Review Article

The Effect of Bariatric Surgery on Circulating Levels of Oxidized Low-Density Lipoproteins Is Apparently Independent of Changes in Body Mass Index: A Systematic Review and Meta-Analysis

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Background. Obesity is related to dyslipidemia and increased circulating oxidized LDL (ox-LDL) concentrations that may predispose to atherosclerosis. Bariatric surgery may lower the risk of cardiovascular mortality. Elevated plasma ox-LDL has been associated with atherogenesis and atherosclerotic cardiovascular disease (ASCVD) events. The aim of this meta-analysis was to investigate the impact of bariatric surgery on proatherogenic circulating ox-LDL levels in patients with severe obesity.

Methods. Four databases were systematically searched from inception to May 1, 2021. Also, to clarify the heterogeneity of studies with regard to treatment duration, research design, and the demographic features, a random-effects model and the generic inverse variance weighting approach were utilized. To determine the association with the estimated effect size, a random-effect meta-regression approach was performed. Finally, a meta-regression analysis was conducted to explore the influence of, respectively, baseline and changes in body mass index (BMI), baseline ox-LDL, and postsurgery follow-up period with the estimated effect size of surgery on ox-LDL levels.

Results. Meta-analysis of 11 studies including 470 subjects showed a significant decline in circulating ox-LDL levels following bariatric surgery (SMD: -0.971, 95% CI: -1.317, -0.626, p < 0.001, I²: 89.43%). The results of meta-regression did not show any significant association between the changes in ox-LDL after bariatric surgery and baseline BMI, duration of follow-up or baseline ox-LDL values. However, there was a significant association between ox-LDL alteration and percentage of BMI change.

Conclusion. Bariatric surgery in patients who had severe obesity causes a decrease of circulating ox-LDL that was apparently dependent in BMI changes.
1. Introduction

Obesity is a major risk factor for impaired glucose tolerance, insulin resistance, and type 2 diabetes mellitus, particularly atherosclerotic cardiovascular disease (ASCVD) [1]. Obesity is associated with atherogenic dyslipidemia, low-grade inflammation, and an overall prothrombotic state [2–4]. Dyslipidemia, and especially elevated plasma low-density lipoprotein (LDL) cholesterol, is a pivotal risk factor for atherosclerosis [5]. It is well established that atherogenesis begins with endothelial dysfunction or damage. When LDL particles rich in cholesterol are present in plasma in larger quantities, they permeate through the altered endothelium into the subendothelial space entering the intima. Once this occurs, the LDL particles are oxidized by reactive oxygen species (ROS) and modified into oxidized LDL (ox-LDL) [6, 7]. Ox-LDL particles are strong ligands for macrophage scavenger receptors (CD36, SR-AI/II, and SR-BI) allowing them to enter macrophages and transform them into foam cells [8]. Foam cells, when piled up, appear macroscopically as fatty streaks which are an important step towards fibro-lipid atherosclerotic plaques build-up. Therefore, it is not surprising that increased circulating ox-LDL levels are linked to clinical ASCVD events [9].

Bariatric surgery is a durable and effective therapeutic approach in severely obese individuals. Most endocrinology societies recommend surgical therapy for individuals with BMI ≥ 40 kg/m² or for those with a BMI ranging from 35 to 39.9 kg/m² and comorbidities who may benefit from weight reduction, as well as for severe obese individuals with a BMI 30.0–34.9 kg/m² and poorly controlled type 2 diabetes mellitus. The most common types of bariatric surgery are sleeve gastrectomy (SG), Roux-en-Y gastric bypass (RYGBP), laparoscopic adjustable gastric band (LAGB), and biliopancreatic diversion/duodenal switch (BDP/DS) [10]. Weight loss following bariatric surgery can lower ASCVD risk as well as ensuing mortality in severely obese individuals [11–15].

Since severe obesity is associated with dyslipidemia and increased LDL oxidation that may predispose to atherosclerosis and its ominous consequences, it would be important to verify if bariatric surgery would reduce oxidative stress. Therefore, the aim of this systematic review and meta-analysis was to establish the effect of bariatric surgery on levels of circulating ox-LDL.

