Ischemia-reperfusion intervention: from enhancements in exercise performance to accelerates performance recovery - a systematic review and meta-analysis

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Abstract: It has been demonstrated that brief cycles of ischemia followed by reperfusion (IR) applied before exercise can improve performance and, IR intervention, applied immediately after exercise (post-exercise ischemic conditioning – PEIC) exerts a potential ergogenic effect to accelerate recovery. Thus, the purpose of this systematic review with meta-analysis was to identify the effects of PEIC on exercise performance, recovery and the responses of associated physiological parameters, such as creatine kinase, perceived recovery and muscle soreness, over 24 h after its application. From 3281 studies, six involving 106 subjects fulfilled the inclusion criteria. Compared to sham (cuff administration with low pressure) and control interventions (no cuff administration), PEIC led to faster performance recovery (p=0.004; ES=0.49) and lower increase in creatine kinase (p<0.001; ES=0.71) and muscle soreness (p<0.001; ES=0.89) over 24 h. The effectiveness of this intervention is more pronounced in subjects with low/moderate fitness level and at least a total time of 10 min of ischemia (e.g. 2 cycles of 5 min) is necessary to promote positive effects.

Keywords: Intermittent occlusion; Blood flow occlusion; Sports; Ergogenic; Ischemic postconditioning

1. Introduction

High-level sports performance is dependent on several factors which require high mechanical [1], psychological [2] and physiological [3] demands. Elite competitors are usually submitted to successive high volume and intensity training sessions and/or to multi-days competitions, with short intervals of recovery. These events can lead to physiological [3] and psychological [4] alterations, impairing sports performance. Thus, to increase the resistance to fatigue and to improve performance, many athletes and coaches search post-exercise recovery strategies [5].

In this context, cycles of ischemia-reperfusion (IR) performed immediately after exercise (Post-exercise ischemic conditioning - PEIC) are an interesting ergogenic aid to accelerate recovery during high intensity exercise sessions [6,7]. This intervention is actually of low cost, non-invasive, easy and quick-to-apply compared with others methods, such as cold water immersion [8,9]. The IR requires the use of a cuff (tourniquet) on the proximal regions of the lower or upper limbs and the perform of repeated bouts of ischemia interspersed with reperfusion periods [10]. Analyzing the studies with IR and exercise performance it is possible to verify that the most common applied IR protocols encompass 3 or 4 bouts of 5 min of ischemia followed by 5 min of reperfusion among bouts [11,12].
IR intervention was initially used before a prolonged ischemic insult that causes myocardial necrosis. This intervention, termed ischemic preconditioning, was able to confer cardiac protection against infarction [13] and it was associated with low increase of tissue necrosis biomarkers [i.e. creatine kinase (CK)] [14] and improved cardiac performance during exercise [15,16]. Afterwards, its use after prolonged ischemic insult, termed ischemic postconditioning, was also demonstrated to confer cardiac protection [14] and reduced oxidative stress [17]. IR intervention to improve exercise performance followed a similar path: firstly, performed before physical exercise promoted a better skeletal muscle capacity [18], improved performance of swimmers [19], runners [20], and cyclists [21]. Then, applied immediately during recovery phase from exercise, prevented the drops in performance 24 h, 48 h, and 72 h after an exercise-induced muscle damage, mitigating increases in CK [7]. It is important to note that both the mentioned tissue biomarkers and oxidative stress are associated with drop in sports performance [22,23]. Therefore, during recovery, PEIC application could attenuate tissue and oxidative injury caused by exercise.

The first study that evaluated PEIC effect during recovery phase [24], employed the intervention immediately after an exercise protocol that involved jumps and repeated sprints. After PEIC, the participants repeated the exercise protocol, and again 24 h later. Beneficial effects were observed both immediately and 24 h post intervention. Specifically, recovery of power production and sprint performance were improved. Compared with other IR studies [25,26], the authors used a short protocol (two bout of 3 min of ischemia followed by 3 min of reperfusion; 2 cycles x 3 min). Based on this protocol, Northey et al. [27] evaluated the velocity of recovery applying the PEIC immediately after a fatiguing resistance exercise protocol. PEIC was not able to attenuate the loss in muscle force during jumps and the torque during concentric isokinetic contractions 1 h and 24 h later. Similar lack of beneficial results using the same IR protocol were observed in academy rugby players 2 h and 24 h after PEIC application [28] but longer protocols (e.g. 3 cycles x 5 min) were efficient to prevent decrements in maximal voluntary contraction, jumps and sprints performance [7,29]. However, it is important to highlight that training status of participants were different among studies as well as exercise types used for produce fatigue and assess performance.

