Exercise and bariatric surgery: A systematic review and meta-analysis of the feasibility and acceptability of exercise and controlled trial methods

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Abstract

This systematic review and meta-analysis assessed the feasibility and acceptability of

exercise and controlled trial methods in adults awaiting or having undergone bariatric

surgery (BS). Search methods used to identify relevant articles were: inclusion of articles

identified in a systematic review, new database search of articles published 2019-21, and

hand searching reference lists. Titles/abstracts and full-texts were screened by two

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reviewers independently against inclusion criteria: adults awaiting or having undergone BS, controlled trial, exercise group compared to a comparison group without exercise. Twenty-eight articles were reviewed; most interventions were supervised, performed after BS, and lasted ≤13 weeks. Pooled data for exercise intervention attendance and dropout rates were 84% (k=10) and 5% (k=19), respectively, though possibly misestimated due to poor/selective reporting. Median study and recruitment duration were 18 weeks and 24 months, respectively, with a pooled enrollment rate of 2.5 participants/month. Pooled data for refusal to participate, enrollment, and retention rates were 23% (k=16), 43% (k=18), and 87% (k=26), respectively. Exercise and controlled trial methods seem feasible and acceptable for adults awaiting or having undergone BS. However, improved reporting of feasibility and acceptability indicators is needed to better identify methodological or practical challenges, and assess bias.

Introduction

The prevalence of severe obesity, defined by a body mass index ≥35 kg/m², is constantly increasing in higher income countries.¹ Worldwide, the number of adults with obesity increased from 100 million in 1975 to 671 million in 2016.¹ Severe obesity increases the risk of morbidity and mortality, and reduces quality of life.²-5 Treatment of severe obesity by bariatric surgery is generally effective for long-term weight loss, and can offer additional benefits (e.g., reduced relative risk of morbidity and mortality, improved quality

of life).^{6, 7} As a result, the number of bariatric surgery procedures around the world has risen dramatically in recent years, with 604,223 surgeries performed in 2018.⁸

Controlled trials provide evidence that exercise is an important part of care before and after bariatric surgery, as it can help to maintain or enhance the short- and long-term benefits of surgery. Previous reviews showed that exercise positively impacts cardiorespiratory fitness, muscle strength, cardiometabolic health, and weight and fat loss after bariatric surgery. However, general questions regarding the feasibility and acceptability of exercise for adults awaiting or having undergone bariatric surgery remain unanswered as previous reviews focused on summarizing the effects of exercise on outcomes of interest.

Summarizing the evidence for the feasibility and acceptability of exercise is needed for guidance of professionals and to help implementation in the field.²⁰ In addition, this information may serve to identify potential aspects of the intervention in need of refinement during implementation. By extension, whether the methods used in controlled trials to assess the effects of exercise are both feasible and acceptable is critical as it can impact the outcomes, provide a higher level of evidence for future guideline updates, and provide a basis for selecting certain methods in future research (e.g., opt for some recruitment strategies over others).²⁰ As well, it can help elucidate potential methodological or practical challenges warranting attention (e.g., time, resources, data management) early on during the developmental phase of the research to ensure successful trial completion. This is especially needed given the limited resources generally available.

Thus, the objective of this systematic review and meta-analysis of published controlled trials were to: i) assess the evidence of the feasibility and acceptability of exercise

(intervention) and controlled trial methods in adults awaiting or having undergone bariatric surgery, and ii) identify factors associated with feasibility and acceptability outcomes. Whilst there are no universally accepted definitions for feasibility and acceptability adopted by authors of such controlled trials, feasibility was considered to reflect whether the exercise intervention can be delivered to participants as planned and whether the methods (e.g., assessment protocol) can be successfully executed by the researchers, ²¹ and acceptability was considered to reflect the suitability of the exercise intervention and methods from the perspective of intended users (e.g., adults awaiting or having undergone bariatric surgery) or those responsible for implementation (e.g., healthcare providers delivering the intervention, research staff recruiting participants). ²¹

Methods

Protocol and registration

The protocol for this systematic review and meta-analysis was registered in the Prospective Register of Systematic Reviews (PROSPERO) international registry (CRD42021255048, 07/2021) and it is reported according to the Preferred Reporting Items for Systematic Reviews and Meta-Analysis (PRISMA) guidelines.²² The completed PRISMA checklist is presented in Supplemental file 1. All supplemental files, and R data are available on Open Science Framework (https://osf.io/h4p27/).

Data sources and searches

Three search methods were used to identify relevant articles. The first method was to include all articles found in a systematic review and meta-analysis published in June 2021

on exercise training and bariatric surgery. As the authors database search was limited to articles published prior to January 2019, the second method was to update the database search, and search for articles published from January 1, 2019 onwards. This search was performed by two reviewers (MSP, ABa) in April 2021 using three databases: PubMed, Web of Science, and EMBASE. The search strategies for each database were developed using the same strategies as in Bellicha et al. (see Supplemental file with search strategy in Bellicha et al. 9). "Feasibility" and "acceptability" were not added as terms to be able to retrieve as many articles as possible. The third method was to hand-search reference lists from eligible articles and relevant reviews 10-19 to identify other potentially relevant articles; this search was completed by one reviewer (MSP).

Eligibility criteria

To be included, articles had to: i) be an English-language full-text article reporting on primary research published in a peer-reviewed journal; ii) involve a controlled trial, either randomized or non-randomized, comparing an exercise group to a control group without exercise, and; iii) include adults (≥ 18 years) awaiting or having undergone bariatric surgery. Articles were excluded if they only presented a study focused on behavioral interventions to promote exercise engagement. Systematic reviews (with or without metaanalyses), study protocols without results, grey literature, case reports, letters/commentaries, animal studies, observational quantitative, qualitative, or mixedmethods studies, and published abstracts were also excluded.

Study selection of the updated search

Once the database searches were completed, records of the articles retrieved were imported into Endnote X9.3.3. After removing duplicates, titles and abstracts were screened concurrently against selection criteria by two reviewers (ABa, MSP). Full-texts were obtained for potentially relevant articles and screened against selection criteria by two reviewers (ABa, MSP). Where there was disagreement on eligibility, a third reviewer (JB) assessed the full-texts of the articles in detail against the selection criteria and discussions took place to determine whether the articles should be included. Next, the reference lists of the eligible articles and relevant reviews were reviewed.

Data extraction

Data pertaining to study details, participants' characteristics, exercise intervention and comparison group(s), and outcomes were extracted and entered into a Microsoft Excel table developed for this review by one reviewer (WC), and checked by two other reviewers (LB, MSP). See Supplemental file 2 for more details.

Feasibility and acceptability data were also extracted and entered into a Microsoft Excel table by one reviewer (MSP), and checked by another reviewer (ABa). Disagreements were discussed and resolved by a third reviewer (JB). To ensure a comprehensive synthesis of feasibility and acceptability evidence, the following approach was used. Data presented in the CONSORT flow chart (where applicable) were extracted for each group. Then, all data reflecting feasibility or acceptability outcomes were extracted from each article including: i) number of participants who discontinued the intervention and reasons ¹; ii) participants' satisfaction ratings/scores; iii) reported attendance rate related to session *frequency* (i.e.,

¹ For the purpose of this review, participants reported as having discontinued the intervention because of lack of attendance or compliance were considered as excluded from analysis.

among participants analyzed, percentage of participants who attended all or a certain percentage of prescribed sessions according to the definition given by the authors of included articles²³; **iv**) reported compliance rate related to session *duration* and *intensity* (i.e., among participants analyzed, percentage of participants who met all or a certain percentage of prescribed session duration and intensity according to the definition given by the authors of included articles)²³; **v**) number and type of adverse events related to exercise intervention only (e.g., fall, injury), and; **vi**) other quantitative or qualitative data showing feasibility and acceptability of the intervention.

Additional data for feasibility and acceptability of controlled trial methods were extracted, and included: i) number of participants assessed for eligibility (i.e., number recruited); ii) number of participants excluded and reasons for exclusion (e.g., did not meet eligibility criteria, declined to participate, other reasons); iii) number of participants randomized, number of participants who refused randomization and refusal reasons; iv) number of participants allocated to each group; v) number of participants who received allocated intervention (i.e., number who started the intervention), and number of participants who did not received allocated intervention (i.e., number randomized to the intervention but did not start the intervention) and reasons; vi) number of participants lost to follow-up (i.e., number who started assessments, but did not complete them until the end of the study) and reasons²; vii) number of participants analyzed, and number excluded from analysis and reasons; viii) recruitment strategies (description and classification into active (contact with participants was made by the research team (e.g., clinician referral, phone calls)) or passive (contact was made by potential participants (e.g., self-referral by poster, newspaper

² For the purpose of this review, participants reported as lost to follow-up because of lack of attendance were considered rather as excluded from analysis.

advertisements)),²⁴ ix) recruitment duration in months, x) missing data (total (i.e., across time points and collectively for all outcomes), per time point, and/or per outcome), and xi) other quantitative or qualitative data showing feasibility and acceptability of the study protocol.

Corresponding authors were contacted two times by email to obtain missing data or clarification, if necessary. Data unavailable or unclear despite the authors responding or when authors were unresponsive after two contact attempts were marked as 'not available' (NA) or unclear (?), respectively. When multiple articles were published from a single study, only data from the first published article containing a flow chart were extracted; more recent parent articles were checked to adjust or add additional data if unavailable or unclear in the earlier article. For articles that included a follow-up phase, data were extracted separately for the intervention (i.e., from recruitment to completion of post-intervention assessment) and follow-up phases (i.e., from post-intervention assessment to the end of the study) to allow for easier comparison between studies without follow-up.

Quality assessment

As in Bellicha et al.'s review,⁹ the quality of included articles was assessed using the National Heart, Lung and Blood Institute quality assessment tool for controlled intervention studies.²⁵ Two reviewers (ABe, ABa) completed this task. Study quality was defined as good, fair or poor when 0, 1 or \geq 2 fatal flaws (i.e., non-randomized study; dropout rate \geq 20%; no intention-to-treat analyses) were identified by both reviewers.

Disagreements were discussed between reviewers.

