The efficacy and safety of triple inhaled treatment in patients with chronic obstructive pulmonary disease: a systematic review and meta-analysis using Bayesian methods

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Purpose: Although tiotropium (TIO) and inhaled corticosteroid (ICS)/long-acting β-agonists are frequently prescribed together, the efficacy of “triple therapy” has not been scientifically demonstrated. We conducted a systematic review and meta-analysis using Bayesian methods to compare triple therapy and TIO monotherapy.

Methods: We searched the MEDLINE, EMBASE, and Cochrane Library databases for randomized controlled trials comparing the efficacy and safety of triple therapy and TIO monotherapy in patients with chronic obstructive pulmonary disease (COPD). We conducted a meta-analysis to compare the effectiveness and safety of triple therapy and TIO monotherapy using Bayesian random effects models.

Results: Seven trials were included, and the risk of bias in the majority of the studies was acceptable. There were no statistically significant differences in the incidence of death and acute exacerbation of disease in the triple therapy and TIO monotherapy groups. Triple therapy improved the prebronchodilator forced expiratory volume in 1 second (mean difference [MD], 63.68 mL; 95% credible interval [CrI], 45.29–82.73), and patients receiving triple therapy showed more improvement in St George Respiratory Questionnaire scores (MD, −3.11 points; 95% CrI, −6.00 to −0.80) than patients receiving TIO monotherapy. However, both of these differences were lower than the minimal clinically important difference (MCID). No excessive adverse effects were reported in triple therapy group.

Conclusion: Triple therapy with TIO and ICSs/long-acting β-agonists was only slightly more efficacious than TIO monotherapy in treating patients with COPD. Further investigations into the efficacy of new inhaled drugs are needed.

Keywords: inhaled long-acting muscarinic antagonists (LAMAs), inhaled corticosteroids (ICSs), inhaled long-acting β2-agonists (LABAs), chronic obstructive pulmonary disease (COPD)

Introduction

Inhaled drugs, including inhaled long-acting muscarinic antagonists (LAMAs), inhaled corticosteroids (ICSs), and inhaled long-acting β2-agonists (LABAs), are the principal therapeutic options for patients with chronic obstructive pulmonary disease (COPD). Until recently, tiotropium (TIO) was the only LAMA available, and it remains the most commonly used LAMA. Currently, these three classes of drugs (TIO, ICS, and LABA) are frequently prescribed together as “triple therapy”; however, there is insufficient scientific evidence demonstrating the efficacy of this combination. Only a few rigorous systematic reviews, supporting the efficacy of triple therapy, including improvements in the health status of the patient and reductions in the future risk of the patient,
have been published. In addition, these reviews have limitations. Only a few clinical trials have been conducted to investigate the benefits of triple therapy, and adverse outcomes, including death, are rarely reported in those studies. The assumption of normality, which may not hold for small studies, is necessary to construct confidence intervals, and continuity correction is required if there are zero events. Bayesian approaches to meta-analysis could overcome some of these issues. Moreover, Bayesian meta-analysis can provide a probabilistic interpretation of the treatment effect of interest and a probability of the effect being larger (or smaller) than a specific value. Bayesian meta-analysis can also be useful when evaluating whether the magnitude of efficacy is greater than the minimal clinically important difference (MCID). Thus, we conducted a systematic review using Bayesian methods to compare the efficacy and safety of triple therapy and TIO monotherapy.

Methods
To conduct this review, we followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines and the BayesWatch guidelines for reporting studies using Bayesian methods.

Data search and selection criteria
We searched the MEDLINE, EMBASE, and Cochrane Library databases (search date: November 2, 2014). The search terms were “COPD” AND “LAMA” AND (“ICS” AND “LABA”) OR ICS/LABA) AND randomized protocol design. LAMAs included TIO, aclidinium, and glycopyrrolate. LABAs included salmeterol (SAL), formoterol (FOR), vilanterol, or indacaterol. ICSs included beclomethasone, budesonide (BUD), fluticasone propionate (FP), fluticasone furoate, triamcinolone, mometasone, and flunisolide. ICS/LABA combination drugs included FP/SAL, BUD/FOR, and fluticasone furoate/vilanterol. Additional details on our search strategy are provided in the Supplementary materials.