2. Methods

2.1. Search Strategy. This systematic review and meta-analysis were done based on the 2009 preferred reporting items for systematic reviews and meta-analysis (PRISMA) guidelines [16]. PubMed, Embase, Scopus, and Web of Science were searched from inception to May 1, 2021, using keywords in abstracts and titles (also in combination with MESH terms) as follows: (“bariatric surgery” OR gastroplasty OR “gastric bypass” OR “Roux-en-Y” OR “gastric band” OR “biliopancreatic diversion” OR gastrectomy OR “duodenal switch” OR “gastrointestinal diversion” OR gastroenterostomy OR “jejunoileal bypass” OR “obesity surgery” OR “weight loss surgery” OR “weight-loss surgery” OR “bariatric procedure” OR “sleeve surgery” OR “metabolic surgery”) AND (“oxidized low density lipoprotein” OR “oxidized LDL” OR “OxLDL” OR “ox-LDL” OR “oxidized Low-Density Lipoprotein” OR “minimally modified oxidized-LDL” OR “MM-LDL” OR “MMLDL” OR “malondialdehyde-low density lipoprotein” OR “malondialdehyde low density lipoprotein” OR “MDA-LDL” OR “MDALDL”).

2.2. Study Selection. For inclusion, only original peer-reviewed studies written in English language were considered. All forms of bariatric surgery procedures were taken into account. Articles must have reported circulating ox-LDL before and after surgery. The exclusion criteria were only abstracts, letters, case reports, comments, meta-analyses, duplicate studies, animal studies, reviews, non-English papers, studies with no surgical intervention, and studies without outcomes.

2.3. Data Extraction. Following the removal of duplicate research, two independent authors reviewed the abstracts and titles of the remaining papers for inclusion. The whole texts of the applicable studies were collected. When two papers with the same research purpose were published by the same organization and/or authors, the study published more recently with a larger sample size was included. Any differences were discussed by authors. The following data was gathered from studies which were eligible for inclusion: (1) the name of the first author, (2) the year of publication, (3) the type of surgery, (4) the study design, (5) the characteristics of the patients, (6) oxidized LDL levels, and (7) the period of follow-up.

2.4. Quality Assessment. The Newcastle-Ottawa scale (NOS) was performed to evaluate the study quality in this meta-analysis [17]. Three features of each eligible study are taken into account: (1) the selection of the studied patients (4 items), (2) the comparability of the studied populations (1 item), and (3) the ascertainment of the exposure (3 items) in case-control studies or outcome of interest in cohort studies.

