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Cold-water immersion (cryotherapy) for preventing and treating muscle soreness after exercise (Review)

Bleakley C, McDonough S, Gardner E, Baxter GD, Hopkins JT, Davison GW

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[Intervention Review]

Cold-water immersion (cryotherapy) for preventing and treating muscle soreness after exercise

Chris Bleakley[1], Suzanne McDonough[1], Evie Gardner[2], G. David Baxter[3], J. Ty Hopkins[4], Gareth W Davison[2]

[1]Health and Rehabilitation Sciences, University of Ulster, Newtownabbey, UK. [2]Faculty of Life and Health Sciences, University of Ulster, Newtownabbey, UK. [3]School of Physiotherapy, University of Otago, Dunedin, New Zealand. [4]Department of Exercise Sciences, Brigham Young University, Provo, Utah, USA

Contact: Chris Bleakley, Health and Rehabilitation Sciences, University of Ulster, School of Health Sciences, Shore Road, Newtownabbey, County Antrim, BT37 0QB, UK. e10204083@uucde.ulst.ac.uk, chrisbleakley@hotmail.com.

A B S T R A C T

Background

Many strategies are in use with the intention of preventing or minimising delayed onset muscle soreness and fatigue after exercise. Cold-water immersion, in water temperatures of less than 15°C, is currently one of the most popular interventional strategies used after exercise.

Objectives

To determine the effects of cold-water immersion in the management of muscle soreness after exercise.

Search methods

In February 2010, we searched the Cochrane Bone, Joint and Muscle Trauma Group Specialised Register, the Cochrane Central Register of Controlled Trials (*The Cochrane Library* (2010, Issue 1), MEDLINE, EMBASE, Cumulative Index to Nursing and Allied Health (CINAHL), British Nursing Index and archive (BNI), and the Physiotherapy Evidence Database (PEDro). We also searched the reference lists of articles, handsearched journals and conference proceedings and contacted experts.

In November 2011, we updated the searches of CENTRAL (2011, Issue 4), MEDLINE (up to November Week 3 2011), EMBASE (to 2011 Week 46) and CINAHL (to 28 November 2011) to check for more recent publications.

Selection criteria

Randomised and quasi-randomised trials comparing the effect of using cold-water immersion after exercise with: passive intervention (rest/no intervention), contrast immersion, warm-water immersion, active recovery, compression, or a different duration/dosage of cold-water immersion. Primary outcomes were pain (muscle soreness) or tenderness (pain on palpation), and subjective recovery (return to previous activities without signs or symptoms).

Data collection and analysis

Three authors independently evaluated study quality and extracted data. Some of the data were obtained following author correspondence or extracted from graphs in the trial reports. Where possible, data were pooled using the fixed-effect model.

Main results

Seventeen small trials were included, involving a total of 366 participants. Study quality was low. The temperature, duration and frequency of cold-water immersion varied between the different trials as did the exercises and settings. The majority of studies failed to report active surveillance of pre-defined adverse events.

Fourteen studies compared cold-water immersion with passive intervention. Pooled results for muscle soreness showed statistically significant effects in favour of cold-water immersion after exercise at 24 hour (standardised mean difference (SMD) -0.55, 95% CI -0.84 to -0.27; 10 trials), 48 hour (SMD -0.66, 95% CI -0.97 to -0.35; 8 trials), 72 hour (SMD -0.93; 95% CI -1.36 to -0.51; 4 trials) and 96 hour (SMD -0.58; 95% CI -1.00 to -0.16; 5 trials) follow-ups. These results were heterogeneous. Exploratory subgroup analyses showed that studies using cross-over designs or running based exercises showed significantly larger effects in favour of cold-water immersion. Pooled results from two studies found cold-water immersion groups had significantly lower ratings of fatigue (MD -1.70; 95% CI -2.49 to -0.90; 10 units scale, best to worst), and potentially improved ratings of physical recovery (MD 0.97; 95% CI -0.10 to 2.05; 10 units scale, worst to best) immediately after the end of cold-water immersion.

Five studies compared cold-water with contrast immersion. Pooled data for pain showed no evidence of differences between the two groups at four follow-up times (immediately, 24, 48 and 72 hours after treatment). Similar findings for pooled analyses at 24, 48 and 72 hour follow-ups applied to the four studies comparing cold-water with warm-water immersion. Single trials only compared cold-water immersion with respectively active recovery, compression and a second dose of cold-water immersion at 24 hours.

Authors' conclusions

There was some evidence that cold-water immersion reduces delayed onset muscle soreness after exercise compared with passive interventions involving rest or no intervention. There was insufficient evidence to conclude on other outcomes or for other comparisons. The majority of trials did not undertake active surveillance of pre-defined adverse events. High quality, well reported research in this area is required.

PLAIN LANGUAGE SUMMARY

Cold-water immersion for preventing and treating muscle soreness after exercise

Delayed onset muscle soreness commonly results after sports and exercise activity. Cold-water immersion (CWI), which involves people immersing themselves in water at temperatures of less than 15°C, is sometimes used to manage muscle soreness after exercise and to speed up recovery time.

Our review included 17 small trials, involving a total of 366 participants. Study quality was low. Fourteen trials compared cold-water immersion applied after exercise with 'passive' treatment involving rest or no treatment. The temperature, duration and frequency of cold-water immersion varied between the different trials as did the exercises and settings. There was some evidence that cold-water immersion reduces muscle soreness at 24, 48, 72 and even at 96 hours after exercise compared with 'passive' treatment. Limited evidence from four trials indicated that participants considered that cold-water immersion improved recovery/reduced fatigue immediately afterwards. Most of the trials did not consider complications relating to cold-water immersion and so we cannot say whether these are a problem. There were only limited data available for other comparisons of cold-water immersion versus warm or contrasting (alternative warm/cold) water immersion, light jogging, and compression stockings. None of these showed important differences between the interventions being compared.

While the evidence shows that cold-water immersion reduces delayed onset muscle soreness after exercise, the optimum method of cold-water immersion and its safety are not clear.

BACKGROUND

The benefits of maintaining an active lifestyle have been well documented, and daily exercise can reduce the risk of serious health problems such as obesity and cardiovascular disease (Van Dam 2008). Depending on intensity levels, exercise can induce various degrees of fatigue to the musculoskeletal, nervous and metabolic systems. Exercise is also associated with microscopic tears in muscle tissue, commonly known as exercise induced muscle damage (EIMD), which may lead to delayed onset muscle soreness (DOMS).

Description of the condition

DOMS commonly peaks between 24 and 48 hours (in some reports up to 72 hours) after exercise and is characterised by muscle shortening, increased passive stiffness, swelling, decreases in strength and power, localised soreness and altered proprioception (Cleak 1992; Proske 2001). DOMS is more likely to occur with unaccustomed exercise or intense exercise involving eccentric muscle activity (i.e. when the muscle is forcibly stretched when active) (Cheung 2003). Although the physiological mechanism underpinning DOMS has not been fully elucidated, it may relate to primary mechanical damage that occurs to muscle cells during exercise. Microscopic disruption of the small muscle fibre units (known as sarcomeres) (Proske 2001) is also related to a number of inflammatory events, and the release of intracellular enzymes such as creatine kinase (CK) (Chatzinikolaou 2010). This response is thought to contribute to the characteristic pain, swelling and decreased muscle function associated with DOMS.

DOMS usually resolves clinically after approximately four to five days, and therefore it is generally regarded as being less severe than other muscle traumas such as strains or complete tears. Notwithstanding this, there are currently a number of prophylactic strategies to prevent or minimise the impact of DOMS thereby allowing athletes to recover faster following exercise. The most common approach is ensuring adequate rest between exercise bouts; however, this is often complemented with other intervention strategies. These strategies include cool down, stretching, massage, non-steroidal anti-inflammatory drugs, hydrotherapy and compression garments. Currently, there is little scientific evidence to support their use. Indeed, a recent Cochrane review found that stretching has no clinically important effect in reducing muscle soreness (Herbert 2011).

Description of the intervention

Recently, cold-water immersion (CWI) has emerged as one of the most popular interventions to prevent DOMS and promote recovery after exercise. Having started in elite level sport, it is becoming increasingly popular amongst amateur athletes. Immediately after exercise, individuals immerse themselves into cold-water baths which vary from custom built temperature controlled spas, to large containers filled with water and ice. In practice there are large variations in the CWI protocols that are employed in terms of: the duration of immersion; water temperature; and the volume of body parts immersed. Sellwood 2007b reported that, anecdotally, contemporary practice among "high-level sports in Australia" was three cycles, each comprising a one-minute immersion in ice water (at approximately 5°C water temperature) followed by one minute out, started immediately after exercise. However, some clinical studies have employed longer durations of application of up to 15 minutes (Banfi 2008; Vaile 2008d), and higher water temperatures of up to 15°C (Vaile 2008e).

How the intervention might work

Despite positive anecdotal reports, the exact physiological rationale for using CWI has not been elucidated. Cold therapy (cryotherapy) has traditionally been reserved for acute soft tissue injuries (e.g. sprains, strains and muscle contusions) to reduce pain, swelling, metabolism and inflammation associated with secondary injury (Knight 2000). It is proposed that cooling tissue immediately after exercise could have the same anti-inflammatory effect, thereby reducing the potential for DOMS. Another common concept is that CWI causes vasoconstriction (decreased blood vessel diameter) in the immersed musculature, which stimulates blood flow and nutrient and waste transportation through the body after exercise. Additionally CWI may decrease nerve transmission speed (Wilcock 2006) and alter receptor threshold, cumulating in decreased pain perception. There may also be a psychological mechanism, whereby the body simply feels more 'awake' with a reduced sensation of fatigue after exercise (Cochrane 2004).

Why it is important to do this review

Common clinical practices must have a clear rationale and high quality evidence to support their use. Currently, the practice of CWI is based on anecdotal evidence and there are no clear guidelines of an optimal or clinically effective treatment protocol. It is also important to consider that CWI induces a degree of shock on the body; therefore, the potential for short and long term side effects must be fully elucidated. Continued disparity in the rationale for using CWI, coupled with vague guidelines for its use, mean that athletes could risk employing more extreme temperatures or longer immersion times before determining actual benefit or risk.

OBJECTIVES

To determine the effects of cold-water immersion (CWI) in the management (prevention and treatment) of muscle soreness after exercise. The following comparisons were made:

- Cold-water immersion versus no cold-water immersion or placebo
- Cold-water immersion versus contrast water immersion (alternate cold and warm-water immersion)
- Cold-water immersion versus warm-water immersion
- Cold-water immersion versus other interventions (including non water based interventions)
- Different durations or dosages of cold-water immersion

METHODS

Criteria for considering studies for this review

Types of studies

Any randomised and quasi-randomised (e.g. using date of birth, record number or alternation to allocate participants into groups) trials evaluating cold-water immersion (cryotherapy) for preventing and treating muscle soreness after exercise.

Types of participants

We focused on participants using cold-water immersion (CWI) after exercise. No restrictions were placed on age group, gender, type or level of exercise. Trials that included participants with localised injury, whether acute (e.g. sprains, strains, contusions), or due to repetitive strain (e.g. tendinopathy) were excluded. We anticipated that people with vascular problems, such as Raynaud's disease, who are contraindicated to cryotherapy would have been excluded from trials.

Types of interventions

One group in the trial must have comprised participants treated with CWI after exercise. CWI was defined as immersion in water at less than 15°C (Low 2000). No restrictions were made on the duration or frequency of immersions. Interventions described as 'plunge' or 'dip' immersions were included. Immersion depth could be to any level of the body, including isolated immersion of a single body part (e.g. arm or leg), provided that the immersed body part (muscle) had undertaken prior exercise.

Comparisons were made to interventions designed to prevent or treat delayed onset muscle soreness including: passive interventions (rest, placebo or no intervention), warm-water immersion (immersion in water at more than 15°C), contrast water immersion (alternating immersions in hot and cold water); cool down, stretching, massage, and compression garments. Studies comparing different durations or dosages of CWI were included. Trials where the same CWI protocol was used in both arms as a co-intervention were not included. Comparisons with pharmacological interventions were not included.

Types of outcome measures

We collected data for the following follow-up times: immediately and 24, 48, 72 and 96 hours post intervention.

Primary outcomes

- Pain (muscle soreness) or tenderness (pain on palpation)
- Subjective recovery (return to previous activities without signs or symptoms)

Secondary outcomes

- Objective measures of muscle strength or power
- Functional assessment measures
- Swelling
- Range of movement
- Biochemical markers (e.g. lactate dehydrogenase (LDH), creatine kinase (CK), creatine phosphokinase (CPK) or isoform muscle creatine kinase (CKmm), myoglobin, skeletal troponin I, interleukin-6 (IL-6), C-reactive protein (CRP), tumour necrosis factor alpha (TNF-alpha).
- Complications or adverse effects as reported by the individual trials

Search methods for identification of studies

Electronic searches

We searched the Cochrane Bone, Joint and Muscle Trauma Group Specialised Register (to February 2010), the Cochrane Central Register of Controlled Trials (*The Cochrane Library* 2010, Issue 1),

MEDLINE (1950 to February 2010), EMBASE (1988 to February 2010), Cumulative Index to Nursing and Allied Health (CINAHL) (1982 to February 2010), British Nursing Index and archive (BNI) (1985 to February 2010), and the Physiotherapy Evidence Database (PEDro) (1929 to February 2010).

A search update was performed to check the more recent literature in November 2011. These were run from January 2010 onwards for the Cochrane Bone, Joint and Muscle Trauma Group Specialised Register (to November 2011), CENTRAL (*The Cochrane Library* 2011, Issue 4), MEDLINE (2010 to November Week 3 2011), EMBASE (2010 to 2011 Week 46) and CINAHL (2010 to 28 November 2011).

In MEDLINE, the subject-specific search was combined with the Cochrane Highly Sensitive Search Strategy for identifying randomised trials in MEDLINE: sensitivity- and precision-maximizing version (Lefebvre 2009). This strategy was modified for use in other databases. In EMBASE, the subject-specific search was combined with the Scottish Intercollegiate Guidelines Network (SIGN) search filter for randomised controlled trials (*see* Appendix 1 for search strategies).

We also searched Current Controlled Trials and the WHO International Clinical Trials Registry for ongoing and recently completed trials (February 2010).

No language restrictions were applied.

Searching other resources

We searched the reference list of articles, and the table of contents of the following journals not registered as being handsearched by The Cochrane Collaboration (February 2010):

- Australian Journal of Science and Medicine in Sport (1998 to February 2010)
- British Journal of Sports Medicine (1964 to February 2010)
- Clinical Journal of Sport Medicine (1991 to February 2010)
- International Journal of Sports Medicine (2005 to February 2010)
- Journal of Applied Physiology (1948 to February 2010)
- Journal of Sports Medicine and Physical Fitness (1998 to February 2010)
- Journal of Sports Sciences (1985 to 1987; 1990 to 1991; 1994; 1996; 2000 to February 2010)
- Medicine and Science in Sports and Exercise (1980 to February 2010)
- Physical Therapy in Sport (2000 to 2002; 2007 to February 2010)

We also searched the conference proceedings of the following organisations:

- American College of Sports Medicine (1986 to February 2010) (in *Medicine and Science in Sports and Exercise*)
- American Physical Therapy Association (1980 to February 2010) (in *Physical Therapy*)
- British Association of Sport and Exercise Medicine (BASEM) (1964 to February 2010) (in *British Journal of Sports Medicine*)
- British Association of Sport and Exercise Sciences (BASES) (1964 to February 2010) (in *Journal of Sports Sciences*)
- World Confederation for Physical Therapy (2003, 2007) (CD ROM)

Between February 2010 and March 2010, we contacted experts in the field (identified by personal contacts, lead authors in published studies, World Wide Web searching) for relevant data in terms of published, unpublished or ongoing studies.

Data collection and analysis

Selection of studies

Two authors (CB, SMcD) independently selected trials for inclusion. The titles and abstracts of publications obtained by the search strategy were screened. All trials classified as relevant by either of the authors were retrieved. Based on the information within the full reports, we used a standardised form to select the trials eligible for inclusion in the review. Where possible, translation of non-English language studies was undertaken. If necessary, we contacted primary authors for clarification of study characteristics. Disagreement between the authors was resolved by consensus, or third party adjudication (GDB, TH).

Data extraction and management

Data were extracted independently by two review authors (CB, SMcD) using a customised form tested prior to use. This was used to extract relevant data on methodological issues, eligibility criteria, interventions (including detailed characteristics of the exercise protocols and cold-water immersions employed), comparisons and outcome measures. Any disagreement was resolved by consensus, or third party adjudication (GDB, TH). Primary authors were contacted to clarify any omitted data or study characteristics. To perform intention-to-treat analysis, data were extracted according to the original allocation groups, and losses to follow-up were noted where possible. There was no blinding to study author, institution or journal at this stage.

Assessment of risk of bias in included studies

The risk of bias was assessed by two authors independently (CB, GDB), using the tool described (and the criteria outlined) in the Cochrane Handbook (Higgins 2009). To minimise bias in the interpretation of this scale, two review authors (CMB, GDB) initially assessed a small sample of unrelated studies (not included in the current review); disparities in risk of bias judgements were reviewed and discussed, prior to evaluating any of the included studies.

Each study was graded for risk of bias in each of the following domains; sequence generation, allocation concealment, blinding, incomplete outcome data, selective outcome reporting. Two other sources of bias were considered based on the following questions: a) Was the exercise protocol clear and consistent between groups? b) Were no co-interventions used, or if present, were they standardised across groups? For each study, the domains were described as reported in the published study report (or if appropriate based on information from related protocols, published comments, or after discussion with the relevant authors) and judged by the review authors as to their risk of bias. They were assigned 'high risk' or 'low risk'. If insufficient detail of what happened in the study was reported, or if what happened in the study was known, but the risk of bias was unknown; or if an entry was not relevant to the study at hand (for example when the outcome being assessed was not measured in the study) then the risk of bias was deemed 'unclear' for that domain. Disagreements

between authors regarding the risk of bias for domains was resolved by consensus.

Differences in the details of the treatment intervention (e.g. duration of immersion, frequency of immersion, water temperature), and characteristics of participants (professional or highly trained athletes, amateur, sedentary) was regarded as a potential source of bias and addressed in the subgroup analysis.

Measures of treatment effect

For each study, risk ratios and 95% confidence intervals were calculated for dichotomous outcomes, and mean differences and 95% confidence intervals were calculated for continuous outcomes. For continuous outcomes that were pooled on different scales, standardised mean differences were used. We had planned to preferentially extract data based on changes from baseline (mean change scores); however, the majority of studies reported follow-up scores.

Unit of analysis issues

Some studies included multiple observations of the same outcome; therefore we extracted data at clinically relevant time points. When available this was: immediately after the intervention, and then at 24 hour intervals following exercise up to 96 hours. In studies using a randomised cross-over design, we aimed to undertake paired analysis when sufficient data were available; otherwise, data were analysed as if these studies used a parallel group design (Deeks 2008). Studies using within-participant designs where contralateral limbs acted as controls were excluded. Had any study used a cluster randomised design, for example by sports teams, data would have been adjusted for clustering.

Dealing with missing data

If necessary, original investigators were contacted and requests made for any missing data. Depending on the nature of the data, assumptions were made on whether the data were missing at random or missing not at random. Data missing at random were ignored, and we focused on the available data only. If data were deemed to be missing not at random, replacement values were not imputed; sensitivity analyses were not undertaken.

When standard deviations were missing from continuous data, studies were scanned for any other statistics (confidence intervals, standard errors, T values, P values, F values) that allow for their calculation. There were no outcomes where the majority of data were unavailable because of missing standard deviations.

Assessment of heterogeneity

Initially, the clinical diversity across studies was qualitatively assessed. If two or more studies were deemed to be clinically homogeneous in terms of participants' characteristics, intervention, comparison, and outcomes, then data were assessed for statistical heterogeneity using RevMan. The chi-squared (Chi^2) test in conjunction with the I^2 statistic was used. The significance for Chi^2 was set at P < 0.1 (Deeks 2008a). The I^2 statistic was used to quantify inconsistency using the following formula $I^2 = [(Q-df)/Q] \times 100\%$, where Q is the Chi^2 statistic and df its degrees of freedom. I^2 values greater than 50% were considered to represent substantial heterogeneity (Higgins 2003).

Assessment of reporting biases

Should sufficient trials become available in future, we plan to use funnel plots to assess for publication bias based on the effect estimates (horizontal scale) against standard error (on a reversed scale, vertical) using Review Manager, with continuous data represented as standardised mean differences, and dichotomous data represented as risk ratios on a logarithmic scale.

Data synthesis

In the event that there was no evidence of heterogeneity of effect (P > 0.1), a fixed-effect model was used for meta-analysis. In cases where there was evidence of statistical heterogeneity, we checked the results using a random-effects model. We also considered the causes for the heterogeneity in terms of the clinical characteristics and the magnitude and direction of effects.

Subgroup analysis and investigation of heterogeneity

We planned subgroup analysis according to the details of the treatment intervention (duration of immersion, frequency of immersion, water temperature), the type of exercise (normal sporting activity, laboratory controlled muscle damage), exercise intensity (duration, environmental temperature), and the participants' characteristics (professional or highly trained athletes, amateur, sedentary). An additional subgroup analysis was planned according to methodological quality (high risk versus low risk of bias) (Deeks 2008b). This was to take the form of stratified analyses (high risk versus low risk) or in a meta-regression. A 'low risk study' was a study scoring low risk in each of the risk of bias domains (excluding blinding of participants and personnel).

Sensitivity analysis

In the event that no studies could be classified as low risk, we considered but did not perform sensitivity analysis based on individual risk of bias domains. We also planned to undertake additional sensitivity analysis should any other study peculiarities become apparent during the review process.

RESULTS

Description of studies

Results of the search

The searches undertaken in February 2010 identified 58 potentially eligible studies (from 65 reports). No new studies were found after contacting experts in the area. A total of 17 studies were included, 38 were excluded and a further three await classification.

Subsequently, the search was updated to November 2011 to check for more recent publications. This resulted in the identification of four more potentially eligible trials (Ascensao 2011; Pournot 2011; Stacey 2010; Vaile 2011), which have been placed in Studies awaiting classification; and another publication (Rowsell 2011) of an already included trial (Rowsell 2009).

Included studies

The 17 included studies were all published between 1998 and 2009 from centres in Australia (Cassar 2010; Halson 2008; Ingram 2009; King 2009; Montgomery 2008; Rowsell 2009; Sellwood 2007a; Vaile 2008c), Japan (Yanagisawa 2003a; Yanagisawa 2003b), United Kingdom (Bailey 2007; Eston 1999; Goodall 2008; Jakeman 2009)

or other areas of Europe (Buchheit 2009; Skurvydas 2006), and the USA (Kuligowski 1998). All were published in English, and in peer reviewed journals. The majority were identified in MEDLINE, EMBASE or CENTRAL, with one (Cassar 2010) identified from searching conference proceedings.

All 17 studies were randomised controlled trials. Ten were parallel group trials (Bailey 2007; Eston 1999; Goodall 2008; Jakeman 2009; Kuligowski 1998; Montgomery 2008; Rowsell 2009; Sellwood 2007a; Yanagisawa 2003a; Yanagisawa 2003b). Six studies used a randomised cross-over design (Buchheit 2009; Cassar 2010; Halson 2008; Ingram 2009; King 2009; Skurvydas 2006) with the time between intervention arms being three (Halson 2008), five (King 2009), seven (Buchheit 2009; Cassar 2010) and 14 (Ingram 2009) days, with one study using an eight month period (Skurvydas 2006). One study (Vaile 2008c) used a mixed design whereby randomisation was undertaken for the three active treatment groups but not the passive treatment group; data were not used from the passive treatment group in this review. In Buchheit 2009, data for some outcomes were extracted from another report (Peiffer 2010) based on the same experiment and participants.

In total, there were 366 trial participants of which 19% were female; the individual studies reported mean ages between 16 and 29 years. Studies were generally small, with 72% using 20 participants or less. The largest study used 54 participants (Kuligowski 1998). All participants were male in 12 studies (Bailey 2007; Buchheit 2009; Cassar 2010; Goodall 2008; Halson 2008; Ingram 2009; Montgomery 2008; Rowsell 2009; Skurvydas 2006; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b). Three studies involved all female participants (Eston 1999; Jakeman 2009; King 2009), and for the remainder, the male:female ratio was 1:1 (Kuligowski 1998) or 1:2.6 (Sellwood 2007a). Details of included studies can be found in the Characteristics of included studies.

Details of exercise

Exercise type, duration and intensity varied across studies. In nine studies, the exercise was designed to produce delayed onset muscle soreness (DOMS) under laboratory controlled conditions; all involved multiple repetitions (50 to 100 repetitions) of resistance to lengthening (eccentric) (Goodall 2008; Jakeman 2009; Kuligowski 1998; Sellwood 2007a; Skurvydas 2006; Vaile 2008c), or alternative lengthening and shortening exercises (Eston 1999; Yanagisawa 2003a; Yanagisawa 2003b). Five studies (Eston 1999; Kuligowski 1998; Sellwood 2007a; Yanagisawa 2003a; Yanagisawa 2003b) targeted a single muscle group, with other studies targeting a number of related muscle groups (lower limb muscles) using a resistance (leg press) machine (Vaile 2008c) or plyometric exercises (repetitive jumping) (Goodall 2008; Jakeman 2009; Skurvydas 2006). Participants subjected to DOMS were described as untrained (Yanagisawa 2003a; Yanagisawa 2003b); healthy (Eston 1999; Jakeman 2009; Kuligowski 1998; Sellwood 2007a), or active (Goodall 2008; Skurvydas 2006). It is likely that the majority were unaccustomed to this type of exercise; only one study specified that participants had a resistance training history (Vaile 2008c).

The remaining eight studies employed running or cycling based exercise. A single bout of cycling was used in two studies; one was a short duration sprint (Buchheit 2009), and one was a 40 minute steady cycle (75% VO_2 max) plus a time trial (Halson 2008). Both were undertaken by trained cyclists using controlled laboratory conditions and consistent environmental conditions during cycling

(34°C to 35°C ambient temperature, 40% relative humidity). In a further study (Cassar 2010), a 30 minute cycling session was undertaken once per day, over a five day period, by trained cyclists at 70% $V0_2$ max.

Three studies (Bailey 2007; Ingram 2009; King 2009) involved running based with active or trained participants exercising for over an hour (maximum 90 minutes). In Bailey 2007, running was intermittent and based on an average intensity of 75% $V0_2$ max. In the other two studies (Ingram 2009; King 2009), the nature and intensity of the exercise was designed to simulate the running demands of a team sport, with one (Ingram 2009) finishing with an additional shuttle run to exhaustion.

In two studies (Montgomery 2008; Rowsell 2009), exercise involved competing in a team sport (basketball, soccer); in both cases trained/high performance participants played one game per day, over a three (Montgomery 2008) or four (Rowsell 2009) day tournament.

Details of cold-water immersion

All studies employed some form of CWI intervention after exercise. The most popular water immersion temperature was between 10°C and 15°C, which was used by just over 75% of studies; the remainder used lower temperatures of 9°C (King 2009) or 5°C (Sellwood 2007a; Yanagisawa 2003a; Yanagisawa 2003b). In 10 studies (Bailey 2007; Buchheit 2009; Eston 1999; Goodall 2008; Jakeman 2009; Kuligowski 1998; Montgomery 2008; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b), treatment involved continuous immersion for between 5 and 24 minutes; the average treatment duration across these studies was 12.6 minutes. The remaining studies (Halson 2008; Ingram 2009; King 2009; Rowsell 2009; Sellwood 2007a; Skurvydas 2006) undertook CWI in sets where participants got out of the water at pre-determined time points; treatment therefore consisted of: three (Halson 2008; Sellwood 2007a) to five (Rowsell 2009) sets of one minute immersions, two sets of five minute immersions (Ingram 2009; King 2009), or two sets of 15 minute immersions (Skurvydas 2006). Three studies (Bailey 2007; Kuligowski 1998; Rowsell 2009) reported that the water was periodically agitated during immersion.

In the majority of studies, CWI was undertaken to approximately the level of the waist (Bailey 2007; Cassar 2010; Goodall 2008; Ingram 2009; Jakeman 2009; King 2009; Sellwood 2007a; Skurvydas 2006), sternum (Buchheit 2009; Halson 2008; Montgomery 2008; Rowsell 2009) or shoulder (Vaile 2008c). In the remaining four studies, CWI was confined to the arm (Eston 1999; Kuligowski 1998) or lower leg muscles (Yanagisawa 2003a; Yanagisawa 2003b). The timing of initiating CWI after exercise was generally consistent across studies; initiated immediately after (Bailey 2007; Cassar 2010; Eston 1999; Goodall 2008; Ingram 2009; King 2009; Kuligowski 1998; Sellwood 2007a; Skurvydas 2006; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b), or within approximately 10 minutes (Jakeman 2009; Montgomery 2008) or 20 minutes (Halson 2008; Rowsell 2009) after finishing treatment. Seven studies undertook additional CWI interventions after completing a single exercise session: Eston 1999 (12, 24, 36, 48, 60 and 72 hours); Goodall 2008 (24, 48 and 72 hours); Ingram 2009 (24 hours); Kuligowski 1998 (24, 48 and 72 hours); Skurvydas 2006 (4, 8 and 24 hours); Vaile 2008c (24, 48 and 72 hours); and one CWI group of Yanagisawa 2003a (24 hours).

Details of comparisons

The 17 included studies were divided into five different groups: studies comparing CWI with passive intervention (no CWI or rest); studies comparing CWI with contrast immersion; studies comparing CWI with warm-water immersion; studies comparing CWI with active recovery; studies comparing CWI with compression garments; and studies comparing two different dosages of CWI. Six studies (Ingram 2009; King 2009; Kuligowski 1998; Montgomery 2008; Vaile 2008c; Yanagisawa 2003a) used more than one relevant treatment comparison and therefore appear in two different sections.

Cold-water immersion versus passive (no intervention/rest)

This was the most common comparison, which was made by 14 studies (Bailey 2007; Buchheit 2009; Cassar 2010; Eston 1999; Goodall 2008; Halson 2008; Ingram 2009; Jakeman 2009; King 2009; Kuligowski 1998; Montgomery 2008; Skurvydas 2006; Yanagisawa 2003a; Yanagisawa 2003b). CWI was compared with a passive intervention defined as either seated rest (Bailey 2007; Buchheit 2009; Cassar 2010; Goodall 2008; Halson 2008; Ingram 2009; Jakeman 2009; King 2009; Skurvydas 2006) or no intervention (Eston 1999; Kuligowski 1998; Yanagisawa 2003a; Yanagisawa 2003b).

Cold-water immersion versus contrast immersion

Five studies made this comparison (Cassar 2010; Ingram 2009; Kuligowski 1998; King 2009; Vaile 2008c). Contrast immersion involved alternate immersions in cold (between 8°C and 15°C) and warm-water (38°C to 45°C); one study (King 2009) used a hot shower. The overall duration of contrast treatment varied across groups: 12 minutes (2 minutes cold: 2 minutes hot x 3 sets (Ingram 2009); 14 minutes (1 minute cold: 1 minute hot x 7 sets) (Cassar 2010; Vaile 2008c); 15 minutes (1 minute cold: 2 minute hot x 5 sets) King 2009; or 24 minutes (3 minutes hot: 1 minute cold x 6 sets) (Kuligowski 1998).

Cold-water immersion versus warm-water immersion

Four studies used this comparison (Kuligowski 1998; Rowsell 2009; Sellwood 2007a; Vaile 2008c) but the details of warm-water immersion across studies. Three (Kuligowski 1998; Rowsell 2009; Vaile 2008c) used immersion in water between 34°C and 40°C, and one (Sellwood 2007a) used water temperatures of 24°C. The total duration of warm-water immersion was: 3 (Sellwood 2007a), 5 (Rowsell 2009), 14 (Vaile 2008c) and 24 minutes (Kuligowski 1998).

Cold-water immersion versus active recovery

One study (King 2009) compared CWI with an active recovery intervention, which involved 15 minutes of jogging at a predetermined and controlled speed; the exercise in this study was a single bout of netball related running.

Cold-water immersion versus compression

One study (Montgomery 2008) compared CWI with compression therapy over the course of a three day (one game per day) basketball tournament. Participants in the compression group wore full length compression garments (18 mm/Hg) post game and at night (18 hours), over the tournament; CWI involved a single immersion after each game.

Cold-water immersion versus cold-water immersion (different dosage)

One study (Yanagisawa 2003a) specifically compared two different treatment dosages of CWI. Both groups completed a CWI treatment immediately after exercise, and one group undertook an additional treatment 24 hours later.

Primary outcomes

Studies were selected to include at least one primary outcome. Pain was the most commonly reported outcome and 15 studies assessed muscle soreness using a Likert scale or a visual analogue scale (VAS). A five point scale was used in two trials (Yanagisawa 2003a; Yanagisawa 2003b), 10 or 11 point or 10 cm scales were used in 11 trials (Bailey 2007; Cassar 2010; Halson 2008; Ingram 2009; Jakeman 2009; King 2009; Montgomery 2008; Rowsell 2009; Sellwood 2007a; Skurvydas 2006; Vaile 2008c), a 12 cm VAS was used in Kuligowski 1998 and a 20 cm VAS was used in Goodall 2008). The written descriptors used at each end of the scale were specified in all but two studies (Ingram 2009; Rowsell 2009). Seven studies specified that pain was measured during self palpation of muscle (Bailey 2007) or during a functional movement associated with the exercised body part(s) (Goodall 2008; Jakeman 2009; Kuligowski 1998; Sellwood 2007a; Vaile 2008c; Yanagisawa 2003b). We anticipated that the remainder of studies were based on muscle soreness measured at rest.