The study selection criteria were as follows: 1) randomized controlled trials (RCTs); 2) studies with adults aged >18 years and diagnosed with COPD; 3) studies comparing triple therapy with nontriple therapy; 4) trials lasting at least 4 weeks; 5) studies reporting at least one of the following outcomes: mortality, annual rate of decline of the forced expiratory volume in the first second (FEV₁), acute exacerbations, changes in prebronchodilator or trough FEV₁, changes in quality of life (St George Respiratory Questionnaire [SGRQ]), changes in the dyspnea scale using quantitative questionnaires (Chronic Respiratory Disease Questionnaire, Baseline/Transition Dyspnea Index, modified Medical Research Council, visual analog scale, numeric rating scale), and safety data (serious adverse events [SAE] and pneumonia); and 6) studies published in the English language. Studies with duplicate data were excluded.

Data extraction and assessment of risk of bias
Two authors (CHL and MSK) independently reviewed the titles, abstracts, and citations of the studies. After screening potentially relevant studies, they independently evaluated full reports for the eligibility based on the study design, intervention, and outcomes. The authors of five studies with missing required data were contacted to obtain additional information on outcomes; two of these authors provided the desired information. In studies with missing standard deviations for changes from baseline in continuous variables, we imputed standard deviations by calculating a correlation coefficient from a study for which we knew these details.

To assess the risk of bias of each study, the Cochrane risk of bias tool was applied. This assessment included the following: 1) the adequacy of sequence generation; 2) allocation concealment; 3) blinding of the participants, personnel, and outcome assessors; 4) incomplete outcome data; 5) selective outcome reporting; and 6) other biases. Any disagreements were resolved by discussion until a consensus was reached.

Statistical analysis
We used Bayesian random effects models to compare the efficacy and safety of triple therapy and TIO monotherapy. In these models, an assumption of normality and continuity correction factors are not required because these methods are well suited to small studies and rare events. We estimated the relative risk (RR) for the binary outcome data or the mean difference (MD) for continuous variables using the posterior mean and corresponding 95% credible interval (CrI), which is the Bayesian equivalent of a confidence interval in classical analyses. We calculated posterior probabilities of the RR being larger or smaller than 1 (denoted \( P(\text{RR}>1) \) or \( P(\text{RR}<1) \), respectively). We also calculated posterior probabilities of the MD being larger or smaller than 0 (denoted \( P(\text{MD}>0) \) and \( P(\text{MD}<0) \), respectively). Additionally, we calculated Bayesian probabilities that the MD was greater than the MCID. The hypothesis of interest was supported if a posterior probability was greater than 0.9.
Inverse-gamma distributions, normal distributions with a mean of zero and large variance, and uniform distributions were considered noninformative priors for parameters. We used Review Manager 5.3 (Cochrane Collaboration, Oxford, UK), R 3.1.2 and WinBUGS 1.4 (Medical Research Council, Cambridge, and Imperial College School of Medicine, UK) software for our analyses. Three chains were considered to detect convergence. In each chain, the first 10,000 iterations were discarded to remove the influence of the initial value, and sampling from 10,000 additional iterations was used to generate summary statistics such as the posterior mean and 95% CrI. For certain analyses, every 10th or 30th number was extracted from the 10,000 samples to remove autocorrelations as needed. Gelman and Rubin statistics, Monte Carlo error, and autocorrelation plots were used to establish convergence of the Markov Chain Monte Carlo method. We performed sensitivity analyses to assess the impact of using different prior distributions. If the posterior median rather than the posterior mean of the between-study standard deviation was greater than one, then heterogeneity of the effects across studies was considered to exist, as the posterior mean is likely to have a skewed distribution. Publication bias was not formally assessed because each analysis included fewer than ten studies.