Beyond a limited amount of investigations and controversial results, it is important to describe the heterogeneity of PEIC protocols, exercises to induce fatigue (specific and non-specific), as well as sample with fitness level and subjects’ characteristics. Therefore, it would be appropriate to evaluate the current status and the future perspective of PEIC on physical performance recovery including their possible mechanisms. To this aim, we conducted this review and meta-analysis.

2. Materials and Methods

This systematic review with meta-analysis followed the same structure as research articles and conform to the Preferred Reporting Items for Systematic Reviews and Meta-analysis (PRISMA) guidelines.

2.1. Database search

Two independently reviewers, using PubMed, Scopus, SPORTDiscus and Web of Science database, identified potential studies published after January 2012 [After conducting a search in the database, it was identified that in 2012 the first study evaluating PEIC effect on performance recovery was published [24]] with the following key terms i) “intermittent occlusion”; ii) “ischemic conditioning”; iii) “ischemic post-conditioning”; iv) “vascular occlusion” combined with “recovery” and “performance recovery”. Only original research studies written in English were included. The literature search was completed on June the 30th 2020. Potential studies were selected based on strict criteria including: i) original research studies; ii) PEIC performed during post exercise; iii) evaluation of healthy participants iv) conduction of exercise or effort test and v) analysis of recovery performance. Studies with animal model or non-healthy subjects’ articles, systematic reviews and meta-analysis were excluded. Restrictions such as participant age and fitness level were not applied.
2.2 Study selection and quality assessment

It was selected all articles that met the criteria for inclusion and evaluation indicated above. The search revealed a total of 3281 articles. Primarily, the duplicates were removed and then the remaining studies were selected by title. If they matched our inclusion criteria, abstracts were checked. Finally, each study which appeared to respect the criteria of eligibility was reviewed (figure 1).

To estimate the quality of all articles included in this review, it was used a checklist (Table 1) based on our previous experience [12,30]. Two investigators analyzed the methodological quality of the articles assigning among 3 possible scores (Yes = 1 point, Unclear = 1/2 point, No = 0 points) for each item (total of 15) of the checklist. The maximum score was 15 points. The sum of the 15 criteria scores achieved by each paper was used to attribute its general quality. The conclusion of the total scores was taken in agreement with both the investigators. All data collection was conducted by two evaluators separately. Whenever discrepancies were detected, a third-part was also involved in the evaluation.

Figure 1. The procedure to select/inclusion of the studies.
Table 1. Checklist used to analyze the quality of publications

| Reporting                                                                 | 0 | ½ | 1 |
|--------------------------------------------------------------------------|---|----|---|
| 1. Is the hypothesis/aim/objective of the study clearly described?       | No | Unclear | Yes |
| 2. Are the main outcomes to be measured clearly described in the Introduction? | No | Unclear | Yes |
| 3. Are the characteristics of the subjects included in the study clearly described? | No | Unclear | Yes |
| 4. Are the interventions of interest clearly described?                   | No | Unclear | Yes |
| 5. Are the main findings of the study clearly described?                 | No | Unclear | Yes |
| 6. Does the study provide estimates of the random variability in the data for the main outcomes? | No | Unclear | Yes |
| 7. Were the instruments of testing reliable?                             | No | Unclear | Yes |
| 8. Was a follow-up duration sufficiently described and consistent within the study? | No | Unclear | Yes |
| 9. Number of participants included in study findings                     | < 5 | 6 - 15 | > 16 |

**Analysis and presentation**

|                                                                 | 0 | ½ | 1 |
|----------------------------------------------------------------|---|----|---|
| 10. Have actual probability values been reported (e.g. 0.035 rather than < 0.05) for the main outcomes except, where the probability value is less than 0.001? | No | Unclear | Yes |
| 11. Was there a statement adequately describing or referencing all statistical procedures used? | No | Unclear | Yes |
| 12. Were the statistical analyses used appropriate?                | No | Unclear | Yes |
| 13. Was the presentation of results satisfactory?                 | No | Unclear | Yes |
| 14. Were confidence intervals given for the main results?         | No | Unclear | Yes |
| 15. Was the conclusion drawn from the statistical analysis justified? | No | Unclear | Yes |
2.3. Data Analysis

Each study was read, and data extracted. Descriptive information (e.g., sample size, age, training level and PEIC sets), performance, perceived recovery, muscle soreness (MS), and CK data were collected from articles or requested to authors via e-mail. When the data were shown only by figures, the webplotdigitizer 4.3 software (apps.automeris.io/wpd) was used to extract them. Within-group change of these variables were determined by calculation of the difference between pre- and post-interventions. The mean relative percentage change was determined by post-minus pre-interventions values divided by pre-intervention value multiplied by 100. To compare the relative percentage change of performance between PEIC and sham/control groups, the Shapiro-Wilk test was performed to verify the normality of data followed by Mann-Whitney test. The GraphPad® (Prism 6.0, San Diego, CA, USA) was used for this analysis.