Two studies (Eston 1999; Sellwood 2007a) measured pain on pressure (tenderness) using a hand held algometer device; one recorded the force in Newtons (N) at which discomfort turned to pain (Eston 1999) and the other measured pain levels on a VAS (10 cm) during application of a standard pressure of 6 lb/cm² (Sellwood 2007a). Only one study (Buchheit 2009) did not measure any component of pain.

Five studies (Buchheit 2009; Cassar 2010; Halson 2008; Montgomery 2008; Rowsell 2009) measured subjective recovery based on a VAS (10 points or 10 cm). In each study, the participants rated different correlates of subjective recovery which we put into the following subcategories: general fatigue (Cassar 2010; Halson 2008; Montgomery 2008; Rowsell 2009), general recovery (Buchheit 2009); or separate ratings of physical and mental recovery (Halson 2008; Rowsell 2009).

Secondary outcomes

There were a range of secondary outcomes reported. Strength was assessed by 10 studies. The majority used a dynamometer or isokinetic dynamometer (Bailey 2007; Buchheit 2009; Eston 1999; Goodall 2008; Jakeman 2009; Kuligowski 1998; Sellwood 2007a), with others using a cable tensiometer (Ingram 2009), force platform (Vaile 2008c), or an unspecified force measuring device (Skurvydas 2006). Two focused on concentric muscle strength (Buchheit 2009; Jakeman 2009) with the remainder measuring isometric strength at a specific mid-range joint angle.

Eight studies reported on power. In five studies, power was assessed by measuring vertical jump performance (centimetres (cm)) either with (King 2009; Rowsell 2009; Skurvydas 2006) or without a counter movement (Bailey 2007; Montgomery 2008). Sellwood 2007a used a related measure based on hopping distance (cm). Using a specialist device, Vaile 2008c measured the power produced in Watts (W) during a weighted squat jump, and Buchheit 2009 measured power output on a cycle ergometer.

Functional performance outcomes were reported by seven studies. Five were based on the time (in seconds) participants took to complete various exercise tasks; these were put into subcategories according to their nature or distance in metres (m) as follows: cycle sprint (Buchheit 2009; Cassar 2010), single running sprint (10 to 20 m) (Bailey 2007; King 2009), multiple running sprint (20 m x 10 to 12 sets) (Ingram 2009; Montgomery 2008; Rowsell 2009), and agility based or sports specific running (King 2009; Montgomery 2008).

Range of movement (ROM) was reported by four studies. Four studies measured active range of movement using a hand held goniometer; we focused on joint movements which tended to lengthen the exercised muscle group (Goodall 2008; Kuligowski 1998; Yanagisawa 2003a). Three reported total range of movement in degrees (Goodall 2008; Kuligowski 1998; Yanagisawa 2003a) and one (Montgomery 2008) reported range of movement at a range of joints based on the sit and reach flexibility test (cm).

Five studies reported swelling using lower limb girth measured with a tape measure (Eston 1999; Goodall 2008; Montgomery 2008; Sellwood 2007a; Vaile 2008c), with values reported in centimetres of limb girth.

Biochemical markers were reported in 12 studies (Bailey 2007; Cassar 2010; Eston 1999; Goodall 2008; Halson 2008; Ingram 2009; Jakeman 2009; Rowsell 2009; Sellwood 2007a; Skurvydas 2006; Vaile 2008c; Yanagisawa 2003a); laboratory procedures were well described with a number of different biomarker molecules assessed. These outcomes were divided into two subcategories: biomarkers of inflammation (IL-1b; IL-6; IL-10; CRP) and muscle damage (creatine kinase (CK); lactate dehydrogenase (LDH); myoglobin).

The majority of studies did not monitor adverse events or complications relating to the interventions. Two studies (Buchheit 2009; Halson 2008) measured core temperature changes associated with CWI; in both studies, participants' mean core temperature (extracted from graphs) did not drop below 37°C.

Missing data

In five included studies (Eston 1999; Halson 2008; Ingram 2009; Kuligowski 1998; Montgomery 2008), the information provided in the trial report was sufficient for analysis of between group differences using Review Manager. In three studies (Bailey 2007; Eston 1999; Jakeman 2009), group dispersion was based on standard errors, with values converted to standard deviations by two independent review authors. In 12 included studies, additional data were requested from study authors; six (Bailey 2007; Cassar 2010; Goodall 2008; Jakeman 2009; King 2009; Skurvydas 2006) were successfully contacted, with each providing the requested information. In five studies (Buchheit 2009; Rowsell 2009; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b), we were able to extract data from graphs; this was undertaken by two independent review authors. In one study (Sellwood 2007a) only median and interquartile range figures were available.

Follow-up

All studies undertook multiple follow-up observations for each outcome. Eleven studies (Bailey 2007; Buchheit 2009; Eston 1999; Goodall 2008; Halson 2008; Ingram 2009; Jakeman 2009; King 2009; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b) reported multiple follow-ups in the first few hours succeeding

the exercise (e.g. pre-exercise, post exercise, pre intervention and post intervention); in such cases, we considered outcomes reported immediately after completing the treatment intervention, to be most clinically relevant. In three studies (Skurvydas 2006; Yanagisawa 2003a; Yanagisawa 2003b) the timing of the first follow-up could not be configured from the trial report and data were extracted from the next clearly defined follow-up time. In such cases (Skurvydas 2006; Yanagisawa 2003a; Yanagisawa 2003b), additional follow-ups were reported within four hours of completing the intervention, which, for the purposes of comparison, was also defined as an 'immediate' follow-up. In all but two studies (Buchheit 2009; Halson 2008) follow-ups were undertaken at 24 hours, with further follow-ups repeated at 48 (Bailey 2007; Eston 1999; Goodall 2008; Ingram 2009; Jakeman 2009; Kuligowski 1998; Sellwood 2007a; Skurvydas 2006; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b), 72 (Eston 1999; Goodall 2008; Jakeman 2009; Sellwood 2007a; Skurvydas 2006; Vaile 2008c) or 96 hours (Goodall 2008; Jakeman 2009; Kuligowski 1998; Yanagisawa 2003a; Yanagisawa 2003b).

In three studies, competitive sports tournaments (Montgomery 2008; Rowsell 2009) or multiple cycling exercises (Cassar 2010) were undertaken over three to five days (one game or cycling bout per day); outcomes were generally reported before and after each game or cycling bout, and repeated for up to 24 hours after the last day of exercise. In all these studies, we considered the 'exercise' to be the entire tournament or days of exercise, and therefore only extracted outcome data collected in the post exercise time period (e.g. after tournament completion or after the last day of exercise).

Excluded studies

After appraisal of the full study reports, we excluded 38 studies. The majority were excluded because they did not use a relevant CWI intervention (n = 19). In a further two studies, the CWI was relevant but used in conjunction with another therapeutic intervention that was not controlled for. Two others were excluded because CWI did not involve the exercised body-part. Ten studies were excluded as they did not report a primary outcome measure (rating of perceived exertion and rating of thermal discomfort were not considered to be measures of subjective recovery). Two trials did not report any form of randomisation, and two used a contralateral body part as a control. One study did not use a relevant time point for follow-up. Detailed reasons for exclusion can be found in the Characteristics of excluded studies.

Studies awaiting classification

Two studies (Fowles 2003; Smith 2008) did not provide sufficient detail to determine their relevance for inclusion, despite contacting the corresponding authors. A further four studies (Ascensao 2011; Pournot 2011; Stacey 2010; Vaile 2011) were identified subsequently to the full search date. Further details of these trials are given in Characteristics of studies awaiting classification.

Risk of bias in included studies

Full details of the quality assessment are given in the Characteristics of included studies. All corresponding authors were contacted, and asked to provide any methodological details which were unclear or missing in the original trial reports. Our requests for information were open ended to avoid any bias resulting from leading questions. Just over 55% of authors responded. Unless an author specifically stated that they did not understand our question, we avoided making multiple requests for information. Risk of bias judgement was made by two independent authors, based on information from trial reports and author correspondence; the results are summarised in Figure 1 and Figure 2. The significance of the variations in risk of bias is unclear as studies could not be subgrouped by high and low risk of bias.

Figure 1. Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

Figure 2. Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.
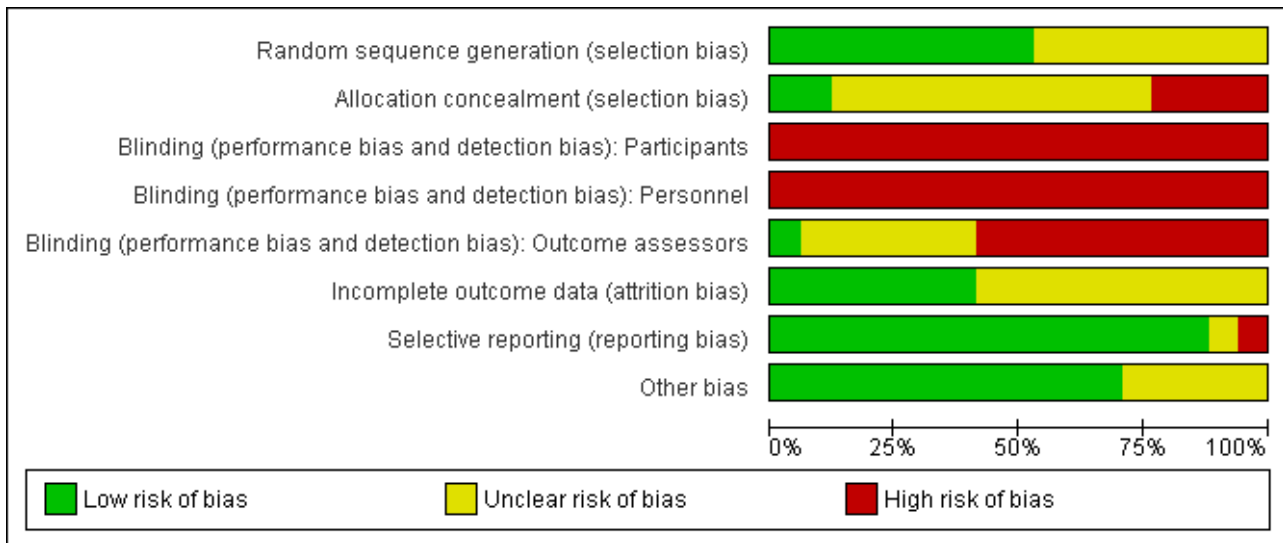
Allocation

Randomisation procedure was described in just nine studies. Sellwood 2007a used a random numbers table; with the remainder clarified after personal correspondence and were based on random numbers table (Bailey 2007; Cassar 2010; Goodall 2008), computer (Ingram 2009) or predetermined sequence generation (Jakeman 2009), coin toss (Buchheit 2009; Eston 1999), or hat draw (King 2009). Allocation concealment was adequately described in just one study (Sellwood 2007a) through the use of opaque sealed envelopes, and is likely in another (King 2009) based on personal correspondence. In the remaining studies, there was no clear indication that the investigators would be unable to predict the prospective group, or in the case of cross-overs, the order of treatments to which participants would be allocated.

Blinding

None of the studies utilised blinding of participants or care givers. One study (Sellwood 2007a) did not inform participants as to which intervention (5°C or 24°C water temperature) was deemed to be therapeutic; however, this was not considered to be true blinding. Only one study (Sellwood 2007a) used blinded outcome assessment. This study also states that participants were requested not to reveal their treatment group to the assessors; however, they did not appear to test the success of their blinding strategy.

Incomplete outcome data

This area of methodology was poorly described. Only one study (Sellwood 2007a) specifically indicated intention-to-treat analysis (with imputation in the event of missing data); however, their results seemed to indicate that there were no missing data or violation from the protocol. After correspondence, six trials (Bailey 2007; Goodall 2008; Ingram 2009; Jakeman 2009; King 2009; Skurvydas 2006) confirmed no losses to follow-up or violation from the study protocol. Six studies (Eston 1999; Kuligowski 1998; Montgomery 2008; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b) provided no information on drop outs, exclusion, missing data or approach to analysis. Four others (Buchheit 2009; Cassar 2010; Halson 2008; Rowsell 2009) were difficult to interpret in

this regard. Buchheit 2009 reported that one participant was dismissed because of incomplete outcomes. Halson 2008 recruited 11 participants who each completed both intervention arms; however, biochemical outcomes were missing for six participants as they were unwilling to provide blood samples. Rowsell 2009 was the only study not to specify the numbers allocated to each study arm; furthermore, just under one third of participants were excluded from the analysis due to injuries sustained at an unspecified time during the exercise protocol (competitive soccer tournament over four days).

Selective reporting

None of the studies made any reference to a published protocol. Buchheit 2009 did not report on two measured outcomes (strength and power) within the trial report, however, these data were available from a secondary publication (Peiffer 2010) based on the same study population and experiment. All studies described outcomes and follow-up times with corresponding results presented by intervention group. In five cases (Bailey 2007; Goodall 2008; Jakeman 2009; King 2009; Skurvydas 2006), additional group summary data were provided by corresponding authors in order to calculate effect size. Data were extracted from graphs in five studies (Buchheit 2009; Rowsell 2009; Vaile 2008c; Yanagisawa 2003a; Yanagisawa 2003b); in one of the studies (Yanagisawa 2003a), summary data from two time points (24 hours, 48 hours) could not be extracted for two outcomes (pain and biomarkers).

Other potential sources of bias

All studies provided in-depth descriptions of the exercise protocols based on exercise type, duration, and intensity. Seven studies (Bailey 2007; Cassar 2010; Goodall 2008; Ingram 2009; Kuligowski 1998; Yanagisawa 2003a; Yanagisawa 2003b) stated that participants were informed to refrain from using co-interventions for the duration of the study; in five cases these interventions were specified: massage (Bailey 2007; Kuligowski 1998), medications/supplements (Bailey 2007; Goodall 2008; Kuligowski 1998), exercise (Goodall 2008; Kuligowski 1998; Yanagisawa 2003a; Yanagisawa

2003b), stretching (Kuligowski 1998), or other physical modalities (Kuligowski 1998; Yanagisawa 2003a; Yanagisawa 2003b). Five studies (Bailey 2007; Halson 2008; King 2009; Montgomery 2008; Rowsell 2009) standardised specific co-interventions across groups: water consumption (Bailey 2007; Halson 2008; King 2009), carbohydrate/food ingestion (Halson 2008; King 2009; Montgomery 2008; Rowsell 2009) and stretching (Montgomery 2008; Rowsell 2009), with two monitoring adherence using participant diaries (Halson 2008; King 2009). Five studies did not provide any details on co-interventions (Buchheit 2009; Jakeman 2009; Sellwood 2007a; Skurvydas 2006; Vaile 2008c).

Effects of interventions

The 17 included studies were divided into five different groups based on comparison. Within each comparison, results are presented in subcategories based on follow-up time (immediate, 24, 48, 72, 96 hours). Although there were no complications or side effects reported within any of the individual studies, it was unclear whether any study actively monitored specific adverse effects as part of their outcomes.

Cold-water immersion versus passive (no intervention/rest)

Primary outcomes

Pain (muscle soreness: VAS, various scales or scores; highest values = worst pain)

Twelve out of the 14 studies in this comparison presented data on muscle soreness based on various visual analogue scores or scales. Pooled results are presented in five subcategories based on follow-up time (see Analysis 1.1; Figure 3). There were no significant differences between groups at immediate follow-up (SMD -0.07, 95% CI -0.43 to 0.28; 7 trials). At all four subsequent times, pooled results showed significantly lower levels of pain in the cold-water immersion group (24 hours: SMD -0.55, 95% CI -0.84 to -0.27, 10 trials); (48 hours: SMD -0.66, 95% CI -0.97 to -0.35; 8 trials); (72 hours: SMD -0.93, 95% CI -1.36 to -0.51; 4 trials); (96 hours: SMD -0.58; 95% CI -1.00 to -0.16; 5 trials). However, there was significant heterogeneity in all four analyses. While increasing the 95% confidence intervals, the findings in favour of CWI were upheld when applying the random-effects model (see Analysis 1.2). In the 24, 48 and 72 hours analyses, cross-over trials have been combined with parallel group trials. Subgroup analysis by study design show statistically significant differences between the pooled results of cross-over trials and parallel group trials at all three follow-up times. For 24 hours, *see* Analysis 1.3 (test for subgroup differences: $Chi^2 = 9.66$, df = 1 (P = 0.002), $I^2 = 89.7\%$); for 48 hours, *see* Analysis 1.4 (test for subgroup differences: $Chi^2 = 6.28$, df = 1 (P = 0.01), $I^2 = 84.1\%$); and for 72 hours (test for subgroup differences: $Chi^2 = 14.34$, df = 1 (P = 0.0002), $I^2 = 93.0\%$). It is clear from all three analyses that the findings in favour of CWI from the three cross-over trials (Ingram 2009; King 2009; Skurvydas 2006) are considerably stronger than those for the parallel group trials.

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Figure 3.   Forest plot of comparison: 1 Cold water immersion (CWI) versus passive, outcome: 1.1 Pain (muscle soreness: various scales Likert and VAS)
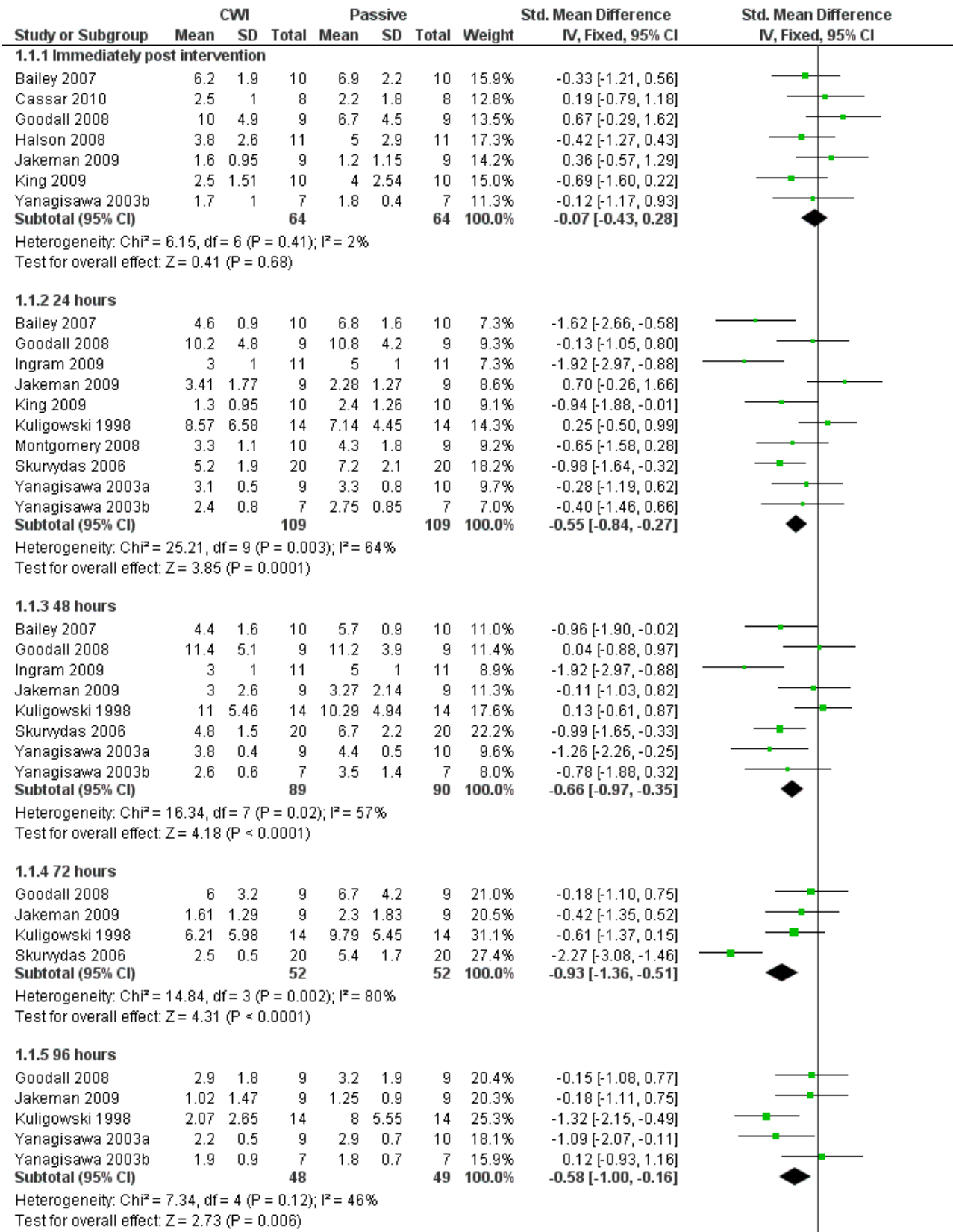
Figure 3. (Continued)

Heterogeneity. Chi² = 7.34, df = 4 (P = 0.12), I² = 46%
Test for overall effect: Z = 2.73 (P = 0.006)

-2 -1 0 1 2
Favours CWI Favours Passive
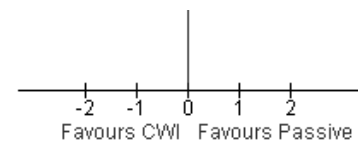
We had planned to perform subgroup analyses according to the specific details of the treatment intervention: e.g. duration of CWI, frequency of CWI, and water temperature. It was only possible to perform subgroup analysis by frequency of immersion (single immersion versus multiple immersions) which showed no statistically significant differences at 24 hours, *see* Analysis 1.6 (test for subgroup differences: Chi² = 0.10, df = 1 (P = 0.75), I² = 0%); and 48 hours, *see* Analysis 1.7 (test for subgroup differences: Chi² = 0.21, df = 1 (P = 0.64), I² = 0%). (Of note is that Yanagisawa 2003a, which compared single versus double CWI (also at 24 hours after exercise) also found no difference between these two groups in muscle soreness at 24 and 48 hour follow-up.)

Further subgroup analysis based on the type of exercise undertaken showed significant differences at 24 hours, *see* Analysis 1.8 (test for subgroup differences: Chi² = 10.82, df = 1 (P = 0.001), I² = 90.8%); and 48 hours, *see* Analysis 1.9 (test for subgroup differences: Chi² = 5.19, df = 1 (P = 0.02), I² = 80.7%). These suggest that effects were stronger when CWI was used after running based exercises (Bailey 2007; Ingram 2009; King 2009; Montgomery 2008) rather than after resistance exercises under laboratory controlled conditions (Goodall 2008; Jakeman 2009; Kuligowski 1998; Skurvydas 2006; Yanagisawa 2003a; Yanagisawa 2003b). However, it should be noted that there was clear heterogeneity in each subgroup as well as the numbers of participants in each subgroup being small.

The majority of participants in the DOMS studies were untrained, and those in the running/cycling studies were athletic or were specifically trained (Ingram 2009; King 2009; Montgomery 2008; Rowsell 2009); we therefore did not perform the planned subgroup analysis on trained versus untrained participants, as it was likely to replicate the subgroup analysis by type of exercise.

Given that all studies were at high risk of bias, we were unable to perform our planned subgroup analysis based on risk of bias.

Pain (Tenderness: algometer)

The one study (Eston 1999) reporting this outcome found no differences between groups at 24, 48 and 72 hour follow-ups (*see* Analysis 1.10).

Subjective recovery (10 point or 10 cm VAS)

Four studies measured different components of subjective recovery at immediate (Buchheit 2009; Cassar 2010; Halson 2008) or 24 hour (Montgomery 2008) follow-up (*see* Analysis 1.11). Two studies (Buchheit 2009; Halson 2008) found that cold-water immersion tended to result in an immediate improvement in ratings of physical recovery after cycling in hot and humid conditions (MD 0.97 units, 95% CI -0.10 to 2.05). However, the results of the two trials are heterogeneous, possibly in part reflecting that Buchheit 2009 involved a short duration cycle (1 km) and Halson 2008, a longer duration cycle (20 minutes). At the same follow-up (immediate),

Halson 2008 found no significant difference in mental recovery (MD 0.60 units, 95% CI -0.86 to 2.06).

Three studies reported subjective rating of fatigue (*see* Analysis 1.12). At immediate follow-up, the pooled results from two studies (Cassar 2010; Halson 2008) found significantly lower ratings of fatigue in the cold-water immersion group (MD -1.70 units, 95% CI -2.49 to -0.90). Montgomery 2008 reported lower, but not significantly lower, levels of fatigue in favour of cold-water immersion, 24 hours after competing in a three day basketball tournament (three interventions undertaken after each match: MD -0.70 units, 95% CI -1.88 to 0.48).

Secondary outcomes

Strength

Studies were placed in two analysis according to the type of continuous data extracted (final outcome value or percentage of baseline (pre-intervention) value) and subgrouped by follow-up time.

Strength (Final outcome values: Newton metres (Nm), N or kilograms (kg))

Maximal strength was reported by five studies (Bailey 2007; Buchheit 2009; Eston 1999; Jakeman 2009; Kuligowski 1998) based on final outcome values, and reported over a range of time points post intervention. There was some variation in the measurement device, contraction type and body part tested. Pooled results are displayed in Analysis 1.13. These tended to favour the passive group but showed no significant differences between groups aside from that at immediate follow-up. However, the lack of robustness of the significant finding for immediate follow-up using the fixed-effect model (MD -19.92 Nm, 95% CI -33.24 to -6.59) is demonstrated by the non-significant effect when using the random-effects model (MD -11.50 Nm, 95% CI -44.17 to 21.17; analysis not shown), reflecting the highly significant heterogeneity.

Strength (% of baseline)

Two studies (Goodall 2008; Skurvydas 2006) reported isometric quadriceps strength at various follow-up times, based on the % of the baseline value (baseline = 100%) (*see* Analysis 1.14). Goodall 2008, which used an isokinetic dynamometer, found no significant differences between groups. The cross-over study (Skurvydas 2006), which did not specify what recording device had been used, found significant differences in favour of cold-water immersion at 24 hours (MD 22.70%, 95% CI 10.66 to 34.74), 48 hours (MD 27.20%, 95% CI 15.73 to 38.67) and 72 hours (MD 25.20%, 95% CI 13.07 to 37.33) follow-up.

Power

Six studies assessed lower limb power, again results were placed in sub-categories according to the type of continuous data extracted

[Final outcome value or percentage (%) decrement in power performance], measurement device, and follow-up time.

Power (jumping performance: final outcome data in centimetres)

At 24 hour follow-up, pooled results from three studies (Bailey 2007; Montgomery 2008; Rowsell 2009) measuring maximal jump height (using a timing mat or similar measuring device) (see Analysis 1.15) had low heterogeneity ($I^2 = 0\%$), and found no difference between groups (MD 0.26 cm, 95% CI -2.04 to 2.56). Only Montgomery 2008 undertook additional follow-up, but found no differences between groups at 48 hours. Skurvydas 2006, a cross-over trial, also measured jump height, and although final outcome values were presented as flight time in centimetres, these were estimated from force plate data. Skurvydas 2006 found no differences between groups immediately post intervention, but jump height was significantly higher in the cold-water immersion group at 24 hours (MD 5.40 cm, 95% CI 2.12 to 8.68), 48 hours (MD 9.60 cm, 95% CI 6.40 to 12.80) and 72 hours (MD 6.60 cm, 95% CI 3.16 to 10.04) follow-up.

Power (jumping performance: percentage decrement)

King 2009 assessed the percentage decrement in jumping performance during five consecutive jumps; this was significantly lower and in favour of cold-water immersion at 24 hours (MD -3.70%, 95% CI -7.17 to -0.23) (see Analysis 1.16).

Power (cycling ergometer: Watts)

Buchheit 2009 found no significant difference between groups in peak power output during a 1 km cycling test (see Analysis 1.17).

Functional performance

Seven studies (Bailey 2007; Buchheit 2009; Cassar 2010; Ingram 2009; King 2009; Montgomery 2008; Rowsell 2009) reported this outcome. Of these, five were based on the time taken to complete a specific performance task over a range of follow-up times (see Analysis 1.18). Results, which are presented as subcategories based on the nature of the performance task and follow-up time showed no significant differences between group for the various measures of functional performance. The largest effect was reported by Cassar 2010, where there was less percentage decrease in performance (from baseline) in the time to volitional fatigue during a cycling task in the cold-water immersion group (MD -13.10%, 95% CI -27.13 to 0.93) at immediate follow-up (see Analysis 1.19).

Range of movement

Four studies measured range of movement, three (Goodall 2008; Kuligowski 1998; Yanagisawa 2003a) of which measured active joint range (in degrees) using a hand held goniometer, based on a movement which lengthened the exercised (primary) muscle group. Studies tested range of movement at various body parts: knee (Goodall 2008), elbow (Kuligowski 1998), ankle (Yanagisawa 2003a), or lumbar flexion (Montgomery 2008), which was based on a sit and reach flexibility test (centimetres). All studies presented final value outcomes. Out of the four trials, only Yanagisawa 2003a reported significant between-group differences, finding higher levels of ankle range of movement in favour of the cold-water immersion group at immediate, 24, 48 and 96 hours follow-up (see Analysis 1.20).

Swelling

Two studies (Eston 1999; Goodall 2008) measured thigh girth using a measuring tape, presenting final value outcomes in centimetres. The pooled results showed no significant differences between groups at 24, 48 or 72 hours follow-up (see Analysis 1.21).

Biomarker (muscle damage)

Creatine kinase (CK) was the most commonly reported biomarker. Pooled results found no significant difference between groups at any follow-up: immediate, 24, 48, 72 and 96 hours follow-up (see Analysis 1.22). Removal of Skurvydas 2006 from the 24 and 48 hour plots, reduces the heterogeneity (respectively, $I^2 = 82\%$ and 84%) to zero.

Results from two studies found no differences in lactate dehydrogenase (LDH) at immediate (Cassar 2010) or at 24, 48 and 96 hours follow-up (Yanagisawa 2003a) (see Analysis 1.23). Data from Yanagisawa 2003a are only presented for 96 hours as we were unable to extract accurate data from graphs at the earlier follow-up points.

Two studies (Bailey 2007; Rowsell 2009) reported on myoglobin levels. While tending to favour cold-water immersion, none of the results for myoglobin levels at immediate, 24 and 48 hours follow-up were statistically significant (see Analysis 1.24).

Biomarker (inflammatory)

There were similar levels of interleukin-6 (IL-6) (Halson 2008) and C-reactive protein (CRP) (Cassar 2010; Halson 2008) between groups in the immediate follow-up (see Analysis 1.25 and Analysis 1.26). Ingram 2009 reported similar patterns with CRP based on follow-up at 24 and 48 hours.

Cold-water immersion versus contrast immersion

Primary outcomes

Pain (muscle soreness: VAS, 10 points or 10 or 12 cm VAS)

Five studies reported this outcome based on a 10 point or 10 or 12 cm visual analogue scale with either unbearable (Kuligowski 1998) or extremely sore (Ingram 2009; King 2009; Vaile 2008c) used as the 'worst' pain descriptor. None of the results at five follow-up times showed a significant difference between the two groups (see Analysis 2.1; random-effects applied due to significant heterogeneity). Only Ingram 2009 showed a significant effect in favour of cold-water immersion at 24 and 48 hours; it should be noted that results of this trial were reported to no decimal places.

Subjective recovery (10 point visual analogue scale)

One study (Cassar 2010) found no differences between groups at immediate follow-up (see Analysis 2.2).

Secondary outcomes

Strength

Three studies (Ingram 2009; Kuligowski 1998; Vaile 2008c) reported strength. The details of the measuring device, and joint movements tested were different across studies. None of the pooled or individual trial results showed significant differences between the two groups at the five follow-up times (see Analysis 2.3).

Power

Two studies measured power (King 2009; Vaile 2008c). Vaile 2008c found no between group differences in jump squat performance at any follow-up (immediate, 24, 48, 72 hours) (see Analysis 2.4). King 2009 found no significant difference between groups in the % decrement in jumping performance (over five consecutive jumps) at 24 hours (MD -3.70%, 95% CI -7.67 to 0.27; see Analysis 2.5).

Functional performance

Time to complete running task in seconds

King 2009 recorded similar between-group running agility times (MD 0.47 seconds, 95% CI -1.69 to 2.63) and running sprint times (MD 0.05 seconds, 95% CI -0.11 to 0.21) at 24 hours (see Analysis 2.6). Ingram 2009 also found that multi-sprint times were similar between groups at 48 hours (MD 0.42 seconds, 95% CI -0.93 to 1.77; see Analysis 2.6).

Time to fatigue

Cassar 2010 found no significant difference between the groups in the percentage decrement in performance (MD 4.20% less decrement in performance, 95% CI -2.53 to 10.93) at immediate follow-up (see Analysis 2.7).

Swelling

This was reported by one study (Vaile 2008c) with no between-group differences found (see Analysis 2.8).

Range of movement

This was reported by one study (Kuligowski 1998) with no between-group differences found (see Analysis 2.9).

Biomarker (muscle damage)

Data for these outcomes are shown in Analysis 2.10 (creatine kinase), Analysis 2.11 (myoglobin) and Analysis 2.12 (lactate dehydrogenase). Reporting for immediate follow-up only, Cassar 2010 found significantly lower levels of creatine kinase but higher levels of myoglobin and lactate dehydrogenase in the CWI group. Ingram 2009 found no difference between the two groups in creatine kinase at 24 and 48 hour follow-ups. The results for Vaile 2008c favoured CWI for all three biomarkers, but the differences between the two groups were only statistically significant for lactate dehydrogenase at 24, 48 and 72 hours (see Analysis 2.12).

Biomarker (inflammatory)

Two trials (Cassar 2010; Vaile 2008c) found no significant differences between groups in inflammatory biomarkers (C-reactive protein and interleukin-6) at various follow-ups (see Analysis 2.13 and Analysis 2.14).

Cold-water immersion versus warm-water immersion

Four studies made this comparison (Kuligowski 1998; Rowsell 2009; Sellwood 2007a; Vaile 2008c).