**Results**

Figure 1 shows a flowchart describing our study selection process. Of 1,777 screened references, 39 studies were reviewed in further detail. Thirty-one of these studies were excluded for various reasons, including duplicate data, non-English language, and a short study duration. Finally, seven studies with a total of 2,122 subjects (triple therapy, n=1,052; TIO only, n=1,070) were included in this meta-analysis. The characteristics of the included studies are presented in Table 1. The duration of the studies ranged from 4 to 52 weeks, with most study durations ranging from 12 to 24 weeks. Four studies were sponsored by a pharmaceutical company. Most of the studies exhibited a low risk of bias according to the six bias assessment scores of the Cochrane Instrument.

**Risk of bias within studies**

Figure 2 shows a graph and summary of the risk of bias assessment. Most of the studies were judged to have a low risk of bias for random sequence generation, incomplete outcome data, and selective reporting. However, evaluations of allocation concealment and blinding of the participants, personnel and outcome assessors were limited, and some of the RCTs were judged to have a high or unclear risk of bias for these parameters.

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**Figure 1** Flowchart for the selection of studies.

**Abbreviations:** COPD, chronic obstructive pulmonary disease; RCT, randomized controlled trial; ICS, inhaled corticosteroid; LABA, long-acting β₂-agonist.
Efficacy outcomes

In the three studies\textsuperscript{4,8,10} that reported the number of deaths, no statistically significant difference in mortality was found between the triple therapy group and the TIO monotherapy group (RR, 1.46; 95% CrI, 0.13–5.17; \( P \) (RR, 1) = 0.47). None of the studies reported rates of lung function decline. Four trials\textsuperscript{4,6,8,10} reported the number of acute exacerbations in patients receiving triple therapy or TIO monotherapy. We found that triple therapy provided a nonsignificant benefit in reducing the incidence of acute exacerbations.

Table 1 Included studies

<table>
<thead>
<tr>
<th>Authors</th>
<th>RCT design</th>
<th>F/U weeks</th>
<th>Age\textsuperscript{a}</th>
<th>FEV\textsubscript{1} %\textsuperscript{a}</th>
<th>Pack-years\textsuperscript{a}</th>
<th>Comparison</th>
<th>Primary outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aaron et al\textsuperscript{4}</td>
<td>Double-blind</td>
<td>52</td>
<td>67.7</td>
<td>41.8</td>
<td>50.3</td>
<td>TIO + FP (1,000 ( \mu )g/d)/SAL (n=145) vs TIO (n=156)</td>
<td>Exacerbation of disease</td>
</tr>
<tr>
<td>Cazzola et al\textsuperscript{5}</td>
<td>Double-blind</td>
<td>12</td>
<td>65.8</td>
<td>38.1</td>
<td>50.9</td>
<td>TIO + FP (1,000 ( \mu )g/d)/SAL (n=29) vs TIO (n=26)</td>
<td>Change in predose FEV\textsubscript{1}</td>
</tr>
<tr>
<td>Welte et al\textsuperscript{10}</td>
<td>Double-blind</td>
<td>12</td>
<td>62.5</td>
<td>37.9</td>
<td>37</td>
<td>TIO + BUD (640 ( \mu )g/d)/FOR (n=329) vs TIO (n=331)</td>
<td>Change in predose FEV\textsubscript{1}</td>
</tr>
<tr>
<td>Hanania et al\textsuperscript{6}</td>
<td>Double-blind</td>
<td>24</td>
<td>61.2</td>
<td>56.7</td>
<td>55.1</td>
<td>TIO + FP (500 ( \mu )g/d)/SAL (n=173) vs TIO (n=169)</td>
<td>Change in predose FEV\textsubscript{1}</td>
</tr>
<tr>
<td>Jung et al\textsuperscript{8}</td>
<td>Open-label</td>
<td>24</td>
<td>67.4</td>
<td>47.5</td>
<td>NR</td>
<td>TIO + FP (500 ( \mu )g/d)/SAL (n=237) vs TIO (n=242)</td>
<td>Change in predose FEV\textsubscript{1}</td>
</tr>
<tr>
<td>Hoshino and Ohtawa\textsuperscript{7}</td>
<td>Open-label</td>
<td>16</td>
<td>71.2</td>
<td>NR</td>
<td>57.7</td>
<td>TIO + FP (500 ( \mu )g/d)/SAL (n=15) vs TIO (n=15)</td>
<td>Airway dimension</td>
</tr>
<tr>
<td>Maltais et al\textsuperscript{9}</td>
<td>Double-blind</td>
<td>4</td>
<td>62.7</td>
<td>54.7</td>
<td>NR</td>
<td>TIO + FP (500 ( \mu )g/d)/SAL (n=124) vs TIO (n=131)</td>
<td>Exercise endurance time</td>
</tr>
</tbody>
</table>

