steinberg GR, Andrews ZB, Kemp BE. AMPK signaling to acetyl-CoA carboxylase is required for fasting- and cold-induced appetite but not thermogenesis. *Elife*. 2018;13:7.
82. Heo MG, Choung SY. Anti-obesity effects of *Spirulina maxima* in high fat diet induced obese rats via the activation of AMPK pathway and SIRT1. *Food Funct*. 2018;9(9):4906–15.
83. Iwabu M, Yamauchi T, Okada-Iwabu M, Sato K, Nakagawa T, Funata M, et al. Adiponectin and AdipoR1 regulate PGC-1 α and mitochondria by Ca(2+) and AMPK/SIRT1. *Nature*. 2010;464(7293):1313–9.
84. Knutti D, Kralli A. PGC-1, a versatile coactivator. *Trends Endocrinol Metab*. 2001;12(8):360–5.
85. Kwon JY, Lee KW, Kim JE, Jung SK, Kang NJ, Hwang MK, et al. Delphinidin suppresses ultraviolet B-induced cyclooxygenases-2 expression through inhibition of MAPKK4 and PI-3 kinase. *Carcinogenesis*. 2009;30(11):1932–40.
86. Ito Y, Ichikawa T, Morohoshi Y, Nakamura T, Saegusa Y, Ishihara K. Effect of tea catechins on body fat accumulation in rats fed a normal diet. *Biomed Res*. 2008;29(1):27–32.
87. Murase T, Nagasawa A, Suzuki J, Hase T, Tokimitsu I. Beneficial effects of tea catechins on diet-induced obesity: stimulation of lipid catabolism in the liver. *Int J Obes Relat Metab Disord*. 2002;26(11):1459–64.
88. Shaik Mohamed Sayed UF, Moshawih S, Goh HP, Kifli N, Gupta G, Singh SK, et al. Natural products as novel anti-obesity agents: insights into mechanisms of action and potential for therapeutic management. *Front Pharmacol*. 2023;14:1182937.
89. Jacobson BH. Effect of Amino Acids on Growth Hormone Release. *Phys Sportsmed*. 1990;18(1):63–70.
90. Knopf RF, Conn JW, Fajans SS, Floyd JC, Guntzsch EM, Rull JA. Plasma Growth Hormone Response to Intravenous Administration of Amino Acids. *J Clin Endocrinol Metab*. 1965;25:1140–4.
91. Burge CM, Carey MF, Payne WR. Rowing performance, fluid balance, and metabolic function following dehydration and rehydration. *Med Sci Sports Exerc*. 1993;25(12):1358–64.
92. Tarnopolsky MA, Cipriano N, Woodcroft C, Pulkkinen WJ, Robinson DC, Henderson JM, et al. Effects of rapid weight loss and wrestling on muscle glycogen concentration. *Clin J Sport Med*. 1996;6(2):78–84.
93. Requena B, Zabala M, Padial P, Feriche B. Sodium bicarbonate and sodium citrate: ergogenic aids? *J Strength Cond Res*. 2005;19(1):213–24.
94. Durkalec-Michalski K, Zawieja EE, Zawieja BE, Michalowska P, Podgorski T. The gender dependent influence of sodium bicarbonate supplementation on anaerobic power and specific performance in female and male wrestlers. *Sci Rep*. 2020;10(1):1878.
95. Porter MM, Stuart S, Boij M, Lexell J. Capillary supply of the tibialis anterior muscle in young, healthy, and moderately active men and women. *J Appl Physiol* (1985). 2002;92(4):1451–7.
96. Simoneau JA, Bouchard C. Human variation in skeletal muscle fiber-type proportion and enzyme activities. *Am J Physiol*. 1989;257(4 Pt 1):E567–72.
97. Russ DW, Lanza IR, Rothman D, Kent-Braun JA. Sex differences in glycolysis during brief, intense isometric contractions. *Muscle Nerve*. 2005;32(5):647–55.

98. Tarnopolsky MA. Gender differences in substrate metabolism during endurance exercise. *Can J Appl Physiol*. 2000;25(4):312–27.