Only variables assessed in more than three studies were taken into account for the meta-analysis. Therefore, the meta-analysis was conducted separately for performance, CK and MS variables using the Review Manager Software, version 5.4, Copenhagen (Denmark). The level of significance adopted was p ≤ 0.05. The effect size (ES) was calculated for each variable using pre-intervention and post-intervention and standard deviations (SDs) to determine the meaningfulness of the difference. A value < 0.2 was considered trivial, > 0.2 - 0.6 small, > 0.6 - 1.2 moderate, > 1.2 - 2.0 large, and > 2.0 - 4.0 very large effect [31]. When a study lacked from the necessary data, the following equation (1) was used to estimate the SD change:

\[
\text{SD change} = \sqrt{([SD \text{ pre}]^2 + [SD \text{ post}]^2 - 2 \times corr \times [pre.post] \times SD \text{ pre} \times SD \text{ post})}
\]

The correlation factor (f.corr) of 0.80 [32] was used for both the PEIC and sham/control groups. To estimate inter-study heterogeneity the I2 statistic was used, where I2 = 25% was considered low, I2 = 50% moderate and I2 = 75% high [33].

3. Results

The search, selection, and inclusion process revealed that only six articles met our inclusion criteria [6,7,24,27–29]. Of these six, four [6,7,24,29] presented favorable results to PEIC and performance recovery, and two studies [27,28] were judged not favorable to PEIC. No study showed a negative effect of PEIC on recovery performance. All studies not favorable to PEIC used cross-over design, while the studies favorable, just two implemented this design [24,29].

The quality scores of the analyzed studies achieved a mean of 12.3 ± 0.84 (81.7 ± 5.0%) points. Only one study obtained a value of 13.5 [6], other studies obtained scores 12.5 [28,29], 12.0 [24] and 11.5 [7,27] (Table S1).

There was a total of 106 participants (102 males and only 4 females) with an average age of 25.1 years (range of standard deviation of 1.0 – 7.0 years). Two studies involved healthy recreational trained subjects [7,24], one study was conducted in trained subjects [6], one in well-trained subjects [27], one in college level participants [28], and one in semi-professional soccer players [29].

The exercise protocols utilized to induce fatigue were different among studies but, in all of them, the authors prescribed only lower body exercises. Three studies performed the same exercises to evaluate performance and induce fatigue (i.e. the subjects performed maximal incremental test to induce fatigue and evaluated recovery performance) [6,7,24], and three included different types of exercise (e.g. one evaluated performance and the other fatigue) [27–29]. It is important highlight that, in the study of Daab et al. [29], some exercises included in the protocol to evaluate performance (e.g. sprints) were also performed in the protocol to induce fatigue, which is specific for soccer. These results are presented in table 2.
Table 2. Characteristics of the PEIC studies

