Effect of Ezetimibe Added to High-Intensity Statin Therapy on Low-Density Lipoprotein Cholesterol Levels: A Meta-Analysis

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Abstract

Background: Adding ezetimibe to high-intensity statin therapy is used for additional lowering of low-density lipoprotein cholesterol (LDL-C); however, there are little data on the efficacy of ezetimibe when combined with a high-intensity statin. A meta-analysis was performed to evaluate the efficacy of ezetimibe added to high-intensity statin therapy on LDL-C levels.

Methods: A literature search from database inception to May 2020 was performed using PubMed, EMBASE and Cochrane Central Register of Controlled Trials. The Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines were used in this meta-analysis, in which the random-effects model was adopted for the calculation of the mean difference (MD). The Cochrane Collaboration’s tool for assessing the risk of bias was used to evaluate the quality of the included trials.

Results: A total of 14 trials with 2,007 patients were included in this study. Compared to the high-intensity statin monotherapy, the MD in LDL-C reduction with high-intensity statin therapy plus ezetimibe was -14.00% (95% confidence interval: -17.78 to -10.22; P < 0.001) with a moderate degree of heterogeneity (P < 0.001, I² = 66%). No significant publication bias among the included trials was identified.

Conclusions: Our study found that adding ezetimibe to high-intensity statin therapy provided a significant but attenuated incremental reduction in LDL-C levels. Whether the magnitude of this additional lowering of LDL-C levels would lead to benefits in clinical cardiovascular outcomes needs further investigation.

Keywords: Ezetimibe; High-intensity statin; LDL-C; Hypercholesterolemia; Coronary heart disease

Introduction

Ezetimibe is the most used non-statin medication for secondary prevention in patients with clinical atherosclerotic cardiovascular disease (ASCVD). It was recommended by the 2018 AHA/ACC Guideline on the Management of Blood Cholesterol [1] (2018 AHA/ACC Cholesterol Guideline) for patients with clinical ASCVD receiving maximally tolerated statin therapy when the low-density lipoprotein cholesterol (LDL-C) level remains ≥ 1.8 mmol/L (70 mg/dL). Ezetimibe was also recommended in patients with severe hypercholesterolemia on maximally tolerated statin therapy and the LDL-C level ≥ 2.6 mmol/L (100 mg/dL).

However, there is a lack of high-quality evidence for these moderate or weak recommendations, with no level A evidence provided in the 2018 AHA/ACC Cholesterol Guideline. In real-world practice, the maximally tolerated statin therapy is a high-intensity statin for majority patients, as evidenced by a prospective cohort study showing that approximately 70% of patients took a high-intensity statin at 1 year after acute coronary syndromes (ACS) [2]. Moreover, 12.6% of patients with ACS in the study were on the combination therapy of a statin and ezetimibe, and among them, about 60% of patients took a high-intensity statin and ezetimibe [2]. Unfortunately, there have been no large-scale randomized double-blinded clinical trials to evaluate the combination of high-intensity statin therapy and ezetimibe. The only available outcome study that was cited in the guidelines was the IMPROVE-IT trial [3], in which a moderate-intensive statin was used and ezetimibe only demonstrated a modest reduction in cardiovascular events. Whether the modest benefit observed in the IMPROVE-IT trial can be extrapolated into the real-world patients taking ezetimibe with a high-intensity statin is unknown. The possibility that the co-administration of high-intensity statin may offset the limited benefits of ezetimibe, making the addition of ezetimibe unnecessary cannot be ruled out.

As outcome trial evaluating the combination of high-intensity statin and ezetimibe was not available, the LDL-C levels as a surrogate endpoint could be used alternatively to...
evaluate the benefits of ezetimibe when adding to the high-intensity statin therapy. Several small trials on this topic have been conducted, but results are conflicting. The 2018 AHA/ACC Cholesterol Guideline stated that “The addition of ezetimibe or bile acid sequestrants to statin therapy typically provides an additional 15% to 25% reduction in LDL-C” [1], but, oddly, no reference regarding this statement was provided in the guideline. Thus, there is a gap of evidence in the medical literature to support this statement.