Note: Mean value.

Abbreviations: RCT, randomized controlled trial; TIO, tiotropium; FP, fluticasone propionate; SAL, salmeterol; BUD, budesonide; FOR, formoterol; NR, not recorded; F/U, follow-up; FEV\textsubscript{1}, forced expiratory volume in the first second.

Figure 2 Risk of bias summary and table.
(RR, 0.80; 95% CrI, 0.35–1.63; P (RR <1) =0.84). All of the trials investigating lung function reported a significant improvement in the prebronchodilator FEV₁ in the patients receiving triple therapy, and the calculated pooled MD was 63.68 mL (95% CrI, 45.29–82.73) (P [MD >0 mL] =1.0). However, the magnitude of this improvement was less than the MCID (100–140 mL)²⁴ (P [MD >100 mL] =0.002). The three studies⁴,⁶,¹⁰ evaluating the SGRQ score reported a significant improvement in the mean change from baseline in patients receiving triple therapy, with a difference of −3.11 points (95% CrI, −6.00 to −0.80; P [MD <0 points] =0.99). However, the magnitude of this change was not over the MCID (−4.0)²⁵ (P [MD <−4.0 points] =0.18) (Table 2 and Figure 3).

Safety outcomes (adverse events)

Patients receiving triple therapy did not experience significantly more adverse events than patients receiving monotherapy (RR, 1.12; 95% CrI, 0.87–1.40; P [RR >1] =0.84). The total number of SAEs was significantly lower in the triple therapy group (RR, 0.62; 95% CrI, 0.17–1.25; P [RR <1] =0.93) than in the TIO monotherapy group. The risk of pneumonia was not significantly higher in the triple therapy group than in the TIO monotherapy group (RR, 1.07; 95% CrI, 0.05–4.28; P [RR >1] =0.27). The incidence of oral candidiasis was not significantly higher in the triple therapy group (RR, 3.63; 95% CrI, 0.46–12.82; P [RR >1] =0.88) (Table 3).

The posterior median of the between-study standard deviation in our meta-analysis was smaller than one, indicating a lack of heterogeneity. We did not find evidence against convergence, and we did not see any substantial differences. However, the CrI and probability changed slightly by changing the specifications of the prior distribution.