| Study                      | N   | Male | Female | Samples | Subjects                              | (Exercise) Fatigue Protocol | Exercise/test to assess performance | Is PEIC favorable to performance? | Other variables analyzed |
|----------------------------|-----|------|--------|---------|---------------------------------------|----------------------------|-------------------------------------|----------------------------------|--------------------------|
| Beaven et al. (2012)       | 14  | 10   | 4      | Paired  | Healthy Recreationally trained         | Jumps / sprints / leg press test | Jumps / sprints / leg press test    | Yes                             | #                        |
| Northey et al. (2016)      | 12  | 12   | 0      | Paired  | Healthy well trained (resistance exercise) | Back Squat (10 sets x 10 repetitions (70% 1RM)) | MVC / Jumps | No                     | MS and PRS               |
| Page et al. (2017)         | 16  | 16   | 0      | No Paired | Healthy Recreationally active          | Jumps (5 sets x 20 repetitions own body weight (box 0.6 m height)) | MIVC / Jumps | Yes                  | MS*, CK*, TC             |
| William et al. (2018)      | 24  | 24   | 0      | Paired  | Rugby Player (college level)           | 6 sets x 50 meters sprints | Jumps | No                     | MS, PRS, CK, lactate, cortisol and testosterone |
| Arriel et al. (2018)       | 28  | 28   | 0      | No Paired | Trained cyclists                       | Maximal Incremental         | Maximal Incremental | Yes                  | MS, PRS, RPE, CK, HR    |
| Daab et al. (2020)         | 12  | 12   | 0      | Paired  | Semi-professional soccer players       | Loughborough intermittent shuttle test & | Jumps / Sprint / MVC | Yes                  | MS, CK, LDH, CRP         |
PEIC, post-exercise ischemic conditioning; MS, muscle soreness; PRS, perceived recovery; RPE, perceived exertion; TC, thigh circumference; CK, creatine kinase; heart rate, HR; LDH, lactate dehydrogenase; CRP, serum C-reactive protein. &, protocol designed to simulate the activities of a real soccer match [six exercise sets lasting approximately 15 min (between 55% and 95% VO2 Max) separated by periods of 3 min] [34]; #, no evaluated. * It was influenced by PEIC.
Table 3 shows information regarding PEIC procedures, interval between fatiguing exercise and PEIC intervention as well as the interval between PEIC application and performance test, information about possibly PEIC effects and type of intervention (i.e. PEIC, sham or control). Only one study performed PEIC or sham intervention 5 min after effort [6] while the five others performed immediately after it [7, 24, 27–29]. At first glance, results do not apparently show a consensus on the PEIC procedure. Most of the studies included sham/placebo intervention but they did not include a control intervention. The study which included a control intervention did not present favorable responses to PEIC [27] and the authors did not include a sham intervention. No studies applied a remote PEIC, i.e. no studies utilized intermittent vascular occlusion applied to remote areas with respect to the exercise muscle group (e.g. apply the PEIC on the lower limbs and performed exercises with the upper limbs or the other way around).
| PEIC sets | Total PEIC and SHAM time (min) | Ischemia pressure (mm Hg) PEIC / SHAM / Limb | Time to test | Groups | Were subjects informed about effects of PEIC? |
|-----------|--------------------------------|---------------------------------------------|--------------|--------|---------------------------------------------|
| Beaven et al. (2012) | 2 x 3 min | 6 | 220 / 15 / thigh | 5 min – 24 h | PEIC / SHAM | No |
| Northey et al. (2016) | 2 x 3 min | 6 | 220 / # / thigh | 1 – 24 h | PEIC / CON | It is not exposed by authors |
| Page et al. (2017) | 3 x 5 min | 15 | 220 / 20 / thigh | 24 – 48 – 72 h | PEIC / SHAM | No |
| William et al. (2018) | 2 x 3 min | 6 | 171 – 266 / 15 / thigh | 2 – 24 h | PEIC / SHAM | Yes |
| Arriel et al. (2018) | 2 x 5 min and 5 x 2 min | 10 and 10 | 50 > SAP / 20/ thigh | 24 h | PEIC / SHAM | Yes |
| Daab et al. (2020) | 3 x 5 min | 15 | 50 > SAP / 20/ thigh | 0 – 24 – 48 – 72 h | PEIC / SHAM | It is not exposed by authors |

PEIC, post-exercise ischemic conditioning; SHAM, cuff administration with low pressure; CON, control; SAP, Systolic arterial pressure; # No SHAM application
Although prevented the drop in exercise performance 24 h post fatigue protocol (Table 4), PEIC did not influence most of the physiological and perceptual variables (Table 5). Only CK, a marker of tissue injury, was attenuated 24 h after the PEIC intervention in well trained subject to resistance exercise [7] and semi-professional soccer players [29], but CK was not attenuated in trained cyclists [6] and in academy rugby union players [28]. However, when we pooled data, the CK percentage change was higher to sham/control compared to PEIC intervention.
Table 4. Comparison of the performance recovery between PEIC and SHAM/CON.