Numerous meta-analysis studies investigating the efficacy of ezetimibe added to ongoing statin therapy on LDL-C levels were reported, but none of them has evaluated the addition of ezetimibe to high-intensity statin therapy. Mikhailidis et al [4] conducted a meta-analysis including five randomized controlled trials involving a total of 5,039 patients; however, only low- to moderate-intensity statins were used in these five trials. Four years later, Mikhailidis et al [5] reported another meta-analysis about ezetimibe of 13 trials including 5,080 patients, but only one included trial employed high-intensity statin. Savarese et al [6] and Ye et al [7] also evaluated the combination of ezetimibe and statin therapy, respectively, but neither of them included trials adding ezetimibe to high-intensity statin therapy. Recently, Yu et al [8] conducted a meta-analysis comparing the efficacy of combination therapy with ezetimibe and statins versus a double dose of statin monotherapy. Among the 11 included studies, only one included trial had a therapy group receiving the combination of ezetimibe and a high-intensity statin. The most comprehensive meta-analysis of ezetimibe was reported by Lorenzi et al [9], and a total of 35 randomized controlled trials were identified and included in their analyses. Of them, only two trials had intervention groups receiving the combination of high-intensity statin therapy and ezetimibe. Besides, ezetimibe has been used as a positive control in several clinical trials [10-13] of proprotein convertase subtilisin/kexin type 9 (PCSK9) inhibitors and cholesteryl ester transfer protein (CETP) inhibitors; however, those trials have not been evaluated in the previous meta-analysis studies.

To the best of our knowledge, there is neither meta-analysis nor large-scale outcome trial examining the addition of ezetimibe to the background high-intensity statin therapy reported yet. Here, we performed a meta-analysis to add some insights into this topic by evaluating the efficacy of ezetimibe in reducing LDL-C levels when added to high-intensity statin therapy, and a discussion was also presented regarding the relationship between the magnitude of LDL-C lowering of ezetimibe and the reduction of ASCVD risk.

Materials and Methods

Literature search

We followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines for reporting the present meta-analysis. A systematic literature search was performed through May 2020 using PubMed (1946-), EMBASE (1947-) and Cochrane Central Register of Controlled Trials with the following key terms: “atorvastatin”, “rosuvastatin”, “high”, “statin” and “ezetimibe” (Supplementary Material 1, www.cardiologyres.org). The screening was restricted to clinical trials in humans published in English. The references were imported into EndNote™ and then Rayyan, which is a systematic reviews web app. Duplicate records were removed automatically.

Article screening and selection

Two authors independently assessed the search result for eligibility by title and/or abstract using Rayyan. Differences were resolved through consensus between the authors. Studies that met the following criteria were included in the meta-analysis: 1) Comparing high-intensity statin plus ezetimibe with the corresponding high-intensity statin monotherapy in patients with a history of clinical ASCVD or hypercholesterolemia; 2) Having comparable baseline LDL-C levels in both groups; 3) Reporting the change in LDL-C levels. Studies containing only a single group (i.e., pre-post treatment) were excluded because they did not contain an adequate comparison group. High-intensity statin therapy was defined as atorvastatin 40 - 80 mg daily or rosuvastatin 20 - 40 mg daily [14]. References passed the initial screening were further evaluated through the full-text review.

Data extraction and statistical analysis

The study design and demographic characteristics of the selected trials, including trial settings, trial locations, sample size, study population, interventions and the changes of LDL-C levels were extracted and assessed. The Cochrane Collaboration’s tool for assessing the risk of bias was used, and two authors independently evaluated the quality of the included trials. Like the article screening process, differences were resolved through consensus between the authors.