2.5. Quantitative Data Synthesis. Meta-analysis was performed using Comprehensive Meta-Analysis (CMA) V2 software (Biostat, NJ) [18]. Information regarding sample size, means, and standard deviations from each group were extracted to calculate the standardized mean differences (SMDs). SMD was applied since several different assays were utilized to determine plasma ox-LDL levels. Random-effects meta-analysis was used to get overall estimate of effect size. Postoperative mean and SD were used to calculate the final effect size. To clarify the heterogeneity of studies regarding treatment duration, design of study, and the characteristics of the studied populations, a random-effects model (owing to interstudy heterogeneity) and the generic inverse variance weighting method were utilized [16]. Clinical heterogeneity was judged by study locations and recruited populations, methods applied for ox-LDL assay, baseline ox-LDL values, and differences in biochemical parameters among studied populations. Statistical heterogeneity was appraised by I²
| Author, year, country | Design of study | Follow-up | Type of surgery | Control (n) | Outcome | BMI change (%) | Patients’ characteristic | No. of patients |
|-----------------------|----------------|-----------|----------------|-------------|---------|---------------|-------------------------|----------------|
| Carmona-Maurici et al., 2020 [24] Spain | Prospective observational cohort study | 6 months 12 months | Laparoscopic RYGB or SG | — | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | -32.27 | Obese patients with atheromatous plaque 51.8 ± 1.8 years old 18 (F)/14 (M) Obese patients without atheromatous plaque 43.5 ± 1.8 years old 29 (F)/5 (M) | 32 |
| Ho et al., 2021 [25] | Prospective, observational study | 6 months 12 months | RYGB, SG, or omega loop bypass | Patients seeking weight management [16] | Unchanged (Mercodia ox-LDL kit) | -29.95 | Morbid obesity patients 50.1 ± 10 years old | 59 |
| Coimbra et al., 2019 [26] | Observational study | 13 months | Laparoscopic adjustable gastric banding (LAGB) | Healthy volunteers [17] | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | -11.85 | Obese patients 49.03 ± 10.71 years old 18 (F)/2 (M) Obese women 46 ± 9 years old Obese women 48 ± 8 years old | 20 |
| Gomez-Martin et al., 2018 [27] | Observational study | 6 months 12 months | SG | Patients matched for age and cardiovascular risk (modified Mediterranean diet) [18] | Significant decrease in ox-LDL levels after 12 months in comparison to baseline and control group (Mercodia ox-LDL kit) | -28.83 | Adolescents with severe obesity 16.5 ± 1.6 years old 10 (M)/29 (F) Adolescents with severe obesity 16.5 ± 1.6 years old 3 (M)/10 (F) | 39 |
| Kelly et al., 2016 [28] | Longitudinal cohort | 3 months 12 months | Laparoscopic RYGB | — | Significant decrease in ox-LDL levels at 12 months (Mercodia ox-LDL kit) | -32.6 | Adolescents with BMI more than 35 kg/m² and insulin-dependent T2DM 58.6 ± 6.1 years old 10 (M)/10 (F) | 20 |
| Müller-Stich et al., 2015 [29] | Prospective cohort | 6 months 12 months | RYGB | — | Unchanged (OxiSelect MDA-LDL-quantitation kit) | -25 | Obese patients 40 ± 14 years old 5 (M)/12 (F) | 17 |
| Van der Schuren et al., 2015 [30] | Observational study | 4 months 7 years | Laparoscopic RYGB | Lean controls [24] | Significant decrease in ox-LDL levels at 7 years (Mercodia ox-LDL kit) | -26.6 | Obese patients 20-60 years old 15 (F)/6 (M) | 21 |
| Julve et al., 2014 [31] | Observational study | 6 months 12 months | RYGB | — | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | — | Obese patients 20-60 years old 15 (F)/6 (M) | 21 |
| Author, year, country | Design of study | Follow-up | Type of surgery | Control (n) | Outcome ox-LDL methods of ox-LDL assessment | BMI change (%) | Patients’ characteristic | No. of patients |
|-----------------------|----------------|-----------|----------------|------------|--------------------------------------------|---------------|------------------------|----------------|
| Martín-Rodríguez et al., 2014 [32] | Prospective cohort | 12 months | Bariatric surgery | Lean Control subjects [30] | Significant decrease in ox-LDL levels (immunodiagnostic system) | -31.37 | Obese patients without metabolic syndrome 40 ± 9 years old 19(F)/4(M) | 23 |
| Garrido-Sánchez et al., 2008 [33] | Observational study | 7 months | Biliopancreatic diversion, or RYGB | Healthy, nonobese persons [11] | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | -30.39 | Obese patients with metabolic syndrome 42 ± 10 years old 34(F)/5(M) | 39 |
| | | | | | | | Morbidly obese patients with: | |
| | | | | | | Normal fasting glucose 39.5 ± 11 years old 14(F)/7(M) | 21 |
| | | | | | | Impaired fasting glucose 44.1 ± 10.6 years old 21(F)/10(M) | 31 |
| | | | | | | Type 2 diabetes 44.5 ± 7.4 years old 14(F)/7(M) | 21 |
| | | | | | | Morbidly obese patients 35.1 ± 13.1 years old 10(F)/10(M) | 20 |
| Uzun et al., 2004 [34] | Observational study | 6 months | Open Swedish adjustable gastric band (SAGB) | — | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | -24.24 | Morbidly obese patients 34.6 ± 9 years old 11(F)/9(M) | 20 |
| | | | Laparoscopic SAGB | — | Significant decrease in ox-LDL levels (Mercodia ox-LDL kit) | -24.27 | | | |
Records identified through database searching include pubmed (12), scopus (27), embase (38), web of science (16) \((n = 93)\)

Records after duplicates removed \((n = 43)\)

Records screened \((n = 43)\)

Full-text articles assessed for eligibility \((n = 11)\)

Studies included in the systematic review and meta-analysis \((n = 11)\)

Excluded \((n = 32)\)

- review articles \((n = 10)\)
- not meeting inclusion criteria \((n = 17)\)
- insufficient data \((n = 5)\)