|                   | PEIC               | SHAM / CON          |
|-------------------|--------------------|---------------------|
|                   | Pre-intervention   | 24-h Post-intervention | Change (%) | Pre-intervention   | 24-h Post-intervention | Change (%) |
| Exercise          |                    |                     |            |                    |                     |            |
| Beaven et al. (2012) | SJₑa (m.s⁻²)      | 20.1 ± 3.9          | 22.8 ± 4.3 | 11.8               | 18.9 ± 3.7           | 17.5 ± 3.6 | -7.4 |
|                   | S 10m (s)          | 12.5 ± 0.8          | 12.4 ± 0.8 | 0.8                | 12.6 ± 0.7           | 12.7 ± 0.8 | -0.8 |
|                   | S 40m (s)          | 42.5 ± 3.4          | 41.8 ± 3.3 | 1.7                | 42.7 ± 3.1           | 42.7 ± 3.2 | 0.0  |
| Northey et al. (2016) | CMJ (cm)          | 41.8 ± 8.8          | 42.1 ± 6.9 | 0.7                | 42.7 ± 7.7           | 42.1 ± 7.0 | -1.4 |
|                   | SJ (cm)            | 37.7 ± 7.8          | 35.3 ± 7.2 | -6.4               | 38.1 ± 5.6           | 36.0 ± 6.1 | -5.5 |
|                   | MD (Kg/f)          | 281.5 ± 46.0        | 256.4 ± 52.0 | -8.9           | 273.7 ± 35.5        | 270.1 ± 39.1 | -1.3 |
| Page et al. (2017) | CMJ (cm)           | 34.0 ± 4.4          | 28.7 ± 1.2 | -15.6              | 38.9 ± 8.1           | 31.1 ± 2.0 | -20.1 |
|                   | MD (Kg/f)          | 611.0 ± 51.0        | 556.0 ± 67.2 | -9.0           | 629.0 ± 136.0       | 515.8 ± 43.3 | -18.0 |
| William et al. (2018) | CMJ (cm)          | 40.4 ± 6.0          | 38.9 ± 6.2 | -3.7               | 39.7 ± 6.0           | 37.6 ± 5.6 | -5.3 |
| Arriel et al. (2018) | IT (s)             | 808.3 ± 122.9       | 811.4 ± 135.1 | 0.4             | 779.9 ± 122.9       | 753.4 ± 110.0 | -3.4 |
| Daab et al. (2020) | CMJ (%)            | 100.0 ± 0.0         | 98.3 ± 1.8 | -1.7               | 100.0 ± 0.0          | 90.6 ± 1.9 | -9.4 |
|                   | SJ (%)             | 100.0 ± 0.0         | 98.8 ± 2.3 | -1.2               | 100.0 ± 0.0          | 90.7 ± 2.4 | -9.3 |
|                   | MD (%)             | 100.0 ± 0.0         | 98.2 ± 4.4 | -1.8               | 100.0 ± 0.0          | 69.8 ± 4.4 | -30.2 |
|                   | S 20m (%)          | 100.0 ± 0.0         | 103.1 ± 0.8 | -3.1           | 100.0 ± 0.0          | 106.7 ± 1.4 | -6.7 |
| **Mean**          |                    |                     | **-2.6***  |                    |                     | **-8.5**  |        |

PEIC, post-exercise ischemic conditioning; SHAM, cuff administration with low pressure; CON, control; SJₑa = squat jump eccentric acceleration; S 10m = 10 meters sprint times over the 6 repeated sprints sprint of 10 meters; S 20m = 20 meters sprint; S 40m = 40 meters sprint times over the 6 repeated sprints; MD = Muscle dynamometry; IT = Incremental test; CMJ = countermovement jump; SJ = Squat Jump.* different from SHAM/CON, p = 0.049
Table 5. Results of the perceived recovery (PRS), muscle soreness (MS) and creatine kinase (CK) between PEIC and SHAM/CON.

| Exercise | PEIC | SHAM / CON |
|----------|------|------------|
|          | Pre-intervention | 24-h Post-intervention | Change (%) | Pre-intervention | 24-h Post-intervention | Change (%) |
| Northey et al. (2016) | PRS (scores) | 8.1 ± 1.5 | 5.6 ± 1.6 | -30.9 | 7.9 ± 0.9 | 5.1 ± 1.9 | -35.4 |
| | MS (scores) | 0.6 ± 0.8 | 3.1 ± 1.9 | 416.7 | 1.0 ± 0.8 | 4.4 ± 2.4 | 340.0 |
| Page et al. (2017) | MS (scores) | 8.9 ± 8.0 | 57.0 ± 24.6 | 540.5 | 15.6 ± 12.5 | 106.1 ± 30.1 | 580.1 |
| | CK (U/L) | 163.5 ± 30.1 | 335.8 ± 243.8 | 105.4 | 178.4 ± 61.4 | 636.4 ± 300.1 | 256.7 |
| William et al. (2018) | CK (U/L) | 218.9 ± 81.9 | 627.1 ± 250.7 | 186.5 | 228.6 ± 81.9 | 731.6 ± 189.7 | 220.0 |
| | PRS (scores) | 8.2 ± 2.2 | 7.2 ± 2.0 | -12.2 | 7.5 ± 2.3 | 7.4 ± 1.7 | -1.3 |
| | MS (scores) | 0.8 ± 1.2 | 0.7 ± 0.9 | -12.5 | 0.6 ± 1.1 | 0.7 ± 1.1 | 16.7 |
| | CK (U/L) | 205.9 ± 138.4 | 244.0 ± 160.2 | 18.5 | 192.5 ± 127.6 | 228.3 ± 138.5 | 18.6 |
| Daab et al. (2020) | MS (Scores) | 0.5 ± 0.1 | 2.5 ± 0.8 | 400.0 | 0.6 ± 0.5 | 3.9 ± 1.1 | 550 |
| | CK (%) | 100.0 ± 0.0 | 200.7 ± 77.9 | 100.7 | 100.0 ± 0.0 | 426.8 ± 75.8 | 326.8 |
| Mean | PRS | -21.6 | 336.2 | 102.8 | | | -18.4 |

PEIC, post-exercise ischemic conditioning; SHAM, cuff administration with low pressure; CON, control.
3.1. Performance recovery, muscle injury markers and muscle soreness analysis