The outcome measure of this analysis was the mean difference (MD) in the reduction of LDL-C levels in patients treated with the combination of a high-intensity statin and ezetimibe vs. the corresponding high-intensity statin monotherapy. The random-effects model with the DerSimonian-Laird method was adopted for the calculation of MD given expected clinical and methodological heterogeneity. Three subgroup analyses were conducted based on the type of high-intensity statins (atorvastatin vs. rosuvastatin), the dose of high-intensity statins (low dose (atorvastatin 40 mg and rosuvastatin 20 mg) vs. high dose (atorvastatin 80 mg and rosuvastatin 40 mg)) and study regions (Asia vs. Western countries). A meta-regression analysis using the random-effects model (method of moments) was performed to evaluate the effect of baseline LDL-C levels. The I² statistic test for assessment of in-between study heterogeneity with values < 25%, 25-50%, 50-75% and > 75% corresponds to no, low, moderate and high degree of heterogeneity, respectively. Small study effects were evaluated through Egger’s test and Begg’s test. We used a confidence interval (CI) of 95% and P < 0.05 as a reflection of statistically significant results in all our analyses. All statistical analyses were conducted by RevMan 5.3 and jamovi 1.1.9.
Results

Literature search and study characteristics

Of the initial 1,037 citations, 14 clinical trials involving 2,007 patients were included in this analysis (Fig. 1) [10, 12, 13, 15-25]. Among them, the study of Nicholls 2017 did not report standard deviations (SDs), and the missing SDs were borrowed from the study of Robinson 2014 due to similarity between these two trials according to the recommendations of Cochrane Handbook for Systematic Reviews of Interventions. The study of Ballantyne 2003 contains two treatment arms of high-intensity statins (atorvastatin 40 mg and 80 mg), and the results of these two arms were reported separately. Thus, in our meta-analysis, the results of these two arms were separately analyzed accordingly though they were from the same study.

The characteristics of the included trials were summarized in Table 1 [10, 12, 13, 15-25]. Four included studies were open-label trials, and two double-blinded studies did not mask ezetimibe use. Three trials were designed to evaluate PCSK9 inhibitors or CETP inhibitors, and the data about ezetimibe were a by-product [9, 11, 12]. A mean LDL-C level of < 1.8 mmol/L (70 mg/dL) was achieved more frequently in the groups of patients receiving the ezetimibe plus high-intensity statin therapy than the groups of patients receiving high-intensity statin monotherapy in the included trials. Of note, in the study of Robinson 2014, a more than 10% increase in the LDL-C levels from baseline in the atorvastatin 80 mg monotherapy group was reported. It is unclear why the LDL-C levels were elevated despite the continuous atorvastatin 80 mg monotherapy after a 4-week lipid stabilization period.
| Studies                  | Design       | Region          | Patients eligibility                                                                 | Treatment, patients number (n) | Baseline LDL-C (mmol/L (mg/dL)) | LDL-C at the end of studies (mmol/L (mg/dL)) |
|-------------------------|--------------|-----------------|---------------------------------------------------------------------------------------|-------------------------------|---------------------------------|---------------------------------------------|
| Azar et al, 2011 [15]   | R, DB, PC, 8 weeks | Western countries | Adults with stable coronary artery disease or coronary artery disease equivalent       | Atorvastatin 40 mg/day, n = 50 | 2.56 ± 0.54 (99 ± 21)          | 2.22 ± 0.36 (86 ± 14)                  |
|                         |              |                 | Atorvastatin 40 mg/d + ezetimibe 10 mg/d, n = 50                                       |                               | 2.64 ± 0.75 (102 ± 29)         | 1.99 ± 0.26 (77 ± 10)                  |
| Ballantyne et al, 2003 [16] | R, DB, PC, 12 weeks | Western countries | Adults with primary hypercholesterolemia, defined as calculated LDL-C of 145 - 250 mg/dL, inclusive, and TG levels ≤ 350 mg/dL. | Atorvastatin 40 mg/day, n = 57 | N/A                             | N/A                                        |
|                         |              |                 | Atorvastatin 40 mg/d + ezetimibe 10 mg/day, n = 57                                     |                               |                                 |                                             |
| Ballantyne et al, 2007 [17] | R, OL, MC, 6 weeks | Western countries | Adults with hypercholesterolemia and a history of CHD or clinical evidence of atherosclerosis or a CHD risk equivalent (10-year CHD risk score > 20%); two most recent fasting LDL-C levels were ≥ 160 mg/dL and < 250 mg/dL and the two measurements were within 15% of each other; a fasting TG concentration of < 400 mg/dL | Rosuvastatin 40 mg/day, n = 230 | 4.94 (191)                      | 2.12 (82)                                 |
|                         |              |                 | Rosuvastatin 40 mg/day + ezetimibe 10 mg/day, n = 235                                 |                               | 4.88 (189)                      | 1.47 (57)                                 |
| Barbosa et al, 2013 [18] | R, OL, 4 weeks | Western countries | Adults with at high risk for CHD, characterized by at least one of the following conditions: established CHD or equivalent; diagnosis of type 2 diabetes; or those whose Framingham risk score was > 20%. | Atorvastatin 40 mg/day, n = 45 | 2.3 (89)                        | 1.91 (74)                                 |
|                         |              |                 | Atorvastatin 40 mg/day + ezetimibe 10 mg/day, n = 37                                |                               | 2.30 (94)                       | 1.60 (62)                                 |