Primary outcomes

Pain (muscle soreness: 10 point or 10 or 12 cm VAS)

There were no significant between-group differences in muscle soreness at immediate (Vaile 2008c), 24 hours (Kuligowski 1998; Rowsell 2009; Vaile 2008c), 48 hours (Kuligowski 1998; Vaile 2008c) or 72 hours (Kuligowski 1998; Vaile 2008c) follow-up (see Analysis

3.1). One study (Kuligowski 1998) found significantly lower levels of pain in favour of CWI at 96 hours (MD -4.64 points, 95% CI -7.72 to -1.56 points on a 12 point scale) (see Analysis 3.1).

Sellwood 2007a recorded various pain outcomes based on: muscle soreness during five different activities (sit to stand, passive stretch, hopping running, isometric contraction) and muscle tenderness at two regions of the thigh at 24, 48 and 72 hours follow-up. The only significant between-group differences were higher levels of muscle soreness in the cold-water immersion group during sit to stand at 24 hours; this amounted to a between-group difference in medians of 6 mm based on a 100 mm VAS.

Subjective recovery

Rowsell 2009 measured two different components of subjective recovery (see Analysis 3.2). At 24 hours, they found no difference between groups using a 10 point VAS for levels of fatigue (MD -0.80 points, 95% CI -3.25 to 1.65). However, also at 24 hours, six out of seven participants of the warm-water immersion group felt that the intervention had no benefit for recovery, compared to one out of seven participants of the cold-water immersion group (risk ratio 0.09, 95% CI 0.01 to 1.30) (see Analysis 3.3).

Secondary outcomes

Strength

Individual and pooled results from Kuligowski 1998 and Vaile 2008c showed no significant differences between the two groups in strength at immediate (Vaile 2008c), 24 hour (both trials), 48 hour (both trials), 72 hour (both trials) and 96 hour (Kuligowski 1998) follow-ups (see Analysis 3.4). Sellwood 2007a also reported no significant differences between the two groups in strength at 24, 48 and 72 hours.

Power

There were no significant differences between groups in terms of counter movement jump height at 24 hours (Rowsell 2009), or peak force produced during a weighted squat jump test at immediate or 24, 48 or 72 hour follow-ups (Vaile 2008c) (see Analysis 3.5). Based on single leg hop distance, Sellwood 2007a also found no significant difference between groups at 24, 48 or 72 hours.

Functional performance

One study (Rowsell 2009) found no differences between the two groups in multi-sprint times at 24 hour follow-up (MD 0.28 seconds, 95% CI -1.55 to 2.11) (see Analysis 3.6).

Range of movement

Kuligowski 1998 found no differences between the two groups in active elbow range of movement at any follow-up (24, 48, 72, 96 hours) (see Analysis 3.7).

Swelling

Similarly both Vaile 2008c and Sellwood 2007a found no between-group differences in thigh girth at any follow-up (immediate, 24, 48, 72 hours) (see Analysis 3.8).

Biomarker (muscle damage)

Three studies (Rowsell 2009; Sellwood 2007a; Vaile 2008c) found similar levels of creatine kinase and myoglobin across groups (see Analysis 3.9, Analysis 3.11). Lactate dehydrogenase (LDH)

levels were also similar at immediate (Vaile 2008c) and 24 hours (Rowsell 2009). However, further follow-ups by Vaile 2008c found significantly lower levels of LDH after cold-water immersion at 24, 48 and 72 hours (see Analysis 3.10).

Biomarker (Inflammatory)

Interleukin-6 levels were similar in the two groups at immediate (Vaile 2008c) and 24 hour follow-up (Rowsell 2009; Vaile 2008c) (see Analysis 3.12). At 24 hours, Rowsell 2009 found that interleukin-10 levels were similar across groups (see Analysis 3.13), but found a tendency towards lower levels of interleukin-1b in the CWI group (see Analysis 3.14).

Cold-water immersion versus active recovery

King 2009, in a cross-over trial involving 10 netballers, found significantly lower levels of muscle soreness in favour of CWI at immediate follow-up (MD -2.65 units, 95% CI -4.05 to -1.25). Although pain levels were still lower in this group at 24 hours (MD -0.90 units, 95% CI -2.00 to 0.20), the difference between the two groups was not statistically significant (see Analysis 4.1).

King 2009 found a significantly smaller decrement in power (measured by the percentage decrement in jump height over five repetitions) in the CWI group at 24 hours (MD -3.30%, 95% CI -6.14% to -0.46%) (see Analysis 4.2). Also at 24 hour follow-up, King 2009 found no differences between groups in terms sprint time (MD -0.03 seconds, 95% CI -0.21 to 0.15) and time taken to complete a multi-sprint test with an agility component (MD -1.12 seconds, 95% CI -3.86 to 1.62) (see Analysis 4.3).

Cold-water immersion versus compression

Only Montgomery 2008 compared CWI with compression therapy (tights). There were no between group differences at 24 hours after completing a three match soccer tournament for any outcome: pain (muscle soreness) (see Analysis 5.1), subjective recovery (rating of fatigue) (see Analysis 5.2), power (vertical jump) (see Analysis 5.3), range of movement (sit and reach) (see Analysis 5.4) and two different functional performance tests (time to complete multi-sprint test and agility test) (see Analysis 5.5).

Cold-water immersion versus multiple cold-water immersion

Only one study (Yanagisawa 2003a) specifically compared two different treatment dosages of CWI. Both groups undertook CWI immediately after exercise; however, one group used an additional treatment 24 hours later. Summary data were reported from range of movement; however, data were extracted from graphs for the remaining outcomes.

Pain was significantly lower in the single immersion group at 96 hours (MD -0.80 units, 95% CI -1.36 to -0.24 points on a 5 point VAS), but there were no differences between groups in the earlier follow-ups (24 and 48 hours) (see Analysis 6.1). Range of movement (ankle dorsiflexion) was slightly higher in the single immersion group at all follow-ups; however, no differences were significant (see Analysis 6.2). Values for both biomarker outcomes (creatine kinase and lactate dehydrogenase) were very similar across groups at 24 and 48 hour follow-ups (we were unable to differentiate between groups on the graph). While both levels of both biomarkers were lower in the single immersion group after 96 hours, there were no statistically significant differences between groups (see Analysis 6.3 and Analysis 6.4).

DISCUSSION

Summary of main results

This review examined the effectiveness of cold-water immersion (CWI) for preventing and treating muscle soreness after exercise. Seventeen studies, involving a total of 366 (mostly young male) participants, were included. For the purposes of the review, we considered exercise to be laboratory induced delayed onset muscle soreness (DOMS) models, running, cycling or team sport activities. We included studies using randomised controlled or randomised cross-over designs. CWI was generally undertaken immediately after exercise, but the temperatures, durations and frequency of CWI varied across studies. Studies were divided into five clearly distinguishable comparison groups; where available, data were extracted at five short term follow-up points: immediate, 24, 48, 72 and 96 hours. Muscle soreness and subjective recovery were the primary outcomes. In many cases it was not possible to draw definite well founded conclusions from the comparisons, due to lack of statistical significance and poor methodological quality. The largest volume of data were extracted for the comparison CWI versus passive intervention (specified as rest or no CWI), which was based on 14 studies.

Cold-water immersion versus passive (no intervention/rest)

Cold-water immersion resulted in significantly greater improvements in muscle soreness. Pooling of muscle soreness data found a significant effect in favour of CWI at four time points (24, 48, 72 and 96 hours); these findings were upheld when a random-effects model was applied. Pooled results from cross-over trials were more favourable for CWI than pooled results from parallel group trials. Subjective recovery and fatigue data were also in favour of CWI; however, these were based on results from just four studies, and mainly at immediate follow-up. Pooled results from two studies found cold-water immersion groups had significantly lower ratings of fatigue, and potentially improved ratings of physical recovery.

The Minimal Important Difference (MID) has been defined as "the smallest difference in score in the domain of interest that patients perceive as important, either beneficial or harmful, and which would lead the clinician to consider a change in the patient's management" (Guyatt 2002). The MID for pain reduction in musculoskeletal conditions of the shoulder has been estimated at 1.4 cm, based on a 10 cm visual analogue scale (Tashjian 2009). One study (Sellwood 2007a) in the current review detailed a sample size calculation which was based on a MID of 25% pain reduction after CWI. The significant findings for muscle soreness reported in favour of CWI (over no intervention/rest) in the current review for 24, 48 and 72 hours all include this MID as shown in Analysis 1.27, where the results for all the trials have been adjusted to fit the same 10 cm VAS (24 hours: MD -1.27 cm, 95% CI -1.70 to -0.84; 48 hours: MD -1.58 cm, 95% CI -2.07 to -1.10; 72 hours: MD -2.16 cm, 95% CI -2.79 to -1.53). While Analysis 1.27 should be considered an exploratory analysis, it illustrates that the statistical significant differences are likely to be clinical relevant also. The clinical relevance of such differences may be dependent also on numerous factors such as the time of the outcome, the capacity for natural recovery after DOMS, and the time and costs associated with CWI. We suggest that a 13% to 22% difference in muscle soreness would be important for many athletes, particularly those in an elite sporting environment.

Due to clinical and methodological diversity, it was not possible to perform extensive pooling of secondary outcome data from studies comparing CWI and passive. In particular, there were variable methods used for reporting similar outcomes across studies. In terms of isokinetic strength, there was conflicting evidence from three studies at the immediate follow-up (Bailey 2007; Buchheit 2009; Jakeman 2009); this is likely to relate to differences in the type of exercise, body part and measurement device employed. A single study (Skurvydas 2006) reported significant differences in favour of CWI for isometric strength at all follow-ups, but provided no details on the measurement device. Pooled data from three studies (Bailey 2007; Montgomery 2008; Rowsell 2009) found no differences in jump height (muscular power) at each follow-up point. This conflicted with the results of a single study (Skurvydas 2006) which reported effects in favour of CWI; this study was not pooled based on differences in exercise type, and measurement device. No differences were reported based on muscular power measured on a cycle ergometer; and although one study (King 2009) found significantly less decrement in jumping performance in favour of CWI, this may not be a clinically important change (MD of 3%). Three out of four studies found no differences in range of movement, with one study (Yanagisawa 2003a) reporting significant differences in favour of CWI at four follow-up points. There were no differences between CWI and rest in terms of swelling, functional performance and a range of biomarker outcomes.

Cold-water immersion versus warm-water immersion or contrast immersion or active recovery or compression

Despite a large number of treatment comparisons, there was little evidence for a superior treatment intervention. Pooled analyses found no differences in terms of muscle soreness between CWI versus contrast immersion, or CWI versus warm-water immersion. In accordance, a single study (Sellwood 2007a), which had a low risk of bias, reported no differences in muscle soreness between CWI versus warm-water immersion. Kuligowski 1998 found a large effect (MD 4.6 points on a 12 cm VAS) in favour of CWI over warm-water immersion, at 96 hour follow-up.

There were very few differences in any of the reported secondary outcomes, when CWI was compared with contrast immersion, warm-water immersion, active recovery or compression tights. One study (King 2009) found a significant effect in favour of active recovery based on a single secondary outcome; but this result was of questionable clinical relevance (MD of 3.3% in muscular power).

Overall completeness and applicability of evidence

We located 17 studies with a total of 366 participants; substantially fewer participants were available for pooling. The majority of participants were young (mean ages between 16 and 29 years), and only 19% were female. Although this probably reflects the current CWI user population in terms of age, we had expected larger numbers of female participants within this review.

We could not find any reports of adverse events or complications in the included studies. One study (Halson 2008) specified that short duration immersion (three minutes) in 11.5°C water temperatures, after exercise in hot conditions, does not alter core temperature to dangerous or hypothermic levels. However, the remainder of studies did not undertake active surveillance for pre-defined adverse events; subsequently the short and long term safety of CWI and comparison interventions remains unknown.

The type of exercise undertaken prior to CWI varied across studies. Eight studies involved trained athletes who used CWI after familiar bouts of cycling or running based exercise. The remaining nine studies used resistance exercises under laboratory controlled conditions to produce DOMS, largely within an untrained population. A subgroup analysis indicated that CWI had a stronger effect on muscle soreness after running based exercises, and was less effective when used after resistance exercises (DOMS models). The significance of these results is difficult to ascertain based on the high risk of bias, the heterogeneity of the trials within each subgroup and the small number of studies and participants within each subgroup. A key difference is that resistance exercises under laboratory controlled conditions are associated with high levels of muscle damage and pain, particularly in an untrained population. In contrast, trained athletes undertaking running based exercises have lower risk of muscle damage and generally have a higher capacity for recovery after exercise.

Trained or elite athletes are most likely to use CWI regularly. Although DOMS is most prevalent in novice athletes, we acknowledge that it may also occur in elite sport: e.g. at the beginning of the sporting season (when returning to training following a period of reduced activity), or after the introduction of a new movement or training method. Rehabilitative eccentric training (e.g. in the management of tendinopathy) may also be associated with DOMS; however, our findings are less applicable to this setting.

It is thought that CWI can reduce muscle soreness through a number of mechanisms including: intracellular-intravascular fluid shifts, reduction of muscle oedema, increased cardiac output (without increasing energy expenditure), enhanced blood flow, nutrient and waste transportation, with an additional psychological benefit (Wilcock 2006). The potential for inducing each of these effects may relate to the type of CWI undertaken. The details of CWI varied in the current review, with water temperatures and immersion time ranging between 5°C and 15°C, and from 3 to 24 minutes respectively. It is not clear whether such variations in treatment parameters affect clinical outcomes. Interestingly, our subgroup analysis found no differences between single and multiple cold-water immersions based on muscle soreness outcome. Further research will ascertain if different dosing schedules are more effective or if there is a specific time frame after exercise when CWI is most effective.

We must also consider that other mechanisms of effect for CWI may exist. Immersion in ice or cold-water is current best practice for managing exertional hyperthermia (McDermott 2009; Smith 2005). CWI may also be of value in cases of mild hyperthermia, as reducing core temperature after exercise should minimise perceived fatigue, enhancing subsequent exercise performance. Three studies (Buchheit 2009; Halson 2008; Rowsell 2009) in the current review, used CWI after exercise in hot conditions (35°C and 40% humidity); however, none reported large effects. We noted that their maximum duration of CWI was five minutes and suggest that longer periods may be required to achieve a clinically relevant effect.

CWI is generally used to enhance recovery after exercise. Reducing muscle soreness is considered to be an important component of recovery, and as such was the most reported primary outcome in this review. Restoration of muscle strength and power, range of movement, and functional performance are also considered

to be important; these were included as secondary outcomes. However, we found few significant differences in these outcomes. We acknowledge that our range of secondary outcomes was not exhaustive and the effect of CWI on full system recovery was not fully addressed. There is evidence also from other systematic reviews (Bleakley 2010) that in certain clinical circumstances, CWI can induce significant biochemical effects including oxidative stress and a potential increase in free radical species formation.

Quality of the evidence

The study quality in this review was low. The majority of studies had a high risk of bias making the validity of most of the results uncertain. The sample size of included studies was also consistently small, raising questions as to the power of individual trials.

We were unable to meaningfully subgroup studies into high and low quality. Just over half of the studies used adequate sequence generation; failure to do this can overestimate treatment effects (Kunz 2007). Concealment of group allocation is also important in this regard (Pildal 2007), particularly when dealing with subjective outcomes (Wood 2008), but was undertaken in just one study (Sellwood 2007a). Blinding was also poorly performed. We acknowledge that, based on the nature of CWI and the comparison interventions, stringent blinding of participants and caregivers is virtually impossible. However, blinding of assessors for most of the outcomes is feasible but was reported in just one study (Sellwood 2007a). The primary outcomes in this review were patient reported. However, a lack of assessor blinding is likely to implicate the secondary outcomes which were mainly objective measures (e.g. strength, swelling, range of movement). Few studies adequately described their approach to analysis; in many cases there was no detail on drop-outs, exclusions, missing data or follow-up points, and we were unable to consider intention-to-treat analysis. Previous Cochrane reviews (Bennett 2005; Herbert 2011) dealing with DOMS studies also found that often missing data were inadequately dealt with, and that few studies indicated intention-to-treat analysis.

Muscle soreness resulting from exercise is usually short lived and self limiting. Cross-over designs are therefore popular in this area of research, with seven included in the current review (however, the cross-over component of Vaile 2008c was not relevant to this review). Cross-over designs can risk certain carry-over effects between treatment periods, which are not present in parallel group designs. Perhaps the most likely source of carry-over is insufficient recovery from the first exercise bout, particularly when studies induce DOMS on an untrained population. This carry-over was minimised in the current review as only one of the cross-over studies in this review used a laboratory based DOMS model and they allowed several months between treatment periods. The remainder of cross-over studies used a shorter time (between 3 and 14 days) between treatment periods; however, it is possible that these times were also appropriate as they used trained individuals completing familiar cycling or running exercises, who were likely to recover faster.

Potential biases in the review process

In this review, we undertook an extensive search of electronic databases and grey literature sources; 17 studies were included, one of which was unpublished. Although we think this was an adequate selection, it is difficult to fully exclude publication bias, study identification bias and study selection bias. We also acknowledge that additional research studies have been undertaken and published since our searches were completed in February 2010. An exploratory search update carried out in November 2011 resulted in the identification of four potentially eligible trials. Based on information provided in the abstracts, two trials (Ascensao 2011; Vaile 2011), which involved a total of 30 participants, are likely to be included in a future update, while the other two trials (Pournot 2011; Stacey 2010), involving a total of 50 participants, may be excluded. Given this, we anticipate that the results of the current review are unlikely to be substantially affected by the inclusion of data from these trials.

Missing data within this review include, in particular, missing outcomes and summary data. As none of the included studies had a registered protocol, bias from selective reporting of results was difficult to fully ascertain. Although we made a concerted attempt to retrieve missing summary data, we were unable to contact some of the authors. To minimise the missing summary data, some of the data presented in the review graphs were estimated from graphs published in the trial reports. While clearly unsatisfactory, this was the best we could do in the absence of information from the investigators. To minimise error, estimations were undertaken by two independent reviewers, with inconsistencies checked through reviewer consensus and a third party.

Six of the included studies used a randomised cross-over design. We were unable to perform any paired analysis, and data were analysed as if these studies used a parallel group design. Cross-over studies were also combined with parallel group trials in the same meta-analysis. Although this approach gives rise to bias through unit of analysis error, this is expected to be conservative as cross-over studies analysed in this way tend to be under-weighted (Deeks 2008). In contrast, our subgroup analysis showed that cross-over designs had a more positive effect on the primary outcome (muscle soreness), compared with parallel group trials. The reasons for this are not clear; however, it may indicate that parallel studies represent the best methodological design for future research in this area.

Three studies used a unique approach based on three to four days of exercising with daily interventions undertaken after each exercise bout. This produced multiple outcome data over a number of days. Within this review, we only focused on outcomes reported after the final exercise session. This provided an indication of the cumulative effect of using CWI throughout the course of the entire three to four day tournament. As we did not extract data after each match, we may have overlooked important intermittent effects.

Finally, we consider that the practical relevance of some of the secondary outcome measures should be questioned and their results interpreted cautiously. Although there were few significant differences reported, many different types of tests were used and the clinical importance of each may be difficult to assess. In some cases (e.g. functional outcomes based on the time taken to complete an exercise test), the clinical significance of the results may depend on the type of exercise, or level of competition. Although we considered a large number of biochemical outcome markers, some were not included as they were not considered to be applicable to our review objectives. These decisions were made based on the current evidence base but may change in light of new evidence in this area.

Agreements and disagreements with other studies or reviews

Recent narrative reviews (Barnett 2006; Cheung 2003) of the best methods of recovery after exercise report little to no evidence for the use of massage, active recovery, contrast water immersion therapy, cryotherapy (including CWI), hyperbaric oxygen therapy, non-steroidal anti-inflammatory drugs, compression garments, stretching, electromyostimulation, combination modalities, homeopathy, ultrasound and electrical current modalities.

Two Cochrane reviews have examined respectively the effectiveness of stretching (Herbert 2011) and hyperbaric oxygen therapy (Bennett 2005) for treating muscle soreness after exercise. Both reviews found little evidence to support the use of each of the respective interventions. Similarly a systematic review (Hing 2008) of randomised controlled studies reported no evidence to support the use of contrast water immersion for aiding recovery after exercise, with the majority of included studies reported as being of limited methodological quality.

Cryotherapy remains a popular intervention for treating muscle strains, joint sprains and contusions. The rationale is somewhat similar to the post exercise environment, and various forms of cooling (including CWI) are used to relieve pain relief, decrease local tissue metabolism and inflammation, and redirect blood flow in the acute phases after injury. A systematic review of randomised controlled trials (Bleakley 2004) found limited evidence that various cooling interventions can induce analgesia after acute soft tissue injury and orthopaedic surgery, and found little to no effect on other important clinical outcomes.

AUTHORS' CONCLUSIONS

Implications for practice

Muscle soreness commonly results after sports and exercise activity. A priority for clinicians and athletes is to encourage full recovery between exercise bouts using an effective intervention programme. This review provides some evidence that cold-water immersion reduces delayed onset muscle soreness after exercise when compared with passive interventions involving rest or no intervention. However, it is not possible to draw definitive conclusions on pain or recovery because of the poor methodological quality, and small sample sizes. The majority of reviewed trials failed to undertake active surveillance of pre-defined adverse events relating to cold-water immersion. Given these findings and the self-limiting nature of most forms of muscle soreness resulting from exercise, there is currently only limited evidence for using cold-water immersion after exercise. There are no data available to determine the optimal method of cold-water immersion.

Implications for research

High quality, well reported research in this area is required. In particular, future studies should: incorporate a randomised controlled parallel group design with adequate sequence generation and allocation concealment; use appropriate sample sizes with power to detect expected differences; use cold-water immersion protocols based on a defined physiological rationale in accordance with the type of exercise and climatic environment; use effective and explicit blinding of outcome with appropriate and full reporting of data; and undertake active surveillance of pre-defined adverse events.

ACKNOWLEDGEMENTS

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* Indicates the major publication for the study

CHARACTERISTICS OF STUDIES

Characteristics of included studies *[ordered by study ID]*

Bailey 2007

Methods	Randomised controlled trial (matched by several anthropometric and physiological characteristics)
Participants	N = 20 males, mean age 22.3 (SD 3.3)
Interventions	CWI (n = 10): Seated immersion to iliac crest in 10°C water for 10 minutes, repeated water agitation
	Passive (n = 10): Seated rest for 10 minutes
Outcomes	**Pain** General whole body soreness (visual analogue scale, 1 "not sore" to 10 "very very sore") (follow-up: immediately, 24, 48 hours) **Strength** Maximal isometric knee extension / flexion (isokinetic dynamometer, % change from baseline) (follow-up: 24, and 48 post exercise) **Power** Vertical jump height (no counter movement) (electronic timing mat, peak height in m) (follow-up: immediately, 1, 24, and 48 hours) **Functional assessment** Sprint performance (15m sprint, photoelectric recording, seconds) (follow-up: 48 hours post exercise) **Biomarker** Muscle damage: Myoglobin (nmol/L), CK (U/L) (Follow-up: immediately, 24 and 48 hours)
Exercise type / intensity	90 minutes of intermittent running (average exercise intensity equal to 75% VO$_2$ max)
Time between completing exercise and initiating intervention	Immediately
Participants' fitness level	Active males, habitually undertaking a variety of sports, weekly number of exercise sessions: passive group: 4 (SD 1); CWI group: 5 (SD 2) VO$_2$ max (ml/kg/min) CWI: 55.2 (SD 4.8); Passive: 56.2 (SD 5.3)

Bailey 2007 *(Continued)*

	Refrained from exercise for at least 2 days prior to the study
	Unfamiliar with 90 minute intermittent running protocol
Notes	Data for pain, strength, power, functional assessment and biomarkers obtained after personal correspondence (converted from SE to SD)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomly allocated (Methods, cryotherapy treatment, pg. 1164)
		Random numbers table used for sequence generation (personal correspondence)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript
		No details on implementation, personal correspondence stated random numbers table only
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript
		No blinding of outcome assessors (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	No details on drop outs, exclusions, missing data or approach to analysis (per protocol / ITT) within the manuscript
		No drops outs or protocol deviations (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available
		Timing of outcome measures stated in methods for two out of five outcomes (pain and repeated sprint)
		No data for power and repeated sprint by intervention group (whole sample data only) within the manuscript, however this was provided after author correspondence
Other bias	Low risk	Exercise protocol clearly described and referenced
		Participants refrained from massage and anti-inflammatory drugs for the duration of the investigation; water consumption standardised

Buchheit 2009

Methods	Randomised cross-over (7 days between interventions)
Participants	N = 10 male cyclists, mean age: 29 (SD 6)

Buchheit 2009 *(Continued)*

Interventions	CWI (n = 10): Seated immersion to mid-sternal level in 14°C water for five minutes. Total recovery 20 minutes, resting in seated position when not in water bath. Passive (n = 10): Seated rest for 20 minutes
Outcomes	**Subjective recovery** Rating of recovery, visual analogue scale (0 'feeling not recovered' to 10 'feeling fully recovered') **Strength** Concentric knee extension (Isokinetic dynamometer, Nm) **Power** Peak power output during exercise test (1 km time trial, cycle ergometer, Watts) **Functional assessment** Time taken to complete exercise test (1 km time trial, cycle ergometer, seconds) [Follow-up for all outcomes: immediate]
Exercise type / intensity	Supramaximal cycling trial (1 km) undertaken at 35°C, 40% humidity
Time between completing exercise and initiating intervention	7.5 minutes
Participants' fitness level	Trained cyclists, mean VO_2 max 56.5 (SD 5.0) ml/kg/min
Notes	Strength and power outcomes were extracted from graphs in a second publication (Peiffer 2010)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	States 'randomly assigned' (pg. H422) Coin toss (personal correspondence)
Allocation concealment (selection bias)	High risk	No details in manuscript No details on implementation, personal correspondence stated random numbers table only
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript "No" (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No details on drop out or missing data within the manuscript N = 11 participants at the start of study, one was dismissed due to incomplete outcome, no details of how missing value was addressed (e.g. intention to treat issues) (personal correspondence)

Buchheit 2009 *(Continued)*

Selective reporting (re-porting bias)	High risk	Data from this trial is presented in a secondary publication (*see* Peiffer 2010)
Other bias	Unclear risk	Exercise protocol clearly described
		No details on co-interventions (immediate follow-up only so may be less associated with bias)

Cassar 2010

Methods	Randomised cross-over study (7 days between conditions)
Participants	N = 8 male cyclists, aged 22 to 35
Interventions	CWI (n = 8): Immersion to waist in water at 8 to 12°C for 14 minutes
	Contrast (n = 8): Immersion to waist in water at 8°C for 1 minute followed by immersion at 40 to 45°C for 1 minute; x 7 sets
	Passive (n = 8): Rest
Outcomes	**Subjective recovery** Rating of fatigue, visual analogue scale (0 to 10 cm) **Pain** Rating of soreness, visual analogue scale (0 to 10 cm) **Functional assessment** Time to fatigue during an exercise test (time maintaining 80 revolutions per minute, cycle ergometer, seconds) **Biomarkers** Muscle damage [CK (U/L), LDH (U/L), Myoglobin (mg/L)] Inflammation [CRP (mg/L)] [Follow-up for all outcomes: immediate]
Exercise type / intensity	30 minutes cycling (70% VO_2 max)
Time between completing exercise and initiating intervention	Immediate
Participants' fitness level	Trained cyclists
Notes	All data provided after personal correspondence

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Randomly assigned (abstract)
		Random numbers table (personal correspondence)

Allocation concealment (selection bias)	Unclear risk	Randomisation order was single blind (personal correspondence)
Blinding (performance bias and detection bias) Participants	High risk	No blinding. Personal correspondence
Blinding (performance bias and detection bias) Personnel	High risk	No blinding. Personal correspondence
Blinding (performance bias and detection bias) Outcome assessors	High risk	Personal correspondence
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	2 participants dropped out of the study (personal correspondence)
Selective reporting (reporting bias)	Unclear risk	Published as an abstract only
Other bias	Low risk	Exercise protocol clearly described
		No co-interventions (personal correspondence)

Eston 1999

Methods	Randomised controlled trial
Participants	N = 15 females, mean age 22 (SE 2)
Interventions	CWI (n = 8): Exercised arm fully submerged in water at 15°C for 15 minutes
	Passive (n = 7): No treatment
Outcomes	**Pain (tenderness)** One site on the biceps (pressure when discomfort changed to pain; algometer, N)
	Strength Maximum isometric elbow flexion (2.36 rad) (dynamometer, N)
	Swelling Thigh circumference (tape measure, cm)
	Biomarker Muscle damage (CK, IU/L)
	[Follow-up for all measures: immediately, 24, 48, and 72 hours after exercise]
Exercise type / intensity	8 sets of 5 maximal elbow flexions (eccentric and concentric), isokinetic dynamometer, 60 seconds rest between each set
Time between completing exercise and initiating intervention	Immediately and every 12 hours for the following 3 days

Eston 1999 *(Continued)*

	Participants' fitness level
Notes	SE converted to SD for: tenderness, strength, swelling and CK

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	Allocated at random to either a cryotherapy or control group (Methods, pg. 232) Coin toss (author correspondence)
Allocation concealment (selection bias)	High risk	No details in manuscript "No" (author correspondence)
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	- No details in manuscript - No (author correspondence)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	- No details in manuscript
Selective reporting (reporting bias)	Low risk	- No published protocol available - Outcomes and follow-ups stated in methods - Means and SD/SE presented by intervention group for all outcomes, at all follow-ups
Other bias	Low risk	- Exercise protocol well described - No evidence of co-interventions

Goodall 2008

Methods	Randomised controlled trial
Participants	n = 18 males, mean age 24 (SD 5)
Interventions	CWI (n = 9): Seated immersion up to iliac crest in water of 15°C for 12 minutes Passive (n = 9): Seated rest
Outcomes	**Pain** Muscle soreness (visual analogue scale; 0 to 200 mm, 'no pain' to 'extremely painful') rating during standing AND during unweighted squat

Goodall 2008 *(Continued)*

> **Strength**
> Maximal voluntary contraction: isometric knee extension at 70 degrees (isokinetic dynamometer, % change from baseline)
>
> **Swelling**
> Thigh circumference (tape measure, cm)
>
> **ROM**
> Active knee flexion (goniometer, degrees)
>
> **Biomarker**
> Muscle damage (CK, IU/L)
>
> [Follow-up for all outcomes: immediately, 24, 48, 72 and 96 hours after exercise]

Exercise type / intensity	5 sets of 20 drop jumps on a concrete based floor, 10 seconds rest between each jump, and 2 minutes rest between each set
Time between completing exercise and initiating intervention	Immediately and every 24 hours for the following 3 days
Participants' fitness level	Physically active males not familiar with eccentric based training; refrained from resistance training or exercise that may induce eccentric muscle damage/soreness for 3 weeks prior to, and for the duration of the study
Notes	Means and SD obtained after personal correspondence

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	States equally but randomly allocated (pg 236) Participants randomised using a random numbers table (personal correspondence)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript "No" (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	No details on drop outs, exclusions, missing data or approach to analysis in manuscript n = 9 participants were randomised to each group and a full data set was obtained at each follow-up (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods

Goodall 2008 *(Continued)*

		Means and SD presented (graphically) by intervention group for all outcomes, at all follow-ups.
Other bias	Low risk	Exercise protocol clearly described
		Participants refrained from exercise and resistance training for three weeks prior to the study, and refrained from exercise, resistance training, NSAIDs and nutritional supplements during the study.