99. Kozak-Collins K, Burke ER, Schoene RB. Sodium bicarbonate ingestion does not improve performance in women cyclists. *Med Sci Sports Exerc*. 1994;26(12):1510–5.
100. Anderson RA, Polansky MM, Bryden NA, Canary JJ. Supplemental-chromium effects on glucose, insulin, glucagon, and urinary chromium losses in subjects consuming controlled low-chromium diets. *Am J Clin Nutr*. 1991;54(5):909–16.
101. Coombes JS, McNaughton LR. Effects of branched-chain amino acid supplementation on serum creatine kinase and lactate dehydrogenase after prolonged exercise. *J Sports Med Phys Fitness*. 2000;40(3):240–6.
102. Koba T, Hamada K, Sakurai M, Matsumoto K, Hayase H, Imaizumi K, et al. Branched-chain amino acids supplementation attenuates the accumulation of blood lactate dehydrogenase during distance running. *J Sports Med Phys Fitness*. 2007;47(3):316–22.
103. Koba T, Hamada K, Sakurai M, Matsumoto K, Higuchi T, Zhao M, et al. Effect Of A Branched-chain Amino Acids Supplementation On Muscle Soreness During Intensive Training Program: 229 Board #136 11:00 AM - 12:30 PM. *Medicine & Science in Sports & Exercise*. 2005;37:S43.
104. Chidi-Ogbolu N, Baar K. Effect of estrogen on musculoskeletal performance and injury risk. *Front Physiol*. 2019;9:421933.
105. Perez-Guisado J, Jakeman PM. Citrulline malate enhances athletic anaerobic performance and relieves muscle soreness. *J Strength Cond Res*. 2010;24(5):1215–22.
106. Figueroa A, Wong A, Jaime SJ, Gonzales JU. Influence of L-citrulline and watermelon supplementation on vascular function and exercise performance. *Curr Opin Clin Nutr Metab Care*. 2017;20(1):92–8.
107. Silva VR, Belozo FL, Micheletti TO, Conrado M, Stout JR, Pimentel GD, et al. beta-hydroxy-beta-methylbutyrate free acid supplementation may improve recovery and muscle adaptations after resistance training: a systematic review. *Nutr Res*. 2017;45:1–9.
108. Gentles JA, Phillips SM. Discrepancies in publications related to HMB-FA and ATP supplementation. *Nutr Metab*. 2017;14(1):42.
109. Gonzalez AM, Stout JR, Jajtner AR, Townsend JR, Wells AJ, Beyer KS, et al. Effects of beta-hydroxy-beta-methylbutyrate free acid and cold water immersion on post-exercise markers of muscle damage. *Amino Acids*. 2014;46(6):1501–11.
110. Hyde PN, Kendall KL, LaFountain RA. Interaction of Beta-Hydroxy-Beta-Methylbutyrate Free Acid and Adenosine Triphosphate on Muscle Mass, Strength, and Power in Resistance-Trained Individuals. *The J Strength Conditioning Res*. 2016;30(10):e10–1.
111. Phillips SM, Aragon AA, Arciero PJ, Arent SM, Close GL, Hamilton DL, et al. Changes in body composition and performance with supplemental HMB-FA+ATP. *J Strength Cond Res*. 2017;31(5):e71–2.
112. Demant TW, Rhodes EC. Effects of creatine supplementation on exercise performance. *Sports Med*. 1999;28(1):49–60.
113. Baker JS, McCormick MC, Robergs RA. Interaction among Skeletal Muscle Metabolic Energy Systems during Intense Exercise. *J Nutr Metab*. 2010;2010:905612.
114. Sahlin K. Muscle energetics during explosive activities and potential effects of nutrition and training. *Sports Med*. 2014;44(Suppl 2):S167–73.
115. Cooper R, Naclerio F, Allgrove J, Jimenez A. Creatine supplementation with specific view to exercise/sports performance: an update. *J Int Soc Sports Nutr*. 2012;9(1):33.