Table 1. Study Characteristics
| Studies                  | Design   | Region          | Patients eligibility                                                                 | Treatment, patients number (n) | Baseline LDL-C (mmol/L (mg/dL)) | LDL-C at the end of studies (mmol/L (mg/dL)) |
|-------------------------|----------|-----------------|--------------------------------------------------------------------------------------|---------------------------------|----------------------------------|---------------------------------------------|
| Blom et al, 2014 [10]   | R, DB, 52 weeks; atorvastatin and ezetimibe were not masked | Western countries | Adults with an LDL-C ≥ 75 mg/dL and a fasting TG level ≤ 400 mg/dL. | Atorvastatin 80 mg/day, n = 73 | 2.49 ± 0.34 (96.2 ± 13.3)          | 2.70 ± 0.82 (104.6 ± 31.61)                |
|                         |          |                 |                                                                                      | Atorvastatin 80 mg/day + ezetimibe 10 mg/day, n = 63 | 3.10 ± 0.84 (119.8 ± 32.4)       | 2.97 ± 1.15 (115.0 ± 44.44)               |
| Hong et al, 2018 [19]   | R, DB, MC, 8 weeks | South Korean | Adults with hypercholesterolemia requiring medical treatment according to NCEP ATP III. | Rosuvastatin 20 mg/day, n = 64 | 3.95 ± 0.81 (152.8 ± 31.4)          | 1.95 ± 0.88 (75.4 ± 34.0)                 |
|                         |          |                 |                                                                                      | Rosuvastatin 20 mg/day + ezetimibe 10 mg/day, n = 64 | 3.97 ± 0.81 (153.5 ± 31.2)       | 1.37 ± 0.57 (53.1 ± 22.0)                 |
| Hougaard et al, 2017 [20]| R, DB, SC, 52 weeks | Western countries | Adults with ST-elevation myocardial infarction                                       | Atorvastatin 80 mg/day, n = 41  | 4.10 ± 0.90 (158.55 ± 34.80)        | 2.00 ± 0.50 (77.34 ± 19.34)               |
|                         |          |                 |                                                                                      | Atorvastatin 80 mg/day + ezetimibe 10 mg/day, n = 39 | 3.70 ± 0.71 (143.08 ± 27.67)     | 1.40 ± 0.80 (54.14 ± 30.94)               |
| Kim et al, 2016 [21]    | R, DB, MC, 8 weeks | South Korean | Adults with initial levels of LDL-C ≤ 250 mg/dL and TG < 400 mg/dL were selected for a lead-in period, then the patients who required lipid-lowering treatment according to the NCEP ATP III guidelines were finally enrolled in the study. | Rosuvastatin 20 mg/day, n = 68  | N/A                               | N/A                                         |
|                         |          |                 |                                                                                      | Rosuvastatin 20 mg/day + ezetimibe 10 mg/day, n = 68 | N/A                              | N/A                                         |
| Kim et al, 2018 [22]    | R, DB, MC, 8 weeks | South Korean | Adults with primary hypercholesterolemia who had an LDL-C level < 250 mg/dL and TG level < 350 mg/dL requiring treatment according to NCEP ATPIII guidelines. | Rosuvastatin 20 mg/day, n = 63  | 4.16 ± 0.79 (160.90 ± 30.57)        | N/A                                         |
|                         |          |                 |                                                                                      | Rosuvastatin 20 mg/day + ezetimibe 10 mg/day, n = 64 | 4.12 ± 0.77 (159.56 ± 29.91)     | N/A                                         |
| Lins et al, 2014 [23]   | R, OL, 4 weeks | Western countries | Adults at high-risk for CHD, characterized by at least one of the following conditions: stable CHD or CHD equivalent; diagnosis of type 2 diabetes; or those whose Framingham risk score was > 20%. | Atorvastatin 40 mg/day, n = 22  | 2.30 (89)                          | 1.94 (75)                                  |
Table 1. Study Characteristics - (continued)

