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Exercise and Adiposity in Overweight and Obese Children and Adolescents: Protocol for a Systematic Review and Network Meta-Analysis of Randomised Trials

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ABSTRACT

Introduction: Overweight and obesity is a worldwide public health problem among children and adolescents. However, the magnitude of effect, as well as hierarchy of exercise interventions (aerobic, strength training, or both), on selected measures of adiposity is not well established despite numerous trials on this issue. The primary purposes of this study are to use the network meta-analytic approach to determine the effects and hierarchy of exercise interventions on selected measures of adiposity in overweight and obese children and adolescents. **Methods and analysis:** Randomised exercise intervention trials ≥ 4 weeks, published in any language between January 1, 1973 and August 31, 2017, and which include direct and/or indirect evidence, will be included. Studies will be located by searching five electronic databases, cross-referencing and expert review. Dual selection and abstraction of data will occur. The primary outcomes will be changes in body mass index (BMI in kg m^2), fat mass and percent body fat. Risk of bias will be assessed using the Cochrane Risk of Bias assessment instrument while confidence in the cumulative evidence will be assessed using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) instrument for network meta-analysis. Network meta-analysis will be performed using multivariate random-effects meta-regression models. The surface under the cumulative ranking curve (SUCRA) will be used to provide a hierarchy of exercise treatments (aerobic, strength, or both). **Dissemination:** The findings of this network meta-analysis will be presented at a professional conference and published in a peer-reviewed journal. **Trial registration number:** PROSPERO #CRD42017073103

INTRODUCTION

Rationale

Overweight and obesity in children and adolescents is a major public health problem worldwide. Between 1980 and 2013, the worldwide prevalence of overweight and obesity in children and adolescents increased by 6.9%, from 16.9% to 23.8%, in boys and by 6.4%, from 16.2% to 22.6%, in girls from developed countries.¹ For developing countries, increases of 4.8%, from 8.1% to 12.9% for boys and 5%, from 8.4% to 13.4% in girls, were reported.¹ In the United States, the prevalence of overweight and obesity, defined as a body mass index (BMI) \geq 85th percentile based on Centers for Disease Control Growth Charts, has been reported to be 31.8% among children and adolescents 2 to 19 years of age, while the prevalence of obesity, defined as a BMI \geq 95th percentile, has been reported as 16.9%.² When compared to 30 years ago, this represents an obesity prevalence that is more than two times higher in US children and more than four times higher in adolescents.^{2 3}

The economic costs associated with overweight and obesity among children and adolescents are also substantial. For example, Finkelstein et al. estimated that the incremental lifetime medical cost of an obese 10-year-old child in the US, relative to a normal weight child who maintained normal weight throughout adulthood, was \$19,000.⁴ Based on the number of obese 10-year-olds in the US, the total direct medical costs associated with obesity were estimated at \$14 billion for this age only.⁴

The negative outcomes associated with obesity in children and adolescents are both immediate and long-term.⁵ For immediacy, a population-based study of US children and adolescents 5 to 17 years of age found that approximately 70% of obese youth had

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a minimum of one cardiovascular disease risk factor (high cholesterol, high blood pressure, etc.).⁶ Obese children and adolescents are also more likely to be diagnosed with prediabetes,⁷ as well as being at an increased risk for bone and joint difficulties, sleep apnea, and social and psychological issues such as stigmatization, poor self-esteem, and poorer health-related quality-of-life.^{8 9}

Long-term, childhood and adolescent overweight and obesity has been demonstrated to track into adulthood,¹⁰⁻¹⁴ thus placing overweight and/or obese adults at a greater risk for cardiovascular disease, type 2 diabetes, stroke, several types of cancer, and osteoarthritis.⁵ These long-term outcomes are important given that overweight and obesity has been reported to be the third leading cause of preventable death in the US, responsible for 216,000 deaths in 2005.¹⁵ In addition, more recent research has shown that up to 18 percent of US deaths between 1986 and 2006 were attributed to obesity.¹⁶ Furthermore, this issue has become so problematic that it is now recognized by the American Medical Association as a disease.¹⁷ Not surprisingly, reducing the prevalence of overweight and obesity among children and adolescents is a major public health priority in the US.¹⁸

One promising intervention in the treatment of overweight and obesity is exercise. However, previous randomised trials that were limited to or included overweight and obese children and adolescents have led to conflicting results,¹⁹⁻⁶⁵ with some reporting statistically significant reductions in adiposity (BMI) as a primary outcome^{19 20 23 24 29 34 35} and others reporting no change.^{21 22 25-28 30-33 36 37 39-47 49-57 64 65 69 71 72} When limited to overweight and obese male and female children and adolescents,^{19 21 24-27 29-33} only 18 (45.0%) have reported statistically significant

104 reductions in BMI.^{19 24 29 35 38 48 58,59-63 65, 50, 52, 54, 56, 57} While this may lead one to the
105 general conclusion that exercise does little to reduce BMI in overweight and obese
106 children and adolescents, this would be shortsighted since it relies on the vote-counting
107 approach,⁷³ an approach that has been shown to be less valid than the meta-analytic
108 approach.^{73 74}

109 Previous systematic reviews with meta-analyses that have focused on the effects of
110 exercise as an independent intervention on BMI as a primary outcome in male and
111 female children and adolescents have reported conflicting findings with five reporting a
112 significant improvement in BMI⁷⁵⁻⁷⁹ and five others reporting no statistically significant
113 improvement.⁸⁰⁻⁸⁴ However, nine of the ten suffer from one or more of the following
114 limitations: (1) inclusion of a small number of studies with exercise as the only
115 intervention,^{78 80-82} (2) inclusion of non-randomised trials,^{75 81} (3) inclusion of children
116 and adolescents who were not overweight or obese.^{77 79 81 83 84} Relevant to this
117 application, all ten suffer from both reliance on pairwise versus network meta-analysis,
118 the latter of which incorporates both direct and indirect evidence. In addition, there was
119 an absence of an established hierarchy for determining which types of exercise
120 (aerobic, strength training, or both) might be best for improving BMI based on both
121 direct and indirect evidence.⁷⁵⁻⁸⁴ To partially address this issue as well as demonstrate
122 feasibility, the investigative team has recently used the network meta-analytic approach
123 to examine the effects of exercise (aerobic, strength training, or both) on BMI z-score in
124 overweight and obese children and adolescents.^{85 86} Statistically significant reductions
125 in BMI z-score were found for aerobic exercise and combined aerobic and strength
126 exercise, but not strength training alone (mean, 95% CI: aerobic, -0.10, -0.15 to -0.05;

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aerobic and strength, -0.11, -0.19 to -0.03; strength, 0.04, -0.07 to 0.15).⁸⁶ Combined aerobic and strength training was ranked best, followed by aerobic exercise and then strength training.⁸⁶ Consistency in evidence and risk of bias did not differ between direct and indirect studies.⁸⁶ It was concluded that combined aerobic exercise and strength training as well as aerobic exercise alone are associated with reductions in BMI z-score.⁸⁶ The lack of effect on BMI z-score in the strength training studies may have been the result of increases in lean muscle mass. However, since BMI in kg·m² continues to be the most frequently assessed and reported measure of adiposity in both the clinical and public health setting, an examination of such using the network meta-analytic approach is needed. In addition, since all types of BMI measures as well as body weight do not capture changes in body composition (fat mass, percent body fat, etc.), the inclusion of such outcomes, as previously suggested,⁸⁶ is also necessary.

Objectives

The primary objectives of the current study are to conduct a systematic review with network meta-analysis of randomised trials to (1) determine the effects of exercise (aerobic, strength training, or both) on adiposity (BMI in kg·m², fat mass, percent body fat) in overweight and obese children and adolescents, and (2) establish a hierarchy of exercise interventions (aerobic, strength training, or both) for treating adiposity (BMI in kg·m², fat mass, percent body fat) in overweight and obese children and adolescents.

METHODS

Overview

This study will follow the guidelines from the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) extension statement for network meta-analyses

of health care interventions.⁸⁷ The protocol for this network meta-analysis is registered in PROSPERO (trial registration number CRD42017073103).

Eligibility criteria

The inclusion criteria for this proposed network meta-analysis will be as follows: (1) direct evidence from randomised trials that compare two or more exercise interventions (aerobic, strength training, both) or indirect evidence from randomised controlled trials that compare an exercise intervention group to a comparative control group (non-intervention, attention control, usual care, wait-list control, placebo), (2) exercise-only intervention (aerobic, strength training, or both), (3) studies lasting ≥ 4 weeks, (4) male and/or female children and adolescents 2 to 18 years of age, (5) participants overweight or obese, as defined by the authors, (6) studies published in any language that include an English language abstract, (7) studies published between January 1, 1973 and August 31, 2017, and (8) data available for BMI in $\text{kg}\cdot\text{m}^2$, fat mass or percent body fat.

Studies will be limited to randomised trials because it is the only way to control for confounders that are not known or measured as well as the observation that nonrandomised controlled trials tend to overestimate the effects of healthcare interventions.^{88 89} Indirect evidence studies will be limited to randomised controlled trials with at least one exercise arm that participates in either aerobic, strength training, or a combination of aerobic and strength training exercise. Direct evidence studies will be limited to randomised trials that include at least two of the following exercise arms: (1) aerobic, (2) strength training, (3) aerobic and strength training exercise.

For the purposes of this study, exercise, aerobic exercise and strength training will be defined according to the 2008 Physical Activity Guidelines for Americans,⁹⁰ defined as

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173 movement that is “planned, structured, and repetitive and purposive in the sense that
174 the improvement or maintenance of one or more components of physical fitness is the
175 objective,”^{90 91} aerobic exercise as “exercise that primarily uses the aerobic energy-
176 producing systems, can improve the capacity and efficiency of these systems, and is
177 effective for improving cardiorespiratory endurance,”⁹⁰ and strength training as “exercise
178 training primarily designed to increase skeletal muscle strength, power, endurance, and
179 mass”.⁹⁰ Four weeks was chosen as the lower cut point for intervention length based
180 on previous research demonstrating improvements in adiposity over this period of time
181 in 11-year olds.²⁸

182 Participants will be limited to overweight and obese children and adolescents, as
183 defined by the original study authors, because it has been shown that this population is
184 at an increased risk for premature morbidity and mortality throughout their lifetime.⁹²

185 Studies will be limited to published articles and examined for potential small-study
186 effects such as publication bias. Unpublished work, defined as master’s theses,
187 dissertations, abstracts from conference proceedings, technical reports, and studies
188 conducted but never reported, will not be included. The rationale for this approach is
189 based on the work of van Driel et al.⁹³ who concluded that (1) the difficulty in retrieving
190 unpublished work could lead to selection bias, (2) many unpublished trials are
191 eventually published, (3) the methodological quality of such studies are poorer than
192 those that are published, and (4) the effort and resources required to obtain unpublished
193 work may not be warranted.⁹³

194 While some research has suggested that studies yielding statistically significant and
195 positive results are more likely to be published in English-language versus non-English

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3 196 language journals,⁹⁴ other research has shown this to not be the case.⁹⁵ Given the
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5 197 former, studies from both English and non-English-language articles will be included
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7 198 with the latter translated into English by the second author using the freely available
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9 199 web-based Babelfish and Bing translators. For those studies that cannot be translated
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11 200 using Babelfish and/or Bing, professional translation services will be utilized.
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15 201 The year 1973 was chosen as the starting point for searching based on a preliminary
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17 202 PubMed search that yielded the first study that met the search, but not necessarily
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19 203 eligibility, criteria.⁹⁶ Body mass index in kg m^2 was included as one of the three primary
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21 204 adiposity outcomes because it is the most commonly used and understood variable by
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23 205 practitioners as well as others and can be easily measured from body weight and
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25 206 height. However, because BMI is an indirect measure of adiposity, fat mass and percent
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27 207 body fat will be included because they are more direct measures of adiposity. The
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29 208 inclusion of fat mass and percent body fat may be especially relevant for studies that
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31 209 include strength training given that decreases in adiposity as measured by BMI may be
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33 210 offset by increases in muscle mass, a secondary outcome that will be coded.
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37 38 211 **Information sources**

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40 212 The following five electronic databases will be searched: (1) PubMed, (2) Web of
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42 213 Science, (3) Cochrane Central Register of Controlled Trials (CENTRAL), (4) Cumulative
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44 214 Index to Nursing and Allied Health Literature (CINAHL), and (5) Sport Discus. In
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46 215 addition to electronic database searches, cross-referencing will be conducted by
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48 216 examining the reference lists of previous review articles as well as each included study
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50 217 for potential articles that meet the inclusion criteria. Upon completion of initial searches,
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the third author will examine the reference list for thoroughness and completeness. Suggested studies will then be retrieved to see if they meet all inclusion criteria.

Search strategy

Search strategies specific to each database will be developed by the investigative team. Major keywords, or forms of keywords to include will be “random”, “children”, “adolescents”, “overweight”, “obese”, “exercise,” “physical fitness”, “body composition”, “fat mass”, “body fat”, “body composition”, “body mass index”, “adiposity”. Searches will be limited to studies published and indexed between January 1, 1973 and August 31, 2017, approximately 34 years. A copy of a preliminary search strategy using PubMed, including limits, can be found in Supplementary file 1. This search strategy will be adapted for other database searches. All database searches and article retrieval will be conducted by the second author with oversight from the first author.

Study records

Study selection

All studies to be screened will be imported into EndNote (version X8; New York, NY: Thomson-Reuters; 2016) and duplicates removed electronically and then manually by the second author. A copy of the database will then be provided to the first author for duplicate screening. To minimize selection bias, the first and second authors will select all studies, independent of each other. They will then review their selections for accuracy and consistency. The full report for each article will be retrieved for all titles and abstracts that appear to meet the inclusion criteria as well as those where uncertainty exists. Multiple reports for the same study will be addressed by including the most recently published article and drawing from prior reports, assuming the same methods

and sample sizes are reported. Based on previous research suggesting neither a clinically nor statistically significant effect on results, blinding to journal titles, study authors, or institutions of the authors will not be employed during the screening and data abstraction processes.⁹⁷ Reasons for excluded studies will be recorded using the following categories: (1) inappropriate population, (2) inappropriate intervention, (3) inappropriate comparison(s), (4) inappropriate outcome(s), (5) inappropriate study design, (6) other. Upon the conclusion of screening, the first and second authors will meet and review their selections. Cohen's kappa statistic (κ) will be used to measure inter-selection agreement.⁹⁸ Any discrepancies will be resolved by consensus. If consensus cannot be reached, the third author will serve as an arbitrator. Upon selecting the final number of studies to include, the overall precision of the searches will be computed by dividing the number of included studies by the total number of studies screened after removing duplicates.⁹⁹ The number needed-to-read (NNR) will then be calculated as the reciprocal of the precision.⁹⁹ A flow diagram that describes the search procedure will be included as well as a supplementary file that includes a reference list of all excluded studies, including the reason(s) for exclusion. Figure 1 illustrates the proposed structure for the flow diagram.

Data abstraction

For this project, Microsoft Excel (version 2016; Redmond, WA: Microsoft Corporation; 2016) will be used to develop comprehensive electronic codebooks that will define the coding process for each of the variables coded. The codebook will be created by the first two authors with feedback from the third author. Consequently, the abstraction of data from the studies in this proposed project should require little subjective judgment

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on the part of the coder. The major groups of variables to code will include (1) study characteristics (author, journal, year of publication, etc.), (2) participant characteristics (age, gender, height, body weight, etc.), and (3) data for primary and secondary outcomes (sample sizes, baseline and post-exercise means and standard deviations, etc.). Table 1 contains a preliminary list of variables that will be coded. Based on previous research by the investigative team,⁸⁶ a codebook capable of including at least 242 items from each study is expected. To avoid data abstraction bias, the first two authors will independently code (dual-coding) all studies to ensure accuracy and consistency. Inter-rater agreement will be assessed using Cohen’s kappa statistic (κ).⁹⁸ Any disagreement in the items coded will be discussed until mutual agreement is reached. If agreement cannot be reached, the third author will serve as an arbitrator.

Outcomes and prioritization

The primary outcomes in this study will be changes BMI in kg m², fat mass, and percent body fat in overweight and obese children and adolescents. Secondary outcomes will include body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, physical activity level, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin.