Feasibility and acceptability data analysis

Feasibility and acceptability data pertaining to the exercise intervention and controlled trial methods were used to calculate several rates/indicators for the intervention phase and for the follow-up phase (where applicable). Table 1 presents the calculations performed according to data extracted from the flow chart and from data in each article.

Statistical analysis

Data were grouped together to provide a tabular summary for narrative synthesis of the included articles, and where possible, quantitative data were pooled for statistical analyses. Prevalence for exercise intervention attendance and compliance, dropout, refusal to participate, recruitment, enrollment, and retention rates, as well as enrollment speed were calculated using random-effect models. Prevalence estimates were computed using the recommended Freeman-Tukey double arcsine transformation. For each outcome of interest, prevalence estimates were synthesized using the double arcsine transformation, and then the pooled estimate was back-transformed to a proportion. Heterogeneity was assessed with Cochran's Q and I² statistics. A low P value (i.e., p<.10) of the Q-statistic indicated that variation in the study-specific effect estimates was due to heterogeneity beyond chance. The I² values ranging from 0% (no observed heterogeneity) to 100% (complete heterogeneity). The risk of publication bias was examined with funnel plots and tested using the Egger's test (p<.10 indicating a publication bias). A trim and fill analysis was also carried out to examine the impact of missing studies by adjusting the

meta-analysis to take into account the theoretically missing studies. ²⁹ Regression residuals were screened to identify potential multivariate outliers using residual Cook distances. Subgroup analyses (i.e., study quality, exercise intervention timing, exercise intervention duration) were carried out for each prevalence estimates. However, no comparison tests were carried out. For study quality, the median (Mdn=8) was used to form higher and lower quality studies (≤8 and >8). For exercise intervention timing, timing was either before or after bariatric surgery. For exercise intervention duration, the median (Mdn=12) was used to form shorter versus longer exercise duration (≤12 and >12 weeks) after excluding one study with a very long intervention duration (i.e., 100 weeks). ³⁰ All analyses were performed in R 4.1 using the 'metafor' package. ³¹

Results

Study selection

The 31 articles from Bellicha et al.'s review⁹ were included (covering articles published prior to January 2019), and 725 additional articles were retrieved during the database search of articles published in January 2019 onwards after removing duplicates (Supplemental file 3). Of the latter, 10 articles met selection criteria. One additional article identified during hand-searching was reviewed; it met selection criteria, yielding a total of 42 articles representing 28 unique studies for review. Thus, 28 articles^{30, 32-58} were included after removing articles coming from the same study.

Participant characteristics

Participant characteristics for each study are described in Supplemental file 4. The 28 articles included 1,250 participants (range 6-220 per study) with a mean age <40 years in

21.4% (k=6) of articles (range 33.3-53.9 years). $^{34, 35, 42, 52, 54, 55}$ Most (k=18, 64.3%) studies had samples comprising \geq 75% women $^{32-34, 36, 37, 39-47, 49, 54, 56, 58}$ and 21.4% (k=6) comprised only women. $^{35, 38, 48, 51-53}$ Beyond providing data on age and sex/gender, 39.3% (k=11) of articles reported other sociodemographic data, $^{30, 32, 33, 35, 39, 44, 45, 49, 51, 52, 56}$ and 42.9% (k=12) reported comorbidities prevalence. $^{32, 33, 35, 36, 39, 46, 49, 51-53, 56, 58}$ Among the 6 articles (21.4%) reporting ethnicity/race, all had samples comprising \geq 50% White participants. $^{36, 37, 40, 41, 49, 56}$ Participants were awaiting or underwent Roux-Y gastric bypass in 32.1% (k=9) of articles, $^{34-36, 38, 45, 48, 50, 51, 53}$ sleeve gastrectomy in 7.1% (k=2), $^{46, 55}$ either RYGB, sleeve gastrectomy, biliopancreatic diversion with duodenal switch or gastric banding in 42.9% (k=12) $^{30, 33, 37, 39, 41-44, 52, 56-58}$; the type of bariatric surgery was not reported in 17.9% (k=5) of articles. $^{32, 40, 46, 49, 54}$

Baseline physical fitness and physical activity levels were reported in 60.7% (k=17)^{32, 33, 35, 37, 40-43, 46-49, 52-54, 56, 57} and 28.6% (k=8) of articles,^{32, 33, 35, 37, 43, 44, 53, 56} respectively. All articles including a 6-minute walking test distance (k=7, 25.0%)^{32, 33, 35, 37, 40, 49, 57} had a mean distance ranging from 273 to 503 m and a mean VO2_{peak} between 16 to 22 ml/kg/min (k=8, 28.6%).^{41, 46-48, 52, 53, 56, 57} Due to the variety of measures/tools, units, and variables reported, data on baseline physical activity levels could not be synthetized. Nevertheless, data reported within articles are presented in Supplemental file 4.

Exercise intervention characteristics

Table 2 presents information on the exercise intervention and comparison groups evaluated within the 28 articles. Only data pertaining to the exercise interventions are synthesized below. A quarter (k=7) of interventions were initiated before bariatric surgery^{32, 33, 40, 41, 46, 49, 54}; the rest (k=21, 75.0%) were initiated after bariatric surgery.^{30, 34-39, 42-45, 47, 48, 50-53, 55-}

⁵⁸ Most (k=16, 57.1%) lasted ≤13 weeks (range 4-104 weeks), $^{32, 33, 35, 38, 40-46, 52, 55-58}$ 89.3% (k=25) were supervised exercise, $^{30, 32-37, 39-41, 43-54, 56-58}$ and 35.7% (k=10) held sessions individually. $^{30, 32, 35, 36, 41-43, 51, 56, 57}$ Exercise sessions were performed: in hospitals, clinics, or medical centers (k=10, 35.7%), $^{33, 40, 43, 49, 51-53, 56-58}$ at home (k=4, 14.3%), $^{30, 36, 41, 42}$ in research centers (k=3, 10.7%), $^{35, 46, 47}$ in community or fitness centers (k=3, 10.7%), $^{44, 50, 54}$ or at a facility (k=3, 10.7%), $^{32, 39, 45}$ location was not specified in 17.9% (k=5) of articles. $^{34, 37, 38, 48, 55}$

Most (k=19, 67.9%) interventions initiated after bariatric surgery started during the first year after the surgery was performed, $^{30, 34-39, 42-44, 48, 50-53, 55-58}$ ranging from 0 to 36 months after bariatric surgery. Only 7.1% (k=2) intervention were initiated \geq 2 years after bariatric surgery. Training consisted of: endurance training (k=6, 21.4%), $^{35, 36, 41, 48, 49, 56}$, resistance training (k=5, 17.9%), $^{38, 39, 44, 45, 53}$ combination of resistance and endurance training (k=16, 57.1%), $^{30, 32-34, 37, 40, 42, 43, 46, 47, 50-52, 54, 57, 58}$ and balance training (k=1, 3.6%). 55

Feasibility and acceptability of the interventions

Table 2 presents information from articles on the feasibility and acceptability of the exercise intervention and comparison groups. Only data pertaining to the exercise interventions are synthetized below. Among the 57.1% (k=16) of articles reporting on whether there were adverse events during exercise, 33, 35-37, 39, 41, 43-45, 48, 51-54, 56, 58 32.1% (k=9) reported none, 36, 41, 43-45, 51-54 14.3% (k=4) reported occasional pain, fatigue, or dyspnea, 33, 35, 39, 56 7.1% (k=2) reported hypoglycemia or hypotension, 33, 58 and 3.6% (k=1) reported back bruise after a fall. 39

Main reasons reported for exercise intervention dropout were: lack of time (12 participants from 3 articles^{36, 37, 56}), work or other commitments (7 participants from 2 articles^{35, 49}), lack of motivation (3 participants from 1 article⁴⁸), dislike of exercise (2 participants from 1 article³⁵), and postponed surgery or complications (2 participants from 2 articles^{43, 49}). Satisfaction with the exercise intervention was only reported in 3.6% (k=1) of articles.³³ In this study,³³ all participants were 'very satisfied' or 'satisfied' with the person delivering the intervention (i.e., a kinesiologist), materials, and exercise modalities, except for one participant who was 'moderately satisfied' with the evening schedule and the training location.

No meta-analysis was performed for exercise intervention compliance rate given the very small number of articles providing these data (10.7%, k=3), more importantly given the varying operationalization of compliance.^{36, 56, 58} The pooled percentage of scheduled exercise session completed (attendance) and the exercise intervention dropout rates are presented Table 3 and Supplemental file 5; related subgroup analysis are available in Table 3 and Supplemental files 6,7, and 8. Briefly, no significant differences in attendance nor dropout rates were found based on studies quality, exercise intervention timing, and exercise intervention duration.

Characteristics of the controlled trial methods

Characteristics pertaining to the methods of the controlled trials are presented in Table 4. Almost two thirds of articles (64.3%, k=18) were published in the last 5 years (2017-21), with the earliest being published in 2010.^{32, 34, 37-39, 41-43, 45-47, 49-54, 58} Most (67.9%, k=19) were randomized controlled studies.^{30, 32, 33, 35-40, 42, 43, 47, 49-51, 53, 55, 56, 58} Studies were

conducted in Europe (35.7%, k=10), $^{30, 39, 43, 46-48, 50, 53, 54, 57}$ North America (32.1%, k=9), $^{33, 36-38, 40, 41, 44, 56, 58}$ Brazil (21.4%, k=6), $^{34, 35, 45, 49, 51, 52}$ and Iran or Turkey (10.7%, k=3). $^{32, 42, 55}$

Study duration per participant (excluding follow-up phase) ranged from 4 to 161 weeks; the median was 18 weeks (Table 4). Only 17.9% (k=5) of studies had a follow-up phase after the exercise intervention, with a duration ranging from 9 to 52 weeks. $^{33, 43, 47, 50, 58}$ Most (85.7%, k=6) studies in which the exercise intervention was delivered before bariatric surgery (k=7, 100%) performed assessments before and after the intervention $^{32, 33, 40, 41, 46, 49, 54}$; the other (14.3%, k=1) performed assessments before, during, and after the intervention. Among the 21 (100%) studies in which the exercise intervention was delivered after bariatric surgery, 33.3% (k=7), $^{30, 35, 38, 39, 42, 55, 57}$ 38.1% (k=8), $^{36, 37, 43, 45, 47, 48, 56}$ and 28.6% (k=6) $^{34, 50, 53, 58}$ performed baseline assessments before bariatric surgery, before the exercise intervention (i.e., after bariatric surgery), or at both these times, respectively. The number of assessments (excluding the follow-up phase) ranged from 2 to 9, with 17.9% (k=5) having \geq 3 assessments. $^{30, 34, 39, 47, 53}$