Halson 2008

Methods	Randomised cross-over (3 days between interventions - personal correspondence)
Participants	N = 11 male cyclists unfamiliar with CWI, mean age 23.8 (SD 1.6)
Interventions	CWI (n = 11): Immersion to mesosternum in water at 11.5°C for 60 seconds x 3, 120 seconds between immersion
	Passive (n = 11): Seated rest, room maintained at 24.2°C and 45.6 relative humidity
Outcomes	**Subjective recovery** - Rating of recovery (visual analogue scale, 1 to 10); physical / mental - Rating of fatigue (visual analogue scale, 1 to 10) **Pain** - Leg soreness (visual analogue scale, 1 to 10) [Immediately after intervention] **Biomarker** Muscle damage (CK,U/L) Inflammation [CRP (mg/dL), IGF-1 (ng/mL), IL-6 pg/mL)] Stress hormones [prolactin (ng/mL), GH (mIU/L), Cortisol (nM), testosterone (ng/dL), adrenaline (pg/mL), noradrenaline (ng/mL)] (Follow-up: Immediately post exercise, and immediately post intervention) **Adverse effects** Core temperature before, during and immediately after immersion (disposable rectal thermometer, °C)
Exercise type / intensity	40 minutes cycling (20 minutes at 75% VO_2 max; time trial of approximately 20 minutes) in an environmental chamber maintained at 34.3°C and 41.2% relative humidity
Time between completing exercise and initiating intervention	20 minutes
Participants' fitness level	Endurance trained cyclists, VO_2 peak: 71.3 (SD 1.2), abstaining from intense exercise for 24 hours prior to each testing session
Notes	Biomarkers were not recorded for N = 4 / 11 (unable to obtain blood sample)

Risk of bias

Bias	Authors' judgement	Support for judgement

Halson 2008 *(Continued)*

Random sequence generation (selection bias)	Unclear risk	Random and counterbalanced order (pg. 333, 2nd paragraph) Participants pair-matched according to peak power obtained from a cycling VO_2 max tests (personal correspondence)
Allocation concealment (selection bias)	High risk	No details in manuscript No measures undertaken to conceal allocation (personal correspondence)
Blinding (performance bias and detection bias) Participants	High risk	No blinding. Confirmed with author contact
Blinding (performance bias and detection bias) Personnel	High risk	No blinding. Confirmed with author contact
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript No blinding (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	All participants completed each intervention arm (personal correspondence) n = 4 participants had missing data for all biochemical outcomes with reasons (unwilling to provide blood sample due to forthcoming racing event)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Low risk	Exercise intervention clearly described Participants provided with instructions for fluid and carbohydrate consumption during the study, and food diaries and training diaries were completed during the study (pg. 333)

Ingram 2009

Methods	Randomised cross-over (14 days between interventions)
Participants	N = 11 male athletes, mean age 27.5 (SD 6)
Interventions	CWI (n = 11): Seated immersion to umbilicus in water at 10°C for 5 minutes x 2 (2.5 minutes standing in room temperature between immersions) Contrast (n = 11): Seated immersion to umbilicus in water at 10°C for 2 minutes, followed by seated immersion to umbilicus in water at 40°C for 2 minutes, x 3 sets (30 second transfer time) Passive (n = 11): Seated rest
Outcomes	**Pain** Muscle soreness (10 point Likert scale) (Follow-up: immediately, 24 and 48 hours post exercise) **Strength**

Ingram 2009 *(Continued)*

Isometric strength at quads, hamstring, hips (cable tensiometer, kgf) (Follow-up: 48 hours post exercise)

Functional assessment
Time to complete exercise (10 x 20 m sprints, seconds) (Follow-up: 48 hours post exercise)

Biomarker
Muscle damage (CK, U/L)

Inflammation (CRP, mg/L)

[Follow-up: 24 and 48 hours post exercise]

Exercise type / intensity	80 minutes of simulated team sports exercise, followed by a 20 m shuttle run test to exhaustion
Time between completing exercise and initiating intervention	Immediately (?) and after 24 hours
Participants' fitness level	Athletes with team games experience
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	'randomly assigned, counterbalanced design' pg. 418) computer generated program (personal correspondence)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript Likely with computer generated program Not confirmed after personal correspondence
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript "No" (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	Manuscript shows n=11 in results (Tables 1 and 2) All participants completed each intervention arm and that there were no missing data (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD/SE presented by intervention group for all outcomes, at all follow-ups

Ingram 2009 *(Continued)*

Other bias	Low risk	Exercise protocol well described
		Manuscript states "no additional exercise or recovery procedures undertaken within 48 hours post exercise" (pg. 418)

Jakeman 2009

Methods	Randomised controlled trial
Participants	N = 18 healthy females, mean age 19.9 (SE 0.97)
Interventions	CWI (n = 9): Seated immersion to iliac crest in 10°C water, for 10 minutes
	Passive (n = 9): Seated rest
Outcomes	**Pain** Leg soreness during unweighted squat (10 cm visual analogue scale, 0 "no pain" to 10 "worst pain ever")
	Strength Concentric knee extension (isokinetic dynamometry at 45°/s, Nm)
	Biomarker Muscle damage (CK, U/L)
	(Follow-up: 1 (immediate), 24, 48, 72 and 96 hours after exercise)
Exercise type / intensity	10 counter movement jumps x 10, adopting a 90 degree knee angle on each landing
Time between completing exercise and initiating intervention	Within 10 minutes
Participants' fitness level	Not engaged in any specific plyometric or lower limb training 6 weeks prior to testing
Notes	Data for pain, strength and biomarkers obtained after personal correspondence (converted from SE to SD)

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	'randomly allocated to a control or treatment group' (Methods, participants and design, pg. 457)
		participants randomised using a predetermined sequence generation (personal correspondence)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript
		Likely with predetermined sequence generation
		Not confirmed after personal correspondence
Blinding (performance bias and detection bias) Participants	High risk	No blinding.

Jakeman 2009 *(Continued)*

Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript No (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	No details in manuscript All participants received intervention as allocation and there was no missing data (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD/SE presented (graphically) by intervention group for all outcomes, at all follow-ups
Other bias	Unclear risk	Exercise protocol is clearly described Few details on the control intervention within the manuscript

King 2009

Methods	Randomised cross-over (5 days between interventions)
Participants	n = 10 female netballers, mean age 19.5 (SD 1.5)
Interventions	CWI (n = 10): Immersion to iliac crest level in water at 9.3°C (+/-1.6°C) for 5 minutes x 2 (2.5 minutes seated at air temperature between immersions) Passive (n = 10): Seated rest for 15 minutes Active (n = 10): Low intensity jogging at predetermined speed for 15 minutes Contrast (n = 10): Immersion to iliac crest level in water at 9.7°C (SD 1.4°C) for 1 minutes followed by a warm shower in water at 39.1°C (SD 2°C) for 2 minute, x 5 sets
Outcomes	**Muscle soreness** (Likert scale; 0 = normal, 10 = extreme soreness) (Follow-up: immediately, 24 hours after intervention) **Power** Countermovement jump (5 jumps in 20 seconds, **%** decrement during 5 reps (follow-up: 24 hrs) **Functional assessment** - Exercise test (10 m sprint, seconds) - Time to complete exercise (netball/agility related circuit, seconds) (Follow-up: 24 hours after intervention)
Exercise type / intensity	Netball related circuit (4 x 15 minutes of intermittent sprinting, jumping, agility, striding) at environmental temperature of 16.9 (SD 2.1°C)

King 2009 *(Continued)*

Time between completing exercise and initiating intervention	Immediate (after post exercise outcomes)
Participants' fitness level	Trained netballers, competing at local and regional representative matches (1 to 2 per week), and training (~3 times per week)
Notes	Data on muscle soreness and subjective recovery obtained after personal correspondence

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	'randomly assigned counterbalanced order' (pg. 1796) drawn from a hat (personal correspondence)
Allocation concealment (selection bias)	Low risk	No details Participants were unaware of which recovery intervention would be used until it was time to complete the recovery. Allocation was completed by the chief investigator alone. This information was then stored in the investigators office and only used for reference by the chief investigator (personal correspondence)
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details No (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	Results Table 1, and Figures 2 and 3 state n=10, so probably no drop outs, however, no other details on exclusions or missing data No missing data, all participants completed as randomised (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Low risk	Exercise protocol clearly described Water consumption was standardised during the testing procedures; participants recorded all food and drink activity in the 24 hours leading up to the first testing session and replicated for the all testing sessions, participants requested to present in a rested state

Kuligowski 1998

Methods	Randomised controlled trial
Participants	N = 56 pain free participants; 28 male, mean age 21.1 (SD 3); 28 female, mean age 20.1 (SD 2.1)
Interventions	CWI (n = 14): Arm immersion in agitated water at 12.8°C to deltoid level for 24 minutes
	Passive (n = 14): no treatment
	Contrast Immersion (n = 14): Arm immersion in agitated water at 38.9°C to deltoid level for 3 minutes, repeated in water at 12.8°C for 1 minute, x 6 sets (24 minutes)
	arm-water immersion (n = 14): Arm immersion in agitated water at 38.9°C to deltoid level for 24 minutes
Outcomes	**Pain** Soreness during active elbow motion (12 cm graphic pain rating scale: dull ache, to unbearable pain) **Strength** Isometric elbow flexion (dynamometer, kg) **ROM** Active elbow flexion (hand held goniometer, degrees) (Follow-up: 24, 48, 96 hours post exercise)
Exercise type / intensity	Eccentric biceps loading, 10 reps at 1 repetition maximum +2.27 kg x 5 sets, 1 minute rest between sets
Time between completing exercise and initiating intervention	Immediately and at 24, 48 and 72 hours
Participants' fitness level	No upper extremity weight training in the previous 9 weeks
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	'Randomly assigned subjects to ...' (pg. 224)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript
Blinding (performance bias and detection bias) Participants	High risk	
Blinding (performance bias and detection bias) Personnel	High risk	
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details in manuscript
Incomplete outcome data (attrition bias)	Unclear risk	No details on drop outs, exclusions, missing data or approach to analysis (per protocol / ITT)

All outcomes

Selective reporting (reporting bias)	Low risk	No published protocol available
		Outcomes and follow-ups stated in methods
		Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Low risk	Exercise protocol clearly described
		Participants were instructed not to use therapeutic modalities, massage, stretching or medications, and refrain from strenuous activity during the course of the study (pg 223)

Montgomery 2008

Methods	Randomised controlled trial (matched by playing position and anthropometrics)
Participants	N = 29 male basketball players, mean age 19.1 (SD: 2.1)
Interventions	All groups: Carbohydrate and stretching
	CWI group (n = 10): Immersion to mesosternale level in 11°C water, 5 x 1 minute. Between immersions participants rested passively for 2 minutes in ambient air (23°C)
	Compression (n = 10): Full length compression garments (18 mm/Hg) post game and overnight (18 hours)
	Control (n = 9): Carbohydrate and stretching only, post game carbohydrate snack (1 g/kg body mass) and 600 ml of fluid in sports drink followed by a standardised programme of 10 stretches (bilateral leg and lower back, each stretch held for 15 seconds, 2 sets)
Outcomes	**Subjective recovery** Rating of general fatigue (visual analogue scale, 1 "not at all" to 10 "extremely tired") **Pain** Leg soreness (visual analogue scale, 1 "normal" to 10 "extremely sore") **Power** Vertical jump (cm) **Functional assessment** Time to complete exercise test (20 sprint; repeated sprint; agility test; all seconds) **Swelling** Limb girth (cm, calf and thigh) **ROM** Sit and reach flexibility test (cm) [Follow-up: 24 hours (after finishing the 3 day tournament)]
Exercise type / intensity	3 day basketball tournament (48 minute game each day), comprising a total of 144 minutes of basketball in mean temperature of 23.2°C
Time between completing exercise and initiating intervention	˜10 minutes (all groups undertook control after each game estimated to take 10 minutes)

Montgomery 2008 *(Continued)*

Participants' fitness level	Regularly competing in state competitions; training load of 8 to 10 hours per week
Notes	Our analysis only included outcomes recorded at the end of the tournament, ~24 hrs after the final game ["pre tournament tests were repeated on the morning of the 4th day" (pg. 1137)]. Outcomes (subjective recovery, pain, muscle power) recorded after each game were not extracted or included in our analysis. Measurement device for vertical jump not clear

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	'randomised controlled trial' (Methods, experimental design, pg. 1137) players matched for positional and anthropometric characteristics and assigned to one of three treatment groups (Methods, experimental procedures, pg. 1137)
Allocation concealment (selection bias)	Unclear risk	No details in manuscript
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details in manuscript
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No details on drop outs, exclusions, missing data or approach to analysis (per protocol / ITT)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Low risk	Cumulative game time is expressed for participants within each treatment group, and is similar (also used as a covariate in statistical analysis) Participants undertook standardised carbohydrate ingestion and stretching between games

Rowsell 2009

Methods	Randomised controlled design (pair-matched according to playing position)
Participants	N = 20 male soccer players, mean age: 15.9 (SD: 0.6)

Rowsell 2009 *(Continued)*

Interventions	CWI (n = 6): Immersion to mesosternale level in 10°C water, 1 minute x 5, manual water agitation throughout. Between immersions participants rested for 60 seconds
	WWI (n = 7): Immersion to mesosternale level in 34°C water, 5 x 1 minute, manual water agitation throughout. Between immersions participants rested for 60 seconds
	Both groups: Standardardised warm up and cool down, carbohydrate, protein and fluid ingestion
Outcomes	**Subjective recovery** Rating of fatigue (visual analogue scale, 1 to 10) Rating or recovery (visual analogue scale, 1 to 10): physical / mental Do you believe that the treatment enhanced recovery from the previous match and improved subsequent physical performance (Yes/No) **Pain** Leg soreness (visual analogue scale, 1 to 10) **Power** Countermovement jump (jump height - standing reach height, cm) **Functional assessment** Time to complete exercise test (repeated sprint (12 x 20 m), seconds) **Biochemical** Inflammatory: IL-1b (pg/ml); IL-6 (pg/ml); IL-10 (pg/ml) Muscle damage: myoglobin (ng/ml), LDH (U/L), CK (U/L) (Follow-up for all outcomes: ~ 22 hours after tournament)
Exercise type / intensity	Soccer tournament, 4 matches in 4 days, similar tactical strategies adopted in each match. Ambient temperature during matches ranged from 25 to 36°C, relative humidity 25 to 58%.
Time between completing exercise and initiating intervention	20 minutes after each match
Participants' fitness level	High performance male junior soccer players
Notes	Data on muscle soreness and subjective recovery (fatigue) extracted from graph. Other data on subjective recovery were not available (rating of physical and mental recovery) N = 7 players sustained injuries throughout the tournament and were not included in the final analysis Our analysis only included outcomes recorded at the end of the tournament. Outcomes undertaken before each of the games were not extracted or included in our analysis. An additional report of the trial (Rowsell 2011) was located in November 2011 - this will be assessed in the update of this review.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	'randomly allocated' (Methods, participants, pg. 566), "players matched for playing position" (pg. 566)
Allocation concealment (selection bias)	Unclear risk	"randomly allocated" (Methods, participants, pg. 566), no further details in manuscript
Blinding (performance bias and detection bias)	High risk	No blinding.

Rowsell 2009 *(Continued)*

Participants

Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details in manuscript
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	The number of participants randomised to each group is not clear n = 7 were excluded from the analysis with reasons (all 7 sustained a soft tissue injury and were unable to complete the study)
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-ups (except physical and mental recovery which stated no between group difference)
Other bias	Low risk	Players in each group aged matched according to playing position so likely to undertake similar exercise intensity; no measure of cumulative game time during the tournament Standardised carbohydrate ingestion and stretching between games

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Sellwood 2007a

Methods	Randomised controlled trial
Participants	N = 40 healthy adults, mean age 21.3 (SD 4.3), 11 male / 29 female
Interventions	Cold-water immersion (n = 20): Immersion in water at 5°C, 1 minute x 3 sets. Warm-water immersion (n = 20): Immersion in water at 24°C, 1 minute x 3 sets Both groups: Immersion to anterior superior iliac spine, participants rested out of the bath for 60 seconds between sets
Outcomes	**Pain** Pain during: sit-stand, passive stretch, hopping, running, isometric contraction (100 mm visual analogue scale, "no pain" to "worst pain possible") **Tenderness** Pain (100 mm visual analogue scale) when pressure applied at two points on the thigh, pressure standardised at 6 lb/cm^2 using an algometer) **Strength** Isometric knee extension (isokinetic dynamometer, 60° flexion, Nm) **Power** Hop for distance (tape measure, m) **Swelling** Thigh circumference (use of tape measure?, mm)

Sellwood 2007a *(Continued)*

Biomarker
Muscle damage (CK, IU/L)

(Follow-up: 24, 48 and 72 hours after exercise)

Exercise type / intensity	5 sets of 10 repetitions of eccentric quadriceps exercise at 120% of 1 repetition maximum, leg extension machine, 1 minute rest between sets
Time between completing exercise and initiating intervention	Immediate
Participants' fitness level	No eccentric quadriceps exercise within past 3 months
Notes	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"Generated using a random numbers table" (Methods, randomisation and masking, pg. 393)
Allocation concealment (selection bias)	Low risk	"Sequentially numbered opaque sealed envelopes held at a central location" (Methods, randomisation and masking, pg. 393)
Blinding (performance bias and detection bias) Participants	High risk	Difficult due to nature of intervention, however participants were not informed as to which intervention was considered therapeutic (Methods, randomisation and masking, pg. 393)
Blinding (performance bias and detection bias) Personnel	High risk	No details
Blinding (performance bias and detection bias) Outcome assessors	Low risk	"Investigator responsible for outcome assessments was blinded to group allocation, and participants advised not to reveal their allocation" (Methods, randomisation and masking, pg. 393)
Incomplete outcome data (attrition bias) All outcomes	Low risk	Data analysis (pg. 395) states that analysis was based on ITT, with imputation using Last Observation Carried Forward (LOCF) in the event of missing data drop outs/missing data is not reported in text however Table 5 which states that n = 20 were followed up in each group; therefore likely that imputation (LOCF) was not required
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD (or median and IQ range) presented by intervention group for all outcomes, at all follow-ups
Other bias	Unclear risk	Exercise protocol clearly described No details on co-interventions

Skurvydas 2006

Methods	Randomised cross-over (9 to 10 months between interventions)
Participants	N = 20 healthy untrained men, aged 20.4 (SD 1.7)
Interventions	CWI (n = 20): Immersion of legs in water at 15°C (SD 1°C) for 15 minutes x 2. 10 minutes between each immersion
	Passive (n = 20): Seated rest
Outcomes	**Pain** Muscle soreness (0 to 10 points) (follow-up: 24, 48, 72 hours post exercise) **Strength** Isometric knee extension (90 degree knee angle, force measuring device, % of baseline) (follow-up: immediate (4 hours), 24, 48, 72 hours post exercise) **Power** Counter movement jump (Kistler force plate, height of jump in cm) (follow-up: immediate (4 hours), 24, 48, 72 hours post exercise) **Biomarker** Muscle damage (CK, IU/L) (Follow-up: 24 hours and 48 hours post exercise)
Exercise type / intensity	100 counter movement jumps from a 0.75 m height
Time between completing exercise and initiating intervention	Immediately and at 4, 8, and 24 hours
Participants' fitness level	Physically active but did not take part in any formal exercise or sports programme
Notes	Data on pain, strength, power and CK were obtained after personal correspondence

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"The choice of their participation in the first and second experiments was made at random" (pg. 142)
Allocation concealment (selection bias)	High risk	No details in manuscript "No" (personal correspondence)
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	High risk	No details in manuscript "No" (personal correspondence)
Incomplete outcome data (attrition bias) All outcomes	Low risk	No details in manuscript

Skurvydas 2006 *(Continued)*

		All 20 participants completed each treatment arm and data were available for all 20 participants (personal correspondence)
Selective reporting (reporting bias)	Low risk	No published protocol available
		Outcomes and follow-ups stated in methods
		Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Unclear risk	Exercise protocol well described
		No details on co-interventions

Vaile 2008c

Methods	Randomised cross-over (8 months between interventions) (*see* Notes)
Participants	N = 38 males
Interventions	CWI (n = 12): Whole body immersion (excluding head and neck) in water at 15°C
	WWI (n = 11): Whole body immersion (excluding head and neck) in water at 38°C
	Contrast (n = 15): Whole body immersion (excluding head and neck) in water at 15°C for 1 minute, followed by whole body immersion (excluding head and neck) in water at 38°C for 1 minute, x 7 sets
	(Passive (n = 38): Seated with minimal movement (*see* Notes))
	All groups: Interventions were 14 minute durations
Outcomes	**Pain** (visual analogue scale, 0 'normal ' to 10 'extremely sore')
	Strength Isometric squat (force platform, peak force, N)
	Power Weighted squat jump (peak force, W)
	Swelling Thigh girth (measuring tape, cm)
	Biomarker Muscle damage [CK (U/L), LDH (U/L), myoglobin (ng/mL)] Inflammation (IL-6, pg/mL)
	[Follow-up: immediately, 24, 48, 72 hours post exercise]
Exercise type / intensity	10 eccentric repetitions on a leg press machine x 7 sets, 3 minutes rest between sets
Time between completing exercise and initiating intervention	Immediately and at 24, 48 and 72 hours
Participants' fitness level	Strength trained
Notes	Data on pain were extracted from graph

* This was a cross-over of hydrotherapy versus passive (it is unclear whether this was randomised). Only the non-cross-over data from the three hydrotherapy groups are presented in the review

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"Subjects were randomly assigned..." (pg. 448)
Allocation concealment (selection bias)	Unclear risk	No details
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No details
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-ups
Other bias	Unclear risk	Exercise protocol was well described No details on co-interventions

Yanagisawa 2003a

Methods	Randomised controlled trial
Participants	N = 28 healthy participants, mean age 23.8 (SD 1.8)
Interventions	CWI (n = 9): Immersion to the head of the fibula in water at 5°C for 15 minutes Double CWI (n = 9): Immersion to the head of the fibula in water at 5°C for 15 minutes Control (n = 10): no intervention (?)
Outcomes	**Pain** Calf soreness (visual analogue scale, 1 (normal) to 5 (very, very sore) (follow-up: 24, 48 and 96 and 168 hours post exercise) **ROM**

Yanagisawa 2003a *(Continued)*

Ankle joint dorsiflexion (goniometer, degrees)

(follow-up: immediate (20 minutes), 24, 48 and 96 and 168 hours post exercise)

Biomarker
Muscle damage [CK, LDH (IU/L)]

(Follow-up: 24, 48 and 96 and 168 hours post exercise)

Exercise type / intensity	5 sets of 20 resisted calf raises at 30% of participants maximal voluntary contraction, 1 minute rest between sets
Time between completing exercise and initiating intervention	Immediate (double CWI group repeated intervention at 24 hours)
Participants' fitness level	Untrained
Notes	No details on control intervention
	Data for pain and biomarkers were extracted from graphs. We were unable to extract data for biomarkers at 24 and 48 hours.

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"Subjects randomly allocated" (Methods, subjects, pg 54)
Allocation concealment (selection bias)	Unclear risk	No details
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No details
Selective reporting (reporting bias)	Low risk	No published protocol available
		Outcomes and follow-ups stated in methods
		Means and SD presented by intervention group for all outcomes, at all follow-up
Other bias	Low risk	Exercise intervention clearly reported
		Participants refrained from physical exercise, other forms of physical therapy, or drinking (alcohol?) for the duration of the study

Yanagisawa 2003b

Methods	Randomised controlled trial
Participants	N = 14 healthy males, mean age 23.8 (SD 2.3)
Interventions	CWI group (n = 7): Immersion to the head of the fibula in water at 5°C for 15 minutes
	Control (n = 7): no intervention (?)
Outcomes	**Pain** Calf soreness during walking (visual analogue scale, 1 (normal) to 5 (very, very sore) [Follow-up: immediate (30 min), 24, 48 and 96 and 168 hours post exercise]
Exercise type / intensity	5 sets of 12 resisted calf raises at 30% of participants maximal voluntary contraction, 1 minute rest between sets
Time between completing exercise and initiating intervention	Immediate
Participants' fitness level	Untrained
Notes	No details on control intervention; data on pain extracted from graph

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	"Subjects were randomly assigned" (Abstract, pg. 1517)
Allocation concealment (selection bias)	Unclear risk	No details
Blinding (performance bias and detection bias) Participants	High risk	No blinding.
Blinding (performance bias and detection bias) Personnel	High risk	No blinding.
Blinding (performance bias and detection bias) Outcome assessors	Unclear risk	No details
Incomplete outcome data (attrition bias) All outcomes	Unclear risk	No details
Selective reporting (reporting bias)	Low risk	No published protocol available Outcomes and follow-ups stated in methods Means and SD presented by intervention group for all outcomes, at all follow-up

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Yanagisawa 2003b *(Continued)*

Other bias	Low risk	Exercise protocol well described
		Participants refrained from physical exercise, and private therapeutic treatment for the duration of the study

CK = creatine kinase
CPK = creatine phosphokinase
IL = interleukin
LDH = lactate dehydrogenase
SD = standard deviation
SE = standard error

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Banfi 2008a	Intervention was CWI combined with active cycle intervention
Chen 2006	Unclear if intervention is CWI (from abstract). Unable to source full text, unable to contact authors
Clements 2002	No relevant outcomes
Coffey 2004	No CWI intervention
Connolly 2003	No CWI intervention
Crowe 2007	No primary outcomes
Dawson 2005	No CWI intervention
Gill 2006	No CWI intervention
Goosey-Tolfrey 2008	CWI did not include muscles/body parts primarily involved in the exercise protocol (isolated wrist immersion only)
Gurovich 2006	No CWI (from abstract)
Hayashi 2004	CWI at 20°C
Heyman 2009	No primary outcomes
Higgins 2010	No relevant follow-up times (abstract unclear however exclusion confirmed after full text version sourced from authors)
Hudson 1999	CWI at 33°C
Khomenok 2008	CWI did not include muscles/body parts primarily involved in the exercise protocol (isolated wrist immersion only)
Kinugasa 2009	Intervention was CWI combined with active cycle intervention
Kokkinidis 1998	No CWI intervention
Lane 2004	No primary outcomes

Study	Reason for exclusion
Lapier 1995	No CWI (abstract unclear, confirmed after correspondence with authors)
Nosaka 1996	No CWI (abstract unclear, confirmed after correspondence with authors and review of full text)
Paddon-Jones 1997	Participants randomised by body site (one arm CWI; other arm control)
Peiffer 2009a	No primary outcomes
Peiffer 2009b	No randomisation; states 'counterbalanced' only
Peiffer 2010b	No randomisation; states 'counterbalanced' only
Peñailillo 2006	No CWI (from abstract)
Price 2004	No CWI intervention
Schniepp 2002	No primary outcomes
Sipaviciene 2007	Not clear if intervention is CWI or cold pack (English abstract only, full text in Lithuanian), no response from authors
Taylor 2008	No relevant outcomes
Tessitore 2007	No CWI intervention
Udermann 1997	No CWI (abstract unclear, confirmed after correspondence with primary author and review of full text)
Vaile 2007	No CWI intervention
Vaile 2008a	No primary outcomes
Vaile 2008b	No primary outcomes
Verducci 2000	No CWI intervention
Webborn 2005	No post exercise CWI intervention
Yamane 2006	No randomisation; each participants had an untreated limb and cooled limb
Yeargin 2006	No primary outcome

CWI = cold-water immersion

Characteristics of studies awaiting assessment [ordered by study ID]

Ascensao 2011

Methods	Randomised controlled trial
Participants	20 male soccer players
Interventions	CWI: 10 min cold water immersion at 10ºC
	Warm water immersion: 10 min thermoneutral water immersion at 35ºC

Ascensao 2011 *(Continued)*

Outcomes	**Pain** DOMS **Strength** Maximal isometric quadriceps strength **Functional assessment** Jump and sprint abilities **Biomarker** Muscle damage [creatine kinase, myoglobin] Inflammation (C-reactive protein) [Follow-up: immediately, 24 and 48 hours post exercise]
Notes	Outside initial search limit.

Fowles 2003

Methods	Randomised cross-over trial
Participants	16 active participants (9 male, 7 female)
Interventions	CWI group: immersion to waist level at 8ºC Control group: light stretching exercises
Outcomes	**Pain** Soreness rating **Power** Vertical jump **Functional assessment** Treadmill run to fatigue (time) Intense 6 km interval run (time to complete) (Follow-up: 48 hours after exercise)
Notes	Exercise was treadmill run to fatigue followed by 6 km run (mean time to completion was 43.5 minutes) Abstract only

Pournot 2011

Methods	Randomised controlled trial
Participants	41 elite athletes (all male)
Interventions	CWI: at 10°C Warm water immersion: at 36°C

Pournot 2011 *(Continued)*

	Contrast: 10 to 42°C
	Passive recovery

Outcomes	**Strength**
	Maximal isometric voluntary contraction (MVC) of the knee extensor muscles
	Power Maximal vertical counter-movement jump
	Functional assessment
	Maximal 30-s rowing test
	Biomarker Muscle damage [creatine kinase, lactate dehydrogenase] Inflammation (IL-6, pg/mL)
	[Follow-up: immediately and 24 hours post exercise]

Notes	Outside initial search limit. May not include primary outcomes.

Smith 2008

Methods	Randomised controlled trial

Participants	31 participants

Interventions	Cryotherapy
	Thermotherapy

Outcomes	**Strength**
	Upper arm 1 repetition maximum

Notes	Exercise was a program of high volume, medium intensity resistance exercises Abstract only

Stacey 2010

Methods	Randomised controlled trial (probably cross-over)

Participants	9 active males (3 cycling bouts)

Interventions	CWI: cold tub with water at 10°C
	Active recovery: cycling

Outcomes	**Pain** Self-assessment
	Subjective measures Perceived exertion and lower extremity sensations
	Biomarker

Stacey 2010 *(Continued)*

	Blood samples (lactate, IL-6, total leukocyte, neutrophil, and lymphocyte cell counts)
	[Follow-up: immediately, post cycling bouts]
Notes	Outside initial search limit. Study design may not be appropriate for inclusion.

Vaile 2011

Methods	Randomised cross-over trial (7 days between interventions)
Participants	10 endurance trained male athletes
Interventions	CWI group: Body immersion (excluding head and neck) in water at 15°C for 15 minutes
	Control group: Active recovery, cycling at 40% of maximum power for 15 minutes
Outcomes	**Functional assessment**
	Total work (kJ) during cycling performance test
Notes	Outside initial search limit

CWI = cold-water immersion

DATA AND ANALYSES

Comparison 1.   Cold-water immersion (CWI) versus passive

Outcome or subgroup title	No. of studies	No. of partici-pants	Statistical method	Effect size
1 Pain (muscle soreness: various scales Likert and VAS)	12		Std. Mean Difference (IV, Fixed, 95% CI)	Subtotals only
1.1 Immediately post intervention	7	128	Std. Mean Difference (IV, Fixed, 95% CI)	-0.07 [-0.43, 0.28]
1.2 24 hours	10	218	Std. Mean Difference (IV, Fixed, 95% CI)	-0.55 [-0.84, -0.27]
1.3 48 hours	8	179	Std. Mean Difference (IV, Fixed, 95% CI)	-0.66 [-0.97, -0.35]
1.4 72 hours	4	104	Std. Mean Difference (IV, Fixed, 95% CI)	-0.93 [-1.36, -0.51]
1.5 96 hours	5	97	Std. Mean Difference (IV, Fixed, 95% CI)	-0.58 [1.00, -0.16]
2 Pain - random effects analysis (muscle soreness: various scales Likert and VAS)	12		Std. Mean Difference (IV, Random, 95% CI)	Subtotals only

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2.1 Immediately post intervention	7	128	Std. Mean Difference (IV, Random, 95% CI)	-0.07 [-0.43, 0.28]
2.2 24 hours	10	218	Std. Mean Difference (IV, Random, 95% CI)	-0.58 [-1.06, -0.10]
2.3 48 hours	8	179	Std. Mean Difference (IV, Random, 95% CI)	-0.70 [-1.18, -0.21]
2.4 72 hours	4	104	Std. Mean Difference (IV, Random, 95% CI)	-0.88 [-1.83, 0.07]
2.5 96 hours	5	97	Std. Mean Difference (IV, Random, 95% CI)	-0.56 [-1.12, 0.01]
3 Subgroup analysis. Study design: Pain at 24 hours (muscle soreness: various scales Likert and VAS)	10	218	Std. Mean Difference (IV, Fixed, 95% CI)	-0.55 [-0.84, -0.27]
3.1 parallel group	7	136	Std. Mean Difference (IV, Fixed, 95% CI)	-0.23 [-0.58, 0.12]
3.2 cross-over	3	82	Std. Mean Difference (IV, Fixed, 95% CI)	-1.17 [-1.65, -0.69]
4 Subgroup analysis. Study design: Pain at 48 hours (muscle soreness: various scales Likert and VAS)	8	179	Std. Mean Difference (IV, Fixed, 95% CI)	-0.66 [-0.97, -0.35]
4.1 parallel group	6	117	Std. Mean Difference (IV, Fixed, 95% CI)	-0.40 [-0.77, -0.02]
4.2 cross-over	2	62	Std. Mean Difference (IV, Fixed, 95% CI)	-1.26 [-1.81, -0.70]
5 Subgroup analysis. Study design: Pain at 72 hours (muscle soreness: various scales Likert and VAS)	4	104	Std. Mean Difference (IV, Fixed, 95% CI)	-0.93 [-1.36, -0.51]
5.1 parallel groups	3	64	Std. Mean Difference (IV, Fixed, 95% CI)	-0.43 [-0.93, 0.07]
5.2 cross-over	1	40	Std. Mean Difference (IV, Fixed, 95% CI)	-2.27 [-3.08, -1.46]
6 Subgroup analysis. Immersion frequency: Pain at 24 hours (muscle soreness: various scales Likert and VAS)	10	218	Std. Mean Difference (IV, Fixed, 95% CI)	-0.55 [-0.84, -0.27]
6.1 Single CWI	6	110	Std. Mean Difference (IV, Fixed, 95% CI)	-0.51 [-0.90, -0.11]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6.2 Multiple CWI	4	108	Std. Mean Difference (IV, Fixed, 95% CI)	-0.60 [-1.00, -0.20]
7 Subgroup analysis. Immersion frequency: Pain at 48 hours (muscle soreness: various scales Likert and VAS)	8	179	Std. Mean Difference (IV, Fixed, 95% CI)	-0.66 [-0.97, -0.35]
7.1 Single CWI	4	71	Std. Mean Difference (IV, Fixed, 95% CI)	-0.75 [-1.25, -0.26]
7.2 Multiple CWI	4	108	Std. Mean Difference (IV, Fixed, 95% CI)	-0.60 [-1.01, -0.20]
8 Subgroup analysis. Exercise type: Pain at 24 hours (muscle soreness: various scales Likert and VAS)	10	218	Std. Mean Difference (IV, Fixed, 95% CI)	-0.55 [-0.84, -0.27]
8.1 Lab controlled muscle damage (DOMS)	6	137	Std. Mean Difference (IV, Fixed, 95% CI)	-0.22 [-0.57, 0.12]
8.2 Other sporting activity	4	81	Std. Mean Difference (IV, Fixed, 95% CI)	-1.23 [-1.72, -0.74]
9 Subgroup analysis. Exercise type: Pain at 48 hours (muscle soreness: various scales Likert and VAS)	8	179	Std. Mean Difference (IV, Fixed, 95% CI)	-0.66 [-0.97, -0.35]
9.1 Lab controlled muscle damage (DOMS)	6	137	Std. Mean Difference (IV, Fixed, 95% CI)	-0.48 [-0.83, -0.14]
9.2 Other sporting activity	2	42	Std. Mean Difference (IV, Fixed, 95% CI)	-1.39 [-2.09, -0.69]
10 Tenderness (pain on palpation)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
10.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.2 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.3 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
11 Subjective recovery (10 point or 10 cm VAS)	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
11.1 Physical recovery (Immediately post intervention)	2	42	Mean Difference (IV, Fixed, 95% CI)	0.97 [-0.10, 2.05]
11.2 Mental recovery (immediately post intervention)	1	22	Mean Difference (IV, Fixed, 95% CI)	0.60 [-0.86, 2.06]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
12 Fatigue (10 point / 10 cm VAS)	3		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
12.1 Fatigue (immediately post intervention)	2	38	Mean Difference (IV, Fixed, 95% CI)	-1.70 [-2.49, -0.90]
12.2 Fatigue (24 hrs post intervention)	1	19	Mean Difference (IV, Fixed, 95% CI)	-0.70 [-1.88, 0.48]
13 Strength (Final value: Nm)	6		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
13.1 Immediate	3	58	Mean Difference (IV, Fixed, 95% CI)	-19.92 [-33.24, -6.59]
13.2 24 hours	4	81	Mean Difference (IV, Fixed, 95% CI)	-1.94 [-6.44, 2.56]
13.3 48 hours	5	103	Mean Difference (IV, Fixed, 95% CI)	-1.44 [-5.46, 2.57]
13.4 72 hours	3	61	Mean Difference (IV, Fixed, 95% CI)	-0.89 [-5.33, 3.56]
13.5 96 hours	2	46	Mean Difference (IV, Fixed, 95% CI)	-0.58 [-4.06, 2.89]
14 Strength (% of baseline)	2		Mean Difference (IV, Random, 95% CI)	Totals not selected
14.1 Immediate	2		Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
14.2 24 hours	2		Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
14.3 48 hours	2		Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
14.4 72 hours	2		Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
14.5 96 hours	1		Mean Difference (IV, Random, 95% CI)	0.0 [0.0, 0.0]
15 Power (jump height: centimetres)	4		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
15.1 Immediate (cross-over trial)	1	40	Mean Difference (IV, Fixed, 95% CI)	-1.80 [-6.29, 2.69]
15.2 24 hours	3	52	Mean Difference (IV, Fixed, 95% CI)	0.26 [-2.04, 2.56]
15.3 24 hours (cross-over trial)	1	40	Mean Difference (IV, Fixed, 95% CI)	5.40 [2.12, 8.68]
15.4 48 hours	1	20	Mean Difference (IV, Fixed, 95% CI)	0.0 [-2.63, 2.63]
15.5 48 hours (cross-over trial)	1	40	Mean Difference (IV, Fixed, 95% CI)	9.60 [6.40, 12.80]