| Studies                  | Design          | Region              | Patients eligibility                                                                 | Treatment, patients number (n) | Baseline LDL-C (mmol/L (mg/dL)) | LDL-C at the end of studies (mmol/L (mg/dL)) |
|--------------------------|-----------------|---------------------|--------------------------------------------------------------------------------------|---------------------------------|----------------------------------|-------------------------------------------|
| Malina et al, 2015 [24]  | R, OL, 4 weeks  | Western countries   | Adults in primary or secondary prevention of CHD, who had an indication for lipid-lowering therapy in accordance with NCEP ATP III | Atorvastatin 40 mg/day + ezetimibe 10 mg/day, n = 21 | 2.45 (95)                     | 1.60 (62)                               |
|                         |                 |                     |                                                                                      | Atorvastatin 40 mg/day, n = 35 | 2.58 ± 0.83 (100 ± 32)         | 2.01 ± 0.67 (78 ± 26)                |
|                         |                 |                     |                                                                                      | Atorvastatin 40 mg/day + ezetimibe 10 mg/day, n = 22 | 2.30 ± 0.72 (89 ± 28)         | 1.42 ± 0.49 (55 ± 19)                 |
| Nicholls et al, 2017 [12]| R, DB, PC, 90 days | Western countries   | Adults with either atherosclerotic cardiovascular disease or type 1 or 2 diabetes mellitus, treated with atorvastatin 40 mg daily for at least 30 days prior to screening and compliant with study drug for the 28-day lead-in phase, had an LDL-C > 70 mg/dL or non-HDL-C > 100 mg/dL at screening and prior to randomization and a TG ≤ 400 mg/dL. | Atorvastatin 40 mg/day, n = 40 | N/A                              | N/A                                      |
|                         |                 |                     |                                                                                      | Atorvastatin 40 mg/day + ezetimibe 10 mg/day, n = 91 |                                  |                                           |
| Robinson et al, 2014 [13]| R, DB, 12 weeks; atorvastatin was not masked | Western countries   | Adults with a screening LDL-C level ≥ 150 mg/dL with no statin, or ≥ 100 mg/dL with non-intensive statin, or ≥ 80 mg/dL with intensive statin, and fasting TG ≤ 400 mg/dL. | Atorvastatin 80 mg/day, n = 110 | 2.52 ± 0.88 (97.5 ± 34.08)       | 2.71 ± 1.08 (104.8 ± 41.66)           |
|                         |                 |                     |                                                                                      | Atorvastatin 80 mg/day + ezetimibe 10 mg/day, n = 110 | 2.47 ± 0.72 (95.5 ± 27.70)       | 2.04 ± 1.31 (78.85 ± 50.89)           |
| Yang et al, 2017 [25]   | R, DB, 12 weeks | South Korean        | Adults with high or moderately high cardiovascular disease as classified by NCEP ATP III | Rosuvastatin 20 mg/day, n = 39 | 3.88 ± 0.98 (150 ± 38)         | 1.68 ± 0.72 (65 ± 28)                |
|                         |                 |                     |                                                                                      | Rosuvastatin 20 mg/day + ezetimibe 10 mg/day, n = 42 | 3.82 ± 0.78 (148 ± 30)         | 1.34 ± 0.62 (52 ± 24)                |