Feasibility and acceptability of the controlled trial methods

Data pertaining to the feasibility and acceptability of the controlled trial methods are presented Table 4. Among the 46.4% (k=13) of articles reporting recruitment duration,^{32, 33, 35, 36, 39, 41-43, 50, 53, 58} the median duration was 24 months (ranged from 6-56 months). Recruitment strategies were reported in most (60.7%, k=17) articles, ^{32, 33, 35, 37, 38, 40-49, 53, 54} among these, active, passive, or mixed approaches were used in 70.6% (k=12),^{32, 35, 37, 40, 42, 43, 46-49, 53, 54} 17.6% (k=3),^{38, 44, 45} and 11.8% (k=2)^{33, 41} studies, respectively. For studies

in which the exercise intervention was delivered after bariatric surgery, 66.7% (k=14) recruited participant before bariatric surgery. 30, 34, 35, 38, 39, 42, 47, 50-53, 55, 57, 58

The reasons for lost to follow-up for the exercise groups were: lack of time and commitment issues (21 participants from 6 articles), 35, 36, 39, 48, 56, 57 surgery-related issues (14 participants from 5 articles), 33, 34, 49, 51, 58 and unreachable (9 participants from 2 articles) (Table 4). 37, 39

Among the 28 articles, 17.9% (k=5)^{33, 39, 41, 51, 53} reported some missing data by outcomes and/or time-points, and 10.7% (k=3)^{36, 43, 50} described the method used to handle missing data.

Pooled data for refusal to participate, recruitment, enrollment, and retention rates, and enrollment speed are presented in Table 3 and Supplemental file 5. Related subgroup analyses are available in Table 3 and Supplemental files 6,7, and 8; no significant differences were found based on study quality, exercise intervention timing, and exercise intervention duration. Randomization refusal rate was not pooled due to the high number of studies without dropout because of dissatisfaction with group allocation (only 4 studies including 8 participants who refused randomization). ^{33, 43, 56, 58}

Studies quality

Study quality was rated as good, fair, and poor for 28.6% (k=8),^{33, 36-38, 43, 50, 51, 53} 35.7% (k=10),^{30, 32, 39, 40, 42, 44, 47, 49, 52, 56} and 35.7% (k=10) of articles,^{34, 35, 41, 45, 46, 48, 54, 55, 57, 58} respectively (Supplemental file 9). Informing these scores, is that most (67.9%, k=19) articles presented a randomized study,^{30, 32, 33, 35-40, 42, 43, 47, 49-51, 53, 55, 56, 58} 46.4% (k=13) reported a dropout rate <20%,^{30, 32, 33, 36-38, 42-44, 47, 50, 52, 53} and 53.6% (k=15) reported intention-to-treat analyses.^{33, 36-40, 43, 44, 46, 49-53, 56} A third or less of articles reported blinding

for outcomes $(10.7\%, k=3)^{32, 36, 49}$ and treatment assignment (0%, k=0), "high attendance rate" (i.e. participation to exercise training sessions $\geq 70\%$ or proportion of completers $\geq 70\%$) (k=10, 35.7%)^{30, 35, 41, 43-45, 47, 49, 51, 54}, and sample size justification (35.7%, k=10).^{32, 35, 39, 42, 43, 45, 47, 50, 51, 53}

Discussion

Summary of evidence

Systematic reviews have summarized evidence on the effects of exercise intervention before and after bariatric surgery. To complement these reviews and make recommendations for practice and future research, the present systematic review summarized evidence on the acceptability and feasibility of exercise interventions and controlled trial methods in adults awaiting or having undergone bariatric surgery.

Overall, available data from 28 articles reviewed show that exercise intervention compliance is very seldom reported (10.7%, k=3), the mean rate for exercise intervention attendance is 84% (based on data available from 28.6% (k=8) of articles), and the mean exercise intervention dropout rate is 5% (based on data available from 64.3% (k=18) of articles). In addition, most adverse events reported noted in the 57.1% (k=16) were relatively minor. Collectively, these indicators suggest that exercise interventions can be feasible and acceptable before and after bariatric surgery, but the relative lack of data available prevents us from making firm conclusions.

Although no consensus exists, attending ≥70% of sessions is often mentioned as "high" for exercise intervention.⁵⁹ With a pooled mean of 84% [95%CI: 77-91%] of sessions completed, the attendance rate calculated could be considered as high, which is important

because attendance rate can impact study results.^{59, 60} This rate is broadly similar to reported rates for people with type 2 diabetes (68-100%)⁶¹ and cancer (70-98%).⁶² However, these findings must be interpreted with caution. Session attendance was not reported in 39.3% (k=11) of included articles (see Table 2), which could overestimate the calculated pooled attendance rate mean, especially if studies with lower attendance were less likely to report these data. As well, session attendance was calculated/reported differently across articles (e.g., % of participants who attended all exercise sessions, % of participants who attended a certain number of exercise sessions). In addition, exercise intervention attendance rate, which relates to frequency, is only one aspect of exercise intervention adherence. Suppose two participants attended all scheduled sessions (100%) attendance), but only one completed the prescribed 30-minute aerobic training at moderate intensity. Focusing solely on attendance can be very misleading if suitable attention is not also given to compliance to the prescribed exercise dosage, which relates to duration and intensity.²³ Authors should give more attention to collecting and reporting data that reflect several key characteristics relating to exercise intervention adherence for transparency and help (potentially) explain lack of efficacity of exercise training. 60

Across 19 exercise groups, the exercise intervention dropout rate was 5%, which is lower than rates observed in adults with depression (15%),⁶³ type 2 diabetes (<20%),⁶¹ and advanced cancer (24%).⁶⁴ The lower rate herein could be explained by the population. Speculatively, adults awaiting or having undergone bariatric surgery may be highly motivated to exercise because of their fear of regaining weight and/or because they receive support via regular medical and nutritional follow-ups. The lower rate could also be explained by the small number of articles that reported this information (64.3%, k=18),

whereby those with higher dropout rate may not have reported these data. Last, it could be because we considered exercise intervention dropout only (not study or follow-up dropout). In other words, participants who completed the intervention, but did not come to the last assessment were not consider in the calculation of the exercise intervention dropout rate. Nevertheless, lack of time, work or other commitments, and lack of motivation were among the main reasons for exercise dropout among the 32.1% (k=9) articles reporting reasons when they had dropouts. These reasons reflect key barriers to physical activity in adults with obesity. and underscore the need to develop strategies to enhance motivation and perhaps offer alternative means for those lacking time or with competing commitments (e.g., online interventions) to reduce dropout and maximize intervention success.

Concerning feasibility and acceptability of controlled trial methods in adults awaiting or having undergone bariatric surgery, results suggest several months (median 24 months (range 6-56 months)) are required to recruit 6 to 220 participants (based on the 46.4% (k=13) articles reporting these data). With a pooled mean of 7 participants assessed for eligibility per month (23% of refusal to participate rate) and 2.5 participants enrolled per month, the pooled enrollment rate was 43%. Recruitment is one of the most challenging steps in research with clinical populations, requiring significant commitment from staff and resources to reach target sample size. As adverse scientific, economic, and ethical impacts can occur with prolonged or inefficient recruitment, and hiring of qualified staff should be considered to optimize recruitment. As well, recruitment strategies should be chosen carefully during the planning phase and reported during the dissemination phase to help others evaluate whether

strategies used were effective.²⁴ Based on available data reported, active recruitment strategies were used most (70.6%, k=12), followed by passive strategies alone (17.6%, k=3), and a mix of active and passive strategies (11.8%, k=2). Active methods of recruitment seem to result in better recruitment, population representativity, and retention than passive methods, but these are more expensive.⁶⁸ This said, the recruitment process was often poorly described or not described at all (39.3%, k=11), making it impossible to determine the most cost-effectiveness strategy/ies for recruiting adults awaiting or having undergone bariatric surgery into exercise trials. Regardless, integrating clinical staff can be helpful for recruitment within the scheduled time frame in bariatric surgery studies⁶⁹; so effective partnerships should be created.

Retention rates for the exercise intervention (87% [95%CI: 80-93%]; k=26) and comparison groups (87% [95%CI: 81-92%]; k=26), as well as randomisation refusal rate (only 8 participants across 4 studies) were good, suggesting the employed methods in the controlled trial methods were feasible and acceptable to participants. There is consensus that retention rates ≥80% are "acceptable" and that such rates help to minimize internal, external, and statistical validity threats. The good pooled retention rate could be explained by the short study duration for participants (median 18 weeks) and small number of assessments (2 or 3 assessments), which are known factors affecting retention. On the other hand, having supervised exercise sessions (89.3%, k=25) required frequent in-person visits, which could have affected enrollment in the first place (i.e., before the intervention/study has started), whereby those who foresaw barriers related to the intervention/study could have refused to participate, resulting in selection bias but good retention rates. This is a serious problem because many adults who might profit from an

exercise intervention may not be included in controlled trials. It is important that we know more about the causes of non-participation and study drop-out, regardless of rates, so as to enable reliable evaluations of exercise interventions and generalization of the results. Thus, authors should provide information about participants versus non-participants, as well as completers and non-completers. As for retention strategies, some guidance is available from the Longitudinal Assessment of Bariatric Surgery study⁷² and the Behavior Change Consortium.⁷³ Potential strategies include flexible schedule, incentives reimbursements, shortening visit length and number, coupling medical to research visits, and participant bonding or identification with the study, though it remains to be seen which strategy (or strategies) is most effective.

To explore factors influencing feasibility and acceptability indicators, as well as heterogeneity, subgroup analyses were performed; no significant differences were noted based on study quality, exercise intervention timing, and intervention duration. The absence of significant differences could be explained by the lack of statistical power herein since subgroup analyses require many more included studies than are needed for the main analyses of the meta-analysis.⁷⁴ The number of sub analysis initially planned (population, intervention and methodology-related characteristics) for this meta-analysis was greatly reduced due to the small number of studies and data available.