Risk of bias assessment in individual studies

Risk of bias for included studies will be assessed using the Cochrane Risk of Bias

Instrument.¹⁰⁰ Assessment is based on judgments of low, high or unclear risk of bias across six defined domains: (1) sequence generation, (2) allocation sequence concealment, (3) blinding of participants and personnel, (4) blinding of outcome assessors, (5) incomplete outcome data, and (6) selective outcome reporting. A seventh domain, whether participants were exercising regularly, as defined by the original study authors prior to taking part in the study, will also be assessed. This risk of bias approach has been recommended over the use of study quality rating scales given the lack of empirical evidence to support the latter.^{89 101 102} Assessment for risk of bias will be limited to the primary outcomes of interest, i.e., changes in BMI in kg·m², fat mass, and percent body fat. All studies will be classified as high risk of bias with respect to the category “blinding of participants and personnel” given that it’s virtually impossible to blind participants to group assignment in exercise intervention protocols. Based on previous research, no study will be excluded based on risk of bias results.¹⁰³

Data Synthesis

Calculation of effect sizes

The primary outcomes for this study will be changes in BMI in kg·m², fat mass (kg), and percent body fat using the original metric. Changes for indirect comparisons will be calculated by subtracting the change outcome difference in the exercise group minus the change outcome difference in the control group. Variances will be computed using the pooled standard deviations of change scores in the exercise and control groups. If change score standard deviations are not available, they will be calculated from 95% confidence intervals (CI) for either change outcome or treatment effect differences as well as pre and post standard deviation values, the latter according to procedures

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developed by Follmann et al.¹⁰⁴ For direct comparisons, i.e., randomised trials with no control group, the same general procedures will be followed except that the control group data will be replaced with one of the exercise interventions as follows: (1) aerobic minus strength training, (2) aerobic and strength training combined minus aerobic training, (3) aerobic and strength training combined minus strength training. Ninety-five percent CI and z-alpha values will be calculated for each outcome from each study. For those studies that include both direct and indirect comparisons, only direct comparison data will be included since a primary purpose of the current meta-analysis is determining which exercise interventions(s) might work best for improving adiposity in children and adolescents. For studies in which adiposity outcomes are assessed at multiple intervention time points, for example, 0, 8, and 16 weeks, only data from the initial and last assessment will be used. If follow-up data are available, results from such will also be analyzed separately to determine the sustainability of changes in adiposity. If any crossover trials are included, treatment effects will be calculated by using all assessments from the intervention and control periods and analyzing them similar to a parallel group trial.¹⁰⁵ While the possibility of a unit-of-analysis error exists as well as studies being under versus over-weighted, this method is believed to be better than alternative approaches, for example, limiting data from the first assessment point or trying to impute standard deviations, especially given the primary and secondary outcomes included and expected distribution of findings.¹⁰⁵

Secondary outcomes (body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, maximum oxygen consumption (relative and absolute), resting systolic and diastolic blood pressure, total cholesterol, high-density

lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin) will be handled using the same approach as for primary outcomes. However, given the different metrics expected and the inability to convert between them, changes in physical activity levels and muscular strength will be calculated using the standardized effect size, adjusted for small sample sizes.¹⁰⁶

Pooled estimates for changes in outcomes

Network (geometry) plots for each outcome will be used to provide a visual representation of the evidence base with nodes (circles) weighted by the number of participants randomised to each treatment and edges (lines) weighted by the number of studies evaluating each pair of treatments.^{107 108} *Contribution plots* for each outcome will be used to determine the most dominant comparisons for each network estimate as well as for the entire network.¹⁰⁷ The weights applied will be a function of the variance of the direct treatment effect and the network structure, the result being a percent contribution of each direct comparison to each network estimate.¹⁰⁷

Network meta-analysis will be performed using *multivariate random-effects meta-regression models* that can be performed within a frequentist setting, allows for the inclusion of potential covariates, and correctly accounts for the correlations from multi-arm trials.^{109 110} A two-tailed alpha value ≤ 0.05 and non-overlapping 95% CI will be considered to represent statistically significant changes. Separate network meta-analysis models will be used to examine for changes in each primary and secondary outcome. Potential *covariates* will be examined by (1) conducting simple meta-

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356 regression for statistically significant associations between covariates and changes in
357 primary outcomes (BMI in kg·m², fat mass, percent fat), (2) examining for
358 multicollinearity between covariates ($r > 0.80$), and (3) building a multiple meta-
359 regression model. A list of potential covariates to examine using simple meta-
360 regression is shown in Table 1. *Transitivity*, i.e., similarity in the distribution of potential
361 effect modifiers across the different pairwise comparisons for each outcome¹¹¹ will
362 include those listed in Table 1. *Inconsistency*, i.e., differences in effect estimates
363 between direct and indirect results for the same comparison,¹¹² will be checked by
364 assessing differences in treatment effects between direct and indirect effect estimates
365 as well as differences between trials with different designs, for example, two-arm versus
366 multi-arm trials.^{110 112 113} However, the probability of inconsistency is considered small
367 given recent research demonstrating that inconsistency was detected in only 2% to 14%
368 of tested loops, depending on the effect measure and heterogeneity estimation
369 method.^{114 115} Finally, *prediction intervals* will be used to enhance interpretation of
370 results with respect to the magnitude of heterogeneity as well as provide an estimate of
371 expected results in a future study.¹¹⁶⁻¹¹⁸ For network meta-analysis, degrees of freedom
372 (*df*) will be set to the number of studies – the number of comparisons – 1.¹¹⁸
373
374 *Meta-biases*
375 *Small-study-effects* (publication bias, etc.) will be assessed using comparison adjusted
376 funnel plots.¹⁰⁷ In the absence of small-study effects, the comparison adjusted funnel
377 plot should be symmetric around the zero line.
378
379 Confidence in cumulative evidence

378 *Quality analysis* of specific pairwise effect estimates in the network meta-analysis will
379 be evaluated using a recently developed modification of the Grading of
380 Recommendations Assessment, Development and Evaluation (GRADE) for network
381 meta-analysis across five domains: (1) study limitations, (2) indirectness, (3)
382 inconsistency, (4) imprecision, and (5) small-study effects.¹¹⁹ Assessment will be
383 conducted using the same procedures as for study selection and data abstraction.

384 To establish a hierarchy of exercise interventions for selected outcomes in the current
385 meta-analysis, *ranking analysis*, i.e., the ability to rank all interventions for a single
386 outcome under study, for example changes in BMI in kg m², will be used based on
387 probabilities. However, because the ranking of treatments based exclusively on the
388 probability of each treatment being the best should be avoided given that it does not
389 account for the uncertainty in the relative treatment effects and the possibility for
390 assigning higher ranks for treatments in which little evidence is available, separate
391 *rankograms and cumulative ranking probability plots* will be used to present ranking
392 probabilities along with their uncertainty for changes in primary and secondary
393 outcomes.^{107 120} The surface under the cumulative ranking curve (SUCRA), a
394 transformation of the mean rank, will be used to establish a hierarchy of exercise
395 interventions (aerobic, strength, both) while accounting for the location and variance of
396 all treatment effects.^{107 120} Larger SUCRA values indicate better ranks for the
397 treatment.^{107 120} Interpretation of all rankings will be approached from the perspective of
398 absolute and relative treatment effects.¹⁰⁸

399 Software used for statistical analysis

All data will be analysed using Stata (V.14.1; Stata/SE for Windows, version 14.0. College Station, TX: Stata Corporation LP; 2015), Microsoft Excel (version 2016; Redmond, WA: Microsoft Corporation; 2016), and two add-ins for Excel, SSC-Stat (V.2.18; SSC-Stat, version 3.0. University of Reading, United Kingdom: Statistical Services Center; 2007), and EZ-Analyze (V.3.0; EZ Analyze, version 3.0. TA Poynton; 2007).

DISSEMINATION

The results of this study will be presented at a professional conference and published in a peer-reviewed journal.

CONTRIBUTORS

GAK is the guarantor. GAK, KSK and RRP drafted the manuscript. All authors contributed to (1) the development of the data sources to search for relevant literature, including search strategy, (2) selection criteria, (3) data extraction criteria and (4) risk of bias assessment strategy. GAK provided statistical expertise while RRP provided content expertise on exercise and adiposity in overweight and obese children and adolescents. All authors read, provided feedback, and approved the final manuscript.

REGISTRATION

In accordance with the Primary Reporting Items for Systematics Reviews and Meta-Analyses Protocols (PRISMA-P) statement, this systematic review with network meta-analysis was registered with the International Prospective Register of Systematic Reviews (PROSPERO) on August 23, 2017 (#CRD42017073103).

AMENDMENTS TO PROTOCOL

422 None to date. If this protocol is amended, the date of each amendment, a description of
423 the change, as well as a rationale for the change, will be provided.

424 **COMPETING INTERESTS**

425 None.

426 **FUNDING**

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429 of the authors and does not necessarily represent the official views of the American
430 Heart Association.

431 **DATA SHARING STATEMENT**

432 All data will be available upon request from the corresponding author.

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REFERENCES

1. Ng M, Fleming T, Robinson M, et al. Global, regional, and national prevalence of overweight and obesity in children and adults during 1980-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2014;384:766-81.

2. Ogden CL, Carroll MD, Kit BK, et al. Prevalence of childhood and adult obesity in the United States, 2011-2012. *JAMA* 2014;311(8):806-14.

3. National Center for Health Statistics. Health, United States, 2011: Socioeconomic Status and Health. Hyattsville, MD, 2012.

4. Finkelstein EA, Graham WCK, Malhotra R. Lifetime direct medical costs of childhood obesity. *Pediatrics* 2014;133(5):854-62.

5. Centers for Disease Control and Prevention. Childhood obesity facts. 8-27-2015. Atlanta, Georgia, US Department of Health & Human Services, Centers for Disease Control and Prevention. 12-3-2015

6. Freedman DS, Mei Z, Srinivasan SR, et al. Cardiovascular risk factors and excess adiposity among overweight children and adolescents: the Bogalusa Heart Study. *J Pediatr* 2007;150(1):12-17.

7. Li C, Ford ES, Zhao G, et al. Prevalence of pre-diabetes and its association with clustering of cardiometabolic risk factors and hyperinsulinemia among U.S. adolescents: National Health and Nutrition Examination Survey 2005-2006. *Diabetes Care* 2009;32(2):342-47.

8. Daniels SR, Arnett DK, Eckel RH, et al. Overweight in children and adolescents: pathophysiology, consequences, prevention, and treatment. *Circulation* 2005;111(15):1999-2012.

9. Dietz WH. Overweight in childhood and adolescence. *N Engl J Med* 2004;350(9):855-57.
10. Singh AS, Mulder C, Twisk JW, et al. Tracking of childhood overweight into adulthood: a systematic review of the literature. *Obes Rev* 2008;9(5):474-88.
11. Guo SS, Chumlea WC. Tracking of body mass index in children in relation to overweight in adulthood. *Am J Clin Nutr* 1999;70(1):145S-48S.
12. Freedman DS, Khan LK, Serdula MK, et al. the relation of childhood bmi to adult adiposity: The Bogalusa Heart Study. *Pediatrics* 2005;115(1):22-27.
13. Freedman DS, Wang J, Thornton JC, et al. Classification of body fatness by body mass index-for-age categories among children. *Arch Pediatr Adolesc Med* 2009;163(9):805-11.
14. Freedman DS, Khan LK, Dietz WH, et al. Relationship of childhood obesity to coronary heart disease risk factors in adulthood: the Bogalusa Heart Study. *Pediatrics* 2001;108:712-18.
15. Danaei G, Ding EL, Mozaffarian D, et al. The preventable causes of death in the United States: comparative risk assessment of dietary, lifestyle, and metabolic risk factors. *PLoS Med* 2009;6(4):e1000058.
16. Masters RK, Reither EN, Powers DA, et al. The impact of obesity on US mortality levels: the importance of age and cohort factors in population estimates. *Am J Public Health* 2013;103(10):1895-901.
17. American Medical Association. Obesity as a Disease, 2013. Document 420 (A13).
18. US Department of Health & Human Services. Healthy People 2020. 2-24-2011. Washington, DC. 2010.

19. Alves JG, Gale CR, Souza E, et al. [Effect of physical exercise on bodyweight in overweight children: a randomized controlled trial in a Brazilian slum]. *Cad Saude Publica* 2008;24(Suppl 2):S353-S59.

20. Benson AC, Torode ME, Fiatarone Singh MA. The effect of high-intensity progressive resistance training on adiposity in children: a randomized controlled trial. *Int J Obes (Lond)* 2008;32(6):1016-27.

21. Daley AJ, Copeland RJ, Wright NP, et al. Exercise therapy as a treatment for psychopathologic conditions in obese and morbidly obese adolescents: a randomized, controlled trial. *Pediatrics* 2006;118(5):2126-34.

22. Donnelly JE, Greene JL, Gibson CA, et al. Physical Activity Across the Curriculum (PAAC): a randomized controlled trial to promote physical activity and diminish overweight and obesity in elementary school children. *Prev Med* 2009;49(4):336-41.

23. Duncan MJ, Al-Nakeeb Y, Nevill AM. Effects of a 6-week circuit training intervention on body esteem and body mass index in British primary school children. *Body Image* 2009;6(3):216-20.

24. Farpour-Lambert NJ, Aggoun Y, Marchand LM, et al. Physical activity reduces systemic blood pressure and improves early markers of atherosclerosis in pre-pubertal obese children. *J Am Coll Cardiol* 2009;54(25):2396-406.

25. Gutin B, Owens S, Slavens G, et al. Effect of physical training on heart-period variability in obese children. *J Pediatr* 1997;130(6):938-43.

26. Gutin B, Owens S, Okuyama T, et al. Effect of physical training and its cessation on percent fat and bone density of children with obesity. *Obes Res* 1999;7(2):208-14.

27. Hagstromer M, Elmberg K, Marild S, et al. Participation in organized weekly physical exercise in obese adolescents reduced daily physical activity. *Acta Paediatr* 2009;98(2):352-54.
28. Jago R, Jonker ML, Missaghian M, et al. Effect of 4 weeks of Pilates on the body composition of young girls. *Prev Med* 2006;42(3):177-80.
29. Karacabey K. The effect of exercise on leptin, insulin, cortisol and lipid profiles in obese children. *J Int Med Res* 2009;37(5):1472-78.
30. Kaufman C, Kelly AS, Kaiser DR, et al. Aerobic-exercise training improves ventilatory efficiency in overweight children. *Pediatr Exerc Sci* 2007;19(1):82-92.
31. Kelly AS, Wetzsteon RJ, Kaiser DR, et al. Inflammation, insulin, and endothelial function in overweight children and adolescents: The role of exercise. *J Pediatr* 2004;145(6):731-36.
32. Kim ES, Im JA, Kim KC, et al. Improved insulin sensitivity and adiponectin level after exercise training in obese Korean youth. *Obesity (Silver Spring)* 2007;15(12):3023-30.
33. Kim HJ, Lee S, Kim TW, et al. Effects of exercise-induced weight loss on acylated and unacylated ghrelin in overweight children. *Clin Endocrinol (Oxf)* 2008;68(3):416-22.
34. Kriemler S, Zahner L, Schindler C, et al. Effect of school based physical activity programme (KISS) on fitness and adiposity in primary schoolchildren: cluster randomised controlled trial. *Br Med J* 2010;340:c785.

35. Li YP, Hu XQ, Schouten EG, et al. Report on childhood obesity in China: effects and sustainability of physical activity intervention on body composition of Chinese youth. *Biomed Environ Sci* 2010;23(3):180-87.

36. Macias-Cervantes MH, Malacara JM, Garay-Sevilla ME, et al. Effect of recreational physical activity on insulin levels in Mexican/Hispanic children. *Eur J Pediatr* 2009;168(10):1195-202.

37. Martinez-Vizcaino V, Salcedo AF, Franquelo GR, et al. Assessment of an after-school physical activity program to prevent obesity among 9- to 10-year-old children: a cluster randomized trial. *Int J Obes (Lond)* 2008;32(1):12-22.

38. Meyer AA, Kundt G, Lenschow U, et al. Improvement of early vascular changes and cardiovascular risk factors in obese children after a six-month exercise program. *J Am Coll Cardiol* 2006;48(9):1865-70.

39. Mo-suwan L, Pongprapai S, Junjana C, et al. Effects of a controlled trial of a school-based exercise program on the obesity indexes of preschool children. *Am J Clin Nutr* 1998;68(5):1006-11.

40. Murphy EC, Carson L, Neal W, et al. Effects of an exercise intervention using Dance Dance Revolution on endothelial function and other risk factors in overweight children. *Int J Pediatr Obes* 2009;4(4):205-14.

41. Neumark-Sztainer D, Story M, Hannan PJ, et al. New Moves: a school-based obesity prevention program for adolescent girls. *Prev Med* 2003;37(1):41-51.

42. Pate RR, Ward DS, Saunders RP, et al. Promotion of physical activity among high-school girls: a randomized controlled trial. *Am J Public Health* 2005;95(9):1582-87.

43. Petty KH, Davis CL, Tkacz J, et al. Exercise effects on depressive symptoms and self-worth in overweight children: a randomized controlled trial. *J Pediatr Psychol* 2009;34(9):929-39.
44. Reilly JJ, Kelly L, Montgomery C, et al. Physical activity to prevent obesity in young children: cluster randomised controlled trial. *Br Med J* 2006;333(7577):1041.
45. Rooney BL, Gritt LR, Havens SJ, et al. Growing healthy families: family use of pedometers to increase physical activity and slow the rate of obesity. *Wis Med J* 2005;104(5):54-60.
46. Shaibi GQ, Cruz ML, Ball GD, et al. Effects of resistance training on insulin sensitivity in overweight Latino adolescent males. *Med Sci Sports Exerc* 2006;38(7):1208-15.
47. Simon C, Schweitzer B, Oujaa M, et al. Successful overweight prevention in adolescents by increasing physical activity: a 4-year randomized controlled intervention. *Int J Obes (Lond)* 2008;32(10):1489-98.
48. Tan S, Yang C, Wang J. Physical training of 9- to 10-year-old children with obesity to lactate threshold intensity. *Pediatr Exerc Sci* 2010;22(3):477-85.
49. Velez A, Golem DL, Arent SM. The impact of a 12-week resistance training program on strength, body composition, and self-concept of Hispanic adolescents. *J Strength Cond Res* 2010;24(4):1065-73.
50. Walther C, Gaede L, Adams V, et al. Effect of increased exercise in school children on physical fitness and endothelial progenitor cells: a prospective randomized trial. *Circulation* 2009;120(22):2251-59.

51. Warren JM, Henry CJ, Lightowler HJ, et al. Evaluation of a pilot school programme aimed at the prevention of obesity in children. *Health Promot Int* 2003;18(4):287-96.

52. Watts K, Beye P, Siafarikas A, et al. Exercise training normalizes vascular dysfunction and improves central adiposity in obese adolescents. *J Am Coll Cardiol* 2004;43(10):1823-27.

53. Watts K, Beye P, Siafarikas A, et al. Effects of exercise training on vascular function in obese children. *J Pediatr* 2004;144(5):620-25.

54. Weintraub DL, Tirumalai EC, Haydel KF, et al. Team sports for overweight children: the Stanford Sports to Prevent Obesity Randomized Trial (SPORT). *Arch Pediatr Adolesc Med* 2008;162(3):232-37.

55. Sigal RJ, Alberga AS, Goldfield GS, et al. Effects of aerobic training, resistance training, or both on percentage body fat and cardiometabolic risk markers in obese adolescents: the healthy eating aerobic and resistance training in youth randomized clinical trial. *JAMA Pediatr* 2014;168(11):1006-14.

56. Meucci M, Cook C, Curry CD, et al. Effects of supervised exercise program on metabolic function in overweight adolescents. *World J Pediatr* 2013;9(4):307-11.

57. Lee S, Deldin AR, White D, et al. Aerobic exercise but not resistance exercise reduces intrahepatic lipid content and visceral fat and improves insulin sensitivity in obese adolescent girls: a randomized controlled trial. *Am J Physiol Endocrinol Metab* 2013;305(10):E1222-E29.

58. Dennis BA, Ergul A, Gower BA, et al. Oxidative stress and cardiovascular risk in overweight children in an exercise intervention program. *Childhood Obesity* 2013;9(1):15-21.

59. Maddison R, Foley L, Ni MC, et al. Effects of active video games on body composition: a randomized controlled trial. *Am J Clin Nutr* 2011;94(1):156-63.
60. Wong PC, Chia MY, Tsou IY, et al. Effects of a 12-week exercise training programme on aerobic fitness, body composition, blood lipids and C-reactive protein in adolescents with obesity. *Ann Acad Med Singapore* 2008;37(4):286-93.
61. Saygin O, Ozturk MA. The effect of twelve week aerobic exercise programme on health related physical fitness components and blood lipids in obese girls. *Afr J Pharm Pharmacol* 2011;5(12):1441-45.
62. Sun MX, Huang XQ, Yan Y, et al. One-hour after-school exercise ameliorates central adiposity and lipids in overweight Chinese adolescents: a randomized controlled trial. *Chin Med J (Engl)* 2011;124(3):323-29.
63. Elloumi M, Makni E, Ounis OB, et al. Six-minute walking test and the assessment of cardiorespiratory responses during weight-loss programmes in obese children. *Physiother Res Int* 2011;16(1):32-42.
64. Lau PWC, Wong dP, Ngo JK, et al. Effects of high-intensity intermittent running exercise in overweight children. *Eur J Sport Sci* 2014:1-9.
65. Alberga AS, Farnesi BC, Lafleche A, et al. The effects of resistance exercise training on body composition and strength in obese prepubertal children. *Phys Sportsmed* 2013;41(3):103-09.
66. Ackel-D'Elia C, Carnier J, Bueno CR, Jr., et al. Effects of different physical exercises on leptin concentration in obese adolescents. *Int J Sports Med* 2014;35(2):164-71.