116. Harkins KM, Mattacola CG, Uhl TL, Malone TR, McCrory JL. Effects of 2 ankle fatigue models on the duration of postural stability dysfunction. *J Athl Train*. 2005;40(3):191–4.
117. Yaggie JA, McGregor SJ. Effects of isokinetic ankle fatigue on the maintenance of balance and postural limits. *Arch Phys Med Rehabil*. 2002;83(2):224–8.
118. Enoka RM, Duchateau J. Muscle fatigue: what, why and how it influences muscle function. *J Physiol*. 2008;586(1):11–23.
119. Bailey SJ, Fulford J, Vanhatalo A, Winyard PG, Blackwell JR, DiMenna FJ, et al. Dietary nitrate supplementation enhances muscle contractile efficiency during knee-extensor exercise in humans. *J Appl Physiol* (1985). 2010;109(1):135–48.
120. Rigamonti E, Touvier T, Clementi E, Manfredi AA, Brunelli S, Rovere-Querini P. Requirement of inducible nitric oxide synthase for skeletal muscle regeneration after acute damage. *J Immunol*. 2013;190(4):1767–77.
121. Wootton-Beard PC, Ryan L. A beetroot juice shot is a significant and convenient source of bioaccessible antioxidants. *J Funct Foods*. 2011;3(4):329–34.
122. Ozkan I, Ibrahim CH. Dehydration, skeletal muscle damage and inflammation before the competitions among the elite wrestlers. *J Phys Ther Sci*. 2016;28(1):162–8.
123. Peeling P, Blee T, Goodman C, Dawson B, Claydon G, Beilby J, et al. Effect of iron injections on aerobic-exercise performance of iron-depleted female athletes. *Int J Sport Nutr Exerc Metab*. 2007;17(3):221–31.
124. Beard JL, Haas JD, Tufts D, Spielvogel H, Vargas E, Rodriguez C. Iron deficiency anemia and steady-state work performance at high altitude. *J Appl Physiol* (1985). 1988;64(5):1878–84.
125. Davies KJ, Maguire JJ, Brooks GA, Dallman PR, Packer L. Muscle mitochondrial bioenergetics, oxygen supply, and work capacity during dietary iron deficiency and repletion. *Am J Physiol*. 1982;242(6):E418–27.
126. Bailey SJ, Winyard PG, Vanhatalo A, Blackwell JR, DiMenna FJ, Wilkerson DP, et al. Acute L-arginine supplementation reduces the O2 cost of moderate-intensity exercise and enhances high-intensity exercise tolerance. *J Appl Physiol* (1985). 2010;109(5):1394–403.
127. Gruetter CA, Barry BK, McNamara DB, Gruetter DY, Kadowitz PJ, Ignarro L. Relaxation of bovine coronary artery and activation of coronary arterial guanylate cyclase by nitric oxide, nitroprusside and a carcinogenic nitrosoamine. *J Cyclic Nucleotide Res*. 1979;5(3):211–24.
128. Takada S, Fumoto Y, Kinugawa S. Ergogenic effects of caffeine are mediated by myokines. *Front Sports Act Living*. 2022;4:969623.
129. Guest N, Corey P, Vescovi J, El-Sohemy A. Caffeine, CYP1A2 Genotype, and Endurance Performance in Athletes. *Med Sci Sports Exerc*. 2018;50(8):1570–8.
130. Rahimi R. The effect of CYP1A2 genotype on the ergogenic properties of caffeine during resistance exercise: a randomized, double-blind, placebo-controlled, crossover study. *Ir J Med Sci*. 2019;188(1):337–45.
131. Ebeling P, Bourey R, Koranyi L, Tuominen JA, Groop LC, Henriksson J, et al. Mechanism of enhanced insulin sensitivity in athletes. Increased blood flow, muscle glucose transport protein (GLUT-4) concentration, and glycogen synthase activity. *J Clin Invest*. 1993;92(4):1623–31.
132. Bellinger BM, Bold A, Wilson GR, Noakes TD, Myburgh KH. Oral creatine supplementation decreases plasma markers of adenine nucleotide degradation during a 1-h cycle test. *Acta Physiol Scand*. 2000;170(3):217–24.