Strengths and limitations of this review

The systematic analysis of the peer-reviewed literature is the main strength of this review. Indeed, the articles identified were the result of an extensive search in different databases, and screening and data extraction were conducted by multiple reviewers to minimize bias.

However, some limitations should be considered when interpreting the results. First, this review contains articles stemming from controlled trials published in peer-reviewed journals with full-texts available in English only. Second, the characteristics of the included articles limits the ability to generalize results. For example, 68% (k=19) of studies were completed in higher income countries, and sociodemographic data were missing in 68% (k=19) of articles. Moreover, men were largely underrepresented. Third, studies were not necessarily designed to explore the feasibility or acceptability of an exercise intervention and/or controlled trial methods, limiting collection of (or reporting) of some data relating to feasibility and acceptability (e.g. satisfaction rate). Fourth, data extraction and interpretation were challenging due to the absence of clear definitions and a lack of consensus on the concepts of "feasibility" and "acceptability," though the CONSORT flow chart presented in most (60.7%, k=17) articles aided. Finally, publication bias could also affect the findings.

Recommendations for exercise intervention implementation, research, and reporting

When interpreting exercise intervention adherence and dropout rates in bariatric surgery population, persons planning to implement exercise interventions should keep in mind the impact of selection bias across included studies due to voluntary basis recruitment and specific selection criteria leading to probable over estimation of the adherence rate and under estimation of the dropout rate.⁵⁹ Based on the voluntary basis, and selection criteria of some studies, it is likely that adults who had a higher disease burden, poorer functional status and motivation were not included, providing only a partial view of what happens in real life conditions; these trials may not address the feasibility and acceptability of exercise

for a subgroup of adults awaiting or having undergone bariatric surgery. Researchers are encouraged to include these adults to examine the effects (as well as the adverse effects) that may result. Relatedly, the lack of data on non-participants raises the possibility that issues of feasibility and acceptability may come about even before participants enroll into trials. Thus, researchers will need to consider how possible differences are likely to impact on the results, as this will influence the certainty of evidence. In addition, the acceptability and feasibility of exercise training in people awaiting or having undergone bariatric surgery may be affected by type of exercise; yet, there was limited variety across studies. Many studies focused on walk, treadmill, or ergocycle (for endurance/aerobic training) or machines or small equipment (for resistance training). Only one study focused on running after bariatric surgery⁴⁸ and another on aquatic training before bariatric surgery⁴⁰. Further investigation into the feasibility and acceptability of different type of exercise is warranted, and whether it matters if it is the researchers who chose the type of exercise or the participants. Because patient-oriented strategies can increase participation and adherence as patients feel included in their decisions, 75-78 and because adults mostly get to choose their own type of exercise in real life, examining whether integrating participants' preferences into interventions improves feasibility and acceptability seems valuable. Also, to reach larger proportion of bariatric surgery patients, several modalities of intervention (e.g. in person, telehealth, individual or group) and setting (hospital, community) should be considered for implementation. Last, based on identified gaps, future studies are necessary to draw conclusions regarding the feasibility and acceptability of multicentric study, long-term data collection (follow-up period after the intervention), exercise intervention in male and visible minorities.

As more research is conducted, the most critical points to address in included articles are the clarity and transparency in reporting feasibility and acceptability data. The major recommendation coming out of this review is that future studies should report consistently these data to better identify which exercise intervention offer the most promising results in terms of efficacy and implementation. The systematic use of the PRISMA flow chart could help in this direction. To increase transparency, researchers should too describe missing data by outcome, by time-point and by group (number and reason), and how missing data were handled. Also, previous and future trials should systematically share their raw data. It facilitates the development of individual patient data meta-analysis (in exercise, e.g., see 10%). This method is particularly recommended to study moderators of intervention effects. For instance, it allows to disentangle subject-level and study-level sources of heterogeneity in adherence rates. Better reports of feasibility and acceptability data could also help future meta-analysis to identify factors associated with better adherence, recruitment, enrollment and retention.

Conclusion

Despite the present lack of data available in controlled trials included, exercise intervention and controlled trial methods in adults awaiting or having undergone bariatric surgery seem to be feasible and acceptable, though there is room for improvement. To improve evidence-based knowledge, identify methodological and practical challenges and bias, better reporting of adherence, dropout, adverse events, recruitment, enrollment, retention data, and strategies used are needed in future controlled trials. This should help implement exercise training protocols in real life conditions that will be specifically adapted to clinical care in the setting of bariatric surgery.

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Table 1. Feasibility and acceptability data analysis

Table 2. Description, feasibility and acceptability of interventions delivered before and after bariatric surgery among studies included (k=28)

Data are presented as mean \pm SD or median (25th-75th percentiles)

ACT = aerobic continuous training; BS = bariatric surgery; HIIT = high-intensity interval training; HR = heart rate; HRR = heart rate reserve; PA = physical activity; NA = not available; OMINI-RES = OMNIResistance Exercise Scale; reps = repetitions; RPE = rating of perceived exertion; RPM = one repetition maximum; VAT-RCP = difference between ventilator anaerobic threshold and respiratory compensation point; wk.= week(s)

*Session duration included warm-up and cool down duration; **Drop out rate = % of participants who discontinued intervention

Table 3. Pooled percentage of feasibility and acceptability indicators of exercise and controlled trial methods in adults awaiting or having undergone bariatric surgery

Study quality and intervention duration subgroup analyses are based on median; Studies considered high quality = score >8/14 and low quality = score \le 8/14; Longer intervention duration = >12 weeks excluding studies with >100 weeks of intervention duration and shorter duration \le 12 weeks. BS = bariatric surgery. Randomization refusal rate (%) was not pooled due to the high number of 0 among studies (only 4 studies including 8 participants who refused randomization).

Table 4. Description, feasibility and acceptability of controlled trial methods in bariatric surgery population among studies included (k=28)

Supplemental file 1. Prisma 2020 checklist

Supplemental file 2. Study details, participants' characteristics, exercise intervention and comparison group(s), and outcomes extracted and entered into a Microsoft Excel table

Supplemental file 3. PRISMA flow diagram

^{*=} except sociodemographic data; BS = bariatric surgery; CBT = cognitive-behavioral therapy; QE = quasi-experimental study; NA = not available; MMTT = Mixed meal tolerance test; PA = physical activity; REE = Resting energy expenditure; RCT = randomized controlled study. S = outcomes self-reported; O = outcomes objectively assessed.

Supplemental file 4. Participant characteristics of controlled trials included (k=28)

Data are presented as mean \pm SD or median (25th-75th percentiles) or mean [range].

12MWRT = 12-minutes walk-run test; 6MWT = 6-minutes walking test; BPD-DS = biliopancreatic diversion with duodenal switch; BMI = body mass index; BS = bariatric surgery; CBT = cognitive-behavioral therapy; FFM = fat free mass; GB = gastric banding; IPAQ-SF = International PA Questionnaire Short Form; HT = hypertension; N = baseline number of participants after randomization; N* = final number of participants (for studies without sociodemographic data for the baseline number of participants after randomization); NA = not available; PA = physical activity; QE = quasi experimental study; RCT = randomized controlled study; RYGB = Roux-Y gastric bypass; SG = sleeve gastrectomy; T2D = type 2 diabetes; y = years.

Supplemental file 5. Forest plots of feasibility and acceptability indicators of exercise and controlled trial methods in adults awaiting or having undergone bariatric surgery

Supplemental file 6. Forest plots of feasibility and acceptability indicators of exercise and controlled trial methods in adults awaiting or having undergone bariatric surgery according to studies quality

Supplemental file 7. Forest plots of feasibility and acceptability indicators of exercise and controlled trial methods in adults awaiting or having undergone bariatric surgery according to exercise training timing

Supplemental file 8. Forest plots of feasibility and acceptability indicators of exercise and controlled trial methods in adults awaiting or having undergone bariatric surgery according to intervention duration

Supplemental file 9. Summary of quality assessment of controlled trials included (k=28)

Criteria for controlled trials: (1) Randomized study; (2) Adequate randomization method; (3) Treatment allocation concealment; (4) Blinding treatment assignment; (5) Blinding outcome assessors; (6) Similar baseline characteristics; (7) Dropout rate <20%; (8) Differential dropout rate between groups <15%; (9) High attendance (i.e. participation to exercise training sessions ≥70% or proportion of completers ≥70 %); (10) Similar background treatments; (11) Valid and reliable outcome measures; (12) Sample size justification; (13) Pre-specified outcomes/subgroups; (14) All randomized participants analyzed (ITT analysis). Three criteria were defined as "fatal flaws" when not met: (1) Randomized study; (7) Dropout rate <20%; (14) Intention-to-Treat analysis). Study quality was defined as good, fair and poor when 0, 1 or ≥2 fatal flaws were identified.

^{*} The reported measurements were taken before BS, because there was no measurement at the beginning of the intervention after BS.

^{**}To note, the ethnicity terms used in this table was those reported by included studies, and control group referred to comparison group without exercise intervention.

Table 1.

