

Appendix 1. PRISMA 2009 Checklist

| Section/topic | # | Checklist item | Reported on page # |
|---------------------------------------|---|---|---------------------------------|
| TITLE | | | |
| Title | 1 | Identify the report as a systematic review, meta-analysis, or both. | 1 |
| ABSTRACT | Π | | |
| Structured summary | 2 | Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number. | 1 |
| INTRODUCTION | | | |
| Rationale | 3 | Describe the rationale for the review in the context of what is already known. | 2 |
| Objectives | 4 | Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS). | 2 |
| METHODS | - | | |
| Protocol and registration | 5 | Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number. | PROSPERO 2020 CRD42020208833 |
| Eligibility criteria | 6 | Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale. | 3 |
| Information sources | 7 | Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched. | 2-3 |
| Search | 8 | Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated. | Appendix 2 |
| Study selection | 9 | State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis). | 3 |
| Data collection process | 10 | Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators. | |
| Data items | 11 | List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made. | 3 |
| Risk of bias in individual studies | 12 | Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis. | |
| Summary measures | 13 | State the principal summary measures (e.g., risk ratio, difference in means). | Not Applied |
| Synthesis of results | Presente and the second second | | Not Applied |



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| Risk of bias across studies | 15 | Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies). | |
| Additional analyses | 16 | Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified. | |
| RESULTS | | · | |
| Study selection | 17 | Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram. | |
| Study characteristics | 18 | For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations. | |
| Risk of bias within studies | 19 | Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12). | |
| Results of individual studies | 20 | For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot. | |
| Synthesis of results | 21 | Present results of each meta-analysis done, including confidence intervals and measures of consistency. | |
| Risk of bias across studies | 22 | Present results of any assessment of risk of bias across studies (see Item 15). | |
| Additional analysis | 23 | Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]). | Not Applied |
| DISCUSSION | | | |
| Summary of evidence | 24 | Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers). | |
| Limitations | 25 | Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias). | |
| Conclusions | 26 | Provide a general interpretation of the results in the context of other evidence, and implications for future research. | 25 |
| FUNDING | | | |
| Funding | 27 | Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review. | Not Applied |

From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(6): e1000097. doi:10.1371/journal.pmed1000097

Appendix 2. Search strings for all databases in the systematic review.

Table 1. Search strings for MEDLINE (by Pubmed)

(("Exercise"[Mesh]) OR "Exercise therapy"[Mesh]) AND "Aged"[Mesh]
 FILTER's:

 -ARTICLE TYPE: Randomized Controlled Trial
 -PUBLICATION DAY: 2015 to JUN/2020
 -AGE: Aged: 65+ years, 80 and over: 80+ years
 -SPECIES: Humans
 -LANGUAGE: English, Spanish, Portuguese

Table 2. Search strings for Elsevier (Scopus)

(Exercise) OR (Exercise therapy) AND (Aged) AND NOT INDEX (medline)
 FILTER's:

 Year: 2015 to JUN/2020
 Document type: Article
 Country/territory: Brazil, Chile, Mexico, Colombia, Argentina, Ecuador, Peru, Cuba, Costa Rica, Uruguay, Venezuela, Jamaica, Puerto Rico, Honduras, El Salvador, Guatemala, Panama, Bolivia
 Language: English, Spanish, Portuguese
 Subject area: Medicine, Health Professions
 Keyword: Aged, Physical Activity, Aging, Resistance Training, Adult, Elderly, Physical Exercise, Training, Strength Training, Very Elderly, Older Adults, Exercise Therapy

Table 3. Search strings for SciELO

(Exercise) OR (Exercise therapy) AND (Aged)
 FILTER's:

 -Collection: Brazil, Chile, Colombia, México, Cuba, Argentina, Costa Rica, Perú, Uruguay
 -Publication Year: 2015 to JUN/2020
 -Type of Literature: Article
 -Language: English, Spanish, Portuguese

| Item category | CERT item description | N. items rated "Yes"/ N. papers with responses (Percentage) |
|----------------|--|---|
| WHAT: | C1: Detailed description of exercise equipment (e.g. weights, | 91/101 |
| materials | treadmill, ergometer, etc.) | 90% |
| WHO: | C2: Detailed description of instructor expertise, qualifications, | 66/101 |
| provider | and/or training | 65% |
| HOW: | C3: Describe whether exercises are performed individually or | 101/101 |
| | in a group | 100% |
| | C4: Describe whether exercises are supervised or | 101/101 |
| | unsupervised; how they are delivered | 100% |
| | C5: Detailed description of how adherence to exercise is | 94/101 |
| | measured and reported | 93% |
| | • | 5/101 |
| | C6: Detailed description of motivation strategies | 5% |
| | C7a: Detailed description of decision rule(s) for determining | *5/10 |
| | exercise progression | 50% |
| delivery | C7b: Detailed description of how exercise program was | *10/10 |
| 5 | progressed | 100% |
| | 1 0 | 22/101 |
| | C8: Detailed description of each exercise to enable replication | 21% |
| | 1 1 | *0/0 |
| | C9: Detailed description of any home program component | 0% |
| | C10: Describe any non-exercise components, e.g. education, | *3/3 |
| | cognitive behavioral therapy, etc. | 100% |
| | C11: Describe the type and number of adverse events that | 5/12 |
| | occur during exercise | 41% |
| WHERE: | 0 | 45/101 |
| location | C12: Describe the setting in which the exercises are performed | 44% |
| WHEN, HOW | | 11/0 |
| MUCH: dosage | C13: Detailed description of exercise intervention, e.g. reps, | 91/101 |
| in accurace | sets, sessions | 90% |
| TAILORING: | C14a: Describe whether the exercises are generic (one size fits | 101/101 |
| what, how | all) or tailored | 100% |
| wildt, How | C14b: Detailed description of how exercises are tailored to the | 58/101 |
| | individual | 57% |
| | C15: Describe the decision rule for determining the starting | 4/101 |
| | level, e.g. beginner, intermediate, advanced, etc. | 3% |
| HOW WELL: | C16a: Describe how adherence or fidelity to the intervention | 94/101 |
| planed, actual | is assessed/measured | 93% |
| Planeu, actual | C16b: Describe the extent to which the intervention was | 35/101 |
| | delivered as planned | 34% |
| * NT. (| carried out this item. Therefore, the total of articles is different fro | |

Appendix 3. Proportions of "Yes" per CERT item for 101 included trials