67. Fazelifar S, Ebrahim K, Sarkisian V. Effect of concurrent training and detraining on anti-inflammatory biomarker and physical fitness levels in obese children. *Rev Bras Med Esporte* 2013;19(5):349-54.

68. Ghorbanian B, Ravassi A, Kordi MR, et al. The effects of rope training on lymphocyte ABCA1 expression, plasma ApoA-I and HDL-c in boy adolescents. *Int J Endocrinol Metab Disord* 2013;11(2):76-81.

69. Lee S, Bacha F, Hannon T, et al. Effects of aerobic versus resistance exercise without caloric restriction on abdominal fat, intrahepatic lipid, and insulin sensitivity in obese adolescent boys: a randomized, controlled trial. *Diabetes* 2012;61(11):2787-95.

70. Song JK, Stebbins CL, Kim TK, et al. Effects of 12 weeks of aerobic exercise on body composition and vascular compliance in obese boys. *J Sports Med Phys Fitness* 2012;52(5):522-29.

71. Cheng HL, Peng P, Zhu R, et al. [Effects of eight weeks exercise prescription intervention on aerobic capacity, body composition, blood lipid and C-reactive protein in obese adolescents.] *J Jilin Univ Med* 2012;38:745-49.

72. Kelly LA, Loza A, Lin X, et al. The effect of a home-based strength training program on type 2 diabetes risk in obese Latino boys. *J Pediatr Endocrinol Metab* 2015;28(3-4):315-22.

73. Hedges LV, Olkin I. Vote-counting methods in research synthesis. *Psychol Bull* 1980;88:359-69.

74. James A, Soler A, Weatherall R. Cognitive behavioural therapy for anxiety disorders in children and adolescents. *Cochrane Database Syst Rev* 2005(4):CD004690.

75. Schranz N, Tomkinson G, Olds T. What is the effect of resistance training on the strength, body composition and psychosocial status of overweight and obese children and adolescents? A systematic review and meta-analysis. *Sports Med* 2013;43(9):893-907.
76. Kelley GA, Kelley KS, Pate RR. Exercise and BMI in overweight and obese children and adolescents: A systematic review with trial sequential meta-analysis. *Biomed Res Int* 2015;2015(Article ID 704539):1-17.
77. Mei H, Xiong Y, Xie S, et al. The impact of long-term school-based physical activity interventions on body mass index of primary school children - a meta-analysis of randomized controlled trials. *BMC Public Health* 2016;16(1):205.
78. Garcia-Hermoso A, Sanchez-Lopez M, Martinez-Vizcaino V. Effects of aerobic plus resistance exercise on body composition related variables in pediatric obesity: A systematic review and meta-analysis of randomized controlled trials. *Pediatr Exerc Sci* 2015;27(4):431-40.
79. Dellert JC, Johnson P. Interventions with children and parents to improve physical activity and body mass index: a meta-analysis. *Am J Health Promot* 2014;28(4):259-67.
80. Atlantis E, Barnes EH, Singh MA. Efficacy of exercise for treating overweight in children and adolescents: a systematic review. *Int J Obes* 2006;30(7):1027-40.
81. Harris KC, Kuramoto LK, Schulzer M, et al. Effect of school-based physical activity interventions on body mass index in children: a meta-analysis. *Can Med Assoc J* 2009;180(7):719-26.

82. McGovern L, Johnson JN, Paulo R, et al. Treatment of pediatric obesity: a systematic review and meta-analysis of randomized trials. *J Clin Endocrinol Metab* 2008;93(12):4600-05.

83. Cesa CC, Sbruzzi G, Ribeiro RA, et al. Physical activity and cardiovascular risk factors in children: meta-analysis of randomized clinical trials. *Prev Med* 2014;69(12):54-62.

84. Guerra PH, Nobre MRC, da Silveira JAC, et al. The effect of school-based physical activity interventions on body mass index: a meta-analysis of randomized trials. *Clinics* 2013;68(9):1263-73.

85. Kelley GA, Kelley KS. Exercise and BMI z-score in overweight and obese children and adolescents: protocol for a systematic review and network meta-analysis of randomised trials. *BMJ Open* 2016;6(e011258):1-7.

86. Kelley GA, Kelley KS, Pate R. Exercise and BMI z-score in overweight and obese children and adolescents: A systematic review and network meta-analysis of randomized trials. *J Evid Based Med* 2016;10(2).

87. Hutton B, Salanti G, Caldwell DM, et al. The PRISMA Extension Statement for Reporting of Systematic Reviews Incorporating Network Meta-analyses of Health Care Interventions: Checklist and Explanations. *Ann Intern Med* 2015;162(11):777-84.

88. Sacks HS, Chalmers TC, Smith H. Randomized versus historical controls for clinical trials. *Am J Med* 1982;72:233-40.

89. Schulz KF, Chalmers I, Hayes R, et al. Empirical evidence of bias: Dimensions of methodological quality associated with estimates of treatment effects in controlled trials. *J Am Med Assoc* 1995;273:4:108-128.
90. Physical activity guidelines advisory report. Washington, DC.: Department of Health and Human Services, 2008.
91. Caspersen CJ, Powell KE, Christenson GM. Physical activity, exercise, and physical fitness: definitions and distinctions for health-related research. *Public Health Rep* 1985;100(2):126-31.
92. Dietz WH. Health consequences of obesity in youth: childhood predictors of adult disease. *Pediatrics* 1998;101(3 Pt 2):518-25.
93. van Driel ML, A. D, J. D, et al. Searching for unpublished trials in Cochrane reviews may not be worth the effort. *J Clin Epidemiol* 2009;62(8):838-44.
94. Egger E, Zellwegerzahner T, Schneider M, et al. Language bias in randomised controlled trials published in English and German. *Lancet* 1997;350:326-29.
95. Moher D, Pham B, Klassen TP, et al. What contributions do languages other than English make on the results of meta-analyses? *J Clin Epidemiol* 2000;53:964-72.
96. Harris MB, Hallbauer ES. Self-directed weight control through dieting and exercise. *Behav Res Therapy* 1973;11:523-29.
97. Berlin JA. Does blinding of readers affect the results of meta-analyses? *Lancet* 1997;350:185-86.
98. Cohen J. Weighted kappa: nominal scale agreement with provision for scaled disagreement or partial credit. *Psychol Bull* 1968;70:213-20.

99. Lee E, Dobbins M, DeCorby K, et al. An optimal search filter for retrieving systematic reviews and meta-analyses. *BMC Med Res Methodol* 2012;12:51.

100. Higgins JP, Altman DG, Gotzsche PC, et al. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. *Br Med J* 2011;343:d5928.

101. Emerson JD, Burdick E, Hoaglin DC, et al. An empirical study of the possible relation of treatment differences to quality scores in controlled randomized clinical trials. *Control Clin Trials* 1990;11:339-52.

102. Juni P, Witschi A, Bloch R, et al. The hazards of scoring the quality of clinical trials for meta-analysis. *J Am Med Assoc* 1999;282:1054-60.

103. Ahn S, Becker BJ. Incorporating quality scores in meta-analysis. *J Educ Behav Stat* 2011;36(5):555-85.

104. Follmann D, Elliot P, Suh I, et al. Variance imputation for overviews of clinical trials with continuous response. *J Clin Epidemiol* 1992;45:769-73.

105. Higgins JPT, Green S, editors. *Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 [updated March 2011]*: The Cochrane Collaboration, 2011.

106. Hedges LV, Olkin I. Statistical methods for meta-analysis. San Diego, CA: Academic Press 1985.

107. Chaimani A, Higgins JPT, Mavridis D, et al. Graphical tools for network meta-analysis in STATA. *PLoS One* 2013;8(10):e76654

108. Catala-Lopez F, Tobias A, Cameron C, et al. Network meta-analysis for comparing treatment effects of multiple interventions: an introduction. *Rheumatol Int* 2014;34(11):1489-96.

109. White IR, Barrett JK, Jackson D, et al. Consistency and inconsistency in network meta-analysis: model estimation using multivariate meta-regression. *Res Synth Methods* 2012;3(2):111-25.
110. White IR. Multivariate random-effects meta-regression: Updates to mvmeta. *Stata J* 2011;11(2):255-70.
111. Jansen JP, Naci H. Is network meta-analysis as valid as standard pairwise meta-analysis? It all depends on the distribution of effect modifiers. *BMC Med* 2013;11
112. Donegan S, Williamson P, D'Alessandro U, et al. Assessing key assumptions of network meta-analysis: a review of methods. *Res Synth Methods* 2013;4(4):291-323.
113. Lu GB, Ades AE. Assessing evidence inconsistency in mixed treatment comparisons. *J Am Stat Assoc* 2006;101(474):447-59.
114. Veroniki AA, Vasiliadis HS, Higgins JP, et al. Evaluation of inconsistency in networks of interventions. *Int J Epidemiol* 2013;42(1):332-45.
115. Song F, Xiong T, Parekh-Bhurke S, et al. Inconsistency between direct and indirect comparisons of competing interventions: meta-epidemiological study. *Br Med J* 2011;343:d4909.
116. Higgins JP, Thompson SG, Spiegelhalter DJ. A re-evaluation of random-effects meta-analysis. *J R Stat Soc Series A* 2009;172(1):137-59.
117. Kelley GA, Kelley KS. Impact of progressive resistance training on lipids and lipoproteins in adults: Another look at a meta-analysis using prediction intervals. *Prev Med* 2009;49:473-75.

1
2
3 118. Cooper HC, Hedges LV, Valentine JF. The handbook of research synthesis. New
4
5 York, New York: Russell Sage 2009.
6
7
8 119. Salanti G, Del Giovane C, Chaimani A, et al. Evaluating the quality of evidence
9
10 from a network meta-analysis. *PLoS One* 2014;9(7): A324.
11
12 120. Salanti G, Ades AE, Ioannidis JP. Graphical methods and numerical summaries for
13
14 presenting results from multiple-treatment meta-analysis: an overview and tutorial.
15
16 *J Clin Epidemiol* 2011;64(2):163-71.
17
18
19
20
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Table 1. Covariates to examine using simple meta-regression.

Characteristics	Variable
Study	Publication year, impact factor of journal, country study conducted, type of control group, bias (sequence generation, allocation concealment, blinding of participants & personnel, blinding of outcome assessors, incomplete outcome data, selective outcome reporting), type of analysis
Participant	Age, gender, race/ethnicity, maturational stage
Exercise	Type (aerobic, strength, both), length, frequency, intensity, duration, total minutes, total minutes (adjusted for compliance), mode, compliance, exercise supervision, setting, number of sets, number of repetitions, rest between sets, number of exercises, type of resistance, equipment used, fidelity (design, training, delivery, receipt, enactment)
Outcome	Baseline values for primary outcomes (BMI in kg m^2 , fat mass, percent fat), method used to assess adiposity, i.e., instrumentation, body weight, lean body mass, waist circumference, waist-to-hip ratio, diet, energy intake, energy expenditure, physical activity level, non-exercise activity, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin

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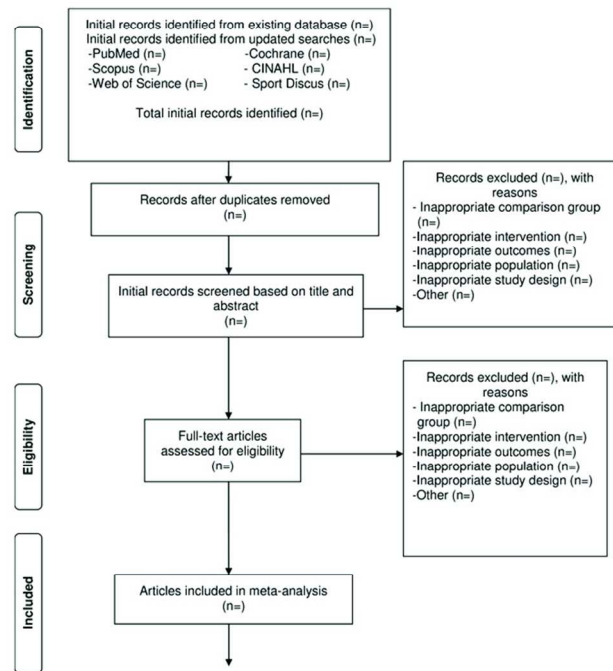
FIGURE LEGEND

Figure 1. Proposed flow diagram to depict the search process.

SUPPLEMENTARY FILE

Supplementary File 1. Preliminary search results in PubMed.

For peer review only



Flow diagram for network meta-analysis.

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PRISMA-P (Preferred Reporting Items for Systematic review and Meta-Analysis Protocols) 2015 checklist: recommended items to address in a systematic review protocol*

Section and topic	Item No	Checklist item	Line #
ADMINISTRATIVE INFORMATION			
Title:			
Identification	1a	Identify the report as a protocol of a systematic review	1-2
Update	1b	If the protocol is for an update of a previous systematic review, identify as such	NA
Registration	2	If registered, provide the name of the registry (such as PROSPERO) and registration number	43; 150-151;416-420
Authors:			
Contact	3a	Provide name, institutional affiliation, e-mail address of all protocol authors; provide physical mailing address of corresponding author	4-7; 9-12;14-17
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review	409-415
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments	421-423
Support:			
Sources	5a	Indicate sources of financial or other support for the review	426-430
Sponsor	5b	Provide name for the review funder and/or sponsor	
Role of sponsor or funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol	
INTRODUCTION			
Rationale	6	Describe the rationale for the review in the context of what is already known	58-138
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to participants, interventions, comparators, and outcomes (PICO)	139-145
METHODS			
Eligibility criteria	8	Specify the study characteristics (such as PICO, study design, setting, time frame) and report characteristics (such as years considered, language, publication status) to be used as criteria for eligibility for the review	152-210
Information sources	9	Describe all intended information sources (such as electronic databases, contact with study authors, trial registers or other grey literature sources) with planned dates of coverage	211-219
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	220-229; Supplementary file 1
Study records:			
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	232-234; 259-261;400-405

Selection process	11b	State the process that will be used for selecting studies (such as two independent reviewers) through each phase of the review (that is, screening, eligibility and inclusion in meta-analysis)	231-257; Figure 1
Data collection process	11c	Describe planned method of extracting data from reports (such as piloting forms, done independently, duplicate), any processes for obtaining and confirming data from investigators	258-274
Data items	12	List and define all variables for which data will be sought (such as PICO items, funding sources) any pre-planned data assumptions and simplifications	264-270;275-284;Table 1
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale	275-284
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis	285-299
Data synthesis	15a	Describe criteria under which study data will be quantitatively synthesised	300-372; 384-398; & Table 1 for 15a-d
	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data and methods of combining data from studies, including any planned exploration of consistency (such as I^2 , Kendall's τ)	
	15c	Describe any proposed additional analyses (such as sensitivity or subgroup analyses, meta-regression)	
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned	
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (such as publication bias across studies, selective reporting within studies)	373-376
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (such as GRADE)	377-383

*** It is strongly recommended that this checklist be read in conjunction with the PRISMA-P Explanation and Elaboration (cite when available) for important clarification on the items. Amendments to a review protocol should be tracked and dated. The copyright for PRISMA-P (including checklist) is held by the PRISMA-P Group and is distributed under a Creative Commons Attribution Licence 4.0.**

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BMJ Open

Exercise and Adiposity in Overweight and Obese Children and Adolescents: Protocol for a Systematic Review and Network Meta-Analysis of Randomised Trials

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ABSTRACT

Introduction: Overweight and obesity is a worldwide public health problem among children and adolescents. However, the magnitude of effect, as well as hierarchy of exercise interventions (aerobic, strength training, or both), on selected measures of adiposity is not well established despite numerous trials on this issue. The primary purposes of this study are to use the network meta-analytic approach to determine the effects and hierarchy of exercise interventions on selected measures of adiposity in overweight and obese children and adolescents. **Methods and analysis:** Randomised exercise intervention trials ≥ 4 weeks, available in any language up to August 31, 2017 and which include direct and/or indirect evidence, will be included. Studies will be located by searching seven electronic databases, cross-referencing and expert review. Dual selection and abstraction of data will occur. The primary outcomes will be changes in body mass index (BMI in kg/m^2), fat mass and percent body fat. Risk of bias will be assessed using the Cochrane Risk of Bias assessment instrument while confidence in the cumulative evidence will be assessed using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) instrument for network meta-analysis. Network meta-analysis will be performed using multivariate random-effects meta-regression models. The surface under the cumulative ranking curve (SUCRA) will be used to provide a hierarchy of exercise treatments (aerobic, strength, or both). **Dissemination:** The findings of this network meta-analysis will be presented at a professional conference and published in a peer-reviewed journal. **Trial registration number:** PROSPERO #CRD42017073103

STRENGTHS AND LIMITATIONS OF THIS STUDY

- To the best of the investigative team’s knowledge, this is the first systematic review to use the network meta-analytic approach to determine the effects as well as hierarchy of exercise interventions (aerobic, strength training, or both) on BMI in kg/m², fat mass and percent body fat in overweight and obese children and adolescents.
- The results of this systematic review with network meta-analysis should be useful to practitioners and policy-makers for making informed decisions about exercise in the treatment of overweight and obesity in children and adolescents.
- The results of this systematic review with network meta-analysis should be useful to researchers with respect to the conduct and reporting of future research on this topic.
- Common to most meta-analyses, the results may yield significant heterogeneity which cannot be explained.
- Like any aggregate data meta-analysis, the possibility of ecological fallacy exists, i.e., that group averages are not reflective of an individual’s values.