Name	Calculation					
Feasibility and acceptability of the exercise intervention						
Attendance rate (relates to session frequency), % (or otherwise)	Not calculated; reported by authors					
Compliance rate (relates to session duration and intensity), % (or otherwise)	Not calculated; reported by authors					
Dropout rate, % = 100 / number of participants allocated to the intervention x number of participants who discont intervention						
Feasibility and acceptability of control	lled trial methods					
Number of participants enrolled, N	= number of participants assessed for eligibility (i.e., recruited) - number of participants excluded					
Refusal rate, %	= 100 / (number of participants enrolled + number of participants who declined to participate) x number of participants who declined to participate					
Recruitment rate, n/month	= number of participants assessed for eligibility /recruitment duration (months)					
Enrollment speed, n/month	= number of participants enrolled / recruitment duration (months)					
Enrollment rate, %	= 100 / number of participants assessed for eligibility x number of participants enrolled					
Retention rate, %	=100/ number of participants allocated to intervention x (number of participants allocated to intervention - number of participants lost to follow-up during the intervention phase)					
	For follow-up phase: = 100 / number of participants at the beginning of follow-up x (number of participants at the beginning of follow-up phase)					
Randomisation refusal rate, %	= 100 / number of participants randomized x number of participants who refused randomization					

Author Year (Country) Brief intervention name	Interventi on starting time	Intervention length Setting	Intervention supervision	Intervent- ion format	Intervention description Exercise type and volume*	Attendance and compliance rate	Dropout rate** Reasons	Adverse events during exercise		
Studies with intervention delivered before BS										
Funderburk 2010 (USA) Aquatic exercise intervention	before BS	13 wk. Rehabilitation center	Supervised by aquatic exercise leaders	NA	Aquatic strength and endurance exercises; Ai Chi exercises for balance, core strengthening and relaxation: 2 x 60 min/wk. + Usual care	Attendance: NA Compliance: NA	NA	NA		
Usual care	before BS	Hospital	Physicians	NA	Proceeding with gastric bypass surgery preparation routine.	Attendance: NA Compliance: NA	NA			
Baillot 2016 (Canada) Endurance and resistance training	3-6 months before BS	Gym in an hospital (possibility sometimes at home)	Supervised by PA specialist + multidisciplinar y team	Group (possibility sometimes individually)	Treadmill, walking circuit, arm-ergocycle, elliptical: 3 x 60 min/wk. (55-85% HRR) Resistance: 2-3 series of 12-15 reps 9 exercises with small equipment (20 min) + Usual care	Attendance: 68.4 (52.8-92.1) % of the total exercise sessions Compliance: NA	6.7% - had stomach pain followed by transportation problem	Occasional muscle or joint pain 3 hypoglycemia (one patient) during exercise sessions		
Usual care	3-6 months before BS	Hospital	Multidisciplinar y team	Group + individual	15 min PA and nutrition counselling meeting every 6 wk. for the first 6 months, then every 8 wk. until the surgery + voluntary group meeting	Attendance: NA Compliance: NA	0%			
Marcon 2017 (Brazil) Walk-simulating endurance training + lifestyle modification	before BS	19 wk. Outpatient clinic	Supervised by NA	NA	Walking-simulating leg and bilateral arm movement: 2 x 25 min/wk. (2-4 Borg RPE) CBT group therapy: 60 min/wk. + Usual care	Attendance: 78.5 ±13.3 % of the total exercise sessions Compliance: NA	22.7% - I surgery was anticipated - 4 were unable to attend intervention	NA		
Walk-simulating endurance training	before BS	Outpatient clinic	Supervised by NA	NA	Walking-simulating leg and bilateral arm movement: 2 x 25 min/wk. (2-4 Borg RPE) + Usual care	Attendance: 80.9 ± 10.8 % of the total exercise sessions Compliance: NA	0%			

Usual care	before BS	Outpatient clinic	Multidisciplinar y team	Individual	Mandatory presence information meetings about the importance of changing eating habits and PA: 5 x 90 min.	Attendance: 100.0 % of the information meetings	0%	
						Compliance: NA		
Picó-Sirvent 2019	before BS	26 wk.	Supervised by	NA	2-4 x 60 to 70 min/wk. Treadmill, ergocycle: 1-2x MICT (60-85%	Attendance: 93.3 ± 6.5 % of	0%	None
(Spain)	ocioic B3	University	sport science	NA	$HRpeak$) + 2 x HIIT (60-95% VO_2peak) / wk.	the total exercise	070	None
MICT-/HIIT-		sport facilities	graduates		Resistance: 2x/wk. 1-4 series of 10-20 reps 4-7 exercises	sessions		
endurance and resistance training					+ Usual care	Compliance: NA		
						Attendance: NA	001	
Usual care	before BS	Hospital	NA	NA	Usual presurgical care indication.	Compliance: NA	0%	
Marc-Hernández 2019	2.6	12 wk.	G : 11	NIA	Ergoycle, arm ergometer, elliptical, and	Attendance: NA	16.70/	NIA
(Spain)	3-6 months	Research	Supervised by NA	NA	treadmill: 2 x 35-50 min continuous/wk. (60-80% HRmax) + 2 x HIIT/wk. (60-80%		16.7%	NA
	before BS	center			VO ₂ max)	Compliance: NA	- 1 left the program	
(HIIT-)Endurance and resistance training					Resistance: 1-4 series of 15-20 reps 4-7 exercises with machines (50-65% RPM) + Usual care			
						Attendance: NA	Not clear	
Usual care	3-6 months before BS	Hospital	Multidisciplinar y team	Individual	Psychological and nutritional counseling. Active lifestyle advice.	Compliance: NA		
Gilbertson 2020	1	4 wk.	Semi-	T., 431 d1	W-11-in 5 20-in/1- ((5 950/ HD)	Attendance: 18.4 ± 3.0 of 20	0%	None
(USA)	1 month before BS	At home	supervised by the research	Individual	Walking: 5 x 30min/wk. (65-85% HRmax) + Usual care	18.4 ± 3.0 01 20 scheduled		
Home walking			team (Use of fitness and activity tracker Weekly texts, emails or			exercise sessions		
			phone calls)			Compliance: NA Attendance: NA		
Usual care	1 month	Medical	Multidisciplinar	Individual	Nutritional instructions and meal replacement		0%	
	before BS	center	y team		shakes.	Compliance: NA		

Arman 2021 (Turkey) Core Stabilization Exercise Program	3-6 months before BS	8 wk. Facility based	Supervised by physiotherapists	Individual	2 x 50-70 min/wk Warm-up: treadmill 10-15 min 50-60%HRmax Core Stabilization Exercise Program (strengthening, endurance and balance exercises) 1-2 series of 7-10 reps 7-12 exercises (50% RPM) + Usual care	Attendance: NA Compliance: NA	0%	NA
Usual care	3-6 months before BS	NA	Supervised by physiotherapists	Individual	PA and sedentary advices and daily PA diary with activities and number of steps. (motivational phone call every 2–3 wk.)	Attendance: NA Compliance: NA	Not clear	
Studies with interv	ention delive	ered after BS						
Castello 2011 (Brazil) Interval endurance training	1 month after BS	12 wk. University laboratory	Supervised by physiotherapist	Individual	Treadmill: 3 x 60min/wk. (50-70% HRmax)	Attendance: 100.0 % of 32 exercise sessions Compliance: NA	31.3% - 3 had trouble balancing work and training - 2 disliked exercise due to muscle or joint pain	Lower limb fatigue, mild dyspnea and mild sweating during exercise sessions
NA	NA	NA	NA	NA	NA	Attendance: NA Compliance: NA	0%	
Shah 2011 (USA) High-volume endurance training + diet intervention	≥ 3 months after BS	12 wk. Gym in a medical center (max. 80% at home)	Semi- supervised by investigator	Individual	Treadmill/walking and ergocycle/rowing: 5 x/wk. (60-70 VO ₂ max) Goal: expend ≥ 2000 kcal/wk. Exercise and diet-related behavioral therapy every two wk	Attendance: NA Compliance: 30.0-53.0 % reached the goal	23.8% - 5 did not have enough time to exercise	Occasional muscle or joint soreness
Diet intervention	≥ 3 months after BS	Research center or by telephone	Investigator	Individual	Behavioral diet-related intervention and feedback on diet every two wk.	Attendance: NA Compliance: NA	33.3 % - 3 preferred to be in the exercise group - 1 had no time	

Stegen 2011 (Belgium) Progressive endurance and resistance training	1 month after BS	12 wk. University rehabilitation facilities	Supervised by movement and rehabilitation sciences master students	Individual	Cycling, walking and stepping: 3 x 75 min/wk. (60-75% HRR) Resistance: 1-3 series of 10-15 reps of knee/elbow flexion/extension exercises using stackweight equipment (60-75% RPM, 25 min)	Attendance: NA Compliance: NA	Not clear	NA
NA	1 month after BS	NA	NA	NA	NA	Attendance: NA Compliance: NA	Not clear	
Coen 2015 (USA) Endurance training	1-3 months after BS	26 wk. Research center or at home or outdoors	Semi- supervised by trained exercise physiologists (at least 1 session/wk.)	Individual	Ergocycle, treadmill, cycling and walking outdoors: 3-5 x 10-30 min/wk. (60-70% HRmax); Goal: expand ≥ 120 min/wk. + Health education	Attendance: NA Compliance: 66.7 % reached the goal	9.1% - 5 had time- commitment issues - 1 was lost to follow-up	None
Health education	1-3 months after BS	NA	NA	Group	Monthly health education sessions (medication, nutrition, PA). PA habit reports.	Attendance: NA Compliance: 90.3 % reported > 30 min PA/wk.	4.8% - 1 moved away - 1 was lost to follow-up - 1 pregnancy	
Huck 2015 (USA) Multifaceted resistance training	1-12 months after BS	12 wk. Community-based training facility	Supervised by certified strength and conditioning specialist	Group	Resistance: 1-3 series of 8-12 reps 8-10 exercises with stack-weight equipment, free weights, resistive bands and body weight. 2-3 x 60 min/wk. (60-75% RPM) + Usual care	Attendance: 84.0 % of training sessions Compliance: NA	Not clear	None
Usual care	1-12 months after BS	NA	Clinicians	NA	Encouragement to increase PA and consume protein shakes.	Attendance: NA Compliance: NA	Not clear	
Marchesi 2015 (Italy) Road running training	12-36 months after BS	44 wk. NA	Supervised by sport physician (1 self-managed session/wk in the last 20 wk.)	NA	Indoor or outdoor road running: 3-4 x 60 min/wk.; goal: participation in a 10.5 km competitive run (55-90% HRmax) Physiological assistance and specialized nutritional counseling.	Attendance: NA Compliance: NA	30.0% - 3 lacked motivation	Symptomatic cholelithiasis (resolved after 3-wk break)
NA	12-13 months	NA	NA	NA	NA	Attendance: NA	0%	

	after BS					Compliance: NA		
Muschitz 2016 (Austria) Endurance and resistance training + specific supplementation	2 wk. after BS	104 wk. At home	Semi- supervised Monitored by physical medicine experts	Individual	Nordic walking: at least 3 x 45 min/wk. Resistance perseverance and equipment training: at least 2 x 30 min/wk. Specific vitamin, mineral and protein supplementation before BS and after BS. + Usual care	Attendance: 88.2 % completed ≥ 80 % exercise requirements Compliance: NA	Not clear	NA
Usual care	2 wk. after BS	NA	Telephone interviews	NA	Advice on PA and balanced nutrition. Basic supplementation.	Attendance: NA Compliance: NA	14.5% - 12 were unwilling to stay in the study - 4 got new jobs outside the area	
Rojhani-Shirazi 2016 (Iran)	5 days after BS	4 wk.	NA	NA	Balance training with periodic exercises (e.g., standing one leg or walking on toes): 4 x 45 min/wk.	Attendance: NA Compliance: NA	NA	NA
NA	after BS	NA	NA	NA	NA	Attendance: NA Compliance: NA	NA	
Campanha- Versiani 2017 (Brazil) Endurance and resistance training	3 months after BS	36 wk.	Supervised by NA	NA	Treadmill: 2 x 25 min/wk. (70-80% HRmax) Resistance: 1-3 series of 10-12 reps at 10RM maximum load 8 exercises (2 x 60 min/wk.) + Usual care	Attendance: 60.0 % attended ≥ 95% scheduled exercise sessions Compliance: NA	Not clear	NA
Usual care	3 months after BS	NA	NA	NA	Dietary counseling. Multivitamin and mineral supplementation.	Attendance: NA Compliance: NA	Not clear	