Outcome or subgroup title	No. of studies	No. of partici-pants	Statistical method	Effect size
15.6 72 hours (cross-over tri-al)	1	40	Mean Difference (IV, Fixed, 95% CI)	6.60 [3.16, 10.04]
16 Power (% decrease in jump height over 5 jumps)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
16.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
17 Power (cycle ergometer power: Watts)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
17.1 Peak power output (Im-mediate)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
18 Functional performance (time to complete exercise test: seconds)	6		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
18.1 1 km cycle time (imme-diate)	1	20	Mean Difference (IV, Fixed, 95% CI)	0.30 [-2.64, 3.24]
18.2 Agility circuit time (24 hours)	1	19	Mean Difference (IV, Fixed, 95% CI)	-0.03 [-0.26, 0.20]
18.3 Sprint time (24 hours)	2	39	Mean Difference (IV, Fixed, 95% CI)	-0.02 [-0.10, 0.07]
18.4 Multisprint / Agility cir-cuit time (24 hours)	1	20	Mean Difference (IV, Fixed, 95% CI)	-0.29 [-2.49, 1.91]
18.5 Multisprint time (24 hours)	2	32	Mean Difference (IV, Fixed, 95% CI)	-0.19 [-1.06, 0.68]
18.6 Sprint time (48 hours)	1	20	Mean Difference (IV, Fixed, 95% CI)	0.02 [-0.11, 0.15]
18.7 Multisprint time (48 hours)	1	22	Mean Difference (IV, Fixed, 95% CI)	-0.07 [-1.37, 1.23]
19 Functional performance (time to fatigue)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
19.1 % decrease from base-line (immediate)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
20 Range of movement (ROM)	4		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
20.1 Degrees (Immediate)	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
20.2 Degrees (24 hours)	3		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
20.3 Centimetres (24 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
20.4 Degrees (48 hours)	3		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
20.5 Degrees (72 hours)	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
20.6 Degrees (96 hours)	3		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
21 Swelling (limb girth in centimetres)	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
21.1 Final value immediate	1	18	Mean Difference (IV, Fixed, 95% CI)	0.10 [-5.16, 5.36]
21.2 Final value 24 hours	2	33	Mean Difference (IV, Fixed, 95% CI)	-1.41 [-5.63, 2.80]
21.3 Final value 48 hours	2	33	Mean Difference (IV, Fixed, 95% CI)	-1.53 [-5.57, 2.51]
21.4 Final value 72 hours	2	33	Mean Difference (IV, Fixed, 95% CI)	-1.54 [-5.65, 2.56]
21.5 Final value 96 hours	1	18	Mean Difference (IV, Fixed, 95% CI)	-0.60 [-5.80, 4.60]
22 Biomarker: creatine kinase	10		Std. Mean Difference (IV, Random, 95% CI)	Subtotals only
22.1 Immediately post	4	68	Std. Mean Difference (IV, Random, 95% CI)	0.01 [-0.66, 0.67]
22.2 24 hours	7	146	Std. Mean Difference (IV, Random, 95% CI)	-0.20 [-1.03, 0.63]
22.3 48 hours	7	152	Std. Mean Difference (IV, Random, 95% CI)	-0.50 [-1.36, 0.37]
22.4 72 hours	3	51	Std. Mean Difference (IV, Random, 95% CI)	0.14 [-0.52, 0.79]
22.5 96 hours	3	55	Std. Mean Difference (IV, Random, 95% CI)	0.01 [-0.70, 0.73]
23 Biomarker: lactate dehydrogenase	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
23.1 U/L (immediate)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
23.2 % change 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
24 Biomarker: myoglobin	3		Std. Mean Difference (IV, Fixed, 95% CI)	Subtotals only
24.1 Immediate	2	36	Std. Mean Difference (IV, Fixed, 95% CI)	-0.51 [-1.18, 0.17]
24.2 24 hours	2	33	Std. Mean Difference (IV, Fixed, 95% CI)	-0.66 [-1.36, 0.05]
24.3 48 hours	1	20	Std. Mean Difference (IV, Fixed, 95% CI)	-0.74 [-1.66, 0.17]
25 Biomarker: interleukin-6 (pg/ml)	1		Std. Mean Difference (IV, Fixed, 95% CI)	Totals not selected

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Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
25.1 Immediate	1		Std. Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
26 Biomarker: C-reactive protein (mg/dL / mg/L)	3		Std. Mean Difference (IV, Fixed, 95% CI)	Subtotals only
26.1 Immediate	2	30	Std. Mean Difference (IV, Fixed, 95% CI)	0.32 [-0.40, 1.04]
26.2 24 hours	1	22	Std. Mean Difference (IV, Fixed, 95% CI)	-0.23 [-1.07, 0.61]
26.3 48 hours	1	22	Std. Mean Difference (IV, Fixed, 95% CI)	-0.21 [-1.05, 0.63]
27 Pain (muscle soreness: all converted to 10 cm VAS)	12		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
27.1 Immediately post intervention	7	128	Mean Difference (IV, Fixed, 95% CI)	0.01 [-0.59, 0.61]
27.2 24 hours	10	218	Mean Difference (IV, Fixed, 95% CI)	-1.27 [-1.70, -0.84]
27.3 48 hours	8	179	Mean Difference (IV, Fixed, 95% CI)	-1.58 [-2.07, -1.10]
27.4 72 hours	4	104	Mean Difference (IV, Fixed, 95% CI)	-2.16 [-2.79, -1.53]
27.5 96 hours	5	97	Mean Difference (IV, Fixed, 95% CI)	-0.64 [-1.21, -0.07]

Analysis 1.1.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 1 Pain (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
1.1.1 Immediately post intervention							
Bailey 2007	10	6.2 (1.9)	10	6.9 (2.2)		15.86%	-0.33[-1.21,0.56]
Cassar 2010	8	2.5 (1)	8	2.2 (1.8)		12.82%	0.19[-0.79,1.18]
Goodall 2008	9	10 (4.9)	9	6.7 (4.5)		13.55%	0.67[-0.29,1.62]
Halson 2008	11	3.8 (2.6)	11	5 (2.9)		17.28%	-0.42[-1.27,0.43]
Jakeman 2009	9	1.6 (1)	9	1.2 (1.2)		14.22%	0.36[-0.57,1.29]
King 2009	10	2.5 (1.5)	10	4 (2.5)		15.02%	-0.69[-1.6,0.22]
Yanagisawa 2003b	7	1.7 (1)	7	1.8 (0.4)		11.26%	-0.12[-1.17,0.93]
Subtotal ***	**64**		**64**			**100%**	**-0.07[-0.43,0.28]**
Heterogeneity: Tau²=0; Chi²=6.15, df=6(P=0.41); I²=2.4%							
Test for overall effect: Z=0.41(P=0.68)							
1.1.2 24 hours							
Bailey 2007	10	4.6 (0.9)	10	6.8 (1.6)		7.32%	-1.62[-2.66,-0.58]
Goodall 2008	9	10.2 (4.8)	9	10.8 (4.2)		9.27%	-0.13[-1.05,0.8]
Ingram 2009	11	3 (1)	11	5 (1)		7.27%	-1.92[-2.97,-0.88]

Favours CWI -2 -1 0 1 2 Favours Passive

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		8.62%	0.7[-0.26,1.66]
King 2009	10	1.3 (1)	10	2.4 (1.3)		9.07%	-0.94[-1.88,-0.01]
Kuligowski 1998	14	8.6 (6.6)	14	7.1 (4.5)		14.33%	0.25[-0.5,0.99]
Montgomery 2008	10	3.3 (1.1)	9	4.3 (1.8)		9.18%	-0.65[-1.58,0.28]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		18.23%	-0.98[-1.64,-0.32]
Yanagisawa 2003a	9	3.1 (0.5)	10	3.3 (0.8)		9.66%	-0.28[-1.19,0.62]
Yanagisawa 2003b	7	2.4 (0.8)	7	2.8 (0.9)		7.04%	-0.4[-1.46,0.66]
Subtotal ***	**109**		**109**			**100%**	**-0.55[-0.84,-0.27]**

Heterogeneity: Tau²=0; Chi²=25.21, df=9(P=0); I²=64.3%

Test for overall effect: Z=3.85(P=0)

1.1.3 48 hours

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
Bailey 2007	10	4.4 (1.6)	10	5.7 (0.9)		11.04%	-0.96[-1.9,-0.02]
Goodall 2008	9	11.4 (5.1)	9	11.2 (3.9)		11.35%	0.04[-0.88,0.97]
Ingram 2009	11	3 (1)	11	5 (1)		8.88%	-1.92[-2.97,-0.88]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		11.34%	-0.11[-1.03,0.82]
Kuligowski 1998	14	11 (5.5)	14	10.3 (4.9)		17.62%	0.13[-0.61,0.87]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		22.22%	-0.99[-1.65,-0.33]
Yanagisawa 2003a	9	3.8 (0.4)	10	4.4 (0.5)		9.57%	-1.26[-2.26,-0.25]
Yanagisawa 2003b	7	2.6 (0.6)	7	3.5 (1.4)		7.98%	-0.78[-1.88,0.32]
Subtotal ***	**89**		**90**			**100%**	**-0.66[-0.97,-0.35]**

Heterogeneity: Tau²=0; Chi²=16.34, df=7(P=0.02); I²=57.15%

Test for overall effect: Z=4.18(P<0.0001)

1.1.4 72 hours

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
Goodall 2008	9	6 (3.2)	9	6.7 (4.2)		20.98%	-0.18[-1.1,0.75]
Jakeman 2009	9	1.6 (1.3)	9	2.3 (1.8)		20.53%	-0.42[-1.35,0.52]
Kuligowski 1998	14	6.2 (6)	14	9.8 (5.5)		31.14%	-0.61[-1.37,0.15]
Skurvydas 2006	20	2.5 (0.5)	20	5.4 (1.7)		27.35%	-2.27[-3.08,-1.46]
Subtotal ***	**52**		**52**			**100%**	**-0.93[-1.36,-0.51]**

Heterogeneity: Tau²=0; Chi²=14.84, df=3(P=0); I²=79.78%

Test for overall effect: Z=4.31(P<0.0001)

1.1.5 96 hours

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
Goodall 2008	9	2.9 (1.8)	9	3.2 (1.9)		20.36%	-0.15[-1.08,0.77]
Jakeman 2009	9	1 (1.5)	9	1.3 (0.9)		20.33%	-0.18[-1.11,0.75]
Kuligowski 1998	14	2.1 (2.7)	14	8 (5.6)		25.33%	-1.32[-2.15,-0.49]
Yanagisawa 2003a	9	2.2 (0.5)	10	2.9 (0.7)		18.13%	-1.09[-2.07,-0.11]
Yanagisawa 2003b	7	1.9 (0.9)	7	1.8 (0.7)		15.86%	0.12[-0.93,1.16]
Subtotal ***	**48**		**49**			**100%**	**-0.58[-1,-0.16]**

Heterogeneity: Tau²=0; Chi²=7.34, df=4(P=0.12); I²=45.52%

Test for overall effect: Z=2.73(P=0.01)

Favours CWI   -2 -1 0 1 2   Favours Passive

Analysis 1.2. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 2 Pain - random effects analysis (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.2.1 Immediately post intervention							
Bailey 2007	10	6.2 (1.9)	10	6.9 (2.2)		15.83%	-0.33[-1.21,0.56]
Cassar 2010	8	2.5 (1)	8	2.2 (1.8)		12.86%	0.19[-0.79,1.18]
Goodall 2008	9	10 (4.9)	9	6.7 (4.5)		13.57%	0.67[-0.29,1.62]
Halson 2008	11	3.8 (2.6)	11	5 (2.9)		17.2%	-0.42[-1.27,0.43]
Jakeman 2009	9	1.6 (1)	9	1.2 (1.2)		14.22%	0.36[-0.57,1.29]
King 2009	10	2.5 (1.5)	10	4 (2.5)		15.01%	-0.69[-1.6,0.22]
Yanagisawa 2003b	7	1.7 (1)	7	1.8 (0.4)		11.32%	-0.12[-1.17,0.93]
Subtotal *	64		64			100%	-0.07[-0.43,0.28]

Heterogeneity: Tau²=0.01; Chi²=6.15, df=6(P=0.41); I²=2.4%

Test for overall effect: Z=0.41(P=0.69)

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.2.2 24 hours							
Bailey 2007	10	4.6 (0.9)	10	6.8 (1.6)		9.04%	-1.62[-2.66,-0.58]
Goodall 2008	9	10.2 (4.8)	9	10.8 (4.2)		9.94%	-0.13[-1.05,0.8]
Ingram 2009	11	3 (1)	11	5 (1)		9.01%	-1.92[-2.97,-0.88]
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		9.67%	0.7[-0.26,1.66]
King 2009	10	1.3 (1)	10	2.4 (1.3)		9.86%	-0.94[-1.88,-0.01]
Kuligowski 1998	14	8.6 (6.6)	14	7.1 (4.5)		11.44%	0.25[-0.5,0.99]
Montgomery 2008	10	3.3 (1.1)	9	4.3 (1.8)		9.9%	-0.65[-1.58,0.28]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		12.16%	-0.98[-1.64,-0.32]
Yanagisawa 2003a	9	3.1 (0.5)	10	3.3 (0.8)		10.09%	-0.28[-1.19,0.62]
Yanagisawa 2003b	7	2.4 (0.8)	7	2.8 (0.9)		8.89%	-0.4[-1.46,0.66]
Subtotal *	109		109			100%	-0.58[-1.06,-0.1]

Heterogeneity: Tau²=0.38; Chi²=25.21, df=9(P=0); I²=64.3%

Test for overall effect: Z=2.36(P=0.02)

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.2.3 48 hours							
Bailey 2007	10	4.4 (1.6)	10	5.7 (0.9)		12.18%	-0.96[-1.9,-0.02]
Goodall 2008	9	11.4 (5.1)	9	11.2 (3.9)		12.33%	0.04[-0.88,0.97]
Ingram 2009	11	3 (1)	11	5 (1)		10.96%	-1.92[-2.97,-0.88]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		12.32%	-0.11[-1.03,0.82]
Kuligowski 1998	14	11 (5.5)	14	10.3 (4.9)		14.67%	0.13[-0.61,0.87]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		15.79%	-0.99[-1.65,-0.33]
Yanagisawa 2003a	9	3.8 (0.4)	10	4.4 (0.5)		11.38%	-1.26[-2.26,-0.25]
Yanagisawa 2003b	7	2.6 (0.6)	7	3.5 (1.4)		10.37%	-0.78[-1.88,0.32]
Subtotal *	89		90			100%	-0.7[-1.18,-0.21]

Heterogeneity: Tau²=0.27; Chi²=16.34, df=7(P=0.02); I²=57.15%

Test for overall effect: Z=2.82(P=0)

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.2.4 72 hours							
Goodall 2008	9	6 (3.2)	9	6.7 (4.2)		24.2%	-0.18[-1.1,0.75]
Jakeman 2009	9	1.6 (1.3)	9	2.3 (1.8)		24.07%	-0.42[-1.35,0.52]
Kuligowski 1998	14	6.2 (6)	14	9.8 (5.5)		26.16%	-0.61[-1.37,0.15]
Skurvydas 2006	20	2.5 (0.5)	20	5.4 (1.7)		25.57%	-2.27[-3.08,-1.46]
Subtotal *	52		52			100%	-0.88[-1.83,0.07]

Heterogeneity: Tau²=0.75; Chi²=14.84, df=3(P=0); I²=79.78%

Test for overall effect: Z=1.82(P=0.07)

Favours CWI   -2 -1 0 1 2   Favours Passive

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.2.5 96 hours							
Goodall 2008	9	2.9 (1.8)	9	3.2 (1.9)		20.31%	-0.15[-1.08,0.77]
Jakeman 2009	9	1 (1.5)	9	1.3 (0.9)		20.3%	-0.18[-1.11,0.75]
Kuligowski 1998	14	2.1 (2.7)	14	8 (5.6)		22.72%	-1.32[-2.15,-0.49]
Yanagisawa 2003a	9	2.2 (0.5)	10	2.9 (0.7)		19.05%	-1.09[-2.07,-0.11]
Yanagisawa 2003b	7	1.9 (0.9)	7	1.8 (0.7)		17.62%	0.12[-0.93,1.16]
Subtotal ***	**48**		**49**			**100%**	**-0.56[-1.12,0.01]**
Heterogeneity: Tau²=0.19; Chi²=7.34, df=4(P=0.12); I²=45.52%							
Test for overall effect: Z=1.92(P=0.06)							

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.3.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 3 Subgroup analysis. Study design: Pain at 24 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
1.3.1 parallel group							
Bailey 2007	10	4.6 (0.9)	10	6.8 (1.6)		7.32%	-1.62[-2.66,-0.58]
Goodall 2008	9	10.2 (4.8)	9	10.8 (4.2)		9.27%	-0.13[-1.05,0.8]
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		8.62%	0.7[-0.26,1.66]
Kuligowski 1998	14	8.6 (6.6)	14	7.1 (4.5)		14.33%	0.25[-0.5,0.99]
Montgomery 2008	10	3.3 (1.1)	9	4.3 (1.8)		9.18%	-0.65[-1.58,0.28]
Yanagisawa 2003a	9	3.1 (0.5)	10	3.3 (0.8)		9.66%	-0.28[-1.19,0.62]
Yanagisawa 2003b	7	2.4 (0.8)	7	2.8 (0.9)		7.04%	-0.4[-1.46,0.66]
Subtotal ***	**68**		**68**			**65.43%**	**-0.23[-0.58,0.12]**
Heterogeneity: Tau²=0; Chi²=13, df=6(P=0.04); I²=53.84%							
Test for overall effect: Z=1.29(P=0.2)							
1.3.2 cross-over							
Ingram 2009	11	3 (1)	11	5 (1)		7.27%	-1.92[-2.97,-0.88]
King 2009	10	1.3 (1)	10	2.4 (1.3)		9.07%	-0.94[-1.88,-0.01]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		18.23%	-0.98[-1.64,-0.32]
Subtotal ***	**41**		**41**			**34.57%**	**-1.17[-1.65,-0.69]**
Heterogeneity: Tau²=0; Chi²=2.55, df=2(P=0.28); I²=21.45%							
Test for overall effect: Z=4.78(P<0.0001)							
Total ***	**109**		**109**			**100%**	**-0.55[-0.84,-0.27]**
Heterogeneity: Tau²=0; Chi²=25.21, df=9(P=0); I²=64.3%							
Test for overall effect: Z=3.85(P=0)							
Test for subgroup differences: Chi²=9.66, df=1 (P=0), I²=89.65%							

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.4.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 4 Subgroup analysis. Study design: Pain at 48 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
1.4.1 parallel group							
Bailey 2007	10	4.4 (1.6)	10	5.7 (0.9)		11.04%	-0.96[-1.9,-0.02]
Goodall 2008	9	11.4 (5.1)	9	11.2 (3.9)		11.35%	0.04[-0.88,0.97]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		11.34%	-0.11[-1.03,0.82]
Kuligowski 1998	14	11 (5.5)	14	10.3 (4.9)		17.62%	0.13[-0.61,0.87]
Yanagisawa 2003a	9	3.8 (0.4)	10	4.4 (0.5)		9.57%	-1.26[-2.26,-0.25]
Yanagisawa 2003b	7	2.6 (0.6)	7	3.5 (1.4)		7.98%	-0.78[-1.88,0.32]
Subtotal ***	**58**		**59**			**68.9%**	**-0.4[-0.77,-0.02]**
Heterogeneity: Tau²=0; Chi²=7.86, df=5(P=0.16); I²=36.39%							
Test for overall effect: Z=2.07(P=0.04)							
1.4.2 cross-over							
Ingram 2009	11	3 (1)	11	5 (1)		8.88%	-1.92[-2.97,-0.88]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		22.22%	-0.99[-1.65,-0.33]
Subtotal ***	**31**		**31**			**31.1%**	**-1.26[-1.81,-0.7]**
Heterogeneity: Tau²=0; Chi²=2.2, df=1(P=0.14); I²=54.49%							
Test for overall effect: Z=4.41(P<0.0001)							
Total ***	**89**		**90**			**100%**	**-0.66[-0.97,-0.35]**
Heterogeneity: Tau²=0; Chi²=16.34, df=7(P=0.02); I²=57.15%							
Test for overall effect: Z=4.18(P<0.0001)							
Test for subgroup differences: Chi²=6.28, df=1 (P=0.01), I²=84.07%							

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.5.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 5 Subgroup analysis. Study design: Pain at 72 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
1.5.1 parallel groups							
Goodall 2008	9	6 (3.2)	9	6.7 (4.2)		20.98%	-0.18[-1.1,0.75]
Jakeman 2009	9	1.6 (1.3)	9	2.3 (1.8)		20.53%	-0.42[-1.35,0.52]
Kuligowski 1998	14	6.2 (6)	14	9.8 (5.5)		31.14%	-0.61[-1.37,0.15]
Subtotal ***	**32**		**32**			**72.65%**	**-0.43[-0.93,0.07]**
Heterogeneity: Tau²=0; Chi²=0.49, df=2(P=0.78); I²=0%							
Test for overall effect: Z=1.69(P=0.09)							
1.5.2 cross-over							
Skurvydas 2006	20	2.5 (0.5)	20	5.4 (1.7)		27.35%	-2.27[-3.08,-1.46]
Subtotal ***	**20**		**20**			**27.35%**	**-2.27[-3.08,-1.46]**
Heterogeneity: Not applicable							
Test for overall effect: Z=5.48(P<0.0001)							
Total ***	**52**		**52**			**100%**	**-0.93[-1.36,-0.51]**
Heterogeneity: Tau²=0; Chi²=14.84, df=3(P=0); I²=79.78%							
Test for overall effect: Z=4.31(P<0.0001)							
Test for subgroup differences: Chi²=14.34, df=1 (P=0), I²=93.03%							

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.6. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 6 Subgroup analysis. Immersion frequency: Pain at 24 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.6.1 Single CWI							
Bailey 2007	10	4.6 (0.9)	10	6.8 (1.6)		7.32%	-1.62[-2.66,-0.58]
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		8.62%	0.7[-0.26,1.66]
King 2009	10	1.3 (1)	10	2.4 (1.3)		9.07%	-0.94[-1.88,-0.01]
Montgomery 2008	10	3.3 (1.1)	9	4.3 (1.8)		9.18%	-0.65[-1.58,0.28]
Yanagisawa 2003a	9	3.1 (0.5)	10	3.3 (0.8)		9.66%	-0.28[-1.19,0.62]
Yanagisawa 2003b	7	2.4 (0.8)	7	2.8 (0.9)		7.04%	-0.4[-1.46,0.66]
Subtotal ***	**55**		**55**			**50.89%**	**-0.51[-0.9,-0.11]**
Heterogeneity: Tau²=0; Chi²=11.69, df=5(P=0.04); I²=57.23%							
Test for overall effect: Z=2.53(P=0.01)							
1.6.2 Multiple CWI							
Goodall 2008	9	10.2 (4.8)	9	10.8 (4.2)		9.27%	-0.13[-1.05,0.8]
Ingram 2009	11	3 (1)	11	5 (1)		7.27%	-1.92[-2.97,-0.88]
Kuligowski 1998	14	8.6 (6.6)	14	7.1 (4.5)		14.33%	0.25[-0.5,0.99]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		18.23%	-0.98[-1.64,-0.32]
Subtotal ***	**54**		**54**			**49.11%**	**-0.6[-1,-0.2]**
Heterogeneity: Tau²=0; Chi²=13.42, df=3(P=0); I²=77.64%							
Test for overall effect: Z=2.93(P=0)							
Total ***	**109**		**109**			**100%**	**-0.55[-0.84,-0.27]**
Heterogeneity: Tau²=0; Chi²=25.21, df=9(P=0); I²=64.3%							
Test for overall effect: Z=3.85(P=0)							
Test for subgroup differences: Chi²=0.1, df=1 (P=0.75), I²=0%							

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.7. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 7 Subgroup analysis. Immersion frequency: Pain at 48 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.7.1 Single CWI							
Bailey 2007	10	4.4 (1.6)	10	5.7 (0.9)		11.04%	-0.96[-1.9,-0.02]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		11.34%	-0.11[-1.03,0.82]
Yanagisawa 2003a	9	3.8 (0.4)	10	4.4 (0.5)		9.57%	-1.26[-2.26,-0.25]
Yanagisawa 2003b	7	2.6 (0.6)	7	3.5 (1.4)		7.98%	-0.78[-1.88,0.32]
Subtotal ***	**35**		**36**			**39.93%**	**-0.75[-1.25,-0.26]**
Heterogeneity: Tau²=0; Chi²=3.02, df=3(P=0.39); I²=0.78%							
Test for overall effect: Z=3(P=0)							
1.7.2 Multiple CWI							
Goodall 2008	9	11.4 (5.1)	9	11.2 (3.9)		11.35%	0.04[-0.88,0.97]
Ingram 2009	11	3 (1)	11	5 (1)		8.88%	-1.92[-2.97,-0.88]
Kuligowski 1998	14	11 (5.5)	14	10.3 (4.9)		17.62%	0.13[-0.61,0.87]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		22.22%	-0.99[-1.65,-0.33]

Favours CWI -2 -1 0 1 2 Favours Passive

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Subtotal ***	54		54			60.07%	-0.6[-1.01,-0.2]

Heterogeneity: Tau²=0; Chi²=13.1, df=3(P=0); I²=77.1%

Test for overall effect: Z=2.94(P=0)

| Total *** | 89 | | 90 | | | 100% | -0.66[-0.97,-0.35] |

Heterogeneity: Tau²=0; Chi²=16.34, df=7(P=0.02); I²=57.15%

Test for overall effect: Z=4.18(P<0.0001)

Test for subgroup differences: Chi²=0.21, df=1 (P=0.64), I²=0%

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.8.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 8 Subgroup analysis. Exercise type: Pain at 24 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.8.1 Lab controlled muscle damage (DOMS)							
Goodall 2008	9	10.2 (4.8)	9	10.8 (4.2)		9.27%	-0.13[-1.05,0.8]
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		8.62%	0.7[-0.26,1.66]
Kuligowski 1998	14	8.6 (6.6)	14	7.1 (4.5)		14.33%	0.25[-0.5,0.99]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		18.23%	-0.98[-1.64,-0.32]
Yanagisawa 2003a	9	3.1 (0.5)	10	3.3 (0.8)		9.66%	-0.28[-1.19,0.62]
Yanagisawa 2003b	7	2.4 (0.8)	7	2.8 (0.9)		7.04%	-0.4[-1.46,0.66]
Subtotal ***	68		69			67.16%	-0.22[-0.57,0.12]

Heterogeneity: Tau²=0; Chi²=10.28, df=5(P=0.07); I²=51.38%

Test for overall effect: Z=1.27(P=0.2)

1.8.2 Other sporting activity							
Bailey 2007	10	4.6 (0.9)	10	6.8 (1.6)		7.32%	-1.62[-2.66,-0.58]
Ingram 2009	11	3 (1)	11	5 (1)		7.27%	-1.92[-2.97,-0.88]
King 2009	10	1.3 (1)	10	2.4 (1.3)		9.07%	-0.94[-1.88,-0.01]
Montgomery 2008	10	3.3 (1.1)	9	4.3 (1.8)		9.18%	-0.65[-1.58,0.28]
Subtotal ***	41		40			32.84%	-1.23[-1.72,-0.74]

Heterogeneity: Tau²=0; Chi²=4.1, df=3(P=0.25); I²=26.83%

Test for overall effect: Z=4.9(P<0.0001)

| Total *** | 109 | | 109 | | | 100% | -0.55[-0.84,-0.27] |

Heterogeneity: Tau²=0; Chi²=25.21, df=9(P=0); I²=64.3%

Test for overall effect: Z=3.85(P=0)

Test for subgroup differences: Chi²=10.82, df=1 (P=0), I²=90.76%

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.9.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 9 Subgroup analysis. Exercise type: Pain at 48 hours (muscle soreness: various scales Likert and VAS).