± indicates standard deviation. CHD: coronary heart disease; DB: double-blinded; HDL-C: high-density lipoprotein cholesterol; LDL-C: low-density lipoprotein cholesterol; MC: multicenter; NCEP ATP III: the third report of the expert panel on detection, evaluation, and treatment of high blood cholesterol in adults; OL: open-label; PC: placebo-controlled; R: randomized; SC: single center; TG: triglyceride.
The overall risk of bias was low or unclear, except for four open-label trials with a high risk of performance bias due to the nature of their study design (Fig. 2). Visual inspection of the funnel plot (Supplementary Material 2, www.cardiologyres.org) suggests some extent of asymmetry, but no statistically significant publication bias among the included trials was identified (Egger’s test, P = 0.702; Begg’s test, P = 0.282).

The MD in LDL-C levels and sensitivity analyses

Compared to the high-intensity statin monotherapy, the MD in LDL-C levels with a high-intensity statin plus ezetimibe was -14.00% (95% CI: -17.78 to -10.22; P < 0.001) (Fig. 3). The result was associated with a moderate degree of heterogeneity (I^2 = 66%, P < 0.001). The exclusion sensitivity analysis showed that the heterogeneity was mainly contributed by the study of Robinson 2014 (Table 2). When this study was omitted, the MD in LDL-C levels was changed to -12.09% (95% CI: -14.64 to -9.55; P < 0.001) with insignificant heterogeneity (I^2 = 21%).

Subgroup analysis and meta-regression

Subgroup analysis yielded non-significant differences between the types of high-intensity statins (P = 0.204), the doses of high-intensity statins (P = 0.838) and the trial regions (P = 0.109). The meta-regression analysis demonstrated a significant effect of baseline LDL-C levels with a coefficient of 0.123 (standard error = 0.044; P = 0.006). However, it should be noted that the significance was eliminated after removing the study of Robinson 2014 (P = 0.055).

Discussion

Here we reported a meta-analysis of 14 clinical trials with 2,007 patients included. The study found that adding ezetimibe to high-intensity statin led to a 14% incremental lowering in LDL-C levels compared to high-intensity statin monotherapy, which was statistically significant. This additional lowering of LDL-C levels was irrespective of the types of high-intensity statin or the trial regions. Also doubling the dose of high-intensity statins did not influence the effect of ezetimibe significantly. Our findings support the statement about the efficacy of ezetimibe in reducing LDL-C levels in 2018 AHA/ACC Cholesterol Guideline.

The trend of our findings was consistent with the results of the IMPROVE-IT trial. In the IMPROVE-IT trial [3], ezetimibe 10 mg daily additionally decreased LDL-C levels by 24% (0.43 mmol/L (16.7 mg/dL), P < 0.001) compared to placebo when added to background therapy of simvastatin 40 mg daily. Unlike the IMPROVE-IT trial, our meta-analysis identified a less percentage change in LDL-C levels, indicating that the impact of ezetimibe on reducing LDL-C levels may be lessened by co-administration with a high-intensity statin.

A similar pattern was observed in the ENHANCE trial as well. In the ENHANCE trial [26], ezetimibe added to background therapy of simvastatin 80 mg daily demonstrated an additional reduction in LDL-C levels by 16.5% (P < 0.01) compared to placebo in patients with familial hypercholesterolemia. The potency of simvastatin 80 mg was slightly less than atorvastatin 40 mg with a roughly 48% LDL-C reduction. The ENHANCE trial resulted in a less percentage change in LDL-C levels compared to the IMPROVE-IT trial, but slightly more percentage change in LDL-C levels compared to our meta-analysis. The differences between these three studies may
also suggest that the impact of ezetimibe on reducing LDL-C levels depends on the intensity of background statin therapy.