INTRODUCTION

Rationale

Overweight and obesity in children and adolescents is a major public health problem worldwide. Between 1980 and 2013, the worldwide prevalence of overweight and obesity in children and adolescents increased by 6.9%, from 16.9% to 23.8%, in boys and by 6.4%, from 16.2% to 22.6%, in girls from developed countries.¹ For developing countries, increases of 4.8%, from 8.1% to 12.9% for boys and 5%, from 8.4% to 13.4% in girls, were reported.¹

The negative outcomes associated with obesity in children and adolescents are both immediate and long-term.² For immediacy, a population-based study of children and adolescents 5 to 17 years of age found that approximately 70% of obese youth had a minimum of one cardiovascular disease risk factor (high cholesterol, high blood pressure, etc.).³ Obese children and adolescents are also more likely to be diagnosed with prediabetes,⁴ as well as being at an increased risk for bone and joint difficulties, sleep apnea, and social and psychological issues such as stigmatization, poor self-esteem, and poorer health-related quality-of-life.^{5 6} Long-term, childhood and adolescent overweight and obesity has been demonstrated to track into adulthood,⁷⁻¹¹ thus placing overweight and/or obese adults at a greater risk for cardiovascular disease, type 2 diabetes, stroke, several types of cancer, and osteoarthritis.²

One promising intervention in the treatment of overweight and obesity is exercise. However, previous randomised trials that were limited to or included overweight and obese children and adolescents have led to conflicting results,¹²⁻⁵⁸ with some reporting statistically significant reductions in adiposity (BMI) as a primary outcome^{12 13 16 17 22 27 28}

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2
3 31 41 51-56 59-63 and others reporting no change.^{14 15 18-21 23-26 29 30 32-40 42-50 57 58 62 64 65} When
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5 limited to overweight and obese male and female children and adolescents,^{12 14 17-20 22-26}
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7 28 31 33 36 38-41 45-57 50, 51, 52, 54, 55, 56, 57 only 18 (45.0%) have reported statistically significant
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9 reductions in BMI.^{12 17 22 28 31 41 51,52-56 58, 50, 52, 54, 56, 57} While this may lead one to the
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11 general conclusion that exercise does little to reduce BMI in overweight and obese
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13 children and adolescents, this would be shortsighted since it relies on the vote-counting
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15 approach,⁶⁶ an approach that has been shown to be less valid than the meta-analytic
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17 approach.^{66 67}

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21 Previous systematic reviews with meta-analyses that have focused on the effects of
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23 exercise as an independent intervention on BMI as a primary outcome in male and
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25 female children and adolescents have reported conflicting findings with five reporting a
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27 significant improvement in BMI⁶⁸⁻⁷² and five others reporting no statistically significant
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29 improvement.⁷³⁻⁷⁷ However, nine of the ten suffer from one or more of the following
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31 limitations: (1) inclusion of a small number of studies with exercise as the only
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33 intervention,^{71 73-75} (2) inclusion of non-randomised trials,^{68 74} (3) inclusion of children
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35 and adolescents who were not overweight or obese.^{70 72 74 76 77} Relevant to this
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37 application, all ten suffer from both reliance on pairwise versus network meta-analysis,
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39 the latter of which incorporates both direct and indirect evidence. In addition, there was
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41 an absence of an established hierarchy for determining which types of exercise
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43 (aerobic, strength training, or both) might be best for improving BMI based on both
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45 direct and indirect evidence.⁶⁸⁻⁷⁷ To partially address this issue as well as demonstrate
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47 feasibility, the investigative team has recently used the network meta-analytic approach
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49 to examine the effects of exercise (aerobic, strength training, or both) on BMI z-score in
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overweight and obese children and adolescents.^{78 79} Statistically significant reductions in BMI z-score were found for aerobic exercise and combined aerobic and strength exercise, but not strength training alone (mean, 95% CI: aerobic, -0.10, -0.15 to -0.05; aerobic and strength, -0.11, -0.19 to -0.03; strength, 0.04, -0.07 to 0.15).⁷⁹ Combined aerobic and strength training was ranked best, followed by aerobic exercise and then strength training.⁷⁹ Consistency in evidence and risk of bias did not differ between direct and indirect studies.⁷⁹ It was concluded that combined aerobic exercise and strength training as well as aerobic exercise alone are associated with reductions in BMI z-score.⁷⁹ The lack of effect on BMI z-score in the strength training studies may have been the result of increases in lean muscle mass. However, since BMI in kg/m² continues to be the most frequently assessed and reported measure of adiposity in both the clinical and public health setting, an examination of such using the network meta-analytic approach is needed. In addition, since all types of BMI measures as well as body weight do not capture changes in body composition (fat mass, percent body fat, etc.), the inclusion of such outcomes, as previously suggested,⁷⁹ is also necessary.

Objectives

The primary objectives of the current study are to conduct a systematic review with network meta-analysis of randomised trials to (1) determine the effects of exercise (aerobic, strength training, or both) on adiposity (BMI in kg/m², fat mass, percent body fat) in overweight and obese children and adolescents, and (2) establish a hierarchy of exercise interventions (aerobic, strength training, or both) for treating adiposity (BMI in kg/m², fat mass, percent body fat) in overweight and obese children and adolescents.

METHODS

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3 **Overview**

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5 This study will follow the guidelines from the Preferred Reporting Items for Systematic

6 Reviews and Meta-Analysis (PRISMA) extension statement for network meta-analyses

7 of health care interventions.⁸⁰ The protocol for this network meta-analysis is registered

8 in PROSPERO (trial registration number CRD42017073103).

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14 **Eligibility criteria**

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16 The inclusion criteria for this proposed network meta-analysis will be as follows: (1)

17 direct evidence from randomised trials that compare two or more exercise interventions

18 (aerobic, strength training, both) or indirect evidence from randomised controlled trials

19 that compare an exercise intervention group to a comparative control group (non-

20 intervention, attention control, usual care, wait-list control, placebo), (2) exercise-only

21 intervention (aerobic, strength training, or both), (3) studies lasting ≥ 4 weeks, (4) male

22 and/or female children and adolescents 2 to 18 years of age, (5) participants overweight

23 or obese, as defined by the authors, (6) studies published in any language up to August

24 31, 2017, (7) data available for BMI in kg/m², fat mass or percent body fat.

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26 Studies will be limited to randomised trials because it is the only way to control for

27 confounders that are not known or measured as well as the observation that

28 nonrandomised controlled trials tend to overestimate the effects of healthcare

29 interventions.^{81 82} Indirect evidence studies will be limited to randomised controlled trials

30 with at least one exercise arm that participates in either aerobic, strength training, or a

31 combination of aerobic and strength training exercise. Direct evidence studies will be

32 limited to randomised trials that include at least two of the following exercise arms: (1)

33 aerobic, (2) strength training, (3) aerobic and strength training exercise.

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For the purposes of this study, exercise, aerobic exercise and strength training will be defined according to the 2008 Physical Activity Guidelines for Americans,⁸³ defined as movement that is “planned, structured, and repetitive and purposive in the sense that the improvement or maintenance of one or more components of physical fitness is the objective,”^{83 84} aerobic exercise as “exercise that primarily uses the aerobic energy-producing systems, can improve the capacity and efficiency of these systems, and is effective for improving cardiorespiratory endurance,”⁸³ and strength training as “exercise training primarily designed to increase skeletal muscle strength, power, endurance, and mass”.⁸³ Four weeks was chosen as the lower cut point for intervention length based on previous research demonstrating improvements in adiposity over this period of time in 11-year olds.²¹

Participants will be limited to overweight and obese children and adolescents, as defined by the original study authors, because it has been shown that this population is at an increased risk for premature morbidity and mortality throughout their lifetime.⁸⁵

While some research has suggested that studies yielding statistically significant and positive results are more likely to be published in English-language versus non-English language journals,⁸⁶ other research has shown this to not be the case.⁸⁷ Given the former, studies from both English and non-English-language articles will be included with the latter translated into English by the second author using the freely available web-based Babelfish and Bing translators. For those studies that cannot be translated using Babelfish and/or Bing, professional translation services will be utilized.

Body mass index in $\text{kg}\cdot\text{m}^2$ was included as one of the three primary adiposity outcomes because it is the most commonly used and understood variable by

practitioners as well as others and can be easily measured from body weight and height. However, because BMI is an indirect measure of adiposity, fat mass and percent body fat will be included because they are more direct measures of adiposity. The inclusion of fat mass and percent body fat may be especially relevant for studies that include strength training given that decreases in adiposity as measured by BMI may be offset by increases in muscle mass, a secondary outcome that will be coded.

Information sources

The following seven electronic databases will be searched: (1) PubMed, (2) Web of Science, (3) Cochrane Central Register of Controlled Trials (CENTRAL), (4) Cumulative Index to Nursing and Allied Health Literature (CINAHL), (5) Sport Discus, (6) Translating Research into Practice (TRIP) and (7) ProQuest Dissertations and Theses. In addition to electronic database searches, cross-referencing will be conducted by examining the reference lists of previous review articles as well as each included study for potential articles that meet the inclusion criteria. Upon completion of initial searches, the third author will examine the reference list for thoroughness and completeness. Suggested studies will then be retrieved to see if they meet all inclusion criteria.

Search strategy

Search strategies specific to each database will be developed by the investigative team. Major keywords, or forms of keywords to include will be “random”, “children”, “adolescents”, “overweight”, “obese”, “exercise,” “physical fitness”, “body composition”, “fat mass”, “body fat”, “body composition”, “body mass index”, “adiposity”. A copy of a preliminary search strategy using PubMed, including limits, can be found in Supplementary file 1. This search strategy will be adapted for other database searches.

All database searches and article retrieval will be conducted by the second author with oversight from the first author.

Study records

Study selection

All studies to be screened will be imported into EndNote (version X8; New York, NY: Thomson-Reuters; 2016) and duplicates removed electronically and then manually by the second author. A copy of the database will then be provided to the first author for duplicate screening. To minimize selection bias, the first and second authors will select all studies, independent of each other. They will then review their selections for accuracy and consistency. The full report for each article will be retrieved for all titles and abstracts that appear to meet the inclusion criteria as well as those where uncertainty exists. Multiple reports for the same study will be addressed by including the most recently published article and drawing from prior reports, assuming the same methods and sample sizes are reported. Based on previous research suggesting neither a clinically nor statistically significant effect on results, blinding to journal titles, study authors, or institutions of the authors will not be employed during the screening and data abstraction processes.⁸⁸ Reasons for excluded studies will be recorded using the following categories: (1) inappropriate population, (2) inappropriate intervention, (3) inappropriate comparison(s), (4) inappropriate outcome(s), (5) inappropriate study design, (6) other. Upon the conclusion of screening, the first and second authors will meet and review their selections. Cohen's kappa statistic (κ) will be used to measure inter-selection agreement.⁸⁹ Any discrepancies will be resolved by consensus. If consensus cannot be reached, the third author will serve as an arbitrator. Upon selecting

the final number of studies to include, the overall precision of the searches will be computed by dividing the number of included studies by the total number of studies screened after removing duplicates.⁹⁰ The number needed-to-read (NNR) will then be calculated as the reciprocal of the precision.⁹⁰ A flow diagram that describes the search procedure will be included as well as a supplementary file that includes a reference list of all excluded studies, including the reason(s) for exclusion. Figure 1 illustrates the proposed structure for the flow diagram.

Data abstraction

For this project, Microsoft Excel (version 2016; Redmond, WA: Microsoft Corporation; 2016) will be used to develop comprehensive electronic codebooks that will define the coding process for each of the variables coded. The codebook will be created by the first two authors with feedback from the third author. Consequently, the abstraction of data from the studies in this proposed project should require little subjective judgment on the part of the coder. The major groups of variables to code will include (1) study characteristics (author, journal, year of publication, etc.), (2) participant characteristics (age, gender, height, body weight, etc.), and (3) data for primary and secondary outcomes (sample sizes, baseline and post-exercise means and standard deviations, etc.). Table 1 contains a preliminary list of variables that will be coded. Based on previous research by the investigative team,⁷⁹ a codebook capable of including at least 242 items from each study is expected. To avoid data abstraction bias, the first two authors will independently code (dual-coding) all studies to ensure accuracy and consistency. Inter-rater agreement will be assessed using Cohen’s kappa statistic (κ).⁸⁹ Any disagreement in the items coded will be discussed until mutual agreement is

reached. If agreement cannot be reached, the third author will serve as an arbitrator.

Outcomes and prioritization

The primary outcomes in this study will be changes BMI in kg/m², fat mass, and percent body fat in overweight and obese children and adolescents. Secondary outcomes will include body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, physical activity level, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin.

Risk of bias assessment in individual studies

Risk of bias for included studies will be assessed using the Cochrane Risk of Bias Instrument.⁹¹ Assessment is based on judgments of low, high or unclear risk of bias across six defined domains: (1) sequence generation, (2) allocation sequence concealment, (3) blinding of participants and personnel, (4) blinding of outcome assessors, (5) incomplete outcome data, and (6) selective outcome reporting. A seventh domain, whether participants were exercising regularly, as defined by the original study authors prior to taking part in the study, will also be assessed using the same approach as for the other six domains. As previously recommended, study-level results will reported for each domain according to risk of bias (low, high, or unclear) while the percentage of low, high, or unclear results across each domain will also be reported.⁹¹ This risk of bias approach has been recommended over the use of study

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quality rating scales given the lack of empirical evidence to support the latter.^{82 92 93}

Assessment for risk of bias will be limited to the primary outcomes of interest, i.e., changes in BMI in kg/m², fat mass, and percent body fat. All studies will be classified as high risk of bias with respect to the category “blinding of participants and personnel” given that it’s virtually impossible to blind participants to group assignment in exercise intervention protocols. Based on previous research, no study will be excluded based on risk of bias results.⁹⁴

Data Synthesis

Calculation of effect sizes

The primary outcomes for this study will be changes in BMI in kg/m², fat mass (kg), and percent body fat using the original metric. Changes for indirect comparisons will be calculated by subtracting the change outcome difference in the exercise group minus the change outcome difference in the control group. Variances will be computed using the pooled standard deviations of change scores in the exercise and control groups. If change score standard deviations are not available, they will be calculated from 95% confidence intervals (CI) for either change outcome or treatment effect differences as well as pre and post standard deviation values, the latter according to procedures developed by Follmann et al.⁹⁵ For direct comparisons, i.e., randomised trials with no control group, the same general procedures will be followed except that the control group data will be replaced with one of the exercise interventions as follows: (1) aerobic minus strength training, (2) aerobic and strength training combined minus aerobic training, (3) aerobic and strength training combined minus strength training. Ninety-five percent CI and z-alpha values will be calculated for each outcome from each study. For

those studies that include both direct and indirect comparisons, only direct comparison data will be included since a primary purpose of the current meta-analysis is determining which exercise interventions(s) might work best for improving adiposity in children and adolescents. For studies in which adiposity outcomes are assessed at multiple intervention time points, for example, 0, 8, and 16 weeks, only data from the initial and last assessment will be used. If follow-up data are available, results from such will also be analyzed separately to determine the sustainability of changes in adiposity. If any crossover trials are included, treatment effects will be calculated by using all assessments from the intervention and control periods and analyzing them similar to a parallel group trial.⁹⁶ While the possibility of a unit-of-analysis error exists as well as studies being under versus over-weighted, this method is believed to be better than alternative approaches, for example, limiting data from the first assessment point or trying to impute standard deviations, especially given the primary and secondary outcomes included and expected distribution of findings.⁹⁶

Secondary outcomes (body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, maximum oxygen consumption (relative and absolute), resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin) will be handled using the same approach as for primary outcomes. However, given the different metrics expected and the inability to convert between them, changes in

physical activity levels and muscular strength will be calculated using the standardized effect size, adjusted for small sample sizes.⁹⁷

Pooled estimates for changes in outcomes

Network (geometry) plots for each outcome will be used to provide a visual representation of the evidence base with nodes (circles) weighted by the number of participants randomised to each treatment and edges (lines) weighted by the number of studies evaluating each pair of treatments.^{98 99} *Contribution plots* for each outcome will be used to determine the most dominant comparisons for each network estimate as well as for the entire network.⁹⁸ The weights applied will be a function of the variance of the direct treatment effect and the network structure, the result being a percent contribution of each direct comparison to each network estimate.⁹⁸

Network meta-analysis will be performed using *multivariate random-effects meta-regression models* that can be performed within a frequentist setting, allows for the inclusion of potential covariates, and correctly accounts for the correlations from multi-arm trials.^{100 101} A two-tailed alpha value ≤ 0.05 and non-overlapping 95% CI will be considered to represent statistically significant changes. Separate network meta-analysis models will be used to examine for changes in each primary and secondary outcome. Potential *covariates* will be examined by (1) conducting simple meta-regression for statistically significant associations between covariates and changes in primary outcomes (BMI in kg/m², fat mass, percent fat), (2) examining for multicollinearity between covariates ($r > 0.80$), and (3) building a multiple meta-regression model. A list of potential covariates to examine using simple meta-regression is shown in Table 1. While we will include all methods used to assess

adiposity, we will also conduct sensitivity analyses to see if results differ according to method of assessment, for example, fat mass assessed using whole body magnetic resonance imaging versus bioelectrical impedance. Secondary outcomes (energy intake and expenditure, physical activity level, muscular strength) will be handled using the same approach. *Transitivity*, i.e., similarity in the distribution of potential effect modifiers across the different pairwise comparisons for each outcome¹⁰² will include those listed in Table 1. *Inconsistency*, i.e., differences in effect estimates between direct and indirect results for the same comparison,¹⁰³ will be checked by assessing differences in treatment effects between direct and indirect effect estimates as well as differences between trials with different designs, for example, two-arm versus multi-arm trials.^{101 103}

¹⁰⁴ However, the probability of inconsistency is considered small given recent research demonstrating that inconsistency was detected in only 2% to 14% of tested loops, depending on the effect measure and heterogeneity estimation method.^{105 106} Finally, *prediction intervals* will be used to enhance interpretation of results with respect to the magnitude of heterogeneity as well as provide an estimate of expected results in a future study.¹⁰⁷⁻¹⁰⁹ For network meta-analysis, degrees of freedom (*df*) will be set to the number of studies – the number of comparisons – 1.¹⁰⁹

Meta-biases

Small-study-effects (publication bias, etc.) will be assessed using comparison adjusted funnel plots.⁹⁸ In the absence of small-study effects, the comparison adjusted funnel plot should be symmetric around the zero line.

Confidence in cumulative evidence

Quality analysis of specific pairwise effect estimates in the network meta-analysis will be evaluated using a recently developed modification of the Grading of Recommendations Assessment, Development and Evaluation (GRADE) for network meta-analysis across five domains: (1) study limitations, (2) indirectness, (3) inconsistency, (4) imprecision, and (5) small-study effects.¹¹⁰ Assessment will be conducted using the same procedures as for study selection and data abstraction.

To establish a hierarchy of exercise interventions for selected outcomes in the current meta-analysis, *ranking analysis*, i.e., the ability to rank all interventions for a single outcome under study, for example changes in BMI in kg/m², will be used based on probabilities. However, because the ranking of treatments based exclusively on the probability of each treatment being the best should be avoided given that it does not account for the uncertainty in the relative treatment effects and the possibility for assigning higher ranks for treatments in which little evidence is available, separate *rankograms and cumulative ranking probability plots* will be used to present ranking probabilities along with their uncertainty for changes in primary and secondary outcomes.^{98 111} The surface under the cumulative ranking curve (SUCRA), a transformation of the mean rank, will be used to establish a hierarchy of exercise interventions (aerobic, strength, both) while accounting for the location and variance of all treatment effects.^{98 111} Larger SUCRA values indicate better ranks for the treatment.^{98 111} Interpretation of all rankings will be approached from the perspective of absolute and relative treatment effects.⁹⁹

Software used for statistical analysis

All data will be analysed using Stata (V.14.1; Stata/SE for Windows, version 14.0. College Station, TX: Stata Corporation LP; 2015), Microsoft Excel (version 2016; Redmond, WA: Microsoft Corporation; 2016), and two add-ins for Excel, SSC-Stat (V.2.18; SSC-Stat, version 3.0. University of Reading, United Kingdom: Statistical Services Center; 2007), and EZ-Analyze (V.3.0; EZ Analyze, version 3.0. TA Poynton; 2007).