133. Oopik V, Paasuke M, Sikku T, Timpmann S, Medijainen L, Ereline J, et al. Effect of rapid weight loss on metabolism and isokinetic performance capacity. A case study of two well trained wrestlers. *J Sports Med Phys Fitness*. 1996;36(2):127–31.
134. Oöpik VTS, Medijainen L. The role and application of dietary creatine supplementation in increasing physical performance capacity. *Biol Sport*. 1995;12:197–212.

135. Meeusen R, Watson P, Hasegawa H, Roelands B, Piacentini MF. Central Fatigue. *Sports Med.* 2006;36(10):881–909.
136. Meeusen R, Watson P, Dvorak J. The brain and fatigue: new opportunities for nutritional interventions? *J Sports Sci.* 2006;24(7):773–82.
137. Westermann CM, Dorland L, Wijnberg ID, de Sain-van der Velden MGM, van Breda E, Barneveld A, et al. Amino acid profile during exercise and training in Standardbreds. *Res Vet Sci.* 2011;91(1):144–9.
138. Sugino T, Aoyagi S, Shirai T, Kajimoto Y, Kajimoto O. Effects of Citric Acid and L-Carnitine on Physical Fatigue. *J Clin Biochem Nutr.* 2007;41(3):224–30.
139. Zielinski J, Kusy K. Hypoxanthine: A Universal Metabolic Indicator of Training Status in Competitive Sports. *Exerc Sport Sci Rev.* 2015;43(4):214–21.
140. Neerunjun JS, Allsop J, Dubowitz V. Hypoxanthine-guanine phosphoribosyltransferase activity of blood and muscle in Duchenne dystrophy. *Muscle Nerve.* 1979;2(1):19–23.
141. Deng B, Zhang F, Wen J, Ye S, Wang L, Yang Y, et al. The function of myostatin in the regulation of fat mass in mammals. *Nutr Metab (Lond).* 2017;14:29.
142. Zhang C, McFarlane C, Lokireddy S, Masuda S, Ge X, Gluckman PD, et al. Inhibition of myostatin protects against diet-induced obesity by enhancing fatty acid oxidation and promoting a brown adipose phenotype in mice. *Diabetologia.* 2012;55(1):183–93.
143. Chen MJ, Han DS, Yang JH, Yang YS, Ho HN, Yang WS. Myostatin and its association with abdominal obesity, androgen and follistatin levels in women with polycystic ovary syndrome. *Hum Reprod.* 2012;27(8):2476–83.
144. Kumar S, Kumar M. Spirulina fusiformis: A Food Supplement against Mercury Induced Hepatic Toxicity. *J Health Sci - J HEALTH SCI.* 2005;51:424–30.
145. Mazokopakis EE, Papadomanolaki MG, Foustieris AA, Kotsiris DA, Lampadakis IM, Ganotakis ES. The hepatoprotective and hypolipidemic effects of Spirulina (*Arthrospira platensis*) supplementation in a Cretan population with non-alcoholic fatty liver disease: a prospective pilot study. *Ann Gastroenterol.* 2014;27(4):387–94.
146. Kalender S, Kalender Y, Ogutcu A, Uzunhisarçikli M, Durak D, Acikgoz F. Endosulfan-induced cardiotoxicity and free radical metabolism in rats: the protective effect of vitamin E. *Toxicology.* 2004;202(3):227–35.
147. Porter NA. Chemistry of lipid peroxidation. *Methods Enzymol.* 1984;105:273–82.
148. Nishitani S, Matsumura T, Fujitani S, Sonaka I, Miura Y, Yagasaki K. Leucine promotes glucose uptake in skeletal muscles of rats. *Biochem Biophys Res Commun.* 2002;299(5):693–6.
149. Mattick JSA, Kamisoglu K, Ierapetritou MG, Androulakis IP, Berthiaume F. Branched-chain amino acid supplementation: impact on signaling and relevance to critical illness. *Wiley Interdiscip Rev Syst Biol Med.* 2013;5(4):449–60.

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