Coleman 2017 (USA) Functional resistance training	6-24 months after BS	52 wk.	Semisupervised by NA (60% self-directed exercise)	Group + Individual	Functional resistance, aerobic and flexibility training with body weight: 2 x 60 min/wk. Self-directed exercise 3 x /wk. Goal: MVPA ≥ 150 min/wk. Weekly phone counseling. Step counting. + maintenance phase with booster exercise sessions and social support	Attendance: NA 32.0-56.0 % of the supervised exercise sessions Compliance: NA Attendance: NA	7.7% - 2 lacked of time	- 44% with initial functionality developed condition during intervention
Usual care	months after BS	Hospital	Nurse care managers	Individual	Dietary and PA counseling 2 wk., 2 months and 6 months after surgery.	Compliance: NA		
Herring 2017 (UK) Endurance and resistance training	12-24 months after BS	12 wk. Outpatient clinic	Supervised by qualified gym instructor	Individual	3 x 60 min/wk., Endurance: 64-77% HRmax, 12-14 Borg RPE Resistance: 3 series of 12 reps 2 exercises (60% RPM, 25 min) Diet information	Attendance: 95.0 % of 36 scheduled exercise sessions Compliance: NA	8.3% - 1 had gastric band deflated	None
Usual care	12-24 months after BS	NA	NA	Individual	Advice session of 30-60 min on PA upon study discharge and diet information.	Attendance: NA Compliance: NA	16.7% - 2 preferred to be in the exercise group	
Onofre 2017 (Brazil) Endurance and resistance training	3 months after BS	12 wk. University hospital	Supervised by physiotherapist	NA	Treadmill: 3 x 60 min/wk. (40–60% HRR and 85–90% HRR) Resistance: upper and lower limp exercises (60-80% RPM, 25 min)	Attendance: NA Compliance: NA	0%	None
Usual care	3 months after BS	NA	NA	NA	General guidelines for the importance of PA	Attendance: NA Compliance: NA	0%	
Daniels 2017 (USA) Progressive resistance training	2 months after BS	12 wk. NA	NA	NA	Resistance: 1-4 series of 8-15 reps on 8-10 exercises, 3 x 60-80 min/wk. (50 - >80% RPM)	Attendance: NA Compliance: NA	0%	NA
Control	2 months after BS	NA	NA	NA	Instruction to continue normal daily activities.	Attendance: NA Compliance: NA	0%	

Hassannejad 2017 (Iran)	Just after BS	12 wk. At home	No supervision	Individual	Resistance: Shoulder and hip exercises with elastic bands, 3 x 20-30 min/wk.	Attendance: NA Compliance: NA	Not clear	NA
Endurance training + resistance training			Daily activity log books		Walking: 3-5 x 20-30 min/wk. (12-14 Borg RPE)			
Endurance training	Just after BS	At home	No supervision Daily activity log books	Individual	Walking: 3-5 x 20-30 min/wk. (12-14 Borg RPE)	Attendance: NA Compliance: NA	Not clear	NA
Diet education	Just after BS	NA	Nutritionist	Individual	Education about standard high-protein diet after BS.	Attendance: NA Compliance: NA	Not clear	
Mundbjerg 2018 (Denmark) Endurance and resistance physical	6 months after BS	26 wk. Fitness center	Supervised by physiologists (additional individual sessions)	NA	2 x 40 min/wk Ergoycle, stair climbing, treadmill, rowing (2x30 min, 50-70% VO ₂ max / 15-17 Borg RPE) Resistance, unconsisting (2x10 min)	Attendance: 59.4 % attended ≥ 50% scheduled exercise sessions	Not clear	NA
training					Resistance: upper extremities (2x10 min) goal: PA ≥ 210 min/wk. + Usual care	Compliance: NA		
Usual care	6 months after BS	NA	NA	NA	Standard dietary recommendations.	Attendance: NA Compliance: NA	Not clear	
Oppert 2018 (France) Resistance training + protein	6 wk. after BS	18 wk. University hospital	Supervised by qualified trainers	Group	Resistance: 4 series of 8-12 reps 6 exercises, 3 x 60 min/wk. (50-75% RPM) Whey protein supplementation, 2 x per day + Usual care	Attendance: 47.8 % completed ≥ 2 exercise sessions/wk. and nutritional requirements	0%	None
Usual care + protein	1 wk. after BS	NA	No supervision	Individual	Whey protein supplementation, 2 x per day. + Usual care	Attendance: NA Attendance: 45.2 % completed all visits and nutritional requirements Compliance: NA	0%	
I	•	•						

Usual care	Before BS	NA	NA	Individual	Nutrition and PA advice. Prescription of iron, multivitamin and mineral supplementation from 15 days before BS.	Attendance: 100.0 % of scheduled visits Compliance: NA	0%	
Murai 2019 (Brazil) Endurance and resistance training	3 months after BS	26 wk. Hospital	Supervised by NA	Individual	Treadmill: 3 x 60-90 min/wk. (50% VAT-RCP) Resistance: 3 series of 8-12 reps on 7 exercises + Health education	Attendance: NA 81.5 ± 13.1 % of total exercise sessions Compliance: NA	0%	None
Health education	3 months after BS	NA	NA	NA	Advice on calcium, vitamin D3 and protein intake. Encouraged to improve PA levels during follow-up	Attendance: NA Compliance: NA	0%	
Diniz-Souza 2020 (Portugal) Multicomponent resistance training +Usual care	1 month after BS	47 wk. Facility based	Supervised by exercise trainer	Group	High-impact training: High ground-reaction force exercises (183-209 gravitational loading peaks > 4.9 g, 20 min) Resistance training: 2-3 series of 4-12 reps 7-8 exercises (65-85% RPM, 35 min) 3 x 75 min/wk. + Usual care	Attendance: 39.0 % attended ≥ 50% scheduled exercise sessions Compliance: NA	Not clear	- 10 required ad hoc exercise program adjustment (4 knee pain, 2 lower back pain, 1 diffuse lower limbs pain) and temporarily interrupted the intervention (1 rolled ankle and 1 hand bruise outside the study, 1 back bruise after fall in exercise session)
Usual care	1 month after BS	NA	NA	Individual	Prescription of proton-pump inhibitors and structured PA. Supplementation advice.	Attendance: NA Compliance: NA	Not clear	
Tardif 2020 (Canada) Moderate endurance and resistance training	3 months after BS	12 wk. Medical center	Supervised by NA	NA	Moderate endurance training (50-75% HRR) and resistance training (25min): 3 x 60 min/wk.	Attendance: 28.0 % attended 85% scheduled exercise sessions Compliance: 100%	0%	11% hypotension
NA	3 months after BS	NA	NA	NA	NA	Attendance: NA Compliance: 100%	0%	

Marc-Hernández 2020 (Spain) ACT- and HIIT- endurance and resistance training	37 months after BS	20 wk. Sport Research Center	Supervised by sports science graduates	NA	Ergocycle, elliptical, treadmill: ACT (60-85% HRmax,) and HIIT (60-95% VO ₂ max), 2-4 x 50 min/wk. Resistance: 1-4 series of 4-7 reps 4-7 exercises (50-75% RPM, 8-28 min)	Attendance: 90.9 % attended ≥ 85% scheduled exercise sessions Compliance: NA	0%	NA
Usual care	37 months after BS	Hospital	NA	NA	NA	Attendance: NA Compliance: NA	20.0% - 2 refused to continue	
Lamarca 2021 (Brazil) Resistance training + protein	2-7 years after BS	12 wk. Facility based	Supervised by qualified professionals	NA	Resistance: 3 series of 8-12 reps on 8 exercises, 3 x 80 min/wk. (6-9 on OMINI-RES) Whey protein supplementation and general training on healthy eating.	Attendance: 80.0 ±7.7% of scheduled exercise sessions Compliance: NA	Not clear	None
Resistance training + placebo	2-7 years after BS	Facility based	Supervised by qualified professionals	NA	Resistance: 3 series of 8-12 reps on 8 exercises, 3 x 80 min/wk. (6-9 on OMINI-RES) Placebo Maltodextrin and general training on healthy eating	Attendance: 84.8 ± 5.4 % of scheduled exercise sessions Compliance: NA	Not clear	NA
Protein	2-7 years after BS	At home	No supervision	Individual	Whey protein supplementation and general training on healthy eating.	Attendance: NA Compliance: NA	Not clear	
Placebo	2-7 years after BS	At home	No supervision	Individual	Placebo Maltodextrin and general training on healthy eating	Attendance: NA Compliance: NA	Not clear	

Table 3.