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.9.1 Lab controlled muscle damage (DOMS)							

Favours CWI -2 -1 0 1 2 Favours Passive

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
Goodall 2008	9	11.4 (5.1)	9	11.2 (3.9)		11.35%	0.04[-0.88,0.97]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		11.34%	-0.11[-1.03,0.82]
Kuligowski 1998	14	11 (5.5)	14	10.3 (4.9)		17.62%	0.13[-0.61,0.87]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		22.22%	-0.99[-1.65,-0.33]
Yanagisawa 2003a	9	3.8 (0.4)	10	4.4 (0.5)		9.57%	-1.26[-2.26,-0.25]
Yanagisawa 2003b	7	2.6 (0.6)	7	3.5 (1.4)		7.98%	-0.78[-1.88,0.32]
Subtotal ***	**68**		**69**			**80.09%**	**-0.48[-0.83,-0.14]**

Heterogeneity: Tau²=0; Chi²=9.33, df=5(P=0.1); I²=46.42%

Test for overall effect: Z=2.72(P=0.01)

1.9.2 Other sporting activity							
Bailey 2007	10	4.4 (1.6)	10	5.7 (0.9)		11.04%	-0.96[-1.9,-0.02]
Ingram 2009	11	3 (1)	11	5 (1)		8.88%	-1.92[-2.97,-0.88]
Subtotal ***	**21**		**21**			**19.91%**	**-1.39[-2.09,-0.69]**

Heterogeneity: Tau²=0; Chi²=1.81, df=1(P=0.18); I²=44.9%

Test for overall effect: Z=3.9(P<0.0001)

| **Total ***** | **89** | | **90** | | | **100%** | **-0.66[-0.97,-0.35]** |

Heterogeneity: Tau²=0; Chi²=16.34, df=7(P=0.02); I²=57.15%

Test for overall effect: Z=4.18(P<0.0001)

Test for subgroup differences: Chi²=5.19, df=1 (P=0.02), I²=80.73%

Favours CWI -2 -1 0 1 2 Favours Passive

Analysis 1.10. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 10 Tenderness (pain on palpation).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
1.10.1 24 hours						
Eston 1999	8	22.6 (20.4)	7	26.9 (10.1)		-4.3[-20.29,11.69]
1.10.2 48 hours						
Eston 1999	8	24.9 (17.9)	7	26.7 (14.6)		-1.8[-18.26,14.66]
1.10.3 72 hours						
Eston 1999	8	22.1 (15.1)	7	21.9 (17.9)		0.2[-16.69,17.09]

Favours CWI -20 -10 0 10 20 Favours Passive

Analysis 1.11. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 11 Subjective recovery (10 point or 10 cm VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
1.11.1 Physical recovery (Immediately post intervention)							
Buchheit 2009	10	6.5 (2.1)	10	4.5 (2)		35.72%	2[0.2,3.8]
Halson 2008	11	6.8 (1.5)	11	6.4 (1.7)		64.28%	0.4[-0.94,1.74]
Subtotal ***	**21**		**21**			**100%**	**0.97[-0.1,2.05]**

Favours passive -2 -1 0 1 2 Favours CWI

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
Heterogeneity: Tau²=0; Chi²=1.96, df=1(P=0.16); I²=48.9%							
Test for overall effect: Z=1.77(P=0.08)							
1.11.2 Mental recovery (immediately post intervention)							
Halson 2008	11	6.7 (1.8)	11	6.1 (1.7)		100%	0.6[-0.86,2.06]
Subtotal ***	**11**		**11**			**100%**	**0.6[-0.86,2.06]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.8(P=0.42)							

Favours passive   -2   -1   0   1   2   Favours CWI

Analysis 1.12. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 12 Fatigue (10 point / 10 cm VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
1.12.1 Fatigue (immediately post intervention)							
Cassar 2010	8	6.9 (0.7)	8	8.8 (1.1)		77.39%	-1.9[-2.8,-1]
Halson 2008	11	5.3 (2)	11	6.3 (2)		22.61%	-1[-2.67,0.67]
Subtotal ***	**19**		**19**			**100%**	**-1.7[-2.49,-0.9]**
Heterogeneity: Tau²=0; Chi²=0.86, df=1(P=0.35); I²=0%							
Test for overall effect: Z=4.18(P<0.0001)							
1.12.2 Fatigue (24 hrs post intervention)							
Montgomery 2008	10	4.5 (0.9)	9	5.2 (1.6)		100%	-0.7[-1.88,0.48]
Subtotal ***	**10**		**9**			**100%**	**-0.7[-1.88,0.48]**
Heterogeneity: Not applicable							
Test for overall effect: Z=1.16(P=0.25)							

Favours CWI   -2   -1   0   1   2   Favours passive

Analysis 1.13. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 13 Strength (Final value: Nm).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
1.13.1 Immediate							
Bailey 2007	10	401 (98)	10	316 (111)		2.11%	85[-6.77,176.77]
Buchheit 2009	10	116 (20)	10	150 (24)		47.34%	-34[-53.36,-14.64]
Jakeman 2009	9	124.5 (17.4)	9	135.6 (22.8)		50.55%	-11.1[-29.84,7.64]
Subtotal ***	**29**		**29**			**100%**	**-19.92[-33.24,-6.59]**
Heterogeneity: Tau²=0; Chi²=7.9, df=2(P=0.02); I²=74.69%							
Test for overall effect: Z=2.93(P=0)							
1.13.2 24 hours							
Bailey 2007	10	402 (107)	10	356 (123)		0.2%	46[-55.04,147.04]
Eston 1999	8	78.3 (59.9)	7	73 (75.4)		0.42%	5.3[-64.29,74.89]
Jakeman 2009	9	130.1 (15)	9	136.5 (25.1)		5.55%	-6.38[-25.48,12.72]
Kuligowski 1998	14	10.6 (4.4)	14	12.4 (7.7)		93.83%	-1.81[-6.46,2.84]
Subtotal ***	**41**		**40**			**100%**	**-1.94[-6.44,2.56]**

Favours passive   -100 -50   0   50   100   Favours CWI

Study or subgroup	CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Heterogeneity: Tau²=0; Chi²=1.12, df=3(P=0.77); I²=0%							
Test for overall effect: Z=0.84(P=0.4)							
1.13.3 48 hours							
Bailey 2007	10	413 (104)	10	371 (117)		0.17%	42[-55.02,139.02]
Eston 1999	8	92.5 (87.4)	7	81.3 (109.5)		0.16%	11.2[-90.03,112.43]
Ingram 2009	11	73.4 (18.5)	11	72.2 (10.7)		10.1%	1.2[-11.43,13.83]
Jakeman 2009	9	122.7 (19)	9	132 (30.8)		2.88%	-9.27[-32.9,14.36]
Kuligowski 1998	14	10.9 (4.4)	14	12.5 (7)		86.69%	-1.6[-5.91,2.71]
Subtotal ***	52		51			100%	-1.44[-5.46,2.57]
Heterogeneity: Tau²=0; Chi²=1.42, df=4(P=0.84); I²=0%							
Test for overall effect: Z=0.71(P=0.48)							
1.13.4 72 hours							
Eston 1999	8	112.3 (76.2)	7	86.4 (117.8)		0.19%	25.9[-76.1,127.9]
Jakeman 2009	9	136 (18.3)	9	139.1 (31.9)		3.44%	-3.07[-27.06,20.92]
Kuligowski 1998	14	11.9 (5.6)	14	12.7 (6.6)		96.37%	-0.86[-5.39,3.67]
Subtotal ***	31		30			100%	-0.89[-5.33,3.56]
Heterogeneity: Tau²=0; Chi²=0.3, df=2(P=0.86); I²=0%							
Test for overall effect: Z=0.39(P=0.7)							
1.13.5 96 hours							
Jakeman 2009	9	138.3 (18.1)	9	153 (25.9)		2.85%	-14.73[-35.33,5.87]
Kuligowski 1998	14	12.9 (5.3)	14	13 (4.1)		97.15%	-0.17[-3.7,3.36]
Subtotal ***	23		23			100%	-0.58[-4.06,2.89]
Heterogeneity: Tau²=0; Chi²=1.86, df=1(P=0.17); I²=46.35%							
Test for overall effect: Z=0.33(P=0.74)							

Favours passive   -100 -50 0 50 100   Favours CWI

Analysis 1.14. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 14 Strength (% of baseline).

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Random, 95% CI	Random, 95% CI
1.14.1 Immediate						
Goodall 2008	9	74.5 (15.9)	9	81.6 (13.8)		-7.1[-20.85,6.65]
Skurvydas 2006	20	84.7 (17)	20	85.8 (10.7)		-1.1[-9.9,7.7]
1.14.2 24 hours						
Goodall 2008	9	87.4 (18.5)	9	82.1 (13.8)		5.3[-9.78,20.38]
Skurvydas 2006	20	97.7 (21.9)	20	75 (16.6)		22.7[10.66,34.74]
1.14.3 48 hours						
Goodall 2008	9	94.3 (22.8)	9	91 (10.6)		3.3[-13.13,19.73]
Skurvydas 2006	20	91.6 (18.1)	20	64.4 (18.9)		27.2[15.73,38.67]
1.14.4 72 hours						
Goodall 2008	9	96.4 (21.8)	9	94.8 (9.6)		1.6[-13.96,17.16]
Skurvydas 2006	20	101.5 (20.6)	20	76.3 (18.5)		25.2[13.07,37.33]

Favours passive   -100 -50 0 50 100   Favours CWI

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Random, 95% CI	Random, 95% CI
1.14.5 96 hours						
Goodall 2008	9	100 (18)	9	97.2 (9.6)		2.8[-10.53,16.13]

Favours passive -100 -50 0 50 100 Favours CWI

Analysis 1.15. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 15 Power (jump height: centimetres).

Study or subgroup	Favours CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.15.1 Immediate (cross-over trial)							
Skurvydas 2006	20	30.7 (9)	20	32.5 (4.9)		100%	-1.8[-6.29,2.69]
Subtotal ***	20		20			100%	-1.8[-6.29,2.69]
Heterogeneity: Not applicable							
Test for overall effect: Z=0.79(P=0.43)							
1.15.2 24 hours							
Bailey 2007	10	35 (3)	10	35 (3)		76.38%	0[-2.63,2.63]
Montgomery 2008	10	61.6 (6.5)	9	58.7 (6.7)		14.92%	2.9[-3.05,8.85]
Rowsell 2009	6	46 (8)	7	48 (6)		8.7%	-2[-9.79,5.79]
Subtotal ***	26		26			100%	0.26[-2.04,2.56]
Heterogeneity: Tau²=0; Chi²=1.12, df=2(P=0.57); I²=0%							
Test for overall effect: Z=0.22(P=0.83)							
1.15.3 24 hours (cross-over trial)							
Skurvydas 2006	20	34.2 (6.9)	20	28.8 (2.9)		100%	5.4[2.12,8.68]
Subtotal ***	20		20			100%	5.4[2.12,8.68]
Heterogeneity: Not applicable							
Test for overall effect: Z=3.23(P=0)							
1.15.4 48 hours							
Bailey 2007	10	34 (3)	10	34 (3)		100%	0[-2.63,2.63]
Subtotal ***	10		10			100%	0[-2.63,2.63]
Heterogeneity: Not applicable							
Test for overall effect: Not applicable							
1.15.5 48 hours (cross-over trial)							
Skurvydas 2006	20	35.2 (6.4)	20	25.6 (3.5)		100%	9.6[6.4,12.8]
Subtotal ***	20		20			100%	9.6[6.4,12.8]
Heterogeneity: Not applicable							
Test for overall effect: Z=5.89(P<0.0001)							
1.15.6 72 hours (cross-over trial)							
Skurvydas 2006	20	35.5 (5.8)	20	28.9 (5.3)		100%	6.6[3.16,10.04]
Subtotal ***	20		20			100%	6.6[3.16,10.04]
Heterogeneity: Not applicable							
Test for overall effect: Z=3.76(P=0)							

Favours passive -10 -5 0 5 10 Favours CWI

Analysis 1.16. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 16 Power (% decrease in jump height over 5 jumps).

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
1.16.1 24 hours						
King 2009	10	4.4 (2.7)	10	8.1 (4.9)		-3.7[-7.17,-0.23]

Favours CWI   -10 -5 0 5 10   Favours passive

Analysis 1.17. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 17 Power (cycle ergometer power: Watts).

Study or subgroup	CWI		Control		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
1.17.1 Peak power output (Immediate)						
Buchheit 2009	10	700 (62)	10	720 (80)		-20[-82.73,42.73]

Favours passive   -100 -50 0 50 100   Favours CWI

Analysis 1.18. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 18 Functional performance (time to complete exercise test: seconds).

Study or subgroup	CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.18.1 1 km cycle time (immediate)							
Buchheit 2009	10	81.5 (3.2)	10	81.2 (3.5)		100%	0.3[-2.64,3.24]
Subtotal ***	10		10			100%	0.3[-2.64,3.24]
Heterogeneity: Tau²=0; Chi²=0, df=0(P<0.0001); I²=100%							
Test for overall effect: Z=0.2(P=0.84)							
1.18.2 Agility circuit time (24 hours)							
Montgomery 2008	10	6.6 (0.2)	9	6.6 (0.3)		100%	-0.03[-0.26,0.2]
Subtotal ***	10		9			100%	-0.03[-0.26,0.2]
Heterogeneity: Not applicable							
Test for overall effect: Z=0.26(P=0.79)							
1.18.3 Sprint time (24 hours)							
King 2009	10	2.3 (0.2)	10	2.4 (0.2)		29.99%	-0.03[-0.19,0.13]
Montgomery 2008	10	3.1 (0.1)	9	3.1 (0.1)		70.01%	-0.01[-0.11,0.09]
Subtotal ***	20		19			100%	-0.02[-0.1,0.07]
Heterogeneity: Tau²=0; Chi²=0.04, df=1(P=0.84); I²=0%							
Test for overall effect: Z=0.36(P=0.72)							
1.18.4 Multisprint / Agilty circuit time (24 hours)							
King 2009	10	34.5 (2.9)	10	34.7 (2)		100%	-0.29[-2.49,1.91]
Subtotal ***	10		10			100%	-0.29[-2.49,1.91]
Heterogeneity: Not applicable							
Test for overall effect: Z=0.26(P=0.8)							
1.18.5 Multisprint time (24 hours)							

Favours CWI   -5 -2.5 0 2.5 5   Favours Passive

Study or subgroup	CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Montgomery 2008	10	27.3 (1)	9	27.6 (1.2)		77.43%	-0.33[-1.32,0.66]
Rowsell 2009	6	43.3 (2.1)	7	43 (1)		22.57%	0.28[-1.55,2.11]
Subtotal ***	**16**		**16**			**100%**	**-0.19[-1.06,0.68]**
Heterogeneity: Tau²=0; Chi²=0.33, df=1(P=0.57); I²=0%							
Test for overall effect: Z=0.43(P=0.67)							
1.18.6 Sprint time (48 hours)							
Bailey 2007	10	2.7 (0.2)	10	2.7 (0.1)		100%	0.02[-0.11,0.15]
Subtotal ***	**10**		**10**			**100%**	**0.02[-0.11,0.15]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.31(P=0.76)							
1.18.7 Multisprint time (48 hours)							
Ingram 2009	11	48.2 (1.8)	11	48.3 (1.3)		100%	-0.07[-1.37,1.23]
Subtotal ***	**11**		**11**			**100%**	**-0.07[-1.37,1.23]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.11(P=0.92)							

Favours CWI -5 -2.5 0 2.5 5 Favours Passive

Analysis 1.19. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 19 Functional performance (time to fatigue).

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
1.19.1 % decrease from baseline (immediate)						
Cassar 2010	8	17.7 (4.7)	8	30.8 (19.7)		-13.1[-27.13,0.93]

Favours CWI -20 -10 0 10 20 Favours passive

Analysis 1.20. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 20 Range of movement (ROM).

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
1.20.1 Degrees (Immediate)						
Goodall 2008	9	127.6 (4.6)	9	128.3 (8.1)		-0.7[-6.79,5.39]
Yanagisawa 2003a	9	30.6 (7.8)	10	19.5 (6.4)		11.13[4.68,17.58]
1.20.2 Degrees (24 hours)						
Goodall 2008	9	128.2 (6.4)	9	125.6 (11.5)		2.6[-6,11.2]
Kuligowski 1998	14	7.2 (7.4)	14	8.4 (5.4)		-1.21[-6.03,3.61]
Yanagisawa 2003a	9	33.9 (6)	10	20.5 (5.9)		13.39[8.02,18.76]
1.20.3 Centimetres (24 hours)						
Montgomery 2008	10	8.6 (11.5)	9	3.2 (6.9)		5.4[-3.03,13.83]
1.20.4 Degrees (48 hours)						
Goodall 2008	9	128.3 (11.4)	9	124.9 (9.8)		3.4[-6.42,13.22]
Kuligowski 1998	14	13.8 (12)	14	13.6 (7)		0.26[-7.03,7.55]

Favours passive -50 -25 0 25 50 Favours CWI

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
Yanagisawa 2003a	9	32.2 (6.2)	10	16.5 (7.1)		15.72[9.75,21.69]
1.20.5 Degrees (72 hours)						
Goodall 2008	9	129.1 (9)	9	125.2 (9.4)		3.9[-4.6,12.4]
Kuligowski 1998	14	14.2 (12.4)	14	17.4 (14.2)		-3.17[-13.03,6.69]
1.20.6 Degrees (96 hours)						
Goodall 2008	9	130 (7.1)	9	125.8 (11.2)		4.2[-4.46,12.86]
Kuligowski 1998	14	9.5 (9.3)	14	16.6 (13.6)		-7.15[-15.76,1.46]
Yanagisawa 2003a	9	32.2 (7.1)	10	22.5 (6.8)		9.7[3.44,15.96]

Favours passive   -50 -25 0 25 50   Favours CWI

Analysis 1.21. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 21 Swelling (limb girth in centimetres).

Study or subgroup	CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.21.1 Final value immediate							
Goodall 2008	9	55.5 (3.8)	9	55.4 (7.1)		100%	0.1[-5.16,5.36]
Subtotal ***	**9**		**9**			**100%**	**0.1[-5.16,5.36]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.04(P=0.97)							
1.21.2 Final value 24 hours							
Eston 1999	8	27.3 (6.2)	7	30.2 (6.2)		44.93%	-2.9[-9.19,3.39]
Goodall 2008	9	55.4 (4.4)	9	55.6 (7.5)		55.07%	-0.2[-5.88,5.48]
Subtotal ***	**17**		**16**			**100%**	**-1.41[-5.63,2.8]**
Heterogeneity: Tau²=0; Chi²=0.39, df=1(P=0.53); I²=0%							
Test for overall effect: Z=0.66(P=0.51)							
1.21.3 Final value 48 hours							
Eston 1999	8	27.3 (5.8)	7	30.4 (6.2)		43.81%	-3.1[-9.2,3]
Goodall 2008	9	55.3 (4.2)	9	55.6 (7.1)		56.19%	-0.3[-5.69,5.09]
Subtotal ***	**17**		**16**			**100%**	**-1.53[-5.57,2.51]**
Heterogeneity: Tau²=0; Chi²=0.45, df=1(P=0.5); I²=0%							
Test for overall effect: Z=0.74(P=0.46)							
1.21.4 Final value 72 hours							
Eston 1999	8	27.1 (6.2)	7	30.4 (7)		37.2%	-3.3[-10.03,3.43]
Goodall 2008	9	55.2 (3.9)	9	55.7 (6.9)		62.8%	-0.5[-5.68,4.68]
Subtotal ***	**17**		**16**			**100%**	**-1.54[-5.65,2.56]**
Heterogeneity: Tau²=0; Chi²=0.42, df=1(P=0.52); I²=0%							
Test for overall effect: Z=0.74(P=0.46)							
1.21.5 Final value 96 hours							
Goodall 2008	9	55 (3.8)	9	55.6 (7)		100%	-0.6[-5.8,4.6]
Subtotal ***	**9**		**9**			**100%**	**-0.6[-5.8,4.6]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.23(P=0.82)							

Favours CWI   -10 -5 0 5 10   Favours passive

Analysis 1.22. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 22 Biomarker: creatine kinase.

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
1.22.1 Immediately post							
Bailey 2007	10	910 (411)	10	1028 (490)		27.94%	-0.25[-1.13,0.63]
Cassar 2010	8	172.3 (31.4)	8	245.5 (129.1)		23.87%	-0.74[-1.76,0.29]
Halson 2008	7	133.1 (33.7)	7	127.1 (48.4)		23.19%	0.13[-0.91,1.18]
Jakeman 2009	9	177.2 (82.2)	9	118.2 (34.5)		25%	0.89[-0.09,1.87]
Subtotal ***	**34**		**34**			**100%**	**0.01[-0.66,0.67]**
Heterogeneity: Tau²=0.21; Chi²=5.54, df=3(P=0.14); I²=45.81%							
Test for overall effect: Z=0.02(P=0.98)							
1.22.2 24 hours							
Bailey 2007	10	1246 (354)	10	1358 (474)		14.61%	-0.26[-1.14,0.62]
Eston 1999	8	5 (1.8)	7	4.8 (1.1)		13.85%	0.12[-0.9,1.13]
Goodall 2008	9	432.7 (253.9)	9	272.2 (198.9)		14.19%	0.67[-0.29,1.63]
Ingram 2009	11	571 (375)	11	681 (720)		14.84%	-0.18[-1.02,0.65]
Jakeman 2009	9	200.8 (125.1)	9	141.5 (88)		14.26%	0.52[-0.42,1.47]
Rowsell 2009	6	1075 (680)	7	881 (627)		13.38%	0.28[-0.82,1.37]
Skurvydas 2006	20	693 (201)	20	1293 (281)		14.87%	-2.41[-3.24,-1.57]
Subtotal ***	**73**		**73**			**100%**	**-0.2[-1.03,0.63]**
Heterogeneity: Tau²=1.02; Chi²=33.28, df=6(P<0.0001); I²=81.97%							
Test for overall effect: Z=0.47(P=0.64)							
1.22.3 48 hours							
Bailey 2007	10	529 (275)	10	766 (461)		14.41%	-0.6[-1.5,0.3]
Eston 1999	8	4.8 (1.7)	7	5.5 (2.2)		13.78%	-0.34[-1.36,0.69]
Goodall 2008	9	267.7 (161.9)	9	190.1 (93.5)		14.18%	0.56[-0.39,1.51]
Ingram 2009	11	337 (218)	11	391 (436)		14.73%	-0.15[-0.99,0.69]
Jakeman 2009	9	129.7 (91.8)	9	104.9 (54.6)		14.26%	0.31[-0.62,1.24]
Skurvydas 2006	20	542 (189)	20	1350 (320)		14.26%	-3.01[-3.95,-2.08]
Yanagisawa 2003a	9	1200 (1700)	10	2500 (6500)		14.39%	-0.25[-1.16,0.65]
Subtotal ***	**76**		**76**			**100%**	**-0.5[-1.36,0.37]**
Heterogeneity: Tau²=1.14; Chi²=36.79, df=6(P<0.0001); I²=83.69%							
Test for overall effect: Z=1.13(P=0.26)							
1.22.4 72 hours							
Eston 1999	8	4.7 (1.5)	7	6 (4.4)		30.67%	-0.37[-1.4,0.65]
Goodall 2008	9	197.7 (103.8)	9	131 (53)		33.52%	0.77[-0.2,1.74]
Jakeman 2009	9	110.3 (50.3)	9	111.6 (74.5)		35.81%	-0.02[-0.94,0.9]
Subtotal ***	**26**		**25**			**100%**	**0.14[-0.52,0.79]**
Heterogeneity: Tau²=0.09; Chi²=2.72, df=2(P=0.26); I²=26.36%							
Test for overall effect: Z=0.41(P=0.68)							
1.22.5 96 hours							
Goodall 2008	9	164.4 (92)	9	126.6 (76.7)		33.25%	0.43[-0.51,1.36]
Jakeman 2009	9	135.7 (85.5)	9	110 (63.7)		33.47%	0.32[-0.61,1.26]

Favours CWI -5 -2.5 0 2.5 5 Favours passive

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Random, 95% CI		Random, 95% CI
Yanagisawa 2003a	9	2500 (3750)	10	12000 (17000)		33.28%	-0.72[-1.65,0.22]
Subtotal ***	**27**		**28**			**100%**	**0.01[-0.7,0.73]**
Heterogeneity: Tau²=0.17; Chi²=3.52, df=2(P=0.17); I²=43.11%							
Test for overall effect: Z=0.03(P=0.98)							

Favours CWI -5 -2.5 0 2.5 5 Favours passive

Analysis 1.23. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 23 Biomarker: lactate dehydrogenase.

Study or subgroup	CWI		Passive		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
1.23.1 U/L (immediate)						
Cassar 2010	8	192 (15)	8	192 (70.7)		0[-50.08,50.08]
1.23.2 % change 96 hours						
Yanagisawa 2003a	9	25 (50)	10	125 (200)		-100[-228.19,28.19]

Favours CWI -200 -100 0 100 200 Favours passive

Analysis 1.24. Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 24 Biomarker: myoglobin.

Study or subgroup	CWI		Passive		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
1.24.1 Immediate							
Bailey 2007	10	29.5 (15.2)	10	45.9 (17.1)		52.15%	-0.97[-1.91,-0.03]
Cassar 2010	8	0 (0)	8	0 (0)		47.85%	0[-0.98,0.98]
Subtotal ***	**18**		**18**			**100%**	**-0.51[-1.18,0.17]**
Heterogeneity: Tau²=0; Chi²=1.97, df=1(P=0.16); I²=49.15%							
Test for overall effect: Z=1.46(P=0.14)							
1.24.2 24 hours							
Bailey 2007	10	5.2 (0.9)	10	8.1 (4.7)		59.09%	-0.82[-1.74,0.1]
Rowsell 2009	6	48 (13)	7	69 (62)		40.91%	-0.42[-1.53,0.69]
Subtotal ***	**16**		**17**			**100%**	**-0.66[-1.36,0.05]**
Heterogeneity: Tau²=0; Chi²=0.3, df=1(P=0.58); I²=0%							
Test for overall effect: Z=1.82(P=0.07)							
1.24.3 48 hours							
Bailey 2007	10	4.2 (1.3)	10	5.6 (2.2)		100%	-0.74[-1.66,0.17]
Subtotal ***	**10**		**10**			**100%**	**-0.74[-1.66,0.17]**
Heterogeneity: Not applicable							
Test for overall effect: Z=1.59(P=0.11)							

Favours CWI -2 -1 0 1 2 Favours passive

Analysis 1.25.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 25 Biomarker: interleukin-6 (pg/ml).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Std. Mean Difference Fixed, 95% CI
1.25.1 Immediate						
Halson 2008	7	4 (1.5)	7	4 (1.6)		0[-1.05,1.05]

Favours CWI -2 -1 0 1 2 Favours passive

Analysis 1.26.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 26 Biomarker: C-reactive protein (mg/dL / mg/L).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Std. Mean Difference Fixed, 95% CI	Weight	Std. Mean Difference Fixed, 95% CI
1.26.1 Immediate							
Cassar 2010	8	0.8 (0.8)	8	0.7 (0.4)		53.88%	0.23[-0.76,1.21]
Halson 2008	7	0.1 (0.1)	7	0.1 (0.1)		46.12%	0.42[-0.64,1.49]
Subtotal ***	**15**		**15**			**100%**	**0.32[-0.4,1.04]**
Heterogeneity: Tau²=0; Chi²=0.07, df=1(P=0.79); I²=0%							
Test for overall effect: Z=0.86(P=0.39)							
1.26.2 24 hours							
Ingram 2009	11	2.5 (1.7)	11	2.9 (1.7)		100%	-0.23[-1.07,0.61]
Subtotal ***	**11**		**11**			**100%**	**-0.23[-1.07,0.61]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.53(P=0.6)							
1.26.3 48 hours							
Ingram 2009	11	1.4 (0.8)	11	1.6 (1)		100%	-0.21[-1.05,0.63]
Subtotal ***	**11**		**11**			**100%**	**-0.21[-1.05,0.63]**
Heterogeneity: Tau²=0; Chi²=0, df=0(P<0.0001); I²=100%							
Test for overall effect: Z=0.5(P=0.62)							

Favours CWI -2 -1 0 1 2 Favours passive

Analysis 1.27.   Comparison 1 Cold-water immersion (CWI) versus passive, Outcome 27 Pain (muscle soreness: all converted to 10 cm VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Passive N	Passive Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
1.27.1 Immediately post intervention							
Bailey 2007	10	5.7 (2.1)	10	6.5 (2.4)		9.2%	-0.77[-2.75,1.21]
Cassar 2010	8	2.5 (1)	8	2.2 (1.8)		17.74%	0.3[-1.13,1.73]
Goodall 2008	9	5 (2.3)	9	3.4 (2.3)		8.36%	1.65[-0.43,3.73]
Halson 2008	11	3.8 (2.6)	11	5 (2.9)		6.82%	-1.2[-3.5,1.1]
Jakeman 2009	9	1.6 (1)	9	1.2 (1.2)		38.04%	0.4[-0.57,1.37]
King 2009	10	2.5 (1.5)	10	4 (2.5)		10.77%	-1.5[-3.33,0.33]
Yanagisawa 2003b	7	1.8 (2.5)	7	2 (1)		9.08%	-0.25[-2.24,1.74]
Subtotal ***	**64**		**64**			**100%**	**0.01[-0.59,0.61]**
Heterogeneity: Tau²=0; Chi²=7.5, df=6(P=0.28); I²=19.98%							

Favours CWI -10 -5 0 5 10 Favours Passive

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Study or subgroup	CWI		Passive		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Test for overall effect: Z=0.02(P=0.98)							
1.27.2 24 hours							
Bailey 2007	10	4 (1)	10	6.4 (1.8)		11.69%	-2.42[-3.67,-1.17]
Goodall 2008	9	5.1 (2.4)	9	5.4 (2.1)		4.24%	-0.3[-2.38,1.78]
Ingram 2009	11	2.2 (1.1)	11	4.4 (1.1)		21.77%	-2.2[-3.12,-1.28]
Jakeman 2009	9	3.4 (1.8)	9	2.3 (1.3)		9.08%	1.13[-0.29,2.55]
King 2009	10	1.3 (1)	10	2.4 (1.3)		19.23%	-1.1[-2.08,-0.12]
Kuligowski 1998	14	7.3 (5.5)	14	6 (3.7)		1.53%	1.34[-2.13,4.81]
Montgomery 2008	10	2.5 (1.2)	9	3.6 (2)		8.23%	-1.1[-2.6,0.4]
Skurvydas 2006	20	5.2 (1.9)	20	7.2 (2.1)		11.94%	-2[-3.24,-0.76]
Yanagisawa 2003a	9	5.3 (1.3)	10	5.8 (2)		8.35%	-0.5[-1.98,0.98]
Yanagisawa 2003b	7	3.5 (2)	7	4.4 (2.1)		3.93%	-0.88[-3.04,1.28]
Subtotal ***	**109**		**109**			**100%**	**-1.27[-1.7,-0.84]**
Heterogeneity: Tau²=0; Chi²=23.75, df=9(P=0); I²=62.1%							
Test for overall effect: Z=5.8(P<0.0001)							
1.27.3 48 hours							
Bailey 2007	10	3.7 (1.8)	10	5.2 (1)		15.19%	-1.43[-2.68,-0.18]
Goodall 2008	9	5.7 (2.6)	9	5.6 (2)		5.44%	0.1[-2,2.2]
Ingram 2009	11	2.2 (1.1)	11	4.4 (1.1)		28.29%	-2.2[-3.12,-1.28]
Jakeman 2009	9	3 (2.6)	9	3.3 (2.1)		4.94%	-0.27[-2.47,1.93]
Kuligowski 1998	14	9.2 (4.6)	14	8.6 (4.1)		2.31%	0.59[-2.63,3.81]
Skurvydas 2006	20	4.8 (1.5)	20	6.7 (2.2)		17.56%	-1.9[-3.07,-0.73]
Yanagisawa 2003a	9	7 (1)	10	8.5 (1.3)		23.28%	-1.5[-2.51,-0.49]
Yanagisawa 2003b	7	4 (1.5)	7	6.3 (3.5)		3%	-2.25[-5.07,0.57]
Subtotal ***	**89**		**90**			**100%**	**-1.58[-2.07,-1.1]**
Heterogeneity: Tau²=0; Chi²=7.91, df=7(P=0.34); I²=11.48%							
Test for overall effect: Z=6.35(P<0.0001)							
1.27.4 72 hours							
Goodall 2008	9	3 (1.6)	9	3.4 (2.1)		13.22%	-0.35[-2.07,1.37]
Jakeman 2009	9	1.6 (1.3)	9	2.3 (1.8)		18.39%	-0.69[-2.15,0.77]
Kuligowski 1998	14	5.2 (5)	14	8.2 (4.5)		3.16%	-2.98[-6.51,0.55]
Skurvydas 2006	20	2.5 (0.5)	20	5.4 (1.7)		65.23%	-2.9[-3.68,-2.12]
Subtotal ***	**52**		**52**			**100%**	**-2.16[-2.79,-1.53]**
Heterogeneity: Tau²=0; Chi²=11.8, df=3(P=0.01); I²=74.59%							
Test for overall effect: Z=6.75(P<0.0001)							
1.27.5 96 hours							
Goodall 2008	9	1.5 (0.9)	9	1.6 (1)		44.68%	-0.15[-1,0.7]
Jakeman 2009	9	1 (1.5)	9	1.3 (0.9)		25.76%	-0.23[-1.36,0.9]
Kuligowski 1998	14	1.7 (2.2)	14	6.7 (4.6)		4.52%	-4.94[-7.63,-2.25]
Yanagisawa 2003a	9	3 (1.3)	10	4.8 (1.8)		17.72%	-1.75[-3.11,-0.39]
Yanagisawa 2003b	7	2.3 (2.3)	7	2 (1.8)		7.32%	0.25[-1.86,2.36]
Subtotal ***	**48**		**49**			**100%**	**-0.64[-1.21,-0.07]**
Heterogeneity: Tau²=0; Chi²=14.86, df=4(P=0.01); I²=73.07%							
Test for overall effect: Z=2.2(P=0.03)							

Favours CWI   -10 -5   0   5   10   Favours Passive

Comparison 2.   Cold-water immersion (CWI) versus contrast immersion

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Pain (muscle soreness: 10 point or 10 or 12 cm VAS)	5		Std. Mean Difference (IV, Random, 95% CI)	Subtotals only
1.1 Immediate	3	63	Std. Mean Difference (IV, Random, 95% CI)	0.04 [-0.66, 0.73]
1.2 24 hours	4	97	Std. Mean Difference (IV, Random, 95% CI)	-0.01 [-0.65, 0.64]
1.3 48 hours	3	77	Std. Mean Difference (IV, Random, 95% CI)	0.02 [-0.87, 0.91]
1.4 72 hours	2	55	Std. Mean Difference (IV, Random, 95% CI)	0.16 [-0.37, 0.69]
1.5 96 hours	1	28	Std. Mean Difference (IV, Random, 95% CI)	-0.12 [-0.86, 0.62]
2 Subjective recovery (10 cm VAS)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 Fatigue (immediate)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Strength	3		Std. Mean Difference (IV, Fixed, 95% CI)	Subtotals only
3.1 Immediate	1	27	Std. Mean Difference (IV, Fixed, 95% CI)	0.04 [-0.72, 0.80]
3.2 24 hours	2	55	Std. Mean Difference (IV, Fixed, 95% CI)	-0.26 [-0.79, 0.28]
3.3 48 hours	3	77	Std. Mean Difference (IV, Fixed, 95% CI)	-0.01 [-0.46, 0.44]
3.4 72 hours	2	55	Std. Mean Difference (IV, Fixed, 95% CI)	-0.17 [-0.70, 0.37]
3.5 96 hours	1	28	Std. Mean Difference (IV, Fixed, 95% CI)	-0.43 [-1.19, 0.32]
4 Power (squat jump in Watts)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
4.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Power (% decrease in jump performance)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
5.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
6 Functional performance (time to complete running task: seconds)	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
6.1 Multisprint/Agility circuit time (24 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.2 Sprint time (24 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
6.3 Multisprint time (48 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
7 Functional performance (time to fatigue)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
7.1 % decrease from baseline (immediate)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8 Swelling (thigh girth: centimetres)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
8.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9 Range of movement (degrees)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
9.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9.2 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9.3 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9.4 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10 Biomarker: creatine kinase (U/L)	3		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
10.1 Immediate	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.2 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.3 48 hours	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
11 Biomarker: myoglobin	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
11.1 Immediate	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of partici-pants	Statistical method	Effect size
11.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
12 Biomarker: lactate de-hydrogenase (U/L)	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
12.1 Immediate	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
12.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
12.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
12.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
13 Biomarker: C-reactive protein (mg/L)	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
13.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
13.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
13.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
14 Biomarker: inter-leukin-6 (pg/mL)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
14.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
14.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 2.1. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 1 Pain (muscle soreness: 10 point or 10 or 12 cm VAS).