In the previous meta-analysis studies, considerably more reduction of LDL-C levels was observed when adding ezetimibe to low- to moderate-intensity statins. In the study of Mikhailidis et al [4], a -23.6% reduction of LDL-C levels was reported when adding ezetimibe to low- to moderate-intensity statin (simvastatin 10 - 20 mg and atorvastatin 10 - 20 mg). In Mikhailidis et al’s later study [5] comparing the efficacy of the addition of ezetimibe to statin vs. doubling statin, the addition of ezetimibe to statin therapy resulted in a roughly -20% greater reduction of LDL-C levels after adjusted the effect of doubling statin dose with the 6% rule. All the included trials used low to moderate statins (atorvastatin 10 - 20 mg and sim-

**Table 2. Exclusion Sensitivity Analysis**

| Study name                  | Mean difference | Standard error | Lower limit | Upper limit | z-value | P-value |
|-----------------------------|-----------------|----------------|-------------|-------------|---------|---------|
| Azar et al, 2011 [15]       | -14.088         | 2.005          | -18.018     | -10.158     | -7.026  | < 0.001 |
| Ballantyne et al, 2003_40 [16] | -14.250         | 2.071          | -18.309     | -10.191     | -6.881  | < 0.001 |
| Ballantyne et al, 2003_80 [16] | -14.457         | 2.067          | -18.508     | -10.406     | -6.995  | < 0.001 |
| Ballantyne et al, 2007 [17] | -14.099         | 2.090          | -18.196     | -10.002     | -6.735  | < 0.001 |
| Barbosa et al, 2013 [18]    | -13.895         | 1.998          | -17.811     | -9.979      | -6.955  | < 0.001 |
| Blom et al, 2014 [10]       | -14.323         | 2.015          | -18.271     | -10.374     | -7.109  | < 0.001 |
| Hong et al, 2018 [19]       | -13.924         | 2.098          | -18.035     | -9.812      | -6.637  | < 0.001 |
| Hougaard et al, 2017 [20]   | -14.400         | 2.076          | -18.469     | -10.330     | -6.935  | < 0.001 |
| Kim et al, 2016 [21]        | -14.562         | 2.060          | -18.600     | -10.524     | -7.068  | < 0.001 |
| Kim et al, 2018 [22]        | -14.047         | 2.144          | -18.249     | -9.844      | -6.552  | < 0.001 |
| Lins et al, 2014 [23]       | -13.784         | 1.999          | -17.702     | -9.866      | -6.895  | < 0.001 |
| Malina et al, 2015 [24]     | -14.101         | 2.014          | -18.048     | -10.154     | -7.002  | < 0.001 |
| Nicholls et al, 2017 [12]   | -13.091         | 1.850          | -16.716     | -9.465      | -7.076  | < 0.001 |
| Robinson et al, 2014 [13]   | -12.094         | 1.299          | -14.641     | -9.547      | -9.307  | < 0.001 |
| Yang et al, 2017 [25]       | -14.559         | 2.010          | -18.499     | -10.620     | -7.244  | < 0.001 |
vastatin 10 - 40 mg), except one trial with atorvastatin 40 mg. In the most comprehensive meta-analysis involving 35 trials [9], the addition of ezetimibe led to an MD in LDL-C of -13.62%, -14.71% and -14.96%, compared to doubling the starting dose of simvastatin, atorvastatin and rosuvastatin, respectively. The results were consistent with Mikhailidis et al’s study, and an approximately 20% additional lowering of LDL-C was achieved if the effect of doubling dose was adjusted with the 6% rule. Notably, among the 35 included trials, only two trials used high-intensity statin as the background therapy (atorvastatin 40 mg [27]; atorvastatin 10 mg or 20 mg or 40 mg or 80 mg [28]). In a pooled analysis of over 21,000 subjects from 27 clinical trials [29], a significant incremental reduction of -23.4% in LDL-C levels was attributable to the addition of ezetimibe compared to the control groups. Similar to the meta-analysis studies, the low- to moderate-intensity statin was found in the majority of the included trials in this pooled analysis. Overall, adding ezetimibe to low- to moderate-intensity statin was associated with an approximately 20-24% extra reduction of LDL-C levels, which was greater than the MD reported in our meta-analysis of adding ezetimibe to high-intensity statin therapy. Taking into account the results of our meta-analysis, the variation in the efficacy of ezetimibe on LDL-C levels could be explained by the different strengths of the background statin therapy.