DISSEMINATION

The results of this study will be presented at a professional conference and published in a peer-reviewed journal.

CONTRIBUTORS

GAK is the guarantor. GAK, KSK and RRP drafted the manuscript. All authors contributed to (1) the development of the data sources to search for relevant literature, including search strategy, (2) selection criteria, (3) data extraction criteria and (4) risk of bias assessment strategy. GAK provided statistical expertise while RRP provided content expertise on exercise and adiposity in overweight and obese children and adolescents. All authors read, provided feedback, and approved the final manuscript.

REGISTRATION

In accordance with the Primary Reporting Items for Systematics Reviews and Meta-Analyses Protocols (PRISMA-P) statement, this systematic review with network meta-analysis was registered with the International Prospective Register of Systematic Reviews (PROSPERO) on August 23, 2017 (#CRD42017073103).

AMENDMENTS TO PROTOCOL

REFERENCES

1. Ng M, Fleming T, Robinson M, *et al.* Global, regional, and national prevalence of overweight and obesity in children and adults during 1980-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2014;384:766-81.
2. CDC. Childhood obesity facts Atlanta, Georgia: U.S. Department of Health & Human Services, Centers for Disease Control and Prevention; 2015 [Available from: <http://www.cdc.gov/healthyschools/obesity/facts.htm> accessed 12/3/2015.
3. Freedman DS, Mei Z, Srinivasan SR, *et al.* Cardiovascular risk factors and excess adiposity among overweight children and adolescents: the Bogalusa Heart Study. *J Pediatr* 2007;150(1):12-17.
4. Li C, Ford ES, Zhao G, *et al.* Prevalence of pre-diabetes and its association with clustering of cardiometabolic risk factors and hyperinsulinemia among U.S. adolescents: National Health and Nutrition Examination Survey 2005-2006. *Diabetes Care* 2009;32(2):342-47.
5. Daniels SR, Arnett DK, Eckel RH, *et al.* Overweight in children and adolescents: pathophysiology, consequences, prevention, and treatment. *Circulation* 2005;111(15):1999-2012.
6. Dietz WH. Overweight in childhood and adolescence. *N Engl J Med* 2004;350(9):855-57.
7. Singh AS, Mulder C, Twisk JW, *et al.* Tracking of childhood overweight into adulthood: a systematic review of the literature. *Obes Rev* 2008;9(5):474-88.
8. Guo SS, Chumlea WC. Tracking of body mass index in children in relation to overweight in adulthood. *Am J Clin Nutr* 1999;70(1):145S-48S.

9. Freedman DS, Khan LK, Serdula MK, *et al.* The Relation of Childhood BMI to Adult Adiposity: The Bogalusa Heart Study. *Pediatrics* 2005;115(1):22-27.

10. Freedman DS, Wang J, Thornton JC, *et al.* Classification of body fatness by body mass index-for-age categories among children. *Arch Pediatr Adolesc Med* 2009;163(9):805-11.

11. Freedman DS, Khan LK, Dietz WH, *et al.* Relationship of childhood obesity to coronary heart disease risk factors in adulthood: the Bogalusa Heart Study. *Pediatrics* 2001;108:712-18.

12. Alves JG, Gale CR, Souza E, *et al.* [Effect of physical exercise on bodyweight in overweight children: a randomized controlled trial in a Brazilian slum]. *Cad Saude Publica* 2008;24(Suppl 2):S353-S59.

13. Benson AC, Torode ME, Fiatarone Singh MA. The effect of high-intensity progressive resistance training on adiposity in children: a randomized controlled trial. *Int J Obes (Lond)* 2008;32(6):1016-27.

14. Daley AJ, Copeland RJ, Wright NP, *et al.* Exercise therapy as a treatment for psychopathologic conditions in obese and morbidly obese adolescents: a randomized, controlled trial. *Pediatrics* 2006;118(5):2126-34.

15. Donnelly JE, Greene JL, Gibson CA, *et al.* Physical Activity Across the Curriculum (PAAC): a randomized controlled trial to promote physical activity and diminish overweight and obesity in elementary school children. *Prev Med* 2009;49(4):336-41.

16. Duncan MJ, Al-Nakeeb Y, Nevill AM. Effects of a 6-week circuit training intervention on body esteem and body mass index in British primary school children. *Body Image* 2009;6(3):216-20.

17. Farpour-Lambert NJ, Aggoun Y, Marchand LM, *et al.* Physical activity reduces systemic blood pressure and improves early markers of atherosclerosis in pre-pubertal obese children. *J Am Coll Cardiol* 2009;54(25):2396-406.
18. Gutin B, Owens S, Slavens G, *et al.* Effect of physical training on heart-period variability in obese children. *J Pediatr* 1997;130(6):938-43.
19. Gutin B, Owens S, Okuyama T, *et al.* Effect of physical training and its cessation on percent fat and bone density of children with obesity. *Obes Res* 1999;7(2):208-14.
20. Hagstromer M, Elmberg K, Marild S, *et al.* Participation in organized weekly physical exercise in obese adolescents reduced daily physical activity. *Acta Paediatr* 2009;98(2):352-54.
21. Jago R, Jonker ML, Missaghian M, *et al.* Effect of 4 weeks of Pilates on the body composition of young girls. *Prev Med* 2006;42(3):177-80.
22. Karacabey K. The effect of exercise on leptin, insulin, cortisol and lipid profiles in obese children. *J Int Med Res* 2009;37(5):1472-78.
23. Kaufman C, Kelly AS, Kaiser DR, *et al.* Aerobic-exercise training improves ventilatory efficiency in overweight children. *Pediatr Exerc Sci* 2007;19(1):82-92.
24. Kelly AS, Wetzsteon RJ, Kaiser DR, *et al.* Inflammation, insulin, and endothelial function in overweight children and adolescents: The role of exercise. *J Pediatr* 2004;145(6):731-36.
25. Kim ES, Im JA, Kim KC, *et al.* Improved insulin sensitivity and adiponectin level after exercise training in obese Korean youth. *Obesity (Silver Spring)* 2007;15(12):3023-30.

26. Kim HJ, Lee S, Kim TW, *et al.* Effects of exercise-induced weight loss on acylated and unacylated ghrelin in overweight children. *Clin Endocrinol (Oxf)* 2008;68(3):416-22.

27. Kriemler S, Zahner L, Schindler C, *et al.* Effect of school based physical activity programme (KISS) on fitness and adiposity in primary schoolchildren: cluster randomised controlled trial. *Br Med J* 2010;340:c785. doi: 10.1136/bmj.c785.c785.

28. Li YP, Hu XQ, Schouten EG, *et al.* Report on childhood obesity in China (8): effects and sustainability of physical activity intervention on body composition of Chinese youth. *Biomed Environ Sci* 2010;23(3):180-87.

29. Macias-Cervantes MH, Malacara JM, Garay-Sevilla ME, *et al.* Effect of recreational physical activity on insulin levels in Mexican/Hispanic children. *Eur J Pediatr* 2009;168(10):1195-202.

30. Martinez-Vizcaino V, Salcedo AF, Franquelo GR, *et al.* Assessment of an after-school physical activity program to prevent obesity among 9- to 10-year-old children: a cluster randomized trial. *Int J Obes (Lond)* 2008;32(1):12-22.

31. Meyer AA, Kundt G, Lenschow U, *et al.* Improvement of early vascular changes and cardiovascular risk factors in obese children after a six-month exercise program. *J Am Coll Cardiol* 2006;48(9):1865-70.

32. Mo-suwan L, Pongprapai S, Junjana C, *et al.* Effects of a controlled trial of a school-based exercise program on the obesity indexes of preschool children. *Am J Clin Nutr* 1998;68(5):1006-11.

33. Murphy EC, Carson L, Neal W, *et al.* Effects of an exercise intervention using Dance Revolution on endothelial function and other risk factors in overweight children. *Int J Pediatr Obes* 2009;4(4):205-14.
34. Neumark-Sztainer D, Story M, Hannan PJ, *et al.* New Moves: a school-based obesity prevention program for adolescent girls. *Prev Med* 2003;37(1):41-51.
35. Pate RR, Ward DS, Saunders RP, *et al.* Promotion of physical activity among high-school girls: a randomized controlled trial. *Am J Public Health* 2005;95(9):1582-87.
36. Petty KH, Davis CL, Tkacz J, *et al.* Exercise effects on depressive symptoms and self-worth in overweight children: a randomized controlled trial. *J Pediatr Psychol* 2009;34(9):929-39.
37. Reilly JJ, Kelly L, Montgomery C, *et al.* Physical activity to prevent obesity in young children: cluster randomised controlled trial. *Br Med J* 2006;333(7577):1041.
38. Rooney BL, Gritt LR, Havens SJ, *et al.* Growing healthy families: family use of pedometers to increase physical activity and slow the rate of obesity. *Wis Med J* 2005;104(5):54-60.
39. Shaibi GQ, Cruz ML, Ball GD, *et al.* Effects of resistance training on insulin sensitivity in overweight Latino adolescent males. *Med Sci Sports Exerc* 2006;38(7):1208-15.
40. Simon C, Schweitzer B, Oujaa M, *et al.* Successful overweight prevention in adolescents by increasing physical activity: a 4-year randomized controlled intervention. *Int J Obes (Lond)* 2008;32(10):1489-98.
41. Tan S, Yang C, Wang J. Physical training of 9- to 10-year-old children with obesity to lactate threshold intensity. *Pediatr Exerc Sci* 2010;22(3):477-85.

42. Velez A, Golem DL, Arent SM. The impact of a 12-week resistance training program on strength, body composition, and self-concept of Hispanic adolescents. *J Strength Cond Res* 2010;24(4):1065-73.

43. Walther C, Gaede L, Adams V, *et al.* Effect of increased exercise in school children on physical fitness and endothelial progenitor cells: a prospective randomized trial. *Circulation* 2009;120(22):2251-59.

44. Warren JM, Henry CJ, Lightowler HJ, *et al.* Evaluation of a pilot school programme aimed at the prevention of obesity in children. *Health Promot Int* 2003;18(4):287-96.

45. Watts K, Beye P, Siafarikas A, *et al.* Exercise training normalizes vascular dysfunction and improves central adiposity in obese adolescents. *J Am Coll Cardiol* 2004;43(10):1823-27.

46. Watts K, Beye P, Siafarikas A, *et al.* Effects of exercise training on vascular function in obese children. *J Pediatr* 2004;144(5):620-25.

47. Weintraub DL, Tirumalai EC, Haydel KF, *et al.* Team sports for overweight children: the Stanford Sports to Prevent Obesity Randomized Trial (SPORT). *Arch Pediatr Adolesc Med* 2008;162(3):232-37.

48. Sigal RJ, Alberga AS, Goldfield GS, *et al.* Effects of aerobic training, resistance training, or both on percentage body fat and cardiometabolic risk markers in obese adolescents: the healthy eating aerobic and resistance training in youth randomized clinical trial. *JAMA Pediatr* 2014;168(11):1006-14.

49. Meucci M, Cook C, Curry CD, *et al.* Effects of supervised exercise program on metabolic function in overweight adolescents. *World J Pediatr* 2013;9(4):307-11.

50. Lee S, Deldin AR, White D, *et al.* Aerobic exercise but not resistance exercise reduces intrahepatic lipid content and visceral fat and improves insulin sensitivity in obese adolescent girls: a randomized controlled trial. *Am J Physiol Endocrinol Metab* 2013;305(10):E1222-E29.
51. Dennis BA, Ergul A, Gower BA, *et al.* Oxidative stress and cardiovascular risk in overweight children in an exercise intervention program. *Childhood Obesity* 2013;9(1):15-21.
52. Maddison R, Foley L, Ni MC, *et al.* Effects of active video games on body composition: a randomized controlled trial. *Am J Clin Nutr* 2011;94(1):156-63.
53. Wong PC, Chia MY, Tsou IY, *et al.* Effects of a 12-week exercise training programme on aerobic fitness, body composition, blood lipids and C-reactive protein in adolescents with obesity. *Ann Acad Med Singapore* 2008;37(4):286-93.
54. Saygin O, Ozturk MA. The effect of twelve week aerobic exercise programme on health related physical fitness components and blood lipids in obese girls. *Afr J Pharm Pharmacol* 2011;5(12):1441-45.
55. Sun MX, Huang XQ, Yan Y, *et al.* One-hour after-school exercise ameliorates central adiposity and lipids in overweight Chinese adolescents: a randomized controlled trial. *Chin Med J (Engl)* 2011;124(3):323-29.
56. Elloumi M, Makni E, Ounis OB, *et al.* Six-minute walking test and the assessment of cardiorespiratory responses during weight-loss programmes in obese children. *Physiother Res Int* 2011;16(1):32-42.
57. Lau PWC, Wong dP, Ngo JK, *et al.* Effects of high-intensity intermittent running exercise in overweight children. *Eur J Sport Sci* 2014:1-9.

58. Alberga AS, Farnesi BC, Lafleche A, *et al.* The effects of resistance exercise training on body composition and strength in obese prepubertal children. *Phys Sportsmed* 2013;41(3):103-09.

59. Ackel-D'Elia C, Carnier J, Bueno CR, Jr., *et al.* Effects of different physical exercises on leptin concentration in obese adolescents. *Int J Sports Med* 2014;35(2):164-71.

60. Fazelifar S, Ebrahim K, Sarkisian V. Effect of concurrent training and detraining on anti-inflammatory biomarker and physical fitness levels in obese children. *Rev Bras Med Esporte* 2013;19(5):349-54.

61. Ghorbanian B, Ravassi A, Kordi MR, *et al.* The effects of rope training on lymphocyte ABCA1 expression, plasma ApoA-I and HDL-c in boy adolescents. *Int J Endocrinol Metab Disord* 2013;11(2):76-81.

62. Lee S, Bacha F, Hannon T, *et al.* Effects of aerobic versus resistance exercise without caloric restriction on abdominal fat, intrahepatic lipid, and insulin sensitivity in obese adolescent boys: a randomized, controlled trial. *Diabetes* 2012;61(11):2787-95.

63. Song JK, Stebbins CL, Kim TK, *et al.* Effects of 12 weeks of aerobic exercise on body composition and vascular compliance in obese boys. *J Sports Med Phys Fitness* 2012;52(5):522-29.

64. Cheng HL, Peng P, Zhu R, *et al.* Effects of eight weeks exercise prescription intervention on aerobic capacity, body composition, blood lipid and C-reactive protein in obese adolescents. [Chinese]. *J Jilin Univ Med* 2012;38:745-49.

65. Kelly LA, Loza A, Lin X, *et al*. The effect of a home-based strength training program on type 2 diabetes risk in obese Latino boys. *J Pediatr Endocrinol Metab* 2015;28(3-4):315-22.
66. Hedges LV, Olkin I. Vote-counting methods in research synthesis. *Psychol Bull* 1980;88:359-69.
67. James A, Soler A, Weatherall R. Cognitive behavioural therapy for anxiety disorders in children and adolescents. *Cochrane Database Syst Rev* 2005(4):CD004690.
68. Schranz N, Tomkinson G, Olds T. What is the effect of resistance training on the strength, body composition and psychosocial status of overweight and obese children and adolescents? A systematic review and meta-analysis. *Sports Med* 2013;43(9):893-907.
69. Kelley GA, Kelley KS, Pate RR. Exercise and BMI in overweight and obese children and adolescents: A systematic review with trial sequential meta-analysis. *Biomed Res Int* 2015;2015(Article ID 704539):1-17.
70. Mei H, Xiong Y, Xie S, *et al*. The impact of long-term school-based physical activity interventions on body mass index of primary school children - a meta-analysis of randomized controlled trials. *BMC Public Health* 2016;16(1):205.
71. Garcia-Hermoso A, Sanchez-Lopez M, Martinez-Vizcaino V. Effects of aerobic plus resistance exercise on body composition related variables in pediatric obesity: A systematic review and meta-analysis of randomized controlled trials. *Pediatr Exerc Sci* 2015;27(4):431-40.

72. Dellert JC, Johnson P. Interventions with children and parents to improve physical activity and body mass index: a meta-analysis. *Am J Health Promot* 2014;28(4):259-67.

73. Atlantis E, Barnes EH, Singh MA. Efficacy of exercise for treating overweight in children and adolescents: a systematic review. *Int J Obes* 2006;30(7):1027-40.

74. Harris KC, Kuramoto LK, Schulzer M, *et al*. Effect of school-based physical activity interventions on body mass index in children: a meta-analysis. *Can Med Assoc J* 2009;180(7):719-26.

75. McGovern L, Johnson JN, Paulo R, *et al*. Treatment of Pediatric Obesity: A Systematic Review and Meta-Analysis of Randomized Trials. *J Clin Endocrinol Metab* 2008;93(12):4600-05.

76. Cesa CC, Sbruzzi G, Ribeiro RA, *et al*. Physical activity and cardiovascular risk factors in children: meta-analysis of randomized clinical trials. *Prev Med* 2014;69(12):54-62.

77. Guerra PH, Nobre MRC, da Silveira JAC, *et al*. The effect of school-based physical activity interventions on body mass index: a meta-analysis of randomized trials. *Clinics* 2013;68(9):1263-73.

78. Kelley GA, Kelley KS. Exercise and BMI z-score in overweight and obese children and adolescents: protocol for a systematic review and network meta-analysis of randomised trials. *BMJ Open* 2016;6(e011258):1-7.

79. Kelley GA, Kelley KS, Pate R. Exercise and BMI z-score in overweight and obese children and adolescents: A systematic review and network meta-analysis of randomized trials. *J Evid Based Med* 2017;10(2):108-28.

80. Hutton B, Salanti G, Caldwell DM, *et al.* The PRISMA Extension Statement for Reporting of Systematic Reviews Incorporating Network Meta-analyses of Health Care Interventions: Checklist and Explanations. *Ann Intern Med* 2015;162(11):777-84.
81. Sacks HS, Chalmers TC, Smith H. Randomized versus historical controls for clinical trials. *Am J Med* 1982;72:233-40.
82. Schulz KF, Chalmers I, Hayes R, *et al.* Empirical evidence of bias: Dimensions of methodological quality associated with estimates of treatment effects in controlled trials. *J Am Med Assoc* 1995;273:408-12.
83. Physical Activity Guidelines Advisory Committee. *Physical Activity Guidelines Advisory Report*. Washington, DC: U.S. Department of Health and Human Services, 2008
84. Caspersen CJ, Powell KE, Christenson GM. Physical activity, exercise, and physical fitness: definitions and distinctions for health-related research. *Public Health Rep* 1985;100(2):126-31.
85. Dietz WH. Health consequences of obesity in youth: childhood predictors of adult disease. *Pediatrics* 1998;101(3 Pt 2):518-25.
86. Egger E, Zellwegerzahner T, Schneider M, *et al.* Language bias in randomised controlled trials published in English and German. *Lancet* 1997;350:326-29.
87. Moher D, Pham B, Klassen TP, *et al.* What contributions do languages other than English make on the results of meta-analyses? *J Clin Epidemiol* 2000;53:964-72.
88. Berlin JA. Does blinding of readers affect the results of meta-analyses? *Lancet* 1997;350:185-86.