Study or subgroup	CWI N	CWI Mean(SD)	Contrast N	Contrast Mean(SD)	Std. Mean Difference Random, 95% CI	Weight	Std. Mean Difference Random, 95% CI
2.1.1 Immediate							
Cassar 2010	8	2.5 (1)	8	1 (2.2)		27.9%	0.83[-0.2,1.86]
King 2009	10	2.5 (1.5)	10	2.7 (1.8)		33.7%	-0.09[-0.96,0.79]
Vaile 2008c	12	3.3 (2)	15	4.1 (1.6)		38.4%	-0.43[-1.2,0.34]
Subtotal *	**30**		**33**			**100%**	**0.04[-0.66,0.73]**
Heterogeneity: Tau²=0.18; Chi²=3.73, df=2(P=0.16); I²=46.35%							
Test for overall effect: Z=0.1(P=0.92)							
2.1.2 24 hours							
Ingram 2009	11	3 (1)	11	4 (1)		23.29%	-0.96[-1.85,-0.07]
King 2009	10	1.3 (1)	10	1.4 (0.7)		23.65%	-0.11[-0.99,0.76]
Kuligowski 1998	14	8.6 (6.6)	14	5.6 (4.1)		26.68%	0.53[-0.23,1.29]
Vaile 2008c	12	5.9 (1.9)	15	5.2 (1.6)		26.38%	0.39[-0.38,1.16]
Subtotal *	**47**		**50**			**100%**	**-0.01[-0.65,0.64]**
Heterogeneity: Tau²=0.26; Chi²=7.38, df=3(P=0.06); I²=59.33%							
Test for overall effect: Z=0.02(P=0.98)							

Favours CWI   -4 -2 0 2 4   Favours Contrast

Study or subgroup	CWI		Contrast		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Random, 95% CI		Random, 95% CI
2.1.3 48 hours							
Ingram 2009	11	3 (1)	11	4 (1)		31.36%	-0.96[-1.85,-0.07]
Kuligowski 1998	14	11 (5.5)	14	8.8 (5.6)		34.62%	0.39[-0.36,1.14]
Vaile 2008c	12	7 (2)	15	5.9 (1.9)		34.02%	0.55[-0.23,1.32]
Subtotal ***	**37**		**40**			**100%**	**0.02[-0.87,0.91]**
Heterogeneity: Tau²=0.44; Chi²=7.29, df=2(P=0.03); I²=72.56%							
Test for overall effect: Z=0.04(P=0.97)							
2.1.4 72 hours							
Kuligowski 1998	14	6.2 (6)	14	5.8 (5)		51.43%	0.07[-0.67,0.81]
Vaile 2008c	12	4.2 (1.1)	15	3.8 (1.8)		48.57%	0.25[-0.51,1.02]
Subtotal ***	**26**		**29**			**100%**	**0.16[-0.37,0.69]**
Heterogeneity: Tau²=0; Chi²=0.11, df=1(P=0.74); I²=0%							
Test for overall effect: Z=0.59(P=0.55)							
2.1.5 96 hours							
Kuligowski 1998	14	2.1 (2.7)	14	2.4 (3)		100%	-0.12[-0.86,0.62]
Subtotal ***	**14**		**14**			**100%**	**-0.12[-0.86,0.62]**
Heterogeneity: Tau²=0; Chi²=0, df=0(P<0.0001); I²=100%							
Test for overall effect: Z=0.32(P=0.75)							

Favours CWI -4 -2 0 2 4 Favours Contrast

Analysis 2.2. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 2 Subjective recovery (10 cm VAS).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.2.1 Fatigue (immediate)						
Cassar 2010	8	6.9 (0.7)	8	6.2 (1.8)		0.7[-0.64,2.04]

Favours CWI -2 -1 0 1 2 Favours contrast

Analysis 2.3. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 3 Strength.

Study or subgroup	CWI		Contrast		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
2.3.1 Immediate							
Vaile 2008c	12	1748 (424)	15	1733 (320)		100%	0.04[-0.72,0.8]
Subtotal ***	**12**		**15**			**100%**	**0.04[-0.72,0.8]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.1(P=0.92)							
2.3.2 24 hours							
Kuligowski 1998	14	10.6 (4.4)	14	13.7 (6.2)		50.12%	-0.56[-1.32,0.2]
Vaile 2008c	12	1877 (418)	15	1857 (405)		49.88%	0.05[-0.71,0.81]
Subtotal ***	**26**		**29**			**100%**	**-0.26[-0.79,0.28]**
Heterogeneity: Tau²=0; Chi²=1.22, df=1(P=0.27); I²=18.28%							

Favours contrast -2 -1 0 1 2 Favours CWI

Study or subgroup	CWI		Contrast		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
Test for overall effect: Z=0.94(P=0.35)							
2.3.3 48 hours							
Ingram 2009	11	73.4 (18.5)	11	70.9 (13.1)		29.09%	0.15[-0.69,0.99]
Kuligowski 1998	14	10.9 (4.4)	14	13.3 (5.9)		36.06%	-0.46[-1.21,0.3]
Vaile 2008c	12	2077 (465)	15	1923 (457)		34.85%	0.32[-0.44,1.09]
Subtotal ***	**37**		**40**			**100%**	**-0.01[-0.46,0.44]**
Heterogeneity: Tau²=0; Chi²=2.23, df=2(P=0.33); I²=10.35%							
Test for overall effect: Z=0.04(P=0.97)							
2.3.4 72 hours							
Kuligowski 1998	14	11.9 (5.6)	14	14.5 (6)		50.57%	-0.44[-1.19,0.31]
Vaile 2008c	12	2074 (487)	15	2018 (477)		49.43%	0.11[-0.65,0.87]
Subtotal ***	**26**		**29**			**100%**	**-0.17[-0.7,0.37]**
Heterogeneity: Tau²=0; Chi²=1.03, df=1(P=0.31); I²=2.65%							
Test for overall effect: Z=0.61(P=0.54)							
2.3.5 96 hours							
Kuligowski 1998	14	12.9 (5.3)	14	15.4 (6)		100%	-0.43[-1.19,0.32]
Subtotal ***	**14**		**14**			**100%**	**-0.43[-1.19,0.32]**
Heterogeneity: Tau²=0; Chi²=0, df=0(P<0.0001); I²=100%							
Test for overall effect: Z=1.13(P=0.26)							

Favours contrast -2 -1 0 1 2 Favours CWI

Analysis 2.4.   Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 4 Power (squat jump in Watts).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.4.1 Immediate						
Vaile 2008c	12	3547 (1033)	15	3328 (806)		219[-493.72,931.72]
2.4.2 24 hours						
Vaile 2008c	12	3735 (872)	15	3675 (741)		60[-559.7,679.7]
2.4.3 48 hours						
Vaile 2008c	12	3939 (877)	15	3805 (821)		134[-513.17,781.17]
2.4.4 72 hours						
Vaile 2008c	914	4080 (914)	15	3937 (808)		143[-270.17,556.17]

Favours CWI -1000 -500 0 500 1000 Favours contrast

Analysis 2.5.   Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 5 Power (% decrease in jump performance).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.5.1 24 hours						
King 2009	10	4.4 (2.7)	10	8.1 (5.8)		-3.7[-7.67,0.27]
				Favours CWI	-20 -10 0 10 20 Favours contrast	
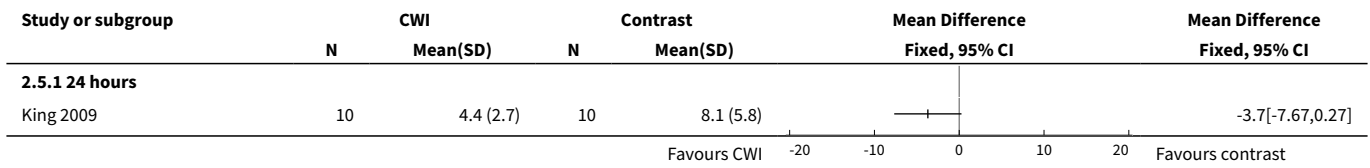
Analysis 2.6.   Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 6 Functional performance (time to complete running task: seconds).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.6.1 Multisprint/Agility circuit time (24 hours)						
King 2009	10	34.5 (2.9)	10	34 (1.9)		0.47[-1.69,2.63]
2.6.2 Sprint time (24 hours)						
King 2009	10	2.3 (0.2)	10	2.3 (0.2)		0.05[-0.11,0.21]
2.6.3 Multisprint time (48 hours)						
Ingram 2009	11	48.2 (1.8)	11	47.8 (1.4)		0.42[-0.93,1.77]
				Favours CWI	-5 -2.5 0 2.5 5 Favours contrast	
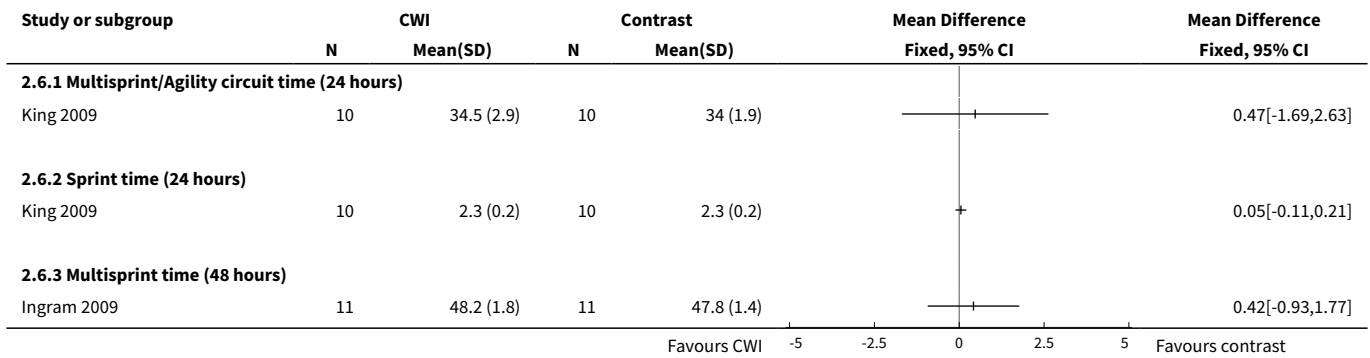
Analysis 2.7.   Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 7 Functional performance (time to fatigue).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.7.1 % decrease from baseline (immediate)						
Cassar 2010	8	17.7 (4.7)	8	13.5 (8.5)		4.2[-2.53,10.93]
				Favours CWI	-10 -5 0 5 10 Favours contrast	
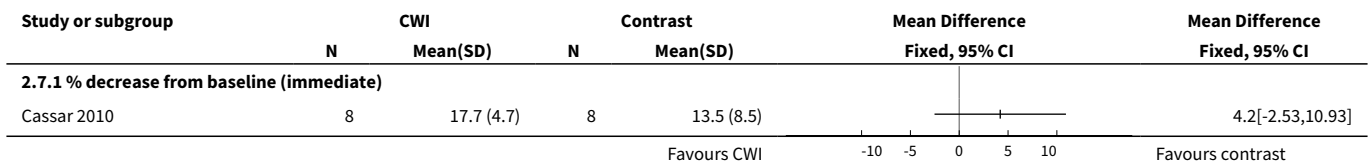
Analysis 2.8.   Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 8 Swelling (thigh girth: centimetres).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.8.1 Immediate						
Vaile 2008c	12	57.4 (3.8)	15	56.8 (4.6)		0.6[-2.57,3.77]
2.8.2 24 hours						
Vaile 2008c	12	57.1 (3.8)	15	56.4 (4.5)		0.7[-2.43,3.83]
2.8.3 48 hours						
Vaile 2008c	12	56.9 (3.8)	15	56.3 (4.6)		0.6[-2.57,3.77]
				Favours CWI	-4 -2 0 2 4 Favours contrast	

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.8.4 72 hours						
Vaile 2008c	12	56.9 (3.8)	15	56.3 (4.5)		0.6[-2.53,3.73]

Favours CWI -4 -2 0 2 4 Favours contrast

Analysis 2.9. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 9 Range of movement (degrees).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.9.1 24 hours						
Kuligowski 1998	14	7.2 (7.4)	14	7.5 (7.6)		-0.31[-5.87,5.25]
2.9.2 48 hours						
Kuligowski 1998	14	13.8 (12)	14	11.7 (10.9)		2.12[-6.37,10.61]
2.9.3 72 hours						
Kuligowski 1998	14	14.2 (12.4)	14	9.6 (9.1)		4.62[-3.42,12.66]
2.9.4 96 hours						
Kuligowski 1998	14	9.5 (9.3)	14	7 (7.4)		2.45[-3.75,8.65]

Favours CWI -20 -10 0 10 20 Favours contrast
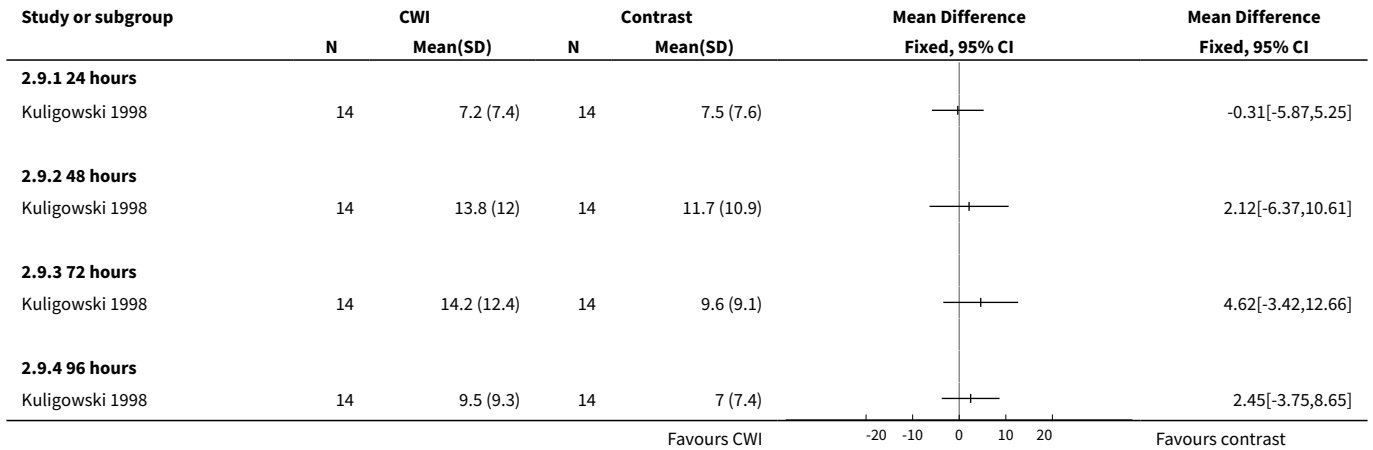
Analysis 2.10. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 10 Biomarker: creatine kinase (U/L).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.10.1 Immediate						
Cassar 2010	8	172.3 (31.4)	8	219 (32.5)		-46.7[-78.02,-15.38]
Vaile 2008c	12	203 (175)	15	229 (147)		-26[-149.85,97.85]
2.10.2 24 hours						
Ingram 2009	11	571 (375)	11	582 (357)		-11[-316.97,294.97]
Vaile 2008c	12	231 (182)	15	736 (1115)		-505[-1078.58,68.58]
2.10.3 48 hours						
Ingram 2009	11	337 (218)	11	332 (168)		5[-157.64,167.64]
Vaile 2008c	12	211 (259)	15	416 (589)		-205[-537.14,127.14]
2.10.4 72 hours						
Vaile 2008c	12	204 (343)	15	359 (433)		-155[-447.71,137.71]

Favours CWI -1000 -500 0 500 1000 Favours contrast

Analysis 2.11. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 11 Biomarker: myoglobin.

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.11.1 Immediate						
Cassar 2010	8	0 (0)	8	0 (0)		0.01[0,0.02]
Vaile 2008c	12	60.7 (30.1)	15	95.4 (76.6)		-34.7[-77.04,7.64]
2.11.2 24 hours						
Vaile 2008c	12	44.9 (25.4)	15	67.2 (51.1)		-22.3[-51.88,7.28]

Favours CWI -100 -50 0 50 100 Favours contrast
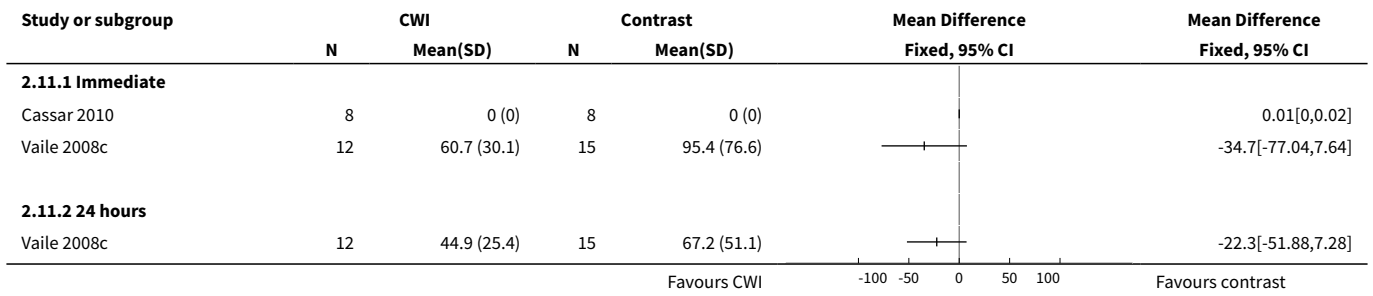
Analysis 2.12. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 12 Biomarker: lactate dehydrogenase (U/L).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.12.1 Immediate						
Cassar 2010	8	192 (15)	8	168 (14.1)		24[9.73,38.27]
Vaile 2008c	12	227 (95)	15	280 (87)		-53[-122.48,16.48]
2.12.2 24 hours						
Vaile 2008c	12	194 (65)	15	291 (132)		-97[-173.25,-20.75]
2.12.3 48 hours						
Vaile 2008c	12	177 (71)	15	264 (117)		-87[-158.55,-15.45]
2.12.4 72 hours						
Vaile 2008c	12	183 (68)	15	254 (109)		-71[-138.25,-3.75]

Favours CWI -200 -100 0 100 200 Favours contrast

Analysis 2.13. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 13 Biomarker: C-reactive protein (mg/L).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.13.1 Immediate						
Cassar 2010	8	0.8 (0.8)	8	0.3 (0.2)		0.46[-0.11,1.03]
2.13.2 24 hours						
Ingram 2009	11	2.5 (1.7)	11	3.3 (3.8)		-0.8[-3.26,1.66]
2.13.3 48 hours						
Ingram 2009	11	1.4 (0.8)	11	2.6 (2.6)		-1.2[-2.81,0.41]

Favours CWI -5 -2.5 0 2.5 5 Favours contrast
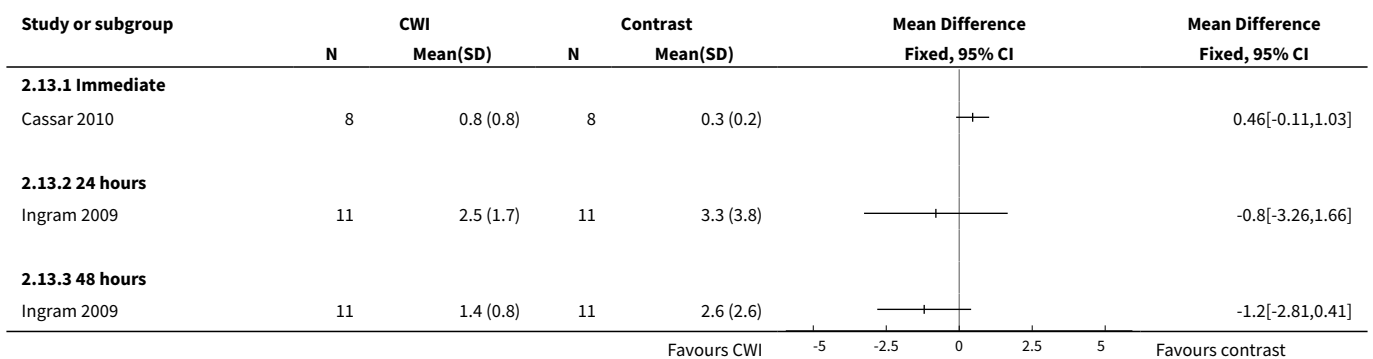
Analysis 2.14. Comparison 2 Cold-water immersion (CWI) versus contrast immersion, Outcome 14 Biomarker: interleukin-6 (pg/mL).

Study or subgroup	CWI		Contrast		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
2.14.1 Immediate						
Vaile 2008c	12	4.5 (6.8)	15	2.2 (0.7)		2.3[-1.56,6.16]
2.14.2 24 hours						
Vaile 2008c	12	3.7 (6.3)	15	1.5 (0.9)		2.2[-1.39,5.79]

Favours CWI -5 -2.5 0 2.5 5 Favours contrast
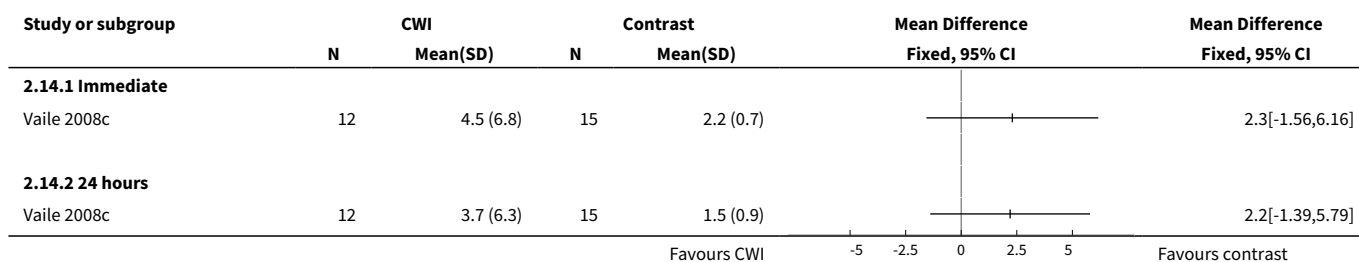
Comparison 3. Cold-water immersion (CWI) versus warm-water immersion (WWI)

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Pain (muscle soreness: 10 point or 10 or 12 cm VAS)	3		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
1.1 Immediate	1	23	Mean Difference (IV, Fixed, 95% CI)	-0.5 [-2.09, 1.09]
1.2 24 hours	3	64	Mean Difference (IV, Fixed, 95% CI)	-0.33 [-1.40, 0.73]
1.3 48 hours	2	51	Mean Difference (IV, Fixed, 95% CI)	0.52 [-0.94, 1.99]
1.4 72 hours	2	51	Mean Difference (IV, Fixed, 95% CI)	-0.40 [-1.83, 1.03]
1.5 96 hours	1	28	Mean Difference (IV, Fixed, 95% CI)	-4.64 [-7.72, -1.56]
2 Subjective recovery	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 Fatigue (24 hours post intervention)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Subjective recovery (Intervention associated with NO benefit for recovery)	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
3.1 24 hours	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Strength	2		Std. Mean Difference (IV, Fixed, 95% CI)	Subtotals only
4.1 Immediate	1	23	Std. Mean Difference (IV, Fixed, 95% CI)	0.42 [-0.41, 1.25]
4.2 24 hours	2	51	Std. Mean Difference (IV, Fixed, 95% CI)	0.28 [-0.27, 0.84]
4.3 48 hours	2	51	Std. Mean Difference (IV, Fixed, 95% CI)	0.30 [-0.27, 0.86]
4.4 72 hours	2	51	Std. Mean Difference (IV, Fixed, 95% CI)	0.20 [-0.36, 0.75]
4.5 96 hours	1	28	Std. Mean Difference (IV, Fixed, 95% CI)	-0.16 [-0.90, 0.58]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
5 Power	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
5.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.2 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
6 Functional performance	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
6.1 Multisprint time at 24 hours (seconds)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
7 Range of movement (ROM)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
7.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
7.2 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
7.3 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
7.4 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8 Swelling	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
8.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
8.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
9 Biomarker: creatine kinase (U/L)	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
9.1 Immediate	1	23	Mean Difference (IV, Fixed, 95% CI)	-66.0 [-328.29, 196.29]
9.2 24 hours	2	36	Mean Difference (IV, Fixed, 95% CI)	-65.27 [-236.38, 105.84]
9.3 48 hours	1	23	Mean Difference (IV, Fixed, 95% CI)	-14.0 [-210.29, 182.29]
9.4 72 hours	1	23	Mean Difference (IV, Fixed, 95% CI)	53.0 [-143.97, 249.97]
10 Biomarker: lactate dehydrogenase (U/L)	2		Mean Difference (IV, Fixed, 95% CI)	Totals not selected

Outcome or subgroup title	No. of studies	No. of partici-pants	Statistical method	Effect size
10.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.2 24 hours	2		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
10.4 72 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
11 Biomarker: myoglo-bin (ng/mL)	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
11.1 Immediate	1	23	Mean Difference (IV, Fixed, 95% CI)	-4.40 [-35.63, 26.83]
11.2 24 hours	2	36	Mean Difference (IV, Fixed, 95% CI)	1.16 [-17.14, 19.46]
12 Biomarker: inter-leukin-6 (pg/ml)	2		Mean Difference (IV, Fixed, 95% CI)	Subtotals only
12.1 Immediate	1	23	Mean Difference (IV, Fixed, 95% CI)	2.2 [-1.72, 6.12]
12.2 24 hours	2	36	Mean Difference (IV, Fixed, 95% CI)	0.07 [-0.90, 1.03]
13 Biomarker: inter-leukin-10 (pg/ml)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
13.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
14 Biomarker: inter-leukin-1b (pg/ml)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
14.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 3.1. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 1 Pain (muscle soreness: 10 point or 10 or 12 cm VAS).

Study or subgroup	CWI		WWI		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
3.1.1 Immediate							
Vaile 2008c	12	3.3 (2)	11	3.8 (1.9)		100%	-0.5[-2.09,1.09]
Subtotal ***	**12**		**11**			**100%**	**-0.5[-2.09,1.09]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.61(P=0.54)							
3.1.2 24 hours							
Kuligowski 1998	14	8.6 (6.6)	14	8.7 (6.7)		4.71%	-0.14[-5.05,4.77]
Rowsell 2009	6	5.6 (2)	7	6.1 (2.4)		19.84%	-0.5[-2.89,1.89]
Vaile 2008c	12	5.9 (1.9)	11	6.2 (1)		75.45%	-0.3[-1.53,0.93]
Subtotal ***	**32**		**32**			**100%**	**-0.33[-1.4,0.73]**
Heterogeneity: Tau²=0; Chi²=0.03, df=2(P=0.99); I²=0%							
Test for overall effect: Z=0.61(P=0.54)							

Favours CWI   -10   -5   0   5   10   Favours WWI

Study or subgroup	CWI		WWI		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
3.1.3 48 hours							
Kuligowski 1998	14	11 (5.5)	14	8.7 (6.3)		11.3%	2.29[-2.06,6.64]
Vaile 2008c	12	7 (2)	11	6.7 (1.8)		88.7%	0.3[-1.25,1.85]
Subtotal ***	**26**		**25**			**100%**	**0.52[-0.94,1.99]**
Heterogeneity: Tau²=0; Chi²=0.71, df=1(P=0.4); I²=0%							
Test for overall effect: Z=0.7(P=0.48)							
3.1.4 72 hours							
Kuligowski 1998	14	6.2 (6)	14	8.9 (5.4)		11.48%	-2.72[-6.93,1.49]
Vaile 2008c	12	4.2 (1.8)	11	4.3 (1.9)		88.52%	-0.1[-1.62,1.42]
Subtotal ***	**26**		**25**			**100%**	**-0.4[-1.83,1.03]**
Heterogeneity: Tau²=0; Chi²=1.32, df=1(P=0.25); I²=24.07%							
Test for overall effect: Z=0.55(P=0.58)							
3.1.5 96 hours							
Kuligowski 1998	14	2.1 (2.7)	14	6.7 (5.2)		100%	-4.64[-7.72,-1.56]
Subtotal ***	**14**		**14**			**100%**	**-4.64[-7.72,-1.56]**
Heterogeneity: Not applicable							
Test for overall effect: Z=2.96(P=0)							

Favours CWI   -10 -5 0 5 10   Favours WWI

Analysis 3.2. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 2 Subjective recovery.

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.2.1 Fatigue (24 hours post intervention)						
Rowsell 2009	6	4.6 (2)	7	5.4 (2.5)		-0.8[-3.25,1.65]

Favours CWI   -5 -2.5 0 2.5 5   Favours WWI

Analysis 3.3. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 3 Subjective recovery (Intervention associated with NO benefit for recovery).

Study or subgroup	CWI	HWI	Risk Ratio	Risk Ratio
	n/N	n/N	M-H, Fixed, 95% CI	M-H, Fixed, 95% CI
3.3.1 24 hours				
Rowsell 2009	0/6	6/7		0.09[0.01,1.3]

Favours CWI   0.001 0.1 1 10 1000   Favours WWI

Analysis 3.4. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 4 Strength.

Study or subgroup	CWI		HWI		Std. Mean Difference	Weight	Std. Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
3.4.1 Immediate							
Vaile 2008c	12	1748 (424)	11	1592 (262)		100%	0.42[-0.41,1.25]
Subtotal * **	**12**		**11**			**100%**	**0.42[-0.41,1.25]**
Heterogeneity: Tau²=0; Chi²=0, df=0(P<0.0001); I²=100%							
Test for overall effect: Z=1(P=0.32)							
3.4.2 24 hours							
Kuligowski 1998	14	10.6 (4.4)	14	10 (7.5)		55.87%	0.1[-0.64,0.84]
Vaile 2008c	12	1877 (418)	11	1685 (286)		44.13%	0.51[-0.32,1.35]
Subtotal * **	**26**		**25**			**100%**	**0.28[-0.27,0.84]**
Heterogeneity: Tau²=0; Chi²=0.52, df=1(P=0.47); I²=0%							
Test for overall effect: Z=1(P=0.32)							
3.4.3 48 hours							
Kuligowski 1998	14	10.9 (4.4)	14	11.6 (7.7)		57.47%	-0.12[-0.86,0.62]
Vaile 2008c	12	2077 (465)	11	1735 (272)		42.53%	0.86[-0.01,1.72]
Subtotal * **	**26**		**25**			**100%**	**0.3[-0.27,0.86]**
Heterogeneity: Tau²=0; Chi²=2.81, df=1(P=0.09); I²=64.46%							
Test for overall effect: Z=1.03(P=0.3)							
3.4.4 72 hours							
Kuligowski 1998	14	11.9 (5.6)	14	12.1 (6.9)		55.82%	-0.04[-0.78,0.7]
Vaile 2008c	12	2074 (487)	11	1868 (291)		44.18%	0.49[-0.34,1.32]
Subtotal * **	**26**		**25**			**100%**	**0.2[-0.36,0.75]**
Heterogeneity: Tau²=0; Chi²=0.86, df=1(P=0.35); I²=0%							
Test for overall effect: Z=0.69(P=0.49)							
3.4.5 96 hours							
Kuligowski 1998	14	12.9 (5.3)	14	14 (8.2)		100%	-0.16[-0.9,0.58]
Subtotal * **	**14**		**14**			**100%**	**-0.16[-0.9,0.58]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.42(P=0.67)							

Favours WWI -2 -1 0 1 2 Favours CWI

Analysis 3.5. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 5 Power.

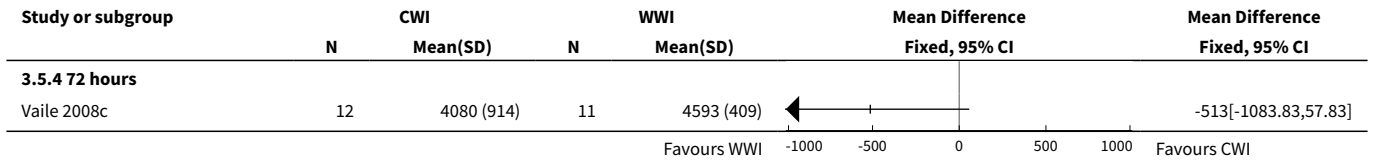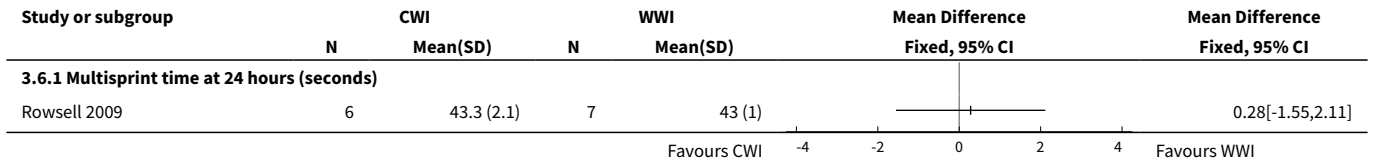| Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.5.1 Immediate						
Vaile 2008c	12	3547 (1033)	11	3446 (351)		101[-519.18,721.18]
3.5.2 24 hours						
Rowsell 2009	6	46 (8)	7	48 (6)		-2[-9.79,5.79]
Vaile 2008c	12	3735 (872)	11	3459 (389)		276[-268.3,820.3]
3.5.3 48 hours						
Vaile 2008c	12	3939 (877)	11	3487 (455)		452[-112.37,1016.37]

Favours WWI -1000 -500 0 500 1000 Favours CWI

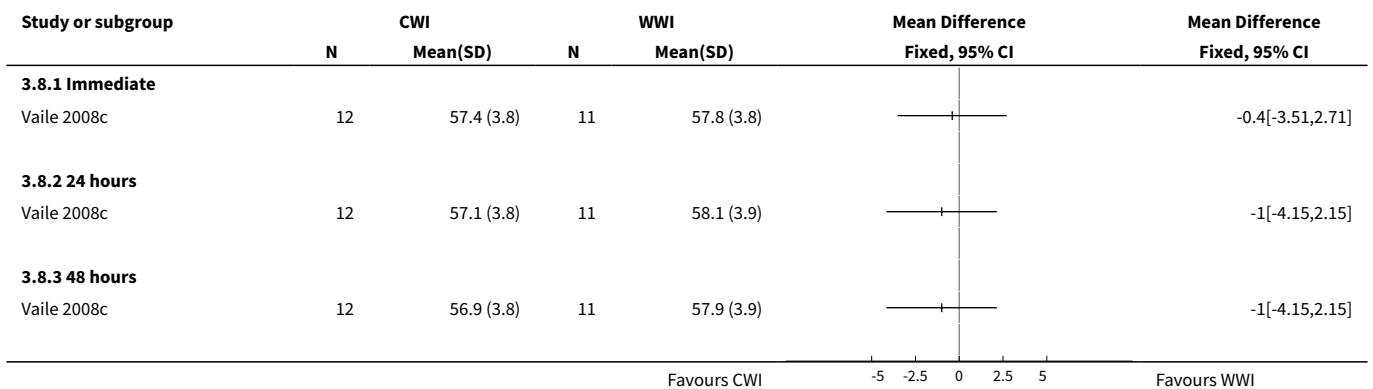| Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.5.4 72 hours						
Vaile 2008c	12	4080 (914)	11	4593 (409)		-513[-1083.83,57.83]

Favours WWI -1000 -500 0 500 1000 Favours CWI

Analysis 3.6. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 6 Functional performance.