The effect of PEIC on performance recovery and CK are shown in figure 2. Concerning recovery performance, there was a significant small effect size of PEIC compared to sham or control group (ES = -0.49, [CI: -0.82, -0.15], p = 0.004). Moderate but non-significant heterogeneity was found amongst studies (I² = 26%, p = 0.17). Regarding CK, there was a significant moderate effect size favoring PEIC compared to sham or control (ES = -0.71, [CI: -1.11, -0.32], p < 0.01). A high significant heterogeneity was found amongst studies (I² = 83%, p < 0.01). Finally, the increase in MS was significantly lowered by PEIC compared to sham or control, with a moderate effect size (ES = -0.89, [CI: -1.34, -0.44], p < 0.01). A moderate significant heterogeneity was found amongst studies (I² = 68%, p = 0.02).
| Study or Subgroup | PEIC Mean | PEIC SD | PEIC Total | SHAM/CON Mean | SHAM/CON SD | SHAM/CON Total | Weight | IV, Fixed, 95% CI | Std. Mean Difference IV, Fixed, 95% CI |
|------------------|-----------|---------|------------|---------------|-------------|----------------|--------|------------------|---------------------------------------|
| Arriel 2018 IT   | -3.1      | 30.8    | 14         | 26.5          | 25.7        | 14             | 17.7%  | -1.01 [1.81, -0.22] |                                      |
| Beaven 2012 SJ   | -2.7      | 2.6     | 5          | 1.4           | 2.3         | 5              | 4.9%   | -1.51 [3.01, -0.01] |                                      |
| Beaven 2012 Sprint 10 | -0.1    | 0.5     | 4          | 0.1           | 0.5         | 4              | 5.6%   | -0.35 [-1.75, 1.06] |                                      |
| Beaven 2012 Sprint 40 | -0.7   | 2.1     | 4          | 0             | 2           | 4              | 5.7%   | -0.30 [-1.70, 1.10] |                                      |
| Daab 2020 CMJ    | 1.7       | 1.8     | 3          | 9.4           | 1.9         | 3              | 0.9%   | -3.33 [-6.02, 0.26] |                                      |
| Daab 2020 MD     | 1.8       | 4.4     | 3          | 30.2          | 4.4         | 3              | 0.4%   | -5.16 [-10.40, 0.07] |                                      |
| Daab 2020 SJ     | 1.2       | 2.3     | 3          | 9.3           | 2.4         | 3              | 1.2%   | -2.76 [-5.06, 0.35] |                                      |
| Daab 2020 Sprint | 3.1       | 0.8     | 3          | 6.7           | 1.4         | 3              | 1.3%   | -2.53 [-5.44, 0.39] |                                      |
| Northey 2016 CMJ | -0.3      | 5.3     | 4          | 0.6           | 4.7         | 4              | 5.8%   | -0.16 [-1.55, 1.23] |                                      |
| Northey 2016 MD  | 25.1      | 31.5    | 4          | 3.6           | 23.0        | 4              | 5.2%   | 0.67 [0.79, 2.13]   |                                      |
| Northey 2016 SJ  | 2.4       | 4.8     | 4          | 2.1           | 3.7         | 4              | 5.8%   | 0.06 [-1.33, 1.45]  |                                      |
| Page 2017 CMJ    | 5.3       | 3.5     | 4          | 7.8           | 6.6         | 4              | 5.6%   | -0.41 [-1.83, 1.00] |                                      |
| Page 2017 MD     | 55.4      | 40.4    | 4          | 113.2         | 104.6       | 4              | 5.3%   | -0.64 [-2.09, 0.82] |                                      |
| Williams 2018 CMJ| 1.5       | 3.9     | 24         | 2.1           | 3.7         | 24             | 34.7%  | -0.16 [-0.72, 0.41] |                                      |

Total (95% CI) 83 83 100.0% -0.49 [-0.82, -0.15]

Heterogeneity: Chisq = 17.56, df = 13 (P = 0.17), I² = 26%
Test for overall effect: Z = 2.85 (P = 0.004)
Figure 2. Forest plot of performance recovery (A) creatine kinase (B) and muscle soreness (C) variables between post-exercise ischemic conditioning (PEIC) and a cuff administration with low pressure (SHAM) or control (CON) interventions. The square is the weight for a given study and is proportional to the weight of the study in the meta-analysis. The horizontal line indicates the 95% confidence interval (CI) for an effect. The diamond at the bottom represents the overall effect calculated using a fixed-effects model. IT = incremental test; SJ = squat jump; Sprint 10 = 10 meters sprint times over the 6 repeated sprints; sprint 40 = 40 meters sprint times over the 6 repeated sprints; CMJ = countermovement jump; MD = muscle dynamometry; Sprint = 20 meters sprint.
4. Discussion