**Figure 1:** Flow chart of included studies.

### Table 2: Quality of bias assessment of the included papers in accordance with the Newcastle-Ottawa scale.

| Study                        | Case definition | Representativeness of the cases | Selection of controls | Definition of controls | Comparability of cases and controls | Ascertainment of exposure | Exposure Same method of ascertainment | Nonresponse rate |
|------------------------------|-----------------|---------------------------------|-----------------------|------------------------|-------------------------------------|--------------------------|--------------------------------------|------------------|
| Ho et al. 2021               | —               | —                               | —                     | —                      | —                                   | *                        | —                                    | —                |
| Carmona-Maurici et al. 2020  | —               | *                               | —                     | —                      | *                                   | —                        | —                                    | —                |
| Coimbra et al. 2019          | —               | —                               | *                     | —                      | —                                   | *                        | —                                    | —                |
| Gómez-Martín et al. 2018     | —               | —                               | —                     | *                      | *                                   | —                        | —                                    | —                |
| Kelly et al. 2016            | —               | —                               | —                     | —                      | *                                   | —                        | —                                    | —                |
| Van der Schueren et al. 2015 | —               | —                               | —                     | *                      | *                                   | *                        | —                                    | —                |
| Müller-Stich et al. 2015     | —               | —                               | —                     | —                      | *                                   | —                        | —                                    | —                |
| Martín-Rodríguez et al. 2014 | —               | —                               | —                     | —                      | *                                   | —                        | —                                    | —                |
| Julve et al. 2014            | —               | —                               | —                     | —                      | *                                   | —                        | —                                    | —                |
| Garrido-Sánchez et al. 2008  | —               | *                               | —                     | *                      | —                                   | —                        | —                                    | —                |
| Uzun et al. 2004             | —               | —                               | —                     | —                      | *                                   | —                        | —                                    | —                |

*Only for comparability a maximum of two stars can be given.*
LDL. (b) Leave-one-out sensitivity analyses for the influence of bariatric surgery on ox-LDL. Findings for each outcome, the magnitude of the confidence intervals for the influence of bariatric surgery on ox-LDL.

2.6. Meta-Regression. A meta-regression analysis was carried out to investigate the impact of, respectively, baseline and changes in BMI, baseline ox-LDL and duration of postsurgery follow-up with the estimated effect size of surgery on ox-LDL concentrations.

2.7. Subgroup Analysis. A subgroup analysis was conducted to investigate the impact of follow-up duration (≥12 months and <12 months) with the estimated effect size of surgery on ox-LDL concentrations.

2.8. GRADE Scoring. We used the grade of recommendations, assessment, development, and evaluation (GRADE) approach to assess the strength of evidence for each outcome [22]. To summarize the findings for each outcome, the GRADEpro GDT software was used. We assigned four points to each outcome and then evaluated factors that reduced the quality of the evidence. For each outcome, points were reduced based on the presence of the following: the overall RoB for each study, inconsistency (significant heterogeneity), indirectness (significant differences in the population, comparisons, and outcomes), and imprecision (the size of the cohort, width, and significance of the confidence intervals (CIs)). As a result, we classified the evidence into four groups depending on the aggregate GRADE ratings for each intervention: high-grade evidence (at least 4 points), moderate grade evidence (3 points), low-grade evidence (2 points), and very low-grade evidence (1 point).

2.9. Publication Bias. In the meta-analysis, the funnel plot was used to realize the presence of publication bias. Hence, Begg’s rank correlation and Egger’s weighted regression tests were often performed to help publication bias detection. When asymmetry in the funnel plot was potentially reduced the quality of the evidence.
missing studies were inserted using the “trim and fill” method. In the event of a significant result, the “fail-safe N” approach was used to compute the number of potentially missing studies required to make the p value nonsignificant. This is another sign of publishing bias [23].

3. Results

A thorough database search identified 93 published papers, 43 of which were directly connected to the issue of this study. After careful consideration, 32 studies were excluded:

**Table 3: Grade assessment.**

| Outcome no. of participants (studies) | Relative effect (95% CI) | Anticipated absolute effects (95% CI) | Certainty | What happens |
|-------------------------------------|--------------------------|--------------------------------------|-----------|--------------|
| Ox-LDL levels (ox-LDL) assessed with: ELISA/Mercodia/ immunodiagnostic/OxiSelect follow-up: range 6 months to 7 years no. of participants: 470 (11 observational studies) | | | | |

*The risk in the intervention group (and its 95% confidence interval) is based on the assumed risk in the comparison group and the relative effect of the intervention (and its 95% CI).