89. Cohen J. Weighted kappa: nominal scale agreement with provision for scaled disagreement or partial credit. *Psychol Bull* 1968;70:213-20.

90. Lee E, Dobbins M, DeCorby K, *et al*. An optimal search filter for retrieving systematic reviews and meta-analyses. *BMC Med Res Methodol* 2012;12:51.

91. Higgins JP, Altman DG, Gotzsche PC, *et al*. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. *Br Med J* 2011;343:d5928.

92. Emerson JD, Burdick E, Hoaglin DC, *et al*. An empirical study of the possible relation of treatment differences to quality scores in controlled randomized clinical trials. *Control Clin Trials* 1990;11:339-52.

93. Juni P, Witschi A, Bloch R, *et al*. The hazards of scoring the quality of clinical trials for meta-analysis. *J Am Med Assoc* 1999;282:1054-60.

94. Ahn S, Becker BJ. Incorporating quality scores in meta-analysis. *J Educ Behav Stat* 2011;36(5):555-85.

95. Follmann D, Elliot P, Suh I, *et al*. Variance imputation for overviews of clinical trials with continuous response. *J Clin Epidemiol* 1992;45:769-73.

96. Higgins JPT, Green S, editors. *Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 [updated March 2011]*: The Cochrane Collaboration, 2011.

97. Hedges LV, Olkin I. *Statistical methods for meta-analysis*. San Diego, CA: Academic Press, 1985.

98. Chaimani A, Higgins JPT, Mavridis D, *et al*. Graphical tools for network meta-analysis in STATA. *PLoS One* 2013;8(10)

99. Catala-Lopez F, Tobias A, Cameron C, *et al*. Network meta-analysis for comparing treatment effects of multiple interventions: an introduction. *Rheumatol Int* 2014;34(11):1489-96.
100. White IR, Barrett JK, Jackson D, *et al*. Consistency and inconsistency in network meta-analysis: model estimation using multivariate meta-regression. *Res Synth Methods* 2012;3(2):111-25.
101. White IR. Multivariate random-effects meta-regression: Updates to mvmeta. *Stata J* 2011;11(2):255-70.
102. Jansen JP, Naci H. Is network meta-analysis as valid as standard pairwise meta-analysis? It all depends on the distribution of effect modifiers. *BMC Med* 2013;11
103. Donegan S, Williamson P, D'Alessandro U, *et al*. Assessing key assumptions of network meta-analysis: a review of methods. *Res Synth Methods* 2013;4(4):291-323.
104. Lu GB, Ades AE. Assessing evidence inconsistency in mixed treatment comparisons. *J Am Stat Assoc* 2006;101(474):447-59.
105. Veroniki AA, Vasiliadis HS, Higgins JP, *et al*. Evaluation of inconsistency in networks of interventions. *Int J Epidemiol* 2013;42(1):332-45.
106. Song F, Xiong T, Parekh-Bhurke S, *et al*. Inconsistency between direct and indirect comparisons of competing interventions: meta-epidemiological study. *Br Med J* 2011;343:d4909.
107. Higgins JP, Thompson SG, Spiegelhalter DJ. A re-evaluation of random-effects meta-analysis. *J R Stat Soc Series A* 2009;172(1):137-59.

Table 1. Covariates to examine using simple meta-regression.

Characteristics	Variable
Study	Publication year, impact factor of journal, country study conducted, type of control group, bias (sequence generation, allocation concealment, blinding of participants & personnel, blinding of outcome assessors, incomplete outcome data, selective outcome reporting), type of analysis
Participant	Age, gender, race/ethnicity, maturational stage
Exercise	Type (aerobic, strength, both), length, frequency, intensity, duration, total minutes, total minutes (adjusted for compliance), mode, compliance, exercise supervision, setting, number of sets, number of repetitions, rest between sets, number of exercises, type of resistance, equipment used, fidelity (design, training, delivery, receipt, enactment)
Outcome	Baseline values for primary outcomes (BMI in kg/m ² , fat mass, percent fat), method used to assess adiposity, i.e., instrumentation, body weight, lean body mass, waist circumference, waist-to-hip ratio, diet, energy intake, energy expenditure, physical activity level, non-exercise activity, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin

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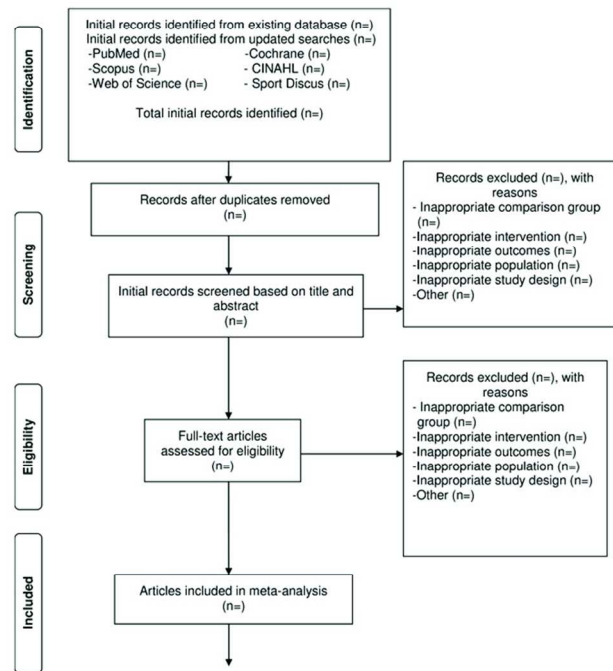
FIGURE LEGEND

Figure 1. Proposed flow diagram to depict the search process.

SUPPLEMENTARY FILE

Supplementary File 1. Preliminary search results in PubMed.

For peer review only



Flow diagram for network meta-analysis.

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PRISMA-P (Preferred Reporting Items for Systematic review and Meta-Analysis Protocols) 2015 checklist: recommended items to address in a systematic review protocol*

Section and topic	Item No	Checklist item	Line #
ADMINISTRATIVE INFORMATION			
Title:			
Identification	1a	Identify the report as a protocol of a systematic review	1-2
Update	1b	If the protocol is for an update of a previous systematic review, identify as such	NA
Registration	2	If registered, provide the name of the registry (such as PROSPERO) and registration number	43; 131-132;394-398
Authors:			
Contact	3a	Provide name, institutional affiliation, e-mail address of all protocol authors; provide physical mailing address of corresponding author	4-7; 9-12;14-17
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review	387-393
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments	399-401
Support:			
Sources	5a	Indicate sources of financial or other support for the review	404-408
Sponsor	5b	Provide name for the review funder and/or sponsor	
Role of sponsor or funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol	
INTRODUCTION			
Rationale	6	Describe the rationale for the review in the context of what is already known	60-119
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to participants, interventions, comparators, and outcomes (PICO)	120-126
METHODS			
Eligibility criteria	8	Specify the study characteristics (such as PICO, study design, setting, time frame) and report characteristics (such as years considered, language, publication status) to be used as criteria for eligibility for the review	133-179
Information sources	9	Describe all intended information sources (such as electronic databases, contact with study authors, trial registers or other grey literature sources) with planned dates of coverage	180-189
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	190-198; Supplementary file 1
Study records:			
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	201-202; 228-230;378-383

Selection process	11b	State the process that will be used for selecting studies (such as two independent reviewers) through each phase of the review (that is, screening, eligibility and inclusion in meta-analysis)	201-226; Figure 1
Data collection process	11c	Describe planned method of extracting data from reports (such as piloting forms, done independently, duplicate), any processes for obtaining and confirming data from investigators	227-243
Data items	12	List and define all variables for which data will be sought (such as PICO items, funding sources) any pre-planned data assumptions and simplifications	233-239;244-253;Table 1
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale	244-253
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis	254-272
Data synthesis	15a	Describe criteria under which study data will be quantitatively synthesised	274-350; 362-376; & Table 1 for 15a-d
	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data and methods of combining data from studies, including any planned exploration of consistency (such as I^2 , Kendall's τ)	
	15c	Describe any proposed additional analyses (such as sensitivity or subgroup analyses, meta-regression)	
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned	
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (such as publication bias across studies, selective reporting within studies)	351-354
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (such as GRADE)	355-361

From: Shamseer L, Moher D, Clarke M, Ghersi D, Liberati A, Petticrew M, Shekelle P, Stewart L, PRISMA-P Group. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015: elaboration and explanation. *BMJ*. 2015 Jan 2;349(jan02 1):g7647.

BMJ Open

Exercise and Adiposity in Overweight and Obese Children and Adolescents: Protocol for a Systematic Review and Network Meta-Analysis of Randomised Trials

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Secondary Subject Heading:	Sports and exercise medicine, Public health, Paediatrics, Epidemiology
Keywords:	exercise, overweight, obesity, children, adolescents, network meta-analysis

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ABSTRACT

Introduction: Overweight and obesity is a worldwide public health problem among children and adolescents. However, the magnitude of effect, as well as hierarchy of exercise interventions (aerobic, strength training, or both), on selected measures of adiposity is not well established despite numerous trials on this issue. The primary purposes of this study are to use the network meta-analytic approach to determine the effects and hierarchy of exercise interventions on selected measures of adiposity in overweight and obese children and adolescents. **Methods and analysis:** Randomised exercise intervention trials ≥ 4 weeks, available in any language up to August 31, 2017 and which include direct and/or indirect evidence, will be included. Studies will be located by searching seven electronic databases, cross-referencing and expert review. Dual selection and abstraction of data will occur. The primary outcomes will be changes in body mass index (BMI in kg/m^2), fat mass and percent body fat. Risk of bias will be assessed using the Cochrane Risk of Bias assessment instrument while confidence in the cumulative evidence will be assessed using the Grading of Recommendations Assessment, Development and Evaluation (GRADE) instrument for network meta-analysis. Network meta-analysis will be performed using multivariate random-effects meta-regression models. The surface under the cumulative ranking curve (SUCRA) will be used to provide a hierarchy of exercise treatments (aerobic, strength, or both). Ethics and **Dissemination:** This study does not require ethics approval. Findings will be presented at a professional conference and published in a peer-reviewed journal. **Trial registration number:** PROSPERO #CRD42017073103

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STRENGTHS AND LIMITATIONS OF THIS STUDY

- To the best of the investigative team’s knowledge, this is the first systematic review to use the network meta-analytic approach to determine the effects as well as hierarchy of exercise interventions (aerobic, strength training, or both) on BMI in kg/m², fat mass and percent body fat in overweight and obese children and adolescents.
- The results of this systematic review with network meta-analysis should be useful to practitioners and policy-makers for making informed decisions about exercise in the treatment of overweight and obesity in children and adolescents.
- The results of this systematic review with network meta-analysis should be useful to researchers with respect to the conduct and reporting of future research on this topic.
- Common to most meta-analyses, the results may yield significant heterogeneity which cannot be explained.
- Like any aggregate data meta-analysis, the possibility of ecological fallacy exists, i.e., that group averages are not reflective of an individual’s values.

INTRODUCTION

Rationale

Overweight and obesity in children and adolescents is a major public health problem worldwide. Between 1980 and 2013, the worldwide prevalence of overweight and obesity in children and adolescents increased by 6.9%, from 16.9% to 23.8%, in boys and by 6.4%, from 16.2% to 22.6%, in girls from developed countries.¹ For developing countries, increases of 4.8%, from 8.1% to 12.9% for boys and 5%, from 8.4% to 13.4% in girls, were reported.¹

The negative outcomes associated with obesity in children and adolescents are both immediate and long-term.² For immediacy, a population-based study of children and adolescents 5 to 17 years of age found that approximately 70% of obese youth had a minimum of one cardiovascular disease risk factor (high cholesterol, high blood pressure, etc.).³ Obese children and adolescents are also more likely to be diagnosed with prediabetes,⁴ as well as being at an increased risk for bone and joint difficulties, sleep apnea, and social and psychological issues such as stigmatization, poor self-esteem, and poorer health-related quality-of-life.^{5 6} Long-term, childhood and adolescent overweight and obesity has been demonstrated to track into adulthood,⁷⁻¹¹ thus placing overweight and/or obese adults at a greater risk for cardiovascular disease, type 2 diabetes, stroke, several types of cancer, and osteoarthritis.²

One promising intervention in the treatment of overweight and obesity is exercise. However, previous randomised trials that were limited to or included overweight and obese children and adolescents have led to conflicting results,¹²⁻⁵⁸ with some reporting statistically significant reductions in adiposity (BMI) as a primary outcome^{12 13 16 17 22 27 28}

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82 31 41 51-56 59-63 and others reporting no change.^{14 15 18-21 23-26 29 30 32-40 42-50 57 58 62 64 65} When
83 limited to overweight and obese male and female children and adolescents,^{12 14 17-20 22-26}
84 28 31 33 36 38-41 45-57 50, 51, 52, 54, 55, 56, 57 only 18 (45.0%) have reported statistically significant
85 reductions in BMI.^{12 17 22 28 31 41 51,52-56 58, 50, 52, 54, 56, 57} While this may lead one to the
86 general conclusion that exercise does little to reduce BMI in overweight and obese
87 children and adolescents, this would be shortsighted since it relies on the vote-counting
88 approach,⁶⁶ an approach that has been shown to be less valid than the meta-analytic
89 approach.^{66 67}
90 Previous systematic reviews with meta-analyses that have focused on the effects of
91 exercise as an independent intervention on BMI as a primary outcome in male and
92 female children and adolescents have reported conflicting findings with five reporting a
93 significant improvement in BMI⁶⁸⁻⁷² and five others reporting no statistically significant
94 improvement.⁷³⁻⁷⁷ However, nine of the ten suffer from one or more of the following
95 limitations: (1) inclusion of a small number of studies with exercise as the only
96 intervention,^{71 73-75} (2) inclusion of non-randomised trials,^{68 74} (3) inclusion of children
97 and adolescents who were not overweight or obese.^{70 72 74 76 77} Relevant to this
98 application, all ten suffer from both reliance on pairwise versus network meta-analysis,
99 the latter of which incorporates both direct and indirect evidence. In addition, there was
100 an absence of an established hierarchy for determining which types of exercise
101 (aerobic, strength training, or both) might be best for improving BMI based on both
102 direct and indirect evidence.⁶⁸⁻⁷⁷ To partially address this issue as well as demonstrate
103 feasibility, the investigative team has recently used the network meta-analytic approach
104 to examine the effects of exercise (aerobic, strength training, or both) on BMI z-score in

overweight and obese children and adolescents.^{78 79} Statistically significant reductions in BMI z-score were found for aerobic exercise and combined aerobic and strength exercise, but not strength training alone (mean, 95% CI: aerobic, -0.10, -0.15 to -0.05; aerobic and strength, -0.11, -0.19 to -0.03; strength, 0.04, -0.07 to 0.15).⁷⁹ Combined aerobic and strength training was ranked best, followed by aerobic exercise and then strength training.⁷⁹ Consistency in evidence and risk of bias did not differ between direct and indirect studies.⁷⁹ It was concluded that combined aerobic exercise and strength training as well as aerobic exercise alone are associated with reductions in BMI z-score.⁷⁹ The lack of effect on BMI z-score in the strength training studies may have been the result of increases in lean muscle mass. However, since BMI in kg/m² continues to be the most frequently assessed and reported measure of adiposity in both the clinical and public health setting, an examination of such using the network meta-analytic approach is needed. In addition, since all types of BMI measures as well as body weight do not capture changes in body composition (fat mass, percent body fat, etc.), the inclusion of such outcomes, as previously suggested,⁷⁹ is also necessary.

Objectives

The primary objectives of the current study are to conduct a systematic review with network meta-analysis of randomised trials to (1) determine the effects of exercise (aerobic, strength training, or both) on adiposity (BMI in kg/m², fat mass, percent body fat) in overweight and obese children and adolescents, and (2) establish a hierarchy of exercise interventions (aerobic, strength training, or both) for treating adiposity (BMI in kg/m², fat mass, percent body fat) in overweight and obese children and adolescents.

METHODS

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3 151 For the purposes of this study, exercise, aerobic exercise and strength training will be
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5 152 defined according to the 2008 Physical Activity Guidelines for Americans,⁸³ defined as
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7 153 movement that is “planned, structured, and repetitive and purposive in the sense that
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9 154 the improvement or maintenance of one or more components of physical fitness is the
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11 155 objective,”^{83 84} aerobic exercise as “exercise that primarily uses the aerobic energy-
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13 156 producing systems, can improve the capacity and efficiency of these systems, and is
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15 157 effective for improving cardiorespiratory endurance,”⁸³ and strength training as “exercise
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17 158 training primarily designed to increase skeletal muscle strength, power, endurance, and
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19 159 mass”.⁸³ Four weeks was chosen as the lower cut point for intervention length based
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21 160 on previous research demonstrating improvements in adiposity over this period of time
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23 161 in 11-year olds.²¹

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25 162 Participants will be limited to overweight and obese children and adolescents, as
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27 163 defined by the original study authors, because it has been shown that this population is
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29 164 at an increased risk for premature morbidity and mortality throughout their lifetime.⁸⁵

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31 165 While some research has suggested that studies yielding statistically significant and
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33 166 positive results are more likely to be published in English-language versus non-English
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35 167 language journals,⁸⁶ other research has shown this to not be the case.⁸⁷ Given the
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37 168 former, studies from both English and non-English-language articles will be included
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39 169 with the latter translated into English by the second author using the freely available
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41 170 web-based Babelfish and Bing translators. For those studies that cannot be translated
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43 171 using Babelfish and/or Bing, professional translation services will be utilized.

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45 172 Body mass index in kg m^2 was included as one of the three primary adiposity
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47 173 outcomes because it is the most commonly used and understood variable by
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197 All database searches and article retrieval will be conducted by the second author with
198 oversight from the first author.

199 **Study records**

200 **Study selection**

201 All studies to be screened will be imported into EndNote (version X8; New York, NY:
202 Thomson-Reuters; 2016) and duplicates removed electronically and then manually by
203 the second author. A copy of the database will then be provided to the first author for
204 duplicate screening. To minimize selection bias, the first and second authors will select
205 all studies, independent of each other. They will then review their selections for accuracy
206 and consistency. The full report for each article will be retrieved for all titles and
207 abstracts that appear to meet the inclusion criteria as well as those where uncertainty
208 exists. Multiple reports for the same study will be addressed by including the most
209 recently published article and drawing from prior reports, assuming the same methods
210 and sample sizes are reported. Based on previous research suggesting neither a
211 clinically nor statistically significant effect on results, blinding to journal titles, study
212 authors, or institutions of the authors will not be employed during the screening and
213 data abstraction processes.⁸⁸ Reasons for excluded studies will be recorded using the
214 following categories: (1) inappropriate population, (2) inappropriate intervention, (3)
215 inappropriate comparison(s), (4) inappropriate outcome(s), (5) inappropriate study
216 design, (6) other. Upon the conclusion of screening, the first and second authors will
217 meet and review their selections. Cohen's kappa statistic (κ) will be used to measure
218 inter-selection agreement.⁸⁹ Any discrepancies will be resolved by consensus. If
219 consensus cannot be reached, the third author will serve as an arbitrator. Upon selecting

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3 220 the final number of studies to include, the overall precision of the searches will be
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5 221 computed by dividing the number of included studies by the total number of studies
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7 222 screened after removing duplicates.⁹⁰ The number needed-to-read (NNR) will then be
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10 223 calculated as the reciprocal of the precision.⁹⁰ A flow diagram that describes the search
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12 224 procedure will be included as well as a supplementary file that includes a reference list
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14 225 of all excluded studies, including the reason(s) for exclusion. Figure 1 illustrates the
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16 226 proposed structure for the flow diagram.