This is the first review and meta-analysis to analyze the effects and interpret the application of PEIC on the performance during recovery. Although a consensus among studies does not exist, our results indicate that PEIC leads to a significantly greater performance recovery and attenuates the CK increase 24 h post fatiguing protocols, especially in subjects with low to moderate levels of physical fitness. Among studies that did not present favorable results to PEIC, one did not include a placebo/sham intervention [27] and another did not include a control intervention [28] in their experimental design. In addition, both investigations adopted a cross-over design. These approaches could generate biased results because it was not possible to blind the subjects to PEIC intervention. Selected ones presented a high quality but only two described statistical significance in the measure of performance and physiological variables [7,29]. Considering that there is no standardization of the present nomenclature and that occlusion-reperfusion interventions are used indifferently prior or post exercise, we would like to suggest that in the future studies researchers use the term post-exercise ischemic conditioning (PEIC) when referring to this type of intervention.

4.1. Quality of the papers

Although a high-quality score has been achieved by studies, some limitations were found. Most of the studies did not described clearly the characteristics of the subjects included in the investigation (criterion 3) [7,27,28]. This can make it difficult to interpret results and to reapply the protocols used. Only two studies reported the exact P-value (criterion 10) for main results [28,29]. The latter one provided information about the strength of the evidence against the null hypothesis, avoiding doubts for the reader to make a decision. We have also concerns about the appropriate use of statistical analysis (criterion 12) [7,24,28], once authors did not perform normality tests of data nor parametric tests for categorical variables (e.g. perceived exertion and pain perception). In addition, they utilized the independent test for paired sample. However, this inappropriate statistical did not influence our meta-analysis, since we have worked with raw data.

Finally, only one study included confidence intervals (CI) for the main results (criterion 14) [24]. The CI is employed to show the dispersion or variability e reliability of an estimate, which likely includes the estimate of the average of populations. This is influenced by the sample size and the homogeneity of the data sample and it can be used to describe how reliable are results of a research. In addition, only two studies estimated the sample size and statistic power [6,29].

4.2. Participants involved

Of 106 participants, only four were female, which reduces the possibility of suggesting the PEIC effects in women. The studies that demonstrated favorable effects of PEIC on performance recovery [6,7,24,29] involved healthy participants, semi-professional soccer players and recreational trained or active subjects, while other studies involved well trained and college level participants [27,28]. There is no study with elite or high-trained athletes. This fact is comprehensible due the difficulty in recruiting athletes available for this kind of research. Therefore, the positive effects of PEIC has not been investigated in this type of population because their physiological and performance responses are different from those found in non-athletes in several aspects. In addition, the responsiveness of IR intervention, when applied before exercise or test to improve performance, has been associated with the training level of the subjects. Specifically, it was demonstrated that participants with low fitness level presented large [35] while high training level small [19] or no response to this intervention [36]. In this context, since PEIC responses have the same pattern, it could be speculated that its responses are also dependent to the fitness level of subjects. However, it is important to highlight that following the quality scores of studies most of them did not clearly describe the characteristics of the enrolled subjects.
4.3. Exercise protocols to induce fatigue and assess performance

The PEIC effects were primarily tested in exercise types that involved vertical jumps, sprints, and resistance exercise [24]. Although one study investigated on incremental cycling test [6], the most common tests were jumps and maximal voluntary contraction [7,27–29]. Although these tests were largely used to evaluate changes in performance [37], they may not be sensitive enough to identify these changes 24 h after exercise [27]. Among studies selected for this review some did not identify significant declines in exercise performance 24 h after the execution of fatiguing protocols in the sham/control, or PEIC groups when compared to pre-intervention. For example, in the Northey et al. [27] study, the performance of countermovement jump and concentric isokinetic peak torque measured in the pre-fatiguing protocol were not different after 24 h among and within groups. The same result was obtained by Williams et al. [28], who investigated peak power output and jump height. They found no difference 24 h after fatiguing protocol for both sham/control and PEIC groups when compared to pre-intervention. On the other hand, in the Arriel et al. [6] study, the sham group presented a drop of performance during incremental cycling 24 h after fatiguing protocol, but after PEIC this phenomenon was not observed. The same happened in the study by Page et al. [7] and by Daab et al. [29] but on the maximal isometric voluntary contraction. Alternatively, this fact may also mean that the physical exercise dose to induce fatigue was not able to cause a drop in performance 24 h after physical exercise. Therefore, these procedures may lead to misleading conclusions on the potential effects of PEIC 24 h post physical exercise.