CI: confidence interval

GRADE working group grades of evidence

High certainty: we are very confident that the true effect lies close to that of the estimate of the effect.

Moderate certainty: we are moderately confident in the effect estimate: the true effect is likely to be close to the estimate of the effect, but there is a possibility that it is substantially different.

Low certainty: our confidence in the effect estimate is limited: the true effect may be substantially different from the estimate of the effect.

Very low certainty: we have very little confidence in the effect estimate: the true effect is likely to be substantially different from the estimate of effect.

**Figure 3: Subgroup analysis to assess the influence follow up duration in ox-LDL alteration.**
Regression of BMI change on std diff in means

Regression of baseline BMI on std diff in means

Figure 4: Continued.
studies were reviews, 17 studies did not meet the inclusion criteria, and 5 studies did not disclose enough data. As a result, 11 studies which evaluated the levels of ox-LDL after bariatric surgery were included (Table 1). The study selection procedure was indicated in Figure 1.

3.1. Quality Assessment of the Included Studies. All selected studies showed insufficient information for case definition, and most of them had lack of information for representativeness of the cases. Because most of the studies did not include a control group, they were not evaluated for selection of controls, definition of controls, comparability, the same method of ascertainment, and nonresponse rate. However, all studies which included met the ascertainment of exposure criteria. Table 2 shows the details of quality assessment.

3.2. Assay Methods. In most of the included studies, serum ox-LDL was assessed using enzyme-linked immunosorbent assay (ELISA) method. Nine studies used Mercodia ox-LDL kit (Mercodia, Uppsala, Sweden) [24–28, 30, 31, 33, 34], one study used immunodiagnostic system (Boldon, UK) [32], and one study used Oxiselect MDA-LDL-Quantitation kit (Cell Biolabs Inc., San Diego, USA) [29].
3.3. Effect of Bariatric Surgery on Circulating Concentrations of Oxidized LDL. Meta-analysis of 11 publications including 470 subjects demonstrated a significant reduction of circulating ox-LDL following bariatric surgery (SMD: -0.971, 95% CI: -1.317, -0.626, \( p < 0.001 \), \( I^2: 89.43\% \)) (Figure 2(a)). In the leave-one-out sensitivity analysis, the reduction in circulating ox-LDL was robust (Figure 2(b)) (low-grade evidence, Table 3).

3.4. Subgroup Analysis. A subgroup analysis was also performed based on follow-up duration (≥12 months and <12 months). Subgroup analyses demonstrated significant reduction of circulating ox-LDL following bariatric surgery in both follow-up periods (≥12 months \( p < 0.001 \) and <12 months \( p < 0.001 \)). However, this analysis did not show significant associations between follow-up duration and change in ox-LDL levels (\( p = 0.309 \)) (Figure 3).

3.5. Meta-Regression. Random-effects meta-regression was used to assess the effect of potential confounders on the ox-LDL reducing effect of bariatric surgery. The results did not designate any significant association between the changes in ox-LDL and baseline BMI (slope: 0.018; 95% CI: -0.041, 0.078; \( p = 0.549 \)), follow-up duration (slope: -0.007; 95% CI: -0.028, 0.012; \( p = 0.444 \)) or baseline ox-LDL (slope: 0.00005; 95% CI: -0.00005, 0.00016; \( p = 0.324 \)). However, there was a significant association between changes in ox-LDL and percentage of BMI change (slope: 0.069; 95% CI: 0.003, 0.135; \( p = 0.039 \)) (Figures 4(a)–4(d)).

3.6. Publication Bias. Egger’s linear regression test (intercept = -7.534, standard error = 0.93; 95%CI = -9.535, -5.533, \( t = 8.026 \), df = 15, two-tailed \( p < 0.001 \)) and Begg’s rank correlation test (Kendall’s Tau with continuity correction = -0.801, \( z = 4.49 \), two-tailed \( p < 0.001 \)) indicated the presence of publication bias in this meta-analysis of bariatric surgery effects on circulating ox-LDL. Trim-and-fill analysis revealed that among all included papers in meta-analysis, there could be five missing studies. The “fail-safe N” test showed that 817 missing studies were required to reduce the effect size to a nonsignificant (\( p < 0.001 \)) value (Figure 5).