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19 227 Data abstraction

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21 228 For this project, Microsoft Excel (version 2016; Redmond, WA: Microsoft Corporation;
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23 229 2016) will be used to develop comprehensive electronic codebooks that will define the
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25 230 coding process for each of the variables coded. The codebook will be created by the
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27 231 first two authors with feedback from the third author. Consequently, the abstraction of
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29 232 data from the studies in this proposed project should require little subjective judgment
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31 233 on the part of the coder. The major groups of variables to code will include (1) study
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33 234 characteristics (author, journal, year of publication, etc.), (2) participant characteristics
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35 235 (age, gender, height, body weight, etc.), and (3) data for primary and secondary
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37 236 outcomes (sample sizes, baseline and post-exercise means and standard deviations,
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39 237 etc.). Table 1 contains a preliminary list of variables that will be coded. Based on
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41 238 previous research by the investigative team,⁷⁹ a codebook capable of including at least
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43 239 242 items from each study is expected. To avoid data abstraction bias, the first two
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45 240 authors will independently code (dual-coding) all studies to ensure accuracy and
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47 241 consistency. Inter-rater agreement will be assessed using Cohen's kappa statistic
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49 242 (κ).⁸⁹ Any disagreement in the items coded will be discussed until mutual agreement is
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reached. If agreement cannot be reached, the third author will serve as an arbitrator.

Outcomes and prioritization

The primary outcomes in this study will be changes BMI in kg/m², fat mass, and percent body fat in overweight and obese children and adolescents. Secondary outcomes will include body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, physical activity level, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin.

Risk of bias assessment in individual studies

Risk of bias for included studies will be assessed using the Cochrane Risk of Bias Instrument.⁹¹ Assessment is based on judgments of low, high or unclear risk of bias across six defined domains: (1) sequence generation, (2) allocation sequence concealment, (3) blinding of participants and personnel, (4) blinding of outcome assessors, (5) incomplete outcome data, and (6) selective outcome reporting. A seventh domain, whether participants were exercising regularly, as defined by the original study authors prior to taking part in the study, will also be assessed using the same approach as for the other six domains. As previously recommended, study-level results will reported for each domain according to risk of bias (low, high, or unclear) while the percentage of low, high, or unclear results across each domain will also be reported.⁹¹ This risk of bias approach has been recommended over the use of study

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quality rating scales given the lack of empirical evidence to support the latter.^{82 92 93}

Assessment for risk of bias will be limited to the primary outcomes of interest, i.e., changes in BMI in kg/m², fat mass, and percent body fat. All studies will be classified as high risk of bias with respect to the category “blinding of participants and personnel” given that it’s virtually impossible to blind participants to group assignment in exercise intervention protocols. Based on previous research, no study will be excluded based on risk of bias results.⁹⁴

Data Synthesis

Calculation of effect sizes

The primary outcomes for this study will be changes in BMI in kg/m², fat mass (kg), and percent body fat using the original metric. Changes for indirect comparisons will be calculated by subtracting the change outcome difference in the exercise group minus the change outcome difference in the control group. Variances will be computed using the pooled standard deviations of change scores in the exercise and control groups. If change score standard deviations are not available, they will be calculated from 95% confidence intervals (CI) for either change outcome or treatment effect differences as well as pre and post standard deviation values, the latter according to procedures developed by Follmann et al.⁹⁵ For direct comparisons, i.e., randomised trials with no control group, the same general procedures will be followed except that the control group data will be replaced with one of the exercise interventions as follows: (1) aerobic minus strength training, (2) aerobic and strength training combined minus aerobic training, (3) aerobic and strength training combined minus strength training. Ninety-five percent CI and z-alpha values will be calculated for each outcome from each study. For

those studies that include both direct and indirect comparisons, only direct comparison data will be included since a primary purpose of the current meta-analysis is determining which exercise interventions(s) might work best for improving adiposity in children and adolescents. For studies in which adiposity outcomes are assessed at multiple intervention time points, for example, 0, 8, and 16 weeks, only data from the initial and last assessment will be used. If follow-up data are available, results from such will also be analyzed separately to determine the sustainability of changes in adiposity. If any crossover trials are included, treatment effects will be calculated by using all assessments from the intervention and control periods and analyzing them similar to a parallel group trial.⁹⁶ While the possibility of a unit-of-analysis error exists as well as studies being under versus over-weighted, this method is believed to be better than alternative approaches, for example, limiting data from the first assessment point or trying to impute standard deviations, especially given the primary and secondary outcomes included and expected distribution of findings.⁹⁶

Secondary outcomes (body weight, lean body mass, waist circumference, waist-to-hip ratio, energy intake, energy expenditure, maximum oxygen consumption (relative and absolute), resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin) will be handled using the same approach as for primary outcomes. However, given the different metrics expected and the inability to convert between them, changes in

adiposity, we will also conduct sensitivity analyses to see if results differ according to method of assessment, for example, fat mass assessed using whole body magnetic resonance imaging versus bioelectrical impedance. Secondary outcomes (energy intake and expenditure, physical activity level, muscular strength) will be handled using the same approach. *Transitivity*, i.e., similarity in the distribution of potential effect modifiers across the different pairwise comparisons for each outcome¹⁰² will include those listed in Table 1. *Inconsistency*, i.e., differences in effect estimates between direct and indirect results for the same comparison,¹⁰³ will be checked by assessing differences in treatment effects between direct and indirect effect estimates as well as differences between trials with different designs, for example, two-arm versus multi-arm trials.^{101 103}¹⁰⁴ However, the probability of inconsistency is considered small given recent research demonstrating that inconsistency was detected in only 2% to 14% of tested loops, depending on the effect measure and heterogeneity estimation method.^{105 106} Finally, *prediction intervals* will be used to enhance interpretation of results with respect to the magnitude of heterogeneity as well as provide an estimate of expected results in a future study.¹⁰⁷⁻¹⁰⁹ For network meta-analysis, degrees of freedom (*df*) will be set to the number of studies – the number of comparisons – 1.¹⁰⁹

Meta-biases

Small-study-effects (publication bias, etc.) will be assessed using comparison adjusted funnel plots.⁹⁸ In the absence of small-study effects, the comparison adjusted funnel plot should be symmetric around the zero line.

Confidence in cumulative evidence

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Quality analysis of specific pairwise effect estimates in the network meta-analysis will be evaluated using a recently developed modification of the Grading of Recommendations Assessment, Development and Evaluation (GRADE) for network meta-analysis across five domains: (1) study limitations, (2) indirectness, (3) inconsistency, (4) imprecision, and (5) small-study effects.¹¹⁰ Assessment will be conducted using the same procedures as for study selection and data abstraction.

To establish a hierarchy of exercise interventions for selected outcomes in the current meta-analysis, *ranking analysis*, i.e., the ability to rank all interventions for a single outcome under study, for example changes in BMI in kg/m², will be used based on probabilities. However, because the ranking of treatments based exclusively on the probability of each treatment being the best should be avoided given that it does not account for the uncertainty in the relative treatment effects and the possibility for assigning higher ranks for treatments in which little evidence is available, separate *rankograms and cumulative ranking probability plots* will be used to present ranking probabilities along with their uncertainty for changes in primary and secondary outcomes.^{98 111} The surface under the cumulative ranking curve (SUCRA), a transformation of the mean rank, will be used to establish a hierarchy of exercise interventions (aerobic, strength, both) while accounting for the location and variance of all treatment effects.^{98 111} Larger SUCRA values indicate better ranks for the treatment.^{98 111} Interpretation of all rankings will be approached from the perspective of absolute and relative treatment effects.⁹⁹

Software used for statistical analysis

378 All data will be analysed using Stata (V.14.1; Stata/SE for Windows, version 14.0.
379 College Station, TX: Stata Corporation LP; 2015), Microsoft Excel (version 2016;
380 Redmond, WA: Microsoft Corporation; 2016), and two add-ins for Excel, SSC-Stat
381 (V.2.18; SSC-Stat, version 3.0. University of Reading, United Kingdom: Statistical
382 Services Center; 2007), and EZ-Analyze (V.3.0; EZ Analyze, version 3.0. TA Poynton;
383 2007).

384 **ETHICS AND DISSEMINATION**

385 This study does not require ethics approval. Findings will be presented at a professional
386 conference and published in a peer-reviewed journal.

387 **CONTRIBUTORS**

388 GAK is the guarantor. GAK, KSK and RRP drafted the manuscript. All authors
389 contributed to (1) the development of the data sources to search for relevant literature,
390 including search strategy, (2) selection criteria, (3) data extraction criteria and (4) risk of
391 bias assessment strategy. GAK provided statistical expertise while RRP provided
392 content expertise on exercise and adiposity in overweight and obese children and
393 adolescents. All authors read, provided feedback, and approved the final manuscript.

394 **REGISTRATION**

395 In accordance with the Primary Reporting Items for Systematics Reviews and Meta-
396 Analyses Protocols (PRISMA-P) statement, this systematic review with network meta-
397 analysis was registered with the International Prospective Register of Systematic
398 Reviews (PROSPERO) on August 23, 2017 (#CRD42017073103).

399 **AMENDMENTS TO PROTOCOL**

REFERENCES

1. Ng M, Fleming T, Robinson M, *et al*. Global, regional, and national prevalence of overweight and obesity in children and adults during 1980-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2014;384:766-81.
2. CDC. Childhood obesity facts Atlanta, Georgia: U.S. Department of Health & Human Services, Centers for Disease Control and Prevention; 2015 [Available from: <http://www.cdc.gov/healthyschools/obesity/facts.htm> accessed 12/3/2015.
3. Freedman DS, Mei Z, Srinivasan SR, *et al*. Cardiovascular risk factors and excess adiposity among overweight children and adolescents: the Bogalusa Heart Study. *J Pediatr* 2007;150(1):12-17.
4. Li C, Ford ES, Zhao G, *et al*. Prevalence of pre-diabetes and its association with clustering of cardiometabolic risk factors and hyperinsulinemia among U.S. adolescents: National Health and Nutrition Examination Survey 2005-2006. *Diabetes Care* 2009;32(2):342-47.
5. Daniels SR, Arnett DK, Eckel RH, *et al*. Overweight in children and adolescents: pathophysiology, consequences, prevention, and treatment. *Circulation* 2005;111(15):1999-2012.
6. Dietz WH. Overweight in childhood and adolescence. *N Engl J Med* 2004;350(9):855-57.
7. Singh AS, Mulder C, Twisk JW, *et al*. Tracking of childhood overweight into adulthood: a systematic review of the literature. *Obes Rev* 2008;9(5):474-88.
8. Guo SS, Chumlea WC. Tracking of body mass index in children in relation to overweight in adulthood. *Am J Clin Nutr* 1999;70(1):145S-48S.

9. Freedman DS, Khan LK, Serdula MK, *et al.* The Relation of Childhood BMI to Adult Adiposity: The Bogalusa Heart Study. *Pediatrics* 2005;115(1):22-27.

10. Freedman DS, Wang J, Thornton JC, *et al.* Classification of body fatness by body mass index-for-age categories among children. *Arch Pediatr Adolesc Med* 2009;163(9):805-11.

11. Freedman DS, Khan LK, Dietz WH, *et al.* Relationship of childhood obesity to coronary heart disease risk factors in adulthood: the Bogalusa Heart Study. *Pediatrics* 2001;108:712-18.

12. Alves JG, Gale CR, Souza E, *et al.* [Effect of physical exercise on bodyweight in overweight children: a randomized controlled trial in a Brazilian slum]. *Cad Saude Publica* 2008;24(Suppl 2):S353-S59.

13. Benson AC, Torode ME, Fiatarone Singh MA. The effect of high-intensity progressive resistance training on adiposity in children: a randomized controlled trial. *Int J Obes (Lond)* 2008;32(6):1016-27.

14. Daley AJ, Copeland RJ, Wright NP, *et al.* Exercise therapy as a treatment for psychopathologic conditions in obese and morbidly obese adolescents: a randomized, controlled trial. *Pediatrics* 2006;118(5):2126-34.

15. Donnelly JE, Greene JL, Gibson CA, *et al.* Physical Activity Across the Curriculum (PAAC): a randomized controlled trial to promote physical activity and diminish overweight and obesity in elementary school children. *Prev Med* 2009;49(4):336-41.

16. Duncan MJ, Al-Nakeeb Y, Nevill AM. Effects of a 6-week circuit training intervention on body esteem and body mass index in British primary school children. *Body Image* 2009;6(3):216-20.

17. Farpour-Lambert NJ, Aggoun Y, Marchand LM, *et al*. Physical activity reduces systemic blood pressure and improves early markers of atherosclerosis in pre-pubertal obese children. *J Am Coll Cardiol* 2009;54(25):2396-406.
18. Gutin B, Owens S, Slavens G, *et al*. Effect of physical training on heart-period variability in obese children. *J Pediatr* 1997;130(6):938-43.
19. Gutin B, Owens S, Okuyama T, *et al*. Effect of physical training and its cessation on percent fat and bone density of children with obesity. *Obes Res* 1999;7(2):208-14.
20. Hagstromer M, Elmberg K, Marild S, *et al*. Participation in organized weekly physical exercise in obese adolescents reduced daily physical activity. *Acta Paediatr* 2009;98(2):352-54.
21. Jago R, Jonker ML, Missaghian M, *et al*. Effect of 4 weeks of Pilates on the body composition of young girls. *Prev Med* 2006;42(3):177-80.
22. Karacabey K. The effect of exercise on leptin, insulin, cortisol and lipid profiles in obese children. *J Int Med Res* 2009;37(5):1472-78.
23. Kaufman C, Kelly AS, Kaiser DR, *et al*. Aerobic-exercise training improves ventilatory efficiency in overweight children. *Pediatr Exerc Sci* 2007;19(1):82-92.
24. Kelly AS, Wetzsteon RJ, Kaiser DR, *et al*. Inflammation, insulin, and endothelial function in overweight children and adolescents: The role of exercise. *J Pediatr* 2004;145(6):731-36.
25. Kim ES, Im JA, Kim KC, *et al*. Improved insulin sensitivity and adiponectin level after exercise training in obese Korean youth. *Obesity (Silver Spring)* 2007;15(12):3023-30.

33. Murphy EC, Carson L, Neal W, *et al.* Effects of an exercise intervention using Dance Revolution on endothelial function and other risk factors in overweight children. *Int J Pediatr Obes* 2009;4(4):205-14.
34. Neumark-Sztainer D, Story M, Hannan PJ, *et al.* New Moves: a school-based obesity prevention program for adolescent girls. *Prev Med* 2003;37(1):41-51.
35. Pate RR, Ward DS, Saunders RP, *et al.* Promotion of physical activity among high-school girls: a randomized controlled trial. *Am J Public Health* 2005;95(9):1582-87.
36. Petty KH, Davis CL, Tkacz J, *et al.* Exercise effects on depressive symptoms and self-worth in overweight children: a randomized controlled trial. *J Pediatr Psychol* 2009;34(9):929-39.
37. Reilly JJ, Kelly L, Montgomery C, *et al.* Physical activity to prevent obesity in young children: cluster randomised controlled trial. *Br Med J* 2006;333(7577):1041.
38. Rooney BL, Gritt LR, Havens SJ, *et al.* Growing healthy families: family use of pedometers to increase physical activity and slow the rate of obesity. *Wis Med J* 2005;104(5):54-60.
39. Shaibi GQ, Cruz ML, Ball GD, *et al.* Effects of resistance training on insulin sensitivity in overweight Latino adolescent males. *Med Sci Sports Exerc* 2006;38(7):1208-15.
40. Simon C, Schweitzer B, Oujaa M, *et al.* Successful overweight prevention in adolescents by increasing physical activity: a 4-year randomized controlled intervention. *Int J Obes (Lond)* 2008;32(10):1489-98.
41. Tan S, Yang C, Wang J. Physical training of 9- to 10-year-old children with obesity to lactate threshold intensity. *Pediatr Exerc Sci* 2010;22(3):477-85.

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2
3 523 42. Velez A, Golem DL, Arent SM. The impact of a 12-week resistance training program
4
5 524 on strength, body composition, and self-concept of Hispanic adolescents. *J Strength*
6
7 525 *Cond Res* 2010;24(4):1065-73.
8
9
10 526 43. Walther C, Gaede L, Adams V, *et al.* Effect of increased exercise in school children
11
12 527 on physical fitness and endothelial progenitor cells: a prospective randomized trial.
13
14 528 *Circulation* 2009;120(22):2251-59.
15
16
17 529 44. Warren JM, Henry CJ, Lightowler HJ, *et al.* Evaluation of a pilot school programme
18
19 530 aimed at the prevention of obesity in children. *Health Promot Int* 2003;18(4):287-96.
20
21 531 45. Watts K, Beye P, Siafarikas A, *et al.* Exercise training normalizes vascular
22
23 532 dysfunction and improves central adiposity in obese adolescents. *J Am Coll Cardiol*
24
25 533 2004;43(10):1823-27.
26
27
28 534 46. Watts K, Beye P, Siafarikas A, *et al.* Effects of exercise training on vascular function
29
30 535 in obese children. *J Pediatr* 2004;144(5):620-25.
31
32
33 536 47. Weintraub DL, Tirumalai EC, Haydel KF, *et al.* Team sports for overweight children:
34
35 537 the Stanford Sports to Prevent Obesity Randomized Trial (SPORT). *Arch Pediatr*
36
37 538 *Adolesc Med* 2008;162(3):232-37.
38
39
40 539 48. Sigal RJ, Alberga AS, Goldfield GS, *et al.* Effects of aerobic training, resistance
41
42 540 training, or both on percentage body fat and cardiometabolic risk markers in obese
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44 541 adolescents: the healthy eating aerobic and resistance training in youth randomized
45
46 542 clinical trial. *JAMA Pediatr* 2014;168(11):1006-14.
47
48
49 543 49. Meucci M, Cook C, Curry CD, *et al.* Effects of supervised exercise program on
50
51 544 metabolic function in overweight adolescents. *World J Pediatr* 2013;9(4):307-11.
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50. Lee S, Deldin AR, White D, *et al.* Aerobic exercise but not resistance exercise reduces intrahepatic lipid content and visceral fat and improves insulin sensitivity in obese adolescent girls: a randomized controlled trial. *Am J Physiol Endocrinol Metab* 2013;305(10):E1222-E29.
51. Dennis BA, Ergul A, Gower BA, *et al.* Oxidative stress and cardiovascular risk in overweight children in an exercise intervention program. *Childhood Obesity* 2013;9(1):15-21.
52. Maddison R, Foley L, Ni MC, *et al.* Effects of active video games on body composition: a randomized controlled trial. *Am J Clin Nutr* 2011;94(1):156-63.
53. Wong PC, Chia MY, Tsou IY, *et al.* Effects of a 12-week exercise training programme on aerobic fitness, body composition, blood lipids and C-reactive protein in adolescents with obesity. *Ann Acad Med Singapore* 2008;37(4):286-93.
54. Saygin O, Ozturk MA. The effect of twelve week aerobic exercise programme on health related physical fitness components and blood lipids in obese girls. *Afr J Pharm Pharmacol* 2011;5(12):1441-45.
55. Sun MX, Huang XQ, Yan Y, *et al.* One-hour after-school exercise ameliorates central adiposity and lipids in overweight Chinese adolescents: a randomized controlled trial. *Chin Med J (Engl)* 2011;124(3):323-29.
56. Elloumi M, Makni E, Ounis OB, *et al.* Six-minute walking test and the assessment of cardiorespiratory responses during weight-loss programmes in obese children. *Physiother Res Int* 2011;16(1):32-42.
57. Lau PWC, Wong dP, Ngo JK, *et al.* Effects of high-intensity intermittent running exercise in overweight children. *Eur J Sport Sci* 2014:1-9.