4.4. PEIC effects on recovery performance, creatine kinase and muscle soreness

Oxidative stress, muscular damage, increase in inflammation, and MS associated with a decreased performance have been all reported after exhaustive exercise performed by different athletes involved in different sports [3,38,39]. This especially over the course of the multi-day races (i.e. 2 to 7 consecutive days of competition) [3,39] and with high training volumes [40]. This phenomenon may be useful for athletes and coaches searching strategies that minimize decrements in performance or to accelerate the recovery from fatiguing efforts. As exposed in figure 2, our statistical analysis showed a favorable PEIC effect on recovery performance, CK (muscle damage markers), and MS. Thus, we can support the use of the PEIC immediately after exercise to reduce these markers and to speed recovery.

It is important to note that all studies analyzed by the present review evaluated only the acute effect of PEIC application. Although there is no study investigated the repeated effect of PEIC application (i.e. several days of application), recent study showed that several days of application of blood flow restriction (just one cycle, and lower pressure than PEIC intervention) after resistance exercise was associated with an impaired muscle adaptation [41], and this fact may be due to magnitude of oxidative stress. The oxidative stress when moderate, play multiple regulatory roles in cells, such as regulation of cell signaling pathways. However, it was speculated that low-to-none levels are not beneficial [42]. Therefore, as cycles of IR were associated with attenuated stress oxidative level [43,44], we suggest that PEIC application should be performed before main competitions or when athletes incorporate a high training volume session. However, further studies are necessary to investigate the PEIC contribution when applied repeatedly on different exercise modes and kinds of sport activities.

4.5. PEIC protocols and possible mechanism

At first glance, there is no a consensus among researchers about the number and duration of IR cycles during PEIC. Furthermore, no consensus exists on period between PEIC application and the return to exercise and whether participants involved were informed about the possible effects of PEIC.

The cycles of ischemia presented a total time ranging from 6 to 15 min and the complete PEIC protocol (occlusion and reperfusion periods) from 12 to 30 min (Table 3). Only one study analyzed two different protocols [6]. While some authors using a total time of ischemia above of 10 min
described positive effects on recovery [7,29,35], others using shorter time did not [27,28]. Although Beaven et al. [24] found positive results for PEIC using total time of 6 min, in this investigation subjects were recreational athletes. Only one study verified two different PEIC protocols with the same total time of ischemia (10 min), but authors did not find significant difference between protocols [6]. A clinical study [45] that demonstrated cardiac injury protection applying intermittent vascular occlusion before prolonged ischemic insult, reported that 4 to 6 ischemic cycles lasting 2 to 5 min yielded significant cardioprotection. Therefore, in addition to training level, it is conceivable that a minimal dose of ischemia is necessary to generate a PEIC response and that highly-trained subjects would need higher dose of PEIC than trained and untrained individuals.

The time between PEIC application and the return to exercise was usually 24 h. However, PEIC effects were also evaluated at 5 min, 1 h, 2 h, 48 h and 72h. While no positive effects were found between 1 and 2 hours after PEIC application [27,28], 24 h has revealed significant changes [6,7,24,29]. Among the few established IR intervention mechanisms, the early (active immediately after reperfusion and lasts 2-3 h) and late (begins 12-24 h after reperfusion) phases of protection, commonly known as the first and second window of protection [16], should be considered. Looking at our results, the second phase appears to be more pronounced. However, some variables (e.g. CK and MS) used for assess the effects of PEIC on recovery have their peak 24 h post exercise. Therefore, it remains unclear whether the first, the second or both phases are effective on performance recovery.

Finally, we can hypothesize that on the early phase, the PEIC could increase nitric oxide and modulate mitochondrial oxygen consumption leading to a decrease of mitochondrial reactive oxygen species (ROS) generation [43] and consequently limit oxidative injury on muscle cell. During the late phase, PEIC could increases iNOS expression (an isoform that synthesizes nitric oxide) [46], and consequently increases nitric oxide production, leading to a possibly improvement in exercise performance by diminishing the level of ROS [43]. These occurrences are also associated with reduced muscle fatigue and damage [47]. In addition, it was speculated that PEIC could lead to an attenuated inflammatory response after physical exercise due to a downregulation of circulating leukocytes [7]. This could be, at least in part, the responsible for the beneficial effects of PEIC on recovery.

5. Conclusions

PEIC intervention has proved to be an effective, non-invasive, inexpensive and easy-to-apply strategy to accelerates performance recovery by the attenuation of creatine kinase and muscle soreness increase in subjects with low-to-moderate fitness level. In highly-trained subjects, a higher dose of PEIC may be administered to elucidate beneficial effects, while the ideal PEIC protocol has not been standardized yet. This intervention is a new approach in the field of ischemic conditioning, which can be applied after exercises that potentially cause muscle damage and soreness such as during multi-day competitions or high training volumes sessions.

Supplementary Materials: The following are available online at www.mdpi.com/xxx/s1, Table S1: Scores assigned to each of the studies for each of the quality (Q) criteria.

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