Feasibility and acceptability data	•				
	Arm numbers	%	95	% CI	I^2
Total attendance rate – exercise intervention	10	84.3	77.0	90.7	0.0
Study quality: High	6	83.9	70.6	94.1	9.4
Low	4	84.2	80.1	88.0	0.0
Exercise intervention timing: Before BS	4	79.4	67.7	89.4	0.0
After BS	6	87.4	76.7	95.6	0.0
Exercise intervention duration: Longer	4	82.2	78.5	85.6	0.0
Shorter	6	86.8	71.4	97.6	27.0
Total compliance rate – exercise intervention	3	No met	a-analy	sis perfo	rmed
Total dropout rate (%) – exercise intervention	19	5.0	1.1	10.5	60.1
Study quality: High	11	5.1	0.6	12.4	61.6
Low	8	5.5	0.0	18.4	58.7
Exercise intervention timing: Before BS	7	3.6	0.0	14.6	44.4
After BS	12	5.6	0.6	13.8	68.5
Exercise intervention duration: Longer	9	3.7	0.0	12.6	64.7
Shorter	10	6.2	0.5	15.8	56.3
Total dropout rate (%) control/comparison groups	18	1.6	0.0	5.3	53.2
Study quality: High	11	2.2	0.0	7.1	60.5
Low	7	1.9	0.0	13.3	41.4
Exercise intervention timing: Before BS	4	0.0	0.0	0.3	0.0
After BS	14	2.8	0.0	8.2	62.3
Exercise intervention duration: Longer	9	0.0	0.0	2.4	6.7
Shorter	8	2.9	0.0	13.1	48.3
Feasibility and acceptability of controlled trial methods					
Total refusal rate (%)	16	22.6	10.0	38.2	94.6
Study quality: High	10	22.7	10.4	38.0	89.5
Low	6	22.9	0.0	69.2	97.5
Exercise intervention timing: Before BS	4	30.7	0.0	81.0	91.7
After BS	12	20.3	6.5	38.7	95.3
Exercise intervention duration: Longer	7	27.2	3.6	60.3	96.8
Shorter	9	18.9	4.7	38.7	91.0
Total recruitment rate (n/month)	12	7.1	3.8	11.3	60.1
Study quality: High	8	6.3	3.8	9.2	0.0
Low	4	10.2	0.1	29.4	81.3
Exercise intervention timing: Before BS	3	8.1	0.7	20.2	0.0
After BS	9	7.0	2.9	12.4	70.4
Exercise intervention duration: Longer	4	7.9	0.0	25.6	85.3
Shorter	8	6.5	3.3	10.5	15.7
Total enrollment speed (n/month)	13	2.5	1.4	3.7	0.0
Study quality: High	9	2.5	1.3	4.0	0.0
Low	4	2.8	0.3	6.8	0.0

Exercise intervention timing: Before BS	3	1.1	0.0	6.2	0.0
After BS	10	2.9	1.6	4.4	0.0
Exercise intervention duration: Longer	5	1.6	0.6	2.9	0.0
Shorter	8	3.3	1.4	5.6	0.0
Total enrollment rate (%)	18	43.3	30.0	57.2	93.6
Study quality: High	12	40.0	26.1	54.7	90.7
Low	6	50.1	14.0	86.1	96.4
Exercise intervention timing: Before BS	4	17.4	0.4	46.5	74.7
After BS	14	50.8	36.4	65.2	93.1
Exercise intervention duration: Longer	9	38.1	17.1	61.4	95.7
Shorter	9	49.0	29.7	68.5	89.1
Total retention rate (%) – exercise intervention	26	87.1	79.6	93.3	79.5
Study quality: High	13	92.9	84.4	98.6	78.7
Low	13	77.2	65.7	87.3	61.3
Exercise intervention timing: Before BS	7	96.4	84.6	100	49.5
After BS	19	83.6	74.1	91.5	82.8
Exercise intervention duration: Longer	12	86.8	75.5	95.5	77.1
Shorter	13	84.3	72.1	93.9	70.8
Total retention rate (%) – control/comparison groups	26	86.8	80.6	92.1	65.5
Study quality: High	13	88.9	81.2	95.1	67.7
Low	13	82.8	71.7	92.0	50.7
Exercise intervention timing: Before BS	6	89.5	77.8	97.8	0.0
After BS	20	85.7	78.1	92.1	72.9
Exercise intervention duration: Longer	12	90.4	62.8	95.7	75.1
Shorter	13	81.7	72.5	89.6	32.0

Author Year (Country) Study design	Study duration for one participant Timing of assessment and indenisation	Outcomes assessed*	Recruitment duration, strategies and timing	Recruit ment rate (number/ mo.)	Refusal rate (%) Reasons to decline to participate	Enrollment speed (participant number/mo.) Enrollment rate (%)	Retention rate (%) Reasons for the lost of follow-up	Randomization refusal rate (%) Reasons to decline randomization
Studies with in	tervention delivere	d before BS						
Funderburk 2010 (USA) Pilot RCT	~13 wk. Pre and post intervention NA	 Weight^O Quality of life^S Psychological distress^S Depression^S Blood pressure^O Physical fitness^O 	NA Active Before BS	NA	NA	NA	NA	NA
Baillot 2016 (Canada) RCT	~12 wk.+ 52 wk. f-up Pre and post intervention + 4x during f-up No	 Anthropometry^O Body composition^O Blood pressure^O Physical fitness^O PAS Quality of life^S Satisfaction^S 	23 months Active and passive Before BS	6.1	18.9 % Reasons: No research interest (n=5) Did not like group exercise (n=2)	1.3 21.4%	For intervention part: Exercise group:100% Control group: 93% Reason: Disagree with her allocation (n=1) For f-up part: Exercise group: 93% Reasons: No surgery (cancer) (n=1) Control group: 93% Reasons: Abandonment (n=1)	Exercise group: 0% Control group: 6.7% Reason: Want to be in exercise group (n=1)
Marcon 2017 (Brazil) RCT	~19 wk. Pre and post intervention NA	1. Anthropometry ^O 2. Functional capacity ^O 3. Physical fitness ^O 4. Cardiovascular risk ^S 5. Lipid profile ^O	NA Active Before BS	NA	7.0 % Reason: NA	NA 13.4%	Exercise group: 100% Exercise group + CBT: 77% Reasons: Surgery was anticipated (n=1) Unable to attend intervention (n=1) Control group: 82% Reasons: Changed address (n=1) Did not attend the assessment (n=3)	Exercise group: 0% Exercise group + CBT: 0% Control group: 0%

Picó-Sirvent	~ 26 wk.	1. Anthrometry ^O	NA					
2019		2. Body composition ^O	1111				Exercise group: 100%	
(Spain)	Pre, mid and	3. Cardiometabolic risk	Active	NA	NA	NA	C 4 1 1000/	Not applicable
(1)	post intervention	factors ^O					Control group: 100%	••
Pilot QE	NA	4.Physical fitness ^O	Before BS					
Marc-		1. Anthopometry ^O					E : 020/	
Hernández	~12 wk.	2. Body composition ^O	NA				Exercise group: 83%	
2019		3. REE ^O			NA	NA	Reasons: Left the program (n=1)	
(Spain)	Pre and post	4. Cardiometabolic risk	Active	NA	Reason: NA		Experienced physical problems (n=1)	Not applicable
	intervention	factors ^O					Control group: 73%	
QE	NA	5. Physical fitness ^O	Before BS			NA	Reason: NA	
		6. Quality of life ^S						
		 Body composition^O Physical fitness^O 	38 months					
Gilbertson	~ 4 wk.	3. Food diary ^S	38 monuis				Exercise group: 90%	
2020		4. MMTT ^O	Active and		73.0%	0.4	Reason:	
(USA)	Pre and post	5. Surgical outcomes ^O	passive	13.8	Reason: NA	2.2.07	Discontinue intervention (n=1)	Not applicable
D'I (OF	intervention	6. Adipose tissue gene	F			3.2 %	G (1 1000/	FF
Pilot QE	NA	expression of adiponectin	Before BS				Control group: 100%	
		and leptin ^O						
Arman 2021	~8 wk.	1. Body composition ^O	6 months				Exercise group: 100%	
(Turkey)	D 1	2. Functional capacity ^O		0.2	39.5%	3.8	Control group: 85%	Exercise group: 0%
(======))	Pre and post	3. Physical fitness ^O	Active	9.3	Reason: NA	41 10/	Reasons:	G 4 1 00/
RCT	intervention NA	4. Fatigue^S5. Quality of life^S	Before BS			41.1%	Sedentary behaviors (n=1) Transportation problem (n=1)	Control group: 0%
			Delore DS				Transportation problem (n-1)	
Studies with in	tervention after ba	ıriatric surgery						
							Exercise group: 69%	
		1. Anthropometry ^O					Reasons:	
Castello 2011	~ 17 wk.	2. Body composition ^O	24 months		12.5.0/	1.2	Did not like to exercise due	F : 00/
(Brazil)		 3. Pulmonary function^O 4. Lipid profile^O 			13.5 % Reason:	1,3	muscle or joint pain (n=2) Trouble balance life and exercise	Exercise group: 0%
RCT	Pre BS and post	5. Heart rate variability ⁰	Active	2.2	Did not consent (n=5)		(n=3)	Control group: 0%
KC1	intervention	6. Physical fitness ^O			Did not consent (n=3)	61.5%	Control group: 63%	Control group. 070
	NA	7. Functional capacity ^O	Before BS			01.570	Reason:	
		1 3					No time to exercise (n=6)	
Shah 2011	~12 wk.	1. Anthropometry ^O	NA				Exercise group: 76%	Exercise group: 0%
(USA)		2. Body composition ^O		NA			Reasons:	<u></u>
	Pre, middle and	3. PA ^O	NA	NA			No time to exercise (n=5)	Control group: 25%
RCT	post	4. Physical fitness ^O			NA	NA	Control group: 67%	Reason:

	intervention NA	 5. REE^O 6. Dietary intake^S 7. Glucose metabolism^O 8. Lipid profile^O 9. Quality of life^S 	After BS				Reasons: Would have preferred to be in exercise group (n=3) No time for the study (n=1)	Want to be in exercise group (n=4)
Stegen 2011 (Belgium) Pilot QE	~ 17 wk. Pre BS and post intervention NA	 Anthropometry^O Body composition^O Physical fitness^O Functional capacity^O 	NA NA Before BS	NA	NA	NA	Exercise group: 80% Reasons: Demanding job, household, or education (n=NA) Control group: 78% Reasons: Demanding job, household, or education(n=NA)	Not applicable
Coen 2015 (USA) RCT	~ 26 wk. Pre and post intervention 200\$ x 2 visits	 Anthropometry^O Body composition^O Glucose metabolism^O Physical fitness^O Blood pressure^O Plasma lipids^O Hepatic enzymes^O 	43 months NA After BS	8.7	NA	3,0 34.4%	Exercise group: 91% Reasons: Commitment issues (n=5) Lost to follow up (n=1) Control group: 95% Reasons: Move in another city (n=1) Pregnancy (n=1) Lost to follow up (n=1)	Exercise group: 0% Control group: 0%
Huck 2015 (USA) Pilot QE	~ 12 wk. Pre and post intervention NA	1. Anthropometry ^O 2. Body composition ^O 3. Heart rate ^O 5. Blood pressure ^O 6. Functional capacity ^O 7. Physical fitness ^O	NA Passive After BS	NA	NA	NA	Exercise group: NA Reason: Economic and personal issues (n=2) Control group: NA	Not applicable
Marchesi 2015 (Italy) Pilot QE	~ 44 wk. Pre, middle and post intervention NA	1. Anthropometry ^O 2. Body composition ^O 3. Physical fitness ^O 4. Cardiological assessment ^O 5. Heart rate variability ^O 6. Psychiatric assessments ^S 7. Blood and urinary metabolic parameters ^O	NA Active After BS	NA	NA	NA	Exercise group: 70% Reason: Motivation problem (n=3) Control group: 100%	Not applicable

4								
Muschitz 2016 (Austria)	~ 113 wk. Pre BS and middle-post intervention (9x)	 Anthropometry^O Bone metabolism^O Bone mineral density^O Quality of life^S 	NA NA	NA	NA	NA NA	Exercise group: 100% Control group: 85% Reasons: Unwilling to stay within the study (n=12)	Exercise group: 0% Control group: 0%
Rei	NA		Before BS			141	Move outside Vienna (n=4)	
Rojhani- Shirazi 2016	~ 4 wk.	1 4 41	NA					
(Iran)	Pre BS and post intervention	 Anthropometry^O Balance^O 	NA	NA	NA	NA	NA	NA
RCT	NA		Before BS					
Campanha- Versiani 2017 (Brazil) QE	~ 52 wk. Pre BS and pre middle-post intervention (5x) NA	 Anthropometry^O Strength^O Bone metabolism^O 	NA NA Before BS	NA	11.8% <i>Reason:</i> NA	NA 84.5%	Exercise group: 60% Reasons: Began climacteric period (n=1) Post-operative complications (n=2) Study dropped (n=9) Control group: 63% Reasons: Began climacteric period (n=1) Post-operative complications (n=2) Study dropped (n=8)	Not applicable
Coleman 2017 (USA) Pilot RCT	~ 52 wk. Pre, middle and post intervention 25 \$x 3 visits + 25 \$/ pedometer return	 Anthrometry^O PA^O Functional capacity^O Sedentary activity, aerobic exercise, flexibility et muscle strength^S 	NA Active After BS	NA	NA	NA 33.6%	Exercise group: 81% Reasons: Do not respond (n=3) Dropout during intervention (n=2) Control group: 92% Reason: Do not respond (n=3)	Exercise group: 0% Control group: 0%
Herring 2017 (UK) RCT	~ 12 wk. + 12 wk. of f-up Pre and post intervention + f- up NA	 Anthropometry^O Physical fitness^O Cardiovascular measurements^O PA^O 	13 months Active After BS	3.6	20% Reasons: Work commitments (n=3) Illnesses preventing exercise (n=2) No surgery (n=2) No reason given (n=3)	1.8% 51.1%	Exercise group: 92% Reasons: Had a gastric bad deflated (n=1) Control group: 83% Reason: Want exercise group (n=2) For f-up part: No lost to f-up	Exercise group: 0% Control group: 8.3% Reason: Want exercise group

~ 26 wk. Pre BS. pre and	1. Anthropometry ^O 2. Body composition ^O	NA	NA		NA	Exercise group: 100%	
post intervention	3. Pulmonary function ^O 4. Physical fitness ^O	NA Before BS	1.11	0%	63.2%	Control group: 100%	Not applicable
~12 wk. PreBS and post intervention NA	Anthropometry ^O Body composition ^O 3. Physical fitness ^O	NA Passive Before BS	NA	NA	NA NA	Exercise group: 100% Control group: 100%	Exercise group: 0% Control group: 0%
~12 wk. Pre BS and post intervention NA	1.Anthropometry ^O 2. Body composition ^O 3. Physical fitness ^O 4. PA ^S 5. Dietary Intake ^S	8 months Active Before BS	11.8	27.7% <i>Reason:</i> NA	7.5 63.8%	NA	Three groups: 0%
~52 wk. + 52 wk. f-up Pre BS, pre and post intervention +f- up NA	1. Anthropometry ^O 2. Body composition ^O 3. Blood pressure and heart rate ^O 4. Glucose and lipid metabolism ^O	25 months NA Before BS	2.8	4.8% Reason: NA	2.4 85.7%	For intervention part: Exercise group: 84% Reasons: Pregnancy (n=1) Declined to participate (n=4) Control group: 89% Reasons: Pregnancy (n=1) Declined to participate (n=2) For f-up part: Exercise group: 82% Reasons: Pregnancy (n=1); Injury (n=1) Declined to participate (n=3) Control group: 80% Reasons: Pregnancy (n=2) Declined to participate (n=3)	Exercise group: 0% Control group: 0%
~ 24 wk. PreBS and 1,3, 6 months after surgery(4x) NA	 Anthropometry^O Body composition^O Physical fitness^O PA^O Quality of life^S Food and beverage consumption^S 	56 months Active Before BS	5.2	62.2% Reason: NA	1.4 26.2%	Exercise + Protein group: 100% Protein group: 100% Control group: 100%	Three groups: 0%
	Pre BS, pre and post intervention NA ~12 wk. PreBS and post intervention NA ~12 wk. Pre BS and post intervention NA ~52 wk. + 52 wk. f-up Pre BS, pre and post intervention +f-up NA ~24 wk. PreBS and 1,3, 6 months after surgery(4x)	1. Anthropometry 2. Body composition 3. Pulmonary function 4. Physical fitness 4. Physical fitness 5. Dietary Intakes 1. Anthropometry 2. Body composition 3. Physical fitness 3. Physical fitness 4. PAs 5. Dietary Intakes 5. Dietary Intakes 5. Dietary Intakes 6. Food and beverage 6. Food a	1. Anthropometry NA	1. Anthropometry NA	1. Anthropometry 2. Body composition 2. Body composition 3. Pulmonary function 4. Physical fitness 5. Before BS	1. Anthropometry 2. Body composition NA	Anthropometry NA NA NA NA NA NA NA N

7. Metabol	lic parameters
and vitami	ns ^O

Murai 2019 (Brazil) RCT	~ 39 wk. Pre BS, pre and post intervention NA	 Anthropometry^O Body composition^O Areal bone mineral density^O Bone parameters^O 	NA NA Before BS	NA	46.2% <i>Reason:</i> NA	NA 30.4%	Exercise group: 69% Reason: Personal reason (n=7) Did not undergo BS (n=4) Control group: 71% Reason: Personal reason (n=7) Did not undergo BS (n=3)	Exercise group: 0% Control group: 0%
Diniz-Sousa 2020 (Portugal) RCT	57-70 wk. Pre BS, middle and post intervention (4x)	 Anthropometry^O Strength^O PA^O Bone parameters^O 	20 months NA Before BS	24	78.8% Reasons: Logistical reasons (n=262) Not interested (n=51)	4.2 17.5%	Exercise group: 73% Reasons: Could not be reached (n=6) Lack of interest (n=1) Travel constriction (n=2) Lack of time (n=6) Control group: 71% Reasons: Could not be reached (n=5) Lack of interest (n=1) Travel constrictions (n=2)	Exercise group: 0% Control group: 0%
Tardif 2020 (Canada) RCT	26 wk. + 26 wk f-up Pre BS, pre and post intervention + f- up NA	 Anthropometry^o Body composition^o Lipids profile^o 	32 months NA Before BS	2.6	0%	1.9 71.4%	For intervention part: Exercise group: 85% Reason: Post-operative complications (n=6) Control group: 85% Reasons: Incomplete data on lipids (n=1) Randomized dissatisfaction (n=2) For f-up part: Exercise group: NA Reason: NA Control group: 85% Reasons: Incomplete data on lipids (n=1) Lack of interest (n=1)	Exercise group: 0% Control group: 10% Reason: Randomized dissatisfaction (n=2)

Marc- Hernández 2020 (Spain) RCT	161 wk. before intervention + 22 wk. + 9 wk. f-up for exercise group 5x before intervention + post intervention + f-up NA	 Anthropometry^O Body composition^O Physical fitness^O Cardio-metabolic risk factors^O Quality of life^S 	36 months Active Before BS	NA	5.0% <i>Reason:</i> NA	1.1 21.1%	Exercise group: 100% For f-up part: No lost to f-up Control group: 80% Reason: Refused to continue (n=2)	Exercise group: 0% Control group: 0%
Lamarca 2021 (Brazil) QE	~ 12 wk. Pre and post intervention (middle for food intake) 3x NA	 Anthropometry^O Body composition^O REE^O Blood parameters^O Dietary intake^S 	18 months Passive After BS	10.0	16.2% <i>Reason:</i> NA	6.6 66.1%	Exercise group + Protein: 56% Reasons: Pregnancy (n=1) External accident/illness (n=1) External exercise training (n=1) Declined (n=9) Exercise group: 47% Reason: External accident/illness (n=5) Declined (n=13) Protein group: 63% Reasons: External accident/illness (n=2) Performed a dermolipectomy (n=1) Declined (n=9) Placebo group: 77% Reasons: Performed a dermolipectomy (n=1) External accident/illness (n=1) Non-specific edema (n=1) Declined (n=3)	Not applicable