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58. Alberga AS, Farnesi BC, Lafleche A, *et al.* The effects of resistance exercise training on body composition and strength in obese prepubertal children. *Phys Sportsmed* 2013;41(3):103-09.

59. Ackel-D'Elia C, Carnier J, Bueno CR, Jr., *et al.* Effects of different physical exercises on leptin concentration in obese adolescents. *Int J Sports Med* 2014;35(2):164-71.

60. Fazelifar S, Ebrahim K, Sarkisian V. Effect of concurrent training and detraining on anti-inflammatory biomarker and physical fitness levels in obese children. *Rev Bras Med Esporte* 2013;19(5):349-54.

61. Ghorbanian B, Ravassi A, Kordi MR, *et al.* The effects of rope training on lymphocyte ABCA1 expression, plasma ApoA-I and HDL-c in boy adolescents. *Int J Endocrinol Metab Disord* 2013;11(2):76-81.

62. Lee S, Bacha F, Hannon T, *et al.* Effects of aerobic versus resistance exercise without caloric restriction on abdominal fat, intrahepatic lipid, and insulin sensitivity in obese adolescent boys: a randomized, controlled trial. *Diabetes* 2012;61(11):2787-95.

63. Song JK, Stebbins CL, Kim TK, *et al.* Effects of 12 weeks of aerobic exercise on body composition and vascular compliance in obese boys. *J Sports Med Phys Fitness* 2012;52(5):522-29.

64. Cheng HL, Peng P, Zhu R, *et al.* Effects of eight weeks exercise prescription intervention on aerobic capacity, body composition, blood lipid and C-reactive protein in obese adolescents. [Chinese]. *J Jilin Univ Med* 2012;38:745-49.

65. Kelly LA, Loza A, Lin X, *et al*. The effect of a home-based strength training program on type 2 diabetes risk in obese Latino boys. *J Pediatr Endocrinol Metab* 2015;28(3-4):315-22.
66. Hedges LV, Olkin I. Vote-counting methods in research synthesis. *Psychol Bull* 1980;88:359-69.
67. James A, Soler A, Weatherall R. Cognitive behavioural therapy for anxiety disorders in children and adolescents. *Cochrane Database Syst Rev* 2005(4):CD004690.
68. Schranz N, Tomkinson G, Olds T. What is the effect of resistance training on the strength, body composition and psychosocial status of overweight and obese children and adolescents? A systematic review and meta-analysis. *Sports Med* 2013;43(9):893-907.
69. Kelley GA, Kelley KS, Pate RR. Exercise and BMI in overweight and obese children and adolescents: A systematic review with trial sequential meta-analysis. *Biomed Res Int* 2015;2015(Article ID 704539):1-17.
70. Mei H, Xiong Y, Xie S, *et al*. The impact of long-term school-based physical activity interventions on body mass index of primary school children - a meta-analysis of randomized controlled trials. *BMC Public Health* 2016;16(1):205.
71. Garcia-Hermoso A, Sanchez-Lopez M, Martinez-Vizcaino V. Effects of aerobic plus resistance exercise on body composition related variables in pediatric obesity: A systematic review and meta-analysis of randomized controlled trials. *Pediatr Exerc Sci* 2015;27(4):431-40.

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72. Dellert JC, Johnson P. Interventions with children and parents to improve physical activity and body mass index: a meta-analysis. *Am J Health Promot* 2014;28(4):259-67.

73. Atlantis E, Barnes EH, Singh MA. Efficacy of exercise for treating overweight in children and adolescents: a systematic review. *Int J Obes* 2006;30(7):1027-40.

74. Harris KC, Kuramoto LK, Schulzer M, *et al*. Effect of school-based physical activity interventions on body mass index in children: a meta-analysis. *Can Med Assoc J* 2009;180(7):719-26.

75. McGovern L, Johnson JN, Paulo R, *et al*. Treatment of Pediatric Obesity: A Systematic Review and Meta-Analysis of Randomized Trials. *J Clin Endocrinol Metab* 2008;93(12):4600-05.

76. Cesa CC, Sbruzzi G, Ribeiro RA, *et al*. Physical activity and cardiovascular risk factors in children: meta-analysis of randomized clinical trials. *Prev Med* 2014;69(12):54-62.

77. Guerra PH, Nobre MRC, da Silveira JAC, *et al*. The effect of school-based physical activity interventions on body mass index: a meta-analysis of randomized trials. *Clinics* 2013;68(9):1263-73.

78. Kelley GA, Kelley KS. Exercise and BMI z-score in overweight and obese children and adolescents: protocol for a systematic review and network meta-analysis of randomised trials. *BMJ Open* 2016;6(e011258):1-7.

79. Kelley GA, Kelley KS, Pate R. Exercise and BMI z-score in overweight and obese children and adolescents: A systematic review and network meta-analysis of randomized trials. *J Evid Based Med* 2017;10(2):108-28.

- 633 80. Hutton B, Salanti G, Caldwell DM, *et al.* The PRISMA Extension Statement for
634 Reporting of Systematic Reviews Incorporating Network Meta-analyses of Health
635 Care Interventions: Checklist and Explanations. *Ann Intern Med* 2015;162(11):777-
636 84.
- 637 81. Sacks HS, Chalmers TC, Smith H. Randomized versus historical controls for clinical
638 trials. *Am J Med* 1982;72:233-40.
- 639 82. Schulz KF, Chalmers I, Hayes R, *et al.* Empirical evidence of bias: Dimensions of
640 methodological quality associated with estimates of treatment effects in controlled
641 trials. *J Am Med Assoc* 1995;273:408-12.
- 642 83. Physical Activity Guidelines Advisory Committee. *Physical Activity Guidelines*
643 *Advisory Report*. Washington, DC: U.S. Department of Health and Human
644 Services, 2008
- 645 84. Caspersen CJ, Powell KE, Christenson GM. Physical activity, exercise, and physical
646 fitness: definitions and distinctions for health-related research. *Public Health Rep*
647 1985;100(2):126-31.
- 648 85. Dietz WH. Health consequences of obesity in youth: childhood predictors of adult
649 disease. *Pediatrics* 1998;101(3 Pt 2):518-25.
- 650 86. Egger E, Zellwegerzahner T, Schneider M, *et al.* Language bias in randomised
651 controlled trials published in English and German. *Lancet* 1997;350:326-29.
- 652 87. Moher D, Pham B, Klassen TP, *et al.* What contributions do languages other than
653 English make on the results of meta-analyses? *J Clin Epidemiol* 2000;53:964-72.
- 654 88. Berlin JA. Does blinding of readers affect the results of meta-analyses? *Lancet*
655 1997;350:185-86.

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89. Cohen J. Weighted kappa: nominal scale agreement with provision for scaled disagreement or partial credit. *Psychol Bull* 1968;70:213-20.

90. Lee E, Dobbins M, DeCorby K, *et al*. An optimal search filter for retrieving systematic reviews and meta-analyses. *BMC Med Res Methodol* 2012;12:51.

91. Higgins JP, Altman DG, Gotzsche PC, *et al*. The Cochrane Collaboration's tool for assessing risk of bias in randomised trials. *Br Med J* 2011;343:d5928.

92. Emerson JD, Burdick E, Hoaglin DC, *et al*. An empirical study of the possible relation of treatment differences to quality scores in controlled randomized clinical trials. *Control Clin Trials* 1990;11:339-52.

93. Juni P, Witschi A, Bloch R, *et al*. The hazards of scoring the quality of clinical trials for meta-analysis. *J Am Med Assoc* 1999;282:1054-60.

94. Ahn S, Becker BJ. Incorporating quality scores in meta-analysis. *J Educ Behav Stat* 2011;36(5):555-85.

95. Follmann D, Elliot P, Suh I, *et al*. Variance imputation for overviews of clinical trials with continuous response. *J Clin Epidemiol* 1992;45:769-73.

96. Higgins JPT, Green S, editors. *Cochrane Handbook for Systematic Reviews of Interventions Version 5.1.0 [updated March 2011]*: The Cochrane Collaboration, 2011.

97. Hedges LV, Olkin I. *Statistical methods for meta-analysis*. San Diego, CA: Academic Press, 1985.

98. Chaimani A, Higgins JPT, Mavridis D, *et al*. Graphical tools for network meta-analysis in STATA. *PLoS One* 2013;8(10)

99. Catala-Lopez F, Tobias A, Cameron C, *et al*. Network meta-analysis for comparing treatment effects of multiple interventions: an introduction. *Rheumatol Int* 2014;34(11):1489-96.
100. White IR, Barrett JK, Jackson D, *et al*. Consistency and inconsistency in network meta-analysis: model estimation using multivariate meta-regression. *Res Synth Methods* 2012;3(2):111-25.
101. White IR. Multivariate random-effects meta-regression: Updates to mvmeta. *Stata J* 2011;11(2):255-70.
102. Jansen JP, Naci H. Is network meta-analysis as valid as standard pairwise meta-analysis? It all depends on the distribution of effect modifiers. *BMC Med* 2013;11
103. Donegan S, Williamson P, D'Alessandro U, *et al*. Assessing key assumptions of network meta-analysis: a review of methods. *Res Synth Methods* 2013;4(4):291-323.
104. Lu GB, Ades AE. Assessing evidence inconsistency in mixed treatment comparisons. *J Am Stat Assoc* 2006;101(474):447-59.
105. Veroniki AA, Vasiliadis HS, Higgins JP, *et al*. Evaluation of inconsistency in networks of interventions. *Int J Epidemiol* 2013;42(1):332-45.
106. Song F, Xiong T, Parekh-Bhurke S, *et al*. Inconsistency between direct and indirect comparisons of competing interventions: meta-epidemiological study. *Br Med J* 2011;343:d4909.
107. Higgins JP, Thompson SG, Spiegelhalter DJ. A re-evaluation of random-effects meta-analysis. *J R Stat Soc Series A* 2009;172(1):137-59.

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55
56
57
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59
60

700 108. Kelley GA, Kelley KS. Impact of progressive resistance training on lipids and
701 lipoproteins in adults: Another look at a meta-analysis using prediction intervals.
702 *Prev Med* 2009;49:473-75.

703 109. Cooper HC, Hedges LV, Valentine JF. The handbook of research synthesis. New
704 York, New York: Russell Sage 2009.

705 110. Salanti G, Del Giovane C, Chaimani A, *et al*. Evaluating the quality of evidence
706 from a network meta-analysis. *PLoS One* 2014;9(7)

707 111. Salanti G, Ades AE, Ioannidis JP. Graphical methods and numerical summaries for
708 presenting results from multiple-treatment meta-analysis: an overview and
709 tutorial. *J Clin Epidemiol* 2011;64(2):163-71.

Table 1. Covariates to examine using simple meta-regression.

Characteristics	Variable
Study	Publication year, impact factor of journal, country study conducted, type of control group, bias (sequence generation, allocation concealment, blinding of participants & personnel, blinding of outcome assessors, incomplete outcome data, selective outcome reporting), type of analysis
Participant	Age, gender, race/ethnicity, maturational stage
Exercise	Type (aerobic, strength, both), length, frequency, intensity, duration, total minutes, total minutes (adjusted for compliance), mode, compliance, exercise supervision, setting, number of sets, number of repetitions, rest between sets, number of exercises, type of resistance, equipment used, fidelity (design, training, delivery, receipt, enactment)
Outcome	Baseline values for primary outcomes (BMI in kg/m ² , fat mass, percent fat), method used to assess adiposity, i.e., instrumentation, body weight, lean body mass, waist circumference, waist-to-hip ratio, diet, energy intake, energy expenditure, physical activity level, non-exercise activity, maximum oxygen consumption (relative and absolute), muscular strength, resting systolic and diastolic blood pressure, total cholesterol, high-density lipoprotein cholesterol, ratio of total cholesterol to high-density lipoprotein cholesterol, non-high density lipoprotein cholesterol, low-density lipoprotein cholesterol, triglycerides, glycosylated hemoglobin, fasting and non-fasting glucose and insulin

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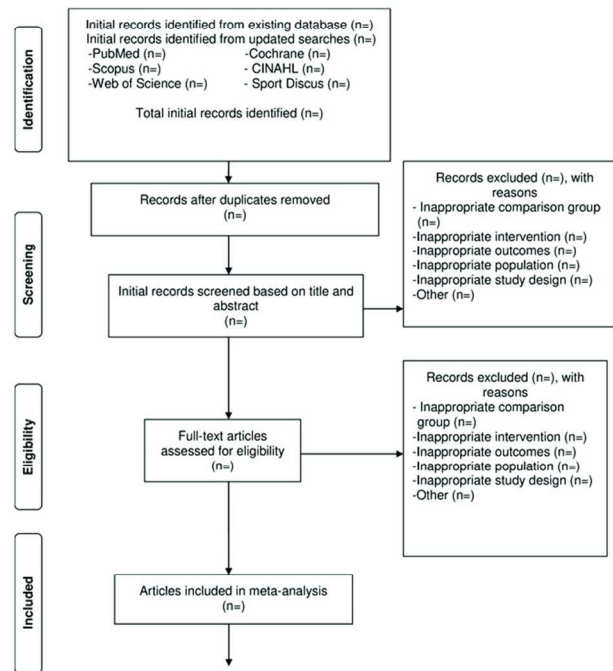
FIGURE LEGEND

Figure 1. Proposed flow diagram to depict the search process.

SUPPLEMENTARY FILE

Supplementary File 1. Preliminary search results in PubMed.

For peer review only



Flow diagram for network meta-analysis.

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PRISMA-P (Preferred Reporting Items for Systematic review and Meta-Analysis Protocols) 2015 checklist: recommended items to address in a systematic review protocol*

Section and topic	Item No	Checklist item	Line #
ADMINISTRATIVE INFORMATION			
Title:			
Identification	1a	Identify the report as a protocol of a systematic review	1-2
Update	1b	If the protocol is for an update of a previous systematic review, identify as such	NA
Registration	2	If registered, provide the name of the registry (such as PROSPERO) and registration number	43; 131-132;394-398
Authors:			
Contact	3a	Provide name, institutional affiliation, e-mail address of all protocol authors; provide physical mailing address of corresponding author	4-7; 9-12;14-17
Contributions	3b	Describe contributions of protocol authors and identify the guarantor of the review	387-393
Amendments	4	If the protocol represents an amendment of a previously completed or published protocol, identify as such and list changes; otherwise, state plan for documenting important protocol amendments	399-401
Support:			
Sources	5a	Indicate sources of financial or other support for the review	404-408
Sponsor	5b	Provide name for the review funder and/or sponsor	
Role of sponsor or funder	5c	Describe roles of funder(s), sponsor(s), and/or institution(s), if any, in developing the protocol	
INTRODUCTION			
Rationale	6	Describe the rationale for the review in the context of what is already known	60-119
Objectives	7	Provide an explicit statement of the question(s) the review will address with reference to participants, interventions, comparators, and outcomes (PICO)	120-126
METHODS			
Eligibility criteria	8	Specify the study characteristics (such as PICO, study design, setting, time frame) and report characteristics (such as years considered, language, publication status) to be used as criteria for eligibility for the review	133-179
Information sources	9	Describe all intended information sources (such as electronic databases, contact with study authors, trial registers or other grey literature sources) with planned dates of coverage	180-189
Search strategy	10	Present draft of search strategy to be used for at least one electronic database, including planned limits, such that it could be repeated	190-198; Supplementary file 1
Study records:			
Data management	11a	Describe the mechanism(s) that will be used to manage records and data throughout the review	201-202; 228-230;378-383

Selection process	11b	State the process that will be used for selecting studies (such as two independent reviewers) through each phase of the review (that is, screening, eligibility and inclusion in meta-analysis)	201-226; Figure 1
Data collection process	11c	Describe planned method of extracting data from reports (such as piloting forms, done independently, duplicate), any processes for obtaining and confirming data from investigators	227-243
Data items	12	List and define all variables for which data will be sought (such as PICO items, funding sources) any pre-planned data assumptions and simplifications	233-239;244-253;Table 1
Outcomes and prioritization	13	List and define all outcomes for which data will be sought, including prioritization of main and additional outcomes, with rationale	244-253
Risk of bias in individual studies	14	Describe anticipated methods for assessing risk of bias of individual studies, including whether this will be done at the outcome or study level, or both; state how this information will be used in data synthesis	254-272
Data synthesis	15a	Describe criteria under which study data will be quantitatively synthesised	274-350; 362-376; & Table 1 for 15a-d
	15b	If data are appropriate for quantitative synthesis, describe planned summary measures, methods of handling data and methods of combining data from studies, including any planned exploration of consistency (such as I^2 , Kendall's τ)	
	15c	Describe any proposed additional analyses (such as sensitivity or subgroup analyses, meta-regression)	
	15d	If quantitative synthesis is not appropriate, describe the type of summary planned	
Meta-bias(es)	16	Specify any planned assessment of meta-bias(es) (such as publication bias across studies, selective reporting within studies)	351-354
Confidence in cumulative evidence	17	Describe how the strength of the body of evidence will be assessed (such as GRADE)	355-361

From: Shamseer L, Moher D, Clarke M, Ghersi D, Liberati A, Petticrew M, Shekelle P, Stewart L, PRISMA-P Group. Preferred reporting items for systematic review and meta-analysis protocols (PRISMA-P) 2015: elaboration and explanation. *BMJ*. 2015 Jan 2;349(jan02 1):g7647.