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.6.1 Multisprint time at 24 hours (seconds)						
Rowsell 2009	6	43.3 (2.1)	7	43 (1)		0.28[-1.55,2.11]

Favours CWI -4 -2 0 2 4 Favours WWI

Analysis 3.7. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 7 Range of movement (ROM).

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.7.1 24 hours						
Kuligowski 1998	14	7.2 (7.4)	14	9.1 (10.2)		-1.88[-8.49,4.73]
3.7.2 48 hours						
Kuligowski 1998	14	13.8 (12)	14	12.5 (9.4)		1.31[-6.69,9.31]
3.7.3 72 hours						
Kuligowski 1998	14	14.2 (12.4)	14	15.5 (13.6)		-1.33[-10.96,8.3]
3.7.4 96 hours						
Kuligowski 1998	14	9.5 (9.3)	14	13.9 (13)		-4.48[-12.85,3.89]

Favours WWI -20 -10 0 10 20 Favours CWI
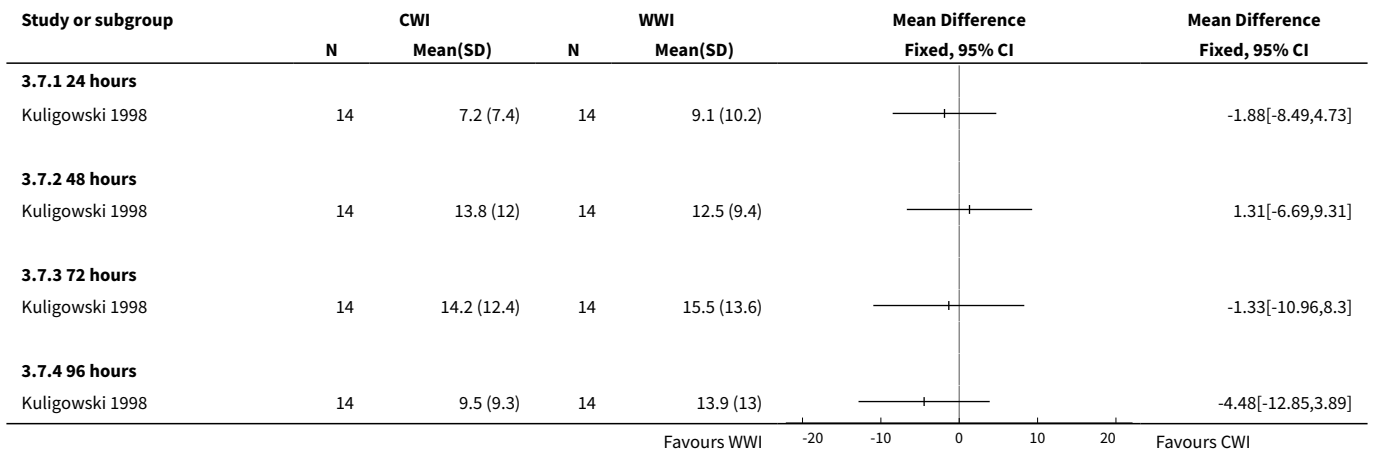
Analysis 3.8. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 8 Swelling.

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.8.1 Immediate						
Vaile 2008c	12	57.4 (3.8)	11	57.8 (3.8)		-0.4[-3.51,2.71]
3.8.2 24 hours						
Vaile 2008c	12	57.1 (3.8)	11	58.1 (3.9)		-1[-4.15,2.15]
3.8.3 48 hours						
Vaile 2008c	12	56.9 (3.8)	11	57.9 (3.9)		-1[-4.15,2.15]

Favours CWI -5 -2.5 0 2.5 5 Favours WWI

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Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.8.4 72 hours						
Vaile 2008c	12	56.9 (3.8)	11	57.6 (3.8)		-0.7[-3.81,2.41]

Favours CWI -5 -2.5 0 2.5 5 Favours WWI

Analysis 3.9.   Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 9 Biomarker: creatine kinase (U/L).

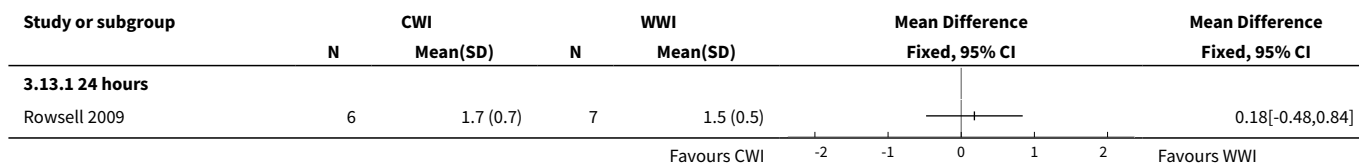| Study or subgroup	CWI		WWI		Mean Difference	Weight	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI		Fixed, 95% CI
3.9.1 Immediate							
Vaile 2008c	12	203 (175)	11	269 (411)		100%	-66[-328.29,196.29]
Subtotal ***	**12**		**11**			**100%**	**-66[-328.29,196.29]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.49(P=0.62)							
3.9.2 24 hours							
Rowsell 2009	6	1075 (680)	7	881 (627)		5.72%	194[-521.39,909.39]
Vaile 2008c	12	231 (182)	11	312 (242)		94.28%	-81[-257.23,95.23]
Subtotal ***	**18**		**18**			**100%**	**-65.27[-236.38,105.84]**
Heterogeneity: Tau²=0; Chi²=0.54, df=1(P=0.46); I²=0%							
Test for overall effect: Z=0.75(P=0.45)							
3.9.3 48 hours							
Vaile 2008c	12	211 (259)	11	225 (221)		100%	-14[-210.29,182.29]
Subtotal ***	**12**		**11**			**100%**	**-14[-210.29,182.29]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.14(P=0.89)							
3.9.4 72 hours							
Vaile 2008c	12	204 (343)	11	151 (57)		100%	53[-143.97,249.97]
Subtotal ***	**12**		**11**			**100%**	**53[-143.97,249.97]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.53(P=0.6)							
Test for subgroup differences: Chi²=0.92, df=1 (P=0.82), I²=0%							

Favours CWI -1000 -500 0 500 1000 Favours WWI

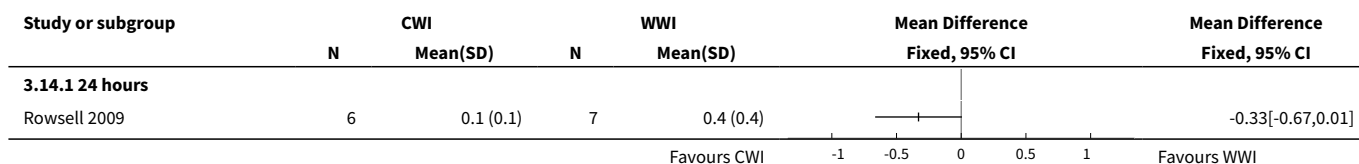## Analysis 3.10.   Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 10 Biomarker: lactate dehydrogenase (U/L).

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.10.1 Immediate						
Vaile 2008c	12	227 (95)	11	278 (85)		-51[-124.57,22.57]
3.10.2 24 hours						
Rowsell 2009	6	482 (65)	7	469 (91)		13[-72.14,98.14]
Vaile 2008c	12	194 (65)	11	271 (90)		-77[-141.66,-12.34]

Favours CWI -100 -50 0 50 100 Favours WWI

Study or subgroup	CWI N	CWI Mean(SD)	WWI N	WWI Mean(SD)	Mean Difference Fixed, 95% CI	Mean Difference Fixed, 95% CI
3.10.3 48 hours						
Vaile 2008c	12	177 (71)	11	260 (69)		-83[-140.24,-25.76]
3.10.4 72 hours						
Vaile 2008c	12	183 (68)	11	254 (83)		-71[-133.34,-8.66]

Favours CWI -100 -50 0 50 100 Favours WWI

Analysis 3.11. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 11 Biomarker: myoglobin (ng/mL).

Study or subgroup	CWI N	CWI Mean(SD)	WWI N	WWI Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
3.11.1 Immediate							
Vaile 2008c	12	60.7 (30.1)	11	65.1 (44.3)		100%	-4.4[-35.63,26.83]
Subtotal ***	12		11			100%	**-4.4[-35.63,26.83]**
Heterogeneity: Not applicable							
Test for overall effect: Z=0.28(P=0.78)							
3.11.2 24 hours							
Rowsell 2009	6	48 (13)	7	69 (62)		15.1%	-21[-68.09,26.09]
Vaile 2008c	12	44.9 (25.4)	11	39.8 (23.2)		84.9%	5.1[-14.76,24.96]
Subtotal ***	18		18			100%	**1.16[-17.14,19.46]**
Heterogeneity: Tau²=0; Chi²=1, df=1(P=0.32); I²=0.18%							
Test for overall effect: Z=0.12(P=0.9)							

Favours CWI -100 -50 0 50 100 Favours WWI

Analysis 3.12. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 12 Biomarker: interleukin-6 (pg/ml).

Study or subgroup	CWI N	CWI Mean(SD)	WWI N	WWI Mean(SD)	Mean Difference Fixed, 95% CI	Weight	Mean Difference Fixed, 95% CI
3.12.1 Immediate							
Vaile 2008c	12	4.5 (6.8)	11	2.3 (1.3)		100%	2.2[-1.72,6.12]
Subtotal ***	12		11			100%	**2.2[-1.72,6.12]**
Heterogeneity: Not applicable							
Test for overall effect: Z=1.1(P=0.27)							
3.12.2 24 hours							
Rowsell 2009	6	1.5 (1)	7	1.6 (0.8)		92.82%	-0.08[-1.08,0.92]
Vaile 2008c	12	3.7 (6.3)	11	1.7 (0.9)		7.18%	2[-1.6,5.6]
Subtotal ***	18		18			100%	**0.07[-0.9,1.03]**
Heterogeneity: Tau²=0; Chi²=1.19, df=1(P=0.28); I²=15.81%							
Test for overall effect: Z=0.14(P=0.89)							

Favours CWI -10 -5 0 5 10 Favours WWI

Analysis 3.13. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 13 Biomarker: interleukin-10 (pg/ml).

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.13.1 24 hours						
Rowsell 2009	6	1.7 (0.7)	7	1.5 (0.5)		0.18[-0.48,0.84]

Favours CWI -2 -1 0 1 2 Favours WWI

Analysis 3.14. Comparison 3 Cold-water immersion (CWI) versus warm-water immersion (WWI), Outcome 14 Biomarker: interleukin-1b (pg/ml).

Study or subgroup	CWI		WWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
3.14.1 24 hours						
Rowsell 2009	6	0.1 (0.1)	7	0.4 (0.4)		-0.33[-0.67,0.01]

Favours CWI -1 -0.5 0 0.5 1 Favours WWI

Comparison 4. Cold-water immersion (CWI) versus active recovery

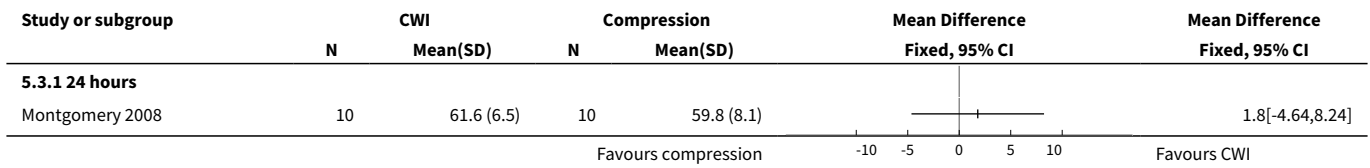| Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Pain (muscle soreness: 10 point VAS)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Power (% decrement in jump performance)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Functional performance (time to complete running test: seconds)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 Sprint time (24 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 Multisprint/Agility time (24 hours)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 4.1. Comparison 4 Cold-water immersion (CWI) versus active recovery, Outcome 1 Pain (muscle soreness: 10 point VAS).

Study or subgroup	CWI		Active		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
4.1.1 Immediate						
King 2009	10	2.5 (1.5)	10	5.2 (1.7)		-2.65[-4.05,-1.25]
4.1.2 24 hours						
King 2009	10	1.3 (1)	10	2.2 (1.5)		-0.9[-2,0.2]
				Favours CWI	-5 -2.5 0 2.5 5	Favours active

Analysis 4.2. Comparison 4 Cold-water immersion (CWI) versus active recovery, Outcome 2 Power (% decrement in jump performance).

Study or subgroup	CWI		Active		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
4.2.1 24 hours						
King 2009	10	4.4 (2.7)	10	7.7 (3.7)		-3.3[-6.14,-0.46]
				Favours CWI	-10 -5 0 5 10	Favours active

Analysis 4.3. Comparison 4 Cold-water immersion (CWI) versus active recovery, Outcome 3 Functional performance (time to complete running test: seconds).

Study or subgroup	CWI		Active		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
4.3.1 Sprint time (24 hours)						
King 2009	10	2.3 (0.2)	10	2.4 (0.2)		-0.03[-0.21,0.15]
4.3.2 Multisprint/Agility time (24 hours)						
King 2009	10	34.5 (2.9)	10	35.6 (3.3)		-1.12[-3.86,1.62]
				Favours CWI	-5 -2.5 0 2.5 5	Favours active

Comparison 5. Cold-water immersion (CWI) versus compression

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Pain (muscle soreness: 10 point VAS)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Subjective recovery (rating of fatigue: 10 point VAS)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected

Outcome or subgroup title	No. of studies	No. of partici-pants	Statistical method	Effect size
2.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Power (vertical jump height: centimetres)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Range of movement (centime-tres)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
4.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5 Functional performance (time to complete running test: sec-onds)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
5.1 Agility circuit time (24 hrs)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
5.2 Multisprint time (24 hrs)	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 5.1. Comparison 5 Cold-water immersion (CWI) versus compression, Outcome 1 Pain (muscle soreness: 10 point VAS).

Study or subgroup	CWI		Compression		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
5.1.1 24 hours						
Montgomery 2008	10	3.3 (1.1)	10	3.2 (1.3)		0.1[-0.96,1.16]

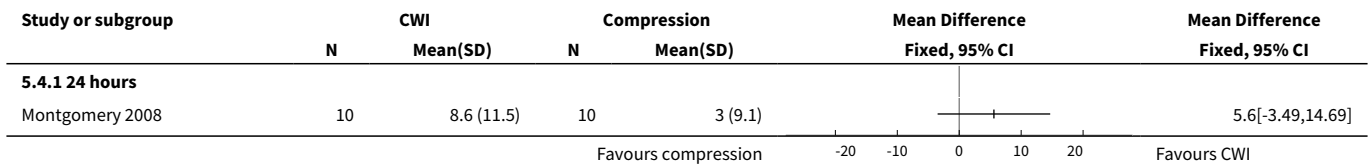Favours CWI   -1   -0.5   0   0.5   1   Favours Compresssion

Analysis 5.2. Comparison 5 Cold-water immersion (CWI) versus compression, Outcome 2 Subjective recovery (rating of fatigue: 10 point VAS).

Study or subgroup	CWI		Compression		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
5.2.1 24 hours						
Montgomery 2008	10	4.5 (0.9)	10	4.3 (1.6)		0.2[-0.94,1.34]

Favours CWI   -2   -1   0   1   2   Favours compression

Analysis 5.3. Comparison 5 Cold-water immersion (CWI) versus compression, Outcome 3 Power (vertical jump height: centimetres).

Study or subgroup	CWI		Compression		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
5.3.1 24 hours						
Montgomery 2008	10	61.6 (6.5)	10	59.8 (8.1)		1.8[-4.64,8.24]

Favours compression -10 -5 0 5 10 Favours CWI

Analysis 5.4. Comparison 5 Cold-water immersion (CWI) versus compression, Outcome 4 Range of movement (centimetres).

Study or subgroup	CWI		Compression		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
5.4.1 24 hours						
Montgomery 2008	10	8.6 (11.5)	10	3 (9.1)		5.6[-3.49,14.69]

Favours compression -20 -10 0 10 20 Favours CWI

Analysis 5.5. Comparison 5 Cold-water immersion (CWI) versus compression, Outcome 5 Functional performance (time to complete running test: seconds).

Study or subgroup	CWI		Compression		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
5.5.1 Agility circuit time (24 hrs)						
Montgomery 2008	10	6.6 (0.2)	10	6.7 (0.3)		-0.06[-0.28,0.16]
5.5.2 Multisprint time (24 hrs)						
Montgomery 2008	10	27.3 (1)	10	27.1 (0.6)		0.22[-0.52,0.96]

Favours CWI -1 -0.5 0 0.5 1 Favours compression

Comparison 6.   Cold-water immersion (CWI) versus double CWI

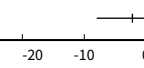| Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Pain (muscle soreness: 5 point VAS)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
1.1 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.2 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.3 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Range of movement (degrees)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
2.1 Immediate	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
2.2 24 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.3 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.4 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3 Biomarker: creatine kinase (U/L)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
3.1 48 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
3.2 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Biomarker: lactate dehydrogenase (U/L)	1		Mean Difference (IV, Fixed, 95% CI)	Totals not selected
4.1 96 hours	1		Mean Difference (IV, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 6.1. Comparison 6 Cold-water immersion (CWI) versus double CWI, Outcome 1 Pain (muscle soreness: 5 point VAS).

Study or subgroup	CWI (single)		Double CWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
6.1.1 24 hours						
Yanagisawa 2003a	9	3.1 (0.5)	9	2.9 (0.2)		0.2[-0.15,0.55]
6.1.2 48 hours						
Yanagisawa 2003a	9	3.8 (0.4)	9	3.8 (0.3)		0[-0.33,0.33]
6.1.3 96 hours						
Yanagisawa 2003a	9	2.2 (0.5)	9	3 (0.7)		-0.8[-1.36,-0.24]

Favours CWI (single) -1 -0.5 0 0.5 1 Favours double CWI
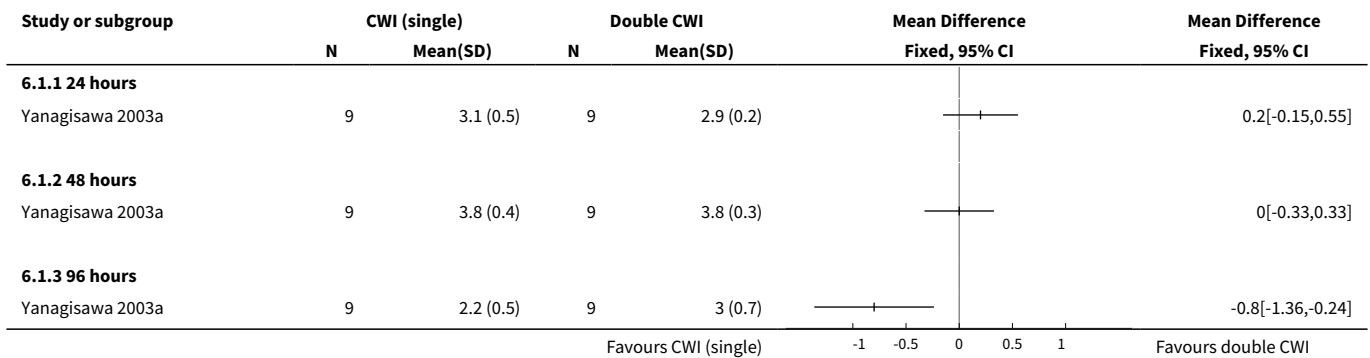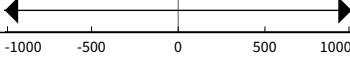
Analysis 6.2. Comparison 6 Cold-water immersion (CWI) versus double CWI, Outcome 2 Range of movement (degrees).
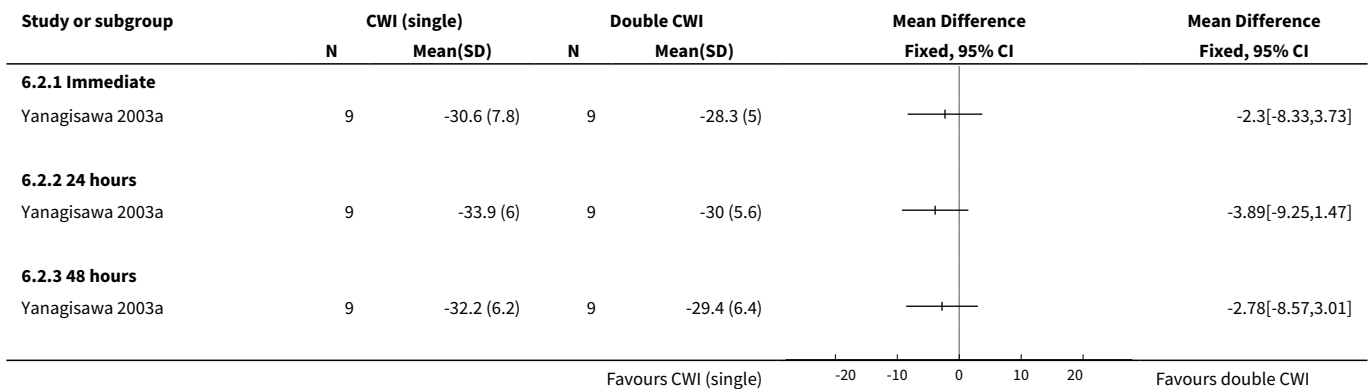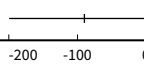
Study or subgroup	CWI (single)		Double CWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
6.2.1 Immediate						
Yanagisawa 2003a	9	-30.6 (7.8)	9	-28.3 (5)		-2.3[-8.33,3.73]
6.2.2 24 hours						
Yanagisawa 2003a	9	-33.9 (6)	9	-30 (5.6)		-3.89[-9.25,1.47]
6.2.3 48 hours						
Yanagisawa 2003a	9	-32.2 (6.2)	9	-29.4 (6.4)		-2.78[-8.57,3.01]

Favours CWI (single) -20 -10 0 10 20 Favours double CWI

Study or subgroup	CWI (single)		Double CWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
6.2.4 96 hours						
Yanagisawa 2003a	9	-31.7 (6.1)	9	-29.4 (6.4)		-2.23[-7.99,3.53]

Favours CWI (single) -20 -10 0 10 20 Favours double CWI

Analysis 6.3. Comparison 6 Cold-water immersion (CWI) versus double CWI, Outcome 3 Biomarker: creatine kinase (U/L).

Study or subgroup	CWI (single)		Double CWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
6.3.1 48 hours						
Yanagisawa 2003a	9	1200 (1700)	9	1200 (800)		0[-1227.48,1227.48]
6.3.2 96 hours						
Yanagisawa 2003a	9	2500 (3750)	9	7500 (8500)		-5000[-11069.65,1069.65]

Favours CWI (single) -1000 -500 0 500 1000 Favours double CWI

Analysis 6.4. Comparison 6 Cold-water immersion (CWI) versus double CWI, Outcome 4 Biomarker: lactate dehydrogenase (U/L).

Study or subgroup	CWI (single)		double CWI		Mean Difference	Mean Difference
	N	Mean(SD)	N	Mean(SD)	Fixed, 95% CI	Fixed, 95% CI
6.4.1 96 hours						
Yanagisawa 2003a	9	25 (50)	9	115 (160)		-90[-199.52,19.52]

Favours CWI (single) -200 -100 0 100 200 Favours double CWI

APPENDICES

Appendix 1. Search strategies

Note: Search strategies for CENTRAL, MEDLINE, EMBASE and CINAHL include line numbers for the search update carried out in November 2011:

Cochrane Central Register of Controlled Trials (Wiley InterScience interface)

#1 MeSH descriptor Cryotherapy, this term only (36)
#2 MeSH descriptor Cold Temperature, this term only (990)
#3 MeSH descriptor Immersion, this term only (203)
#4 (#2 AND #3) (52)
#5 (#1 OR #4) (417)
#6 ((cold or ice or contrast) NEAR/3 (immers* or bath or therapy)):ti,ab,kw (901)
#7 (cryotherapy):ti,ab,kw (727)
#8 (#5 OR #6 OR #7) (1558)
#9 MeSH descriptor Exercise, this term only (7811)
#10 MeSH descriptor Muscle, Skeletal, this term only (4004)
#11 MeSH descriptor Athletic Injuries, this term only (0)
#12 MeSH descriptor Soft Tissue Injuries, this term only (0)
#13 MeSH descriptor Creatine Kinase explode all trees (1111)
#14 MeSH descriptor Physical Exertion, this term only (3090)

#15 MeSH descriptor Muscle Fatigue, this term only (480)
#16 MeSH descriptor Muscle Cramp, this term only (127)
#17 MeSH descriptor Spasm, this term only (171)
#18 MeSH descriptor Muscle Rigidity, this term only (58)
#19 MeSH descriptor Sprains and Strains, this term only (0)
#20 MeSH descriptor Muscle Weakness, this term only (194)
#21 (sore* NEAR/3 musc*):ti,ab,kw (328)
#22 (DOMS):ti,ab,kw (100)
#23 (exercise induced and (muscle* NEAR/2 (damage* or injur*))):ti,ab,kw (124)
#24 MeSH descriptor Lactic Acid, this term only (1497)
#25 (lactate* or lactic):ti,ab,kw (5373)
#26 (#9 OR #10 OR #11 OR #12 OR #13 OR #14 OR #15 OR #16 OR #17 OR #18 OR #19 OR #20 OR #21 OR #22 OR #23 OR #24 OR #25) (18950)
#27 (#8 AND #26), from 2010 to 2011 (12)

MEDLINE (Ovid interface)

1 Cryotherapy/ (3223)
2 Cold Temperature/ (38292)
3 Immersion/ (4157)
4 2 and 3 (682)
5 1 or 4 (3896)
6 ((cold or ice or contrast) adj3 (immers$ or bath or therapy)).tw. (3113)
7 cryotherapy.tw. (4562)
8 5 or 6 or 7 (9925)
9 Exercise/ (57099)
10 Muscle, Skeletal/ (89754)
11 Athletic Injuries/ (18407)
12 Soft Tissue Injuries/ (2989)
13 exp Creatine Kinase/ (22424)
14 Physical Exertion/ (51283)
15 Muscle Fatigue/ (4927)
16 Muscle Cramp/ or Spasm/ or Muscle Rigidity/ or "Sprains and Strains"/ or Muscle Weakness/ (16965)
17 (sore$ adj3 musc$).tw. (964)
18 DOMS.tw. (303)
19 (exercise induced and (muscle$ adj2 (damage$ or injur$))).tw. (488)
20 Lactic Acid/ (27006)
21 (lactate$ or lactic).tw. (95395)
22 or/9-21 (336771)
23 and/8,22 (557)
24 randomized controlled trial.pt. (323380)
25 controlled clinical trial.pt. (84102)
26 randomized.ab. (238622)
27 placebo.ab. (134849)
28 randomly.ab. (174807)
29 trial.ab. (247461)
30 groups.ab. (1147391)
31 29 or 27 or 24 or 26 or 30 or 25 or 28 (1671645)
32 animals/ not (humans/ and animals/) (3631995)
33 31 not 32 (1371077)
34 33 and 23 (157)
35 limit 34 to ed=20100101-20120101 (40)

EMBASE (Ovid Interface)

1 Cryotherapy/ (9985)
2 Low temperature procedures/ or Cold treatment/ or Cooling/ or Cooling water/ or Cold/ (30464)
3 ((cold or ice or contrast) adj3 (immers$ or bath or therapy)).tw. (3510)
4 cryotherapy.tw. (5525)
5 or/1-4 (45294)
6 Exercise/ or Muscle exercise/ or Exercise recovery/ (145790)
7 exp Skeletal Muscle/ (174363)
8 Muscle Injury/ (6215)
9 Soft tissue injury/ or Sport injury/ (25459)

![Cochrane Library logo] Cochrane
Library

Trusted evidence.
Informed decisions.
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Cochrane Database of Systematic Reviews

10   Creatine Kinase/ or Creatine kinase blood level/ (29906)
11   Muscle Fatigue/ (7631)
12   Muscle Stiffness/ or Muscle Cramp/ or Muscle Spasm/ or Muscle Tightness/ or Muscle Rigidity/ or Muscle Strain/ or Muscle Weakness/ (43431)
13   (sore$ adj3 musc$).tw. (1064)
14   DOMS.tw. (359)
15   (exercise induced and (muscle$ adj2 (damage$ or injur$))).tw. (521)
16   Lactic Acid/ (37432)
17   (lactate$ or lactic).tw. (100509)
18   or/6-17 (493490)
19   and/5,18 (2675)
20   Clinical trial/ (821137)
21   Randomized controlled trial/ (292701)
22   Randomization/ (54986)
23   Single blind procedure/ (14442)
24   Double blind procedure/ (101701)
25   Crossover procedure/ (31195)
26   Placebo/ (187497)
27   randomi?ed controlled trial$.tw. (66241)
28   rct.tw. (7997)
29   random allocation.tw. (1065)
30   randomly allocated.tw. (15806)
31   allocated randomly.tw. (1718)
32   (allocated adj2 random).tw. (688)
33   single blind$.tw. (11219)
34   double blind$.tw. (119142)
35   ((treble or triple) adj blind$).tw. (250)
36   placebo$.tw. (161437)
37   Prospective study/ (176527)
38   or/20-37 (1156411)
39   Case study/ (13795)
40   case report.tw. (210181)
41   Abstract report/ or Letter/ (799539)
42   or/39-41 (1019415)
43   38 not 42 (1122896)
44   limit 43 to human (1023188)
45   19 and 44 (330)
46   (2010$ or 2011$).em. (2360483)
47   45 and 46 (75)

CINAHL (EBSCO Interface)

S1   (MH "Cryotherapy") (994)
S2   "water immersion" OR "ice" OR "cold" OR "cooling" (10204)
S3   S1 or S2 (10896)
S4   Muscle or injury or sport or soft tissue or stiffness or cramp or spasm or recovery or exercise or performance (260896)
S5   S3 and S4 (2266)
S6   (MH "Clinical Trials+") (132628)
S7   (MH "Evaluation Research+") (17559)
S8   (MH "Comparative Studies") (64058)
S9   (MH "Crossover Design") (8666)
S10   PT Clinical Trial (68624)
S11   (MH "Random Assignment") (31138)
S12   S6 or S7 or S8 or S9 or S10 or S11 (217079)
S13   TX ((clinical or controlled or comparative or placebo or prospective or randomi?ed) and (trial or study (372280)
S14   TX (random* and (allocat* or allot* or assign* or basis* or divid* or order*)) (54007)
S15   TX ((singl* or doubl* or trebl* or tripl*) and (blind* or mask   (588697)
S16   TX (crossover* or 'cross over') or TX cross n1 over (10971)
S17   TX ((allocat* or allot* or assign* or divid*) and (condition* or experiment* or intervention* or treatment* or therap* or control* or group*)) (67272)
S18   S13 or S14 or S15 or S16 or S17 (890995)
S19   S12 or S18 (945591)

S20   S5 and S19 (1164)
S21 (MH "Animals") (40857)
S22 S20 Not S21 (138)
S23 2010 or 2011 (702465)
S24 S22 and S23 (224)

PEDro

Using the 'Advanced search' option, each term in the 'Abstract & Title' field was combined with each term in the 'Subdiscipline' field in separate searches.

Abstract & Title: cryotherapy/ cold/ immersion/ hydrotherapy/"contrast therapy"/ ice/ water/CWI
Subdiscipline: musculoskeletal/ orthopaedics/ sports/
Method: clinical trial
Match all search terms (AND)

BNI (Ovid Interface)

1 Cryotherapy.mp.
2 immersion.mp.
3 ((cold or ice or contrast) adj3 (immers$ or bath or therapy)).tw.
4 exercise/ or muscle exercise/ or exercise recovery/
5 exp "Musculoskeletal System and Disorders"/
6 soft tissue injury.mp.
7 creatine kinase.mp.
8 muscle.mp.
9 lactic acid.mp.
10 athletic.mp.
11 physical exertion.mp.
12 4 or 5 or 6 or 7 or 8 or 9 or 10 or 11
13 cold.mp.
14 body temperature/ or low temperature.mp.
15 cooling.mp.
16 1 or 2 or 3 or 13 or 14 or 15
17 12 and 16

CONTRIBUTIONS OF AUTHORS

Chris Bleakley developed the research idea, and wrote the original protocol. Evie Gardner provided statistical support.

Selection of studies was carried out by Suzanne McDonough and Chris Bleakley.

Data extraction and interpretation were carried out by Chris Bleakley, Ty Hopkins, Gareth Davison and Philip Glasgow.

Risk of bias assessment and data analysis were carried by Chris Bleakley, David Baxter and Suzanne McDonough.

All authors commented on drafts and approved the final version. Chris Bleakley wrote the final review and is the guarantor.

DECLARATIONS OF INTEREST

None known.

SOURCES OF SUPPORT

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

The order of the two primary outcomes was reversed, with muscle soreness becoming the lead item.

We presented outcome data split into 24 hour instead of 12 hour intervals.

NOTES

The search update in November 2011 to assess recent literature was initiated and carried out by the editorial base of the Cochrane Bone, Joint and Muscle Trauma Group. Study selection was performed by Helen Handoll.

INDEX TERMS

Medical Subject Headings (MeSH)

*Immersion; Cryotherapy [*methods]; Exercise [*physiology]; Muscular Diseases [prevention & control] [*therapy]; Pain [prevention & control]; Pain Management [*methods]; Randomized Controlled Trials as Topic

MeSH check words

Humans