4. Discussion

The results of this meta-analysis revealed a substantial decrease of circulating ox-LDL after bariatric surgery. Of importance, the results of meta-regression did not reveal any significant relationship between the changes in baseline BMI, duration of follow-up or baseline ox-LDL value, and ox-LDL after bariatric surgery. However, there was significant association between the changes in ox-LDL after bariatric surgery and percentage of BMI change.

Some authors have tried to explain the beneficial effect of bariatric surgery on ASCVD by decreasing not only body weight, oxysterols, and ox-LDL but also by decreasing plasminogen activator inhibitor-1 (PAI-1), which is elevated in extremely obese patients and by decreased proliferation of vasa vasorum [35]. Bariatric surgery is a well-documented technique for weight loss can have numerous favorable consequences, such as rise in GLP-1 and its potential function in the metabolism including remission of T2DM [36]. Furthermore, bariatric surgery may be a useful therapy for people who have cardiovascular risk factors [37]. Also, bariatric surgery decreases oxidative stress parameters and glycoproteins and increases antioxidant enzymes paraoxonase-1 and catalase as well, which supports the idea that this procedure decreases oxidation of lipoproteins thus having antiatherogenic effects [34,35].

Others have tried to explain the beneficial effect of bariatric surgery, besides reducing oxysterols and ox-LDL, by increasing not only HDL-cholesterol but also the number of larger HDL particles which are more atheroprotective and reducing the number of small HDL particles which are less protective [9]. Bariatric surgery, apart from decreasing ox-LDL, also reduces the levels of triglycerides and often decreases the number of LDL particles with a decreased proportion of smaller, more atherogenic LDL particles [38]. Already two decades ago, it has been proposed that small dense LDL particles confer greater risk for atherosclerosis and ASCVD than large, buoyant LDL particles, which might be attributed to the greater oxidative vulnerability of small dense LDL [39].

BMI is an indicator of general obesity and demonstrates significant associations with CVDs and ASCVD risk factor. However, as previously suggested, our data support a direct relationship between the magnitude of improvement in cardiovascular risk factors and the amount of BMI reduction [40]. These consequences may be related to positive correlation between change in serum LDL—as a marker of cardiovascular status—and change in BMI [41]. Furthermore, weight loss-induced LDL reduction might decrease the circulating level of the substrate (i.e., LDL particles) for oxidation, and this could partially account for reduction in the generation of ox-LDL [42]. Also, some think that waist circumference, which has been shown to be associated with elevated ASCVD risk, is a better indicator of abdominal obesity [43]. The others suggest that waist-to-hip ratio is even better for measuring abdominal obesity, and it has been proven that this measure corresponds with ASCVD [44].
The favorable effects of bariatric surgery are most likely to be multifactorial [45]. It should be highlighted that weight reduction is not always required to see improvements in some of with ASCVD risks. Indeed, there is considerable evidence to imply that bariatric surgery may cause changes in oxidative stress, inflammation, and adipokines via non-weight loss pathways [28].

Also, this rather surprising issue that bariatric surgery can improve markers of cardiovascular disorders among patients with severe obesity as long as body fat mass is redistributed or replaced by muscle mass independent of significant weight loss needs further investigation.

This study has some advantages. One of them is that this is the first meta-analysis trying to establish the effect of bariatric surgery on circulating ox-LDL in patients with severe obesity. Further investigations should focus on this premise. This meta-analysis has some limitations as well. The studies which were included had an overall relatively small number of patients. Also, there is heterogeneity in types of operations performed, with known differences in metabolic effects between operations. Also, the methods for measuring ox-LDL concentrations in some studies included in this meta-analysis were different and might have explained heterogeneity in our findings. Besides, the quality of the evidence which was evaluated with GRADE approach was low. Additionally, the time intervals and follow-up compliance are not known in some of the reported studies.

In conclusion, bariatric surgery seems to cause a decrease of circulating ox-LDL which is associated with percentage of BMI change, baseline BMI, duration of follow-up, or baseline ox-LDL value. Future studies may focus on the potential neurohormonal effects that could contribute to this reduction, both dependent and independent of weight loss factors.

Data Availability

There is not any raw data associated with this review article.

Additional Points

Key Points. (i) Obesity is related to increased circulating oxidized LDL (ox-LDL) and atherosclerotic risks. (ii) Bariatric surgery may lower the risk of cardiovascular mortality. (iii) Bariatric surgery in severely obese causes a decrease of circulating ox-LDL.

Conflicts of Interest

The authors declare that there is no conflict of interest.

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