Though more patients achieved a target LDL-C level of <1.8 mmol/L (70 mg/dL) when adding ezetimibe to the high-intensity statin therapy in the included trials, it was well established that the ASCVD risk reduction was associated with the magnitude of the lowering of LDL-C levels [30-32]. As described in the IMPROVE-IT trial, adding ezetimibe to a moderate-intensive statin was associated with a 6.4% relative risk reduction in cardiovascular outcomes (absolute risk reduction, 2.0%; hazard ratio, 0.936; 95% CI, 0.89 - 0.99; P = 0.016). While other factors also play an important role in influencing the cardiovascular outcome, the percentage reduction in LDL-C levels from baseline was linearly correlated with ASCVD rates [31]. Hence, it was very unlikely that the proportional benefit of ezetimibe in reducing major vascular events, translated from the 14% reduction in LDL-C levels, could be the same as the modest benefit reported in the IMPROVE-IT trial. We hypothesized that the magnitude of the clinical benefit of adding ezetimibe to a high-intensity statin in reducing cardiovascular events might not be clinically significant. Unfortunately, at the time of writing, there is no good evidence for the effectiveness of ezetimibe in reducing cardiovascular events with the background high-intensity statin therapy. Without any patient-level data, the clinical significance of adding ezetimibe to high-intensity statin therapy in reducing cardiovascular events remains unclear. Though ezetimibe is well tolerated when combined with statin therapy, the cardiovascular benefits of adding ezetimibe to the background high-intensity statin therapy beyond lipids lowering have to be examined to support the use of ezetimibe.

Limitations

Our study has several limitations. First, this meta-analysis is limited by the small number of included trials, the small sample size of each trial and the various trial settings, as shown in Table 1. Second, some of the included trials about PCSK9 inhibitors or CETP inhibitors were not designed to evaluate the effectiveness of ezetimibe on LDL-C levels. Third, the use of statins or ezetimibe was not masked in two trials in addition to four open-label trials, potentially increased risk of performance risk. Fourth, the moderate degree of heterogeneity caused by the study of Robinson 2014 might be explained by the small-study effects, as there were only about 100 patients in each group. Without any additional information or explanation provided by the article, we were unable to find a conclusive reason of the heterogeneity. Thus, we accept the existence of the heterogeneity and the potential greater uncertainty within our results. Taken together, our results should be interpreted with caution.

Conclusions

In summary, adding ezetimibe to high-intensity statin therapy provided a significant but attenuated additional reduction in LDL-C levels. However, whether this further lowering of LDL-C levels achieved by adding ezetimibe to a high-intensity statin would lead to a benefit in cardiovascular outcomes needs further investigation. Our study bridges the gap in the medical literature of the clinic role of ezetimibe with high-intensity statin therapy, and reinforces the statement in the 2018 AHA/ACC Cholesterol Guideline about the efficacy of ezetimibe in reducing LDL-C levels.

Supplementary Material

Suppl 1. Detailed Search Strategy Used in Each Database.
Suppl 2. Funnel plot of the 14 studies included in the meta-analysis.

Acknowledgments

None to declare.

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Dr. Egolum consults/advisory board for Akcea Therapeutics, Alnylam, AstraZeneca and Pfizer. Dr. Ling serves on an advisory board for Alnylam. The remaining authors have nothing to disclose.

Conflict of Interest

None to declare.

Informed Consent

Not applicable.
Author Contributions

HL contributed to the conception and design of the work. JL, HP and MC contributed to the data acquisition, analysis, and interpretation of data for the work. JL and HL drafted the manuscript. UE critically revised the manuscript. All authors gave final approval and agree to be accountable for all aspects of the work ensuring integrity and accuracy.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

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