Discussion

The goals of treating patients with COPD include improving the health status of the patient (lung function, quality of life, and exercise capacity) and decreasing future risks of acute exacerbation, lung function decline, and death.¹ TIO is known to improve the quality of life and lung function of patients with COPD,²⁶ reduce the likelihood of experiencing acute exacerbation,²⁶–²⁸ delay declines in lung function in patients with stage II disease,²⁹ and possibly decrease mortality.³⁰ ICS and LABAs are commonly used in combination in a single device (ICS/LABA). Large clinical trials showed that ICS/LABA combinations, including FP/SAL and BUD/FOR, improve the quality of life and lung function of patients with COPD,³¹–³⁴ reduce the incidence of acute exacerbation,³¹–³⁴ slow declines in lung function,³⁵ and possibly reduce mortality.³¹ However, the efficacy and safety of triple therapy combining TIO and ICS/LABA have rarely been comprehensively investigated, especially in the context of accomplishing these treatment goals. As mentioned in the Introduction, a classical meta-analysis may not be adequate to evaluate the benefits of triple therapy. Because only a few clinical trials⁴–¹⁰ were conducted and rare events were included, the assumption of normality may not hold for the analysis. Therefore, we used Bayesian approaches to reduce some of these problems.¹¹,¹² Moreover, Bayesian meta-analysis can also be useful when evaluating whether the magnitude of efficacy is greater than the MCID³⁴ by using posterior probabilities.¹²,¹³

In our systematic review, we found limited evidence indicating that triple therapy can reduce future risks. Although we used Bayesian methods to analyze the incidence of rare events, we did not find that triple therapy reduced mortality to a greater extent than TIO monotherapy. None of the studies in our review investigated declines in lung function as an outcome. Additionally, we did not find that triple therapy significantly reduced the incidence of acute exacerbation (P [RR <1] =0.84). Welte et al⁸ reported that triple therapy produced a reduction in the incidence of acute exacerbation; however, this effect was not significantly different from that found by other studies in our meta-analysis.

Table 2 Summary of efficacy outcomes

<table>
<thead>
<tr>
<th>Outcome variable</th>
<th>Number of studies</th>
<th>Total number of patients included</th>
<th>RR or MD (95% CrI)</th>
<th>Posterior probability</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Death hectic</td>
<td>3</td>
<td>705/725</td>
<td>1.46 (0.13–5.17)</td>
<td>P (RR &lt;1) =0.47</td>
<td>1.72</td>
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<tr>
<td>Acute exacerbation</td>
<td>4</td>
<td>870/888</td>
<td>0.80 (0.35–1.63)</td>
<td>P (RR &lt;1) =0.84</td>
<td>0.37</td>
</tr>
<tr>
<td>Changes in FEV₁</td>
<td>6</td>
<td>915/928</td>
<td>63.68 mL (45.29–82.73)</td>
<td>P (MD &gt;0 mL) =1.0</td>
<td>9.69</td>
</tr>
<tr>
<td>Changes in SGRQ</td>
<td>3</td>
<td>697/711</td>
<td>−3.11 points (−6.00 to −0.80)</td>
<td>P (MD &lt;0 points) =0.99</td>
<td>2.16</td>
</tr>
</tbody>
</table>

Notes: RR for binary outcomes, MD for continuous outcomes. Abbreviations: FEV₁, forced expiratory volume in the first second; TIO, tiotropium; SGRQ, St George Respiratory Questionnaire; RR, relative risk; CrI, credible interval; MD, mean difference; SD, standard deviation (between studies).

We refer to the full-text article for detailed discussion.
Kwak et al

A

<table>
<thead>
<tr>
<th>Study</th>
<th>TIO + ICS/LABA</th>
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</tr>
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<tr>
<td></td>
<td>Events</td>
<td>Total</td>
</tr>
<tr>
<td>Aaron et al4</td>
<td>6</td>
<td>145</td>
</tr>
<tr>
<td>Jung et al5</td>
<td>0</td>
<td>231</td>
</tr>
<tr>
<td>Welte et al10</td>
<td>1</td>
<td>329</td>
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<tr>
<td>Overall</td>
<td></td>
<td></td>
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Estimate of heterogeneity between studies (SD = 0.87)

B

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<th>TIO monotherapy</th>
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<tr>
<td></td>
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<td>Total</td>
</tr>
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<td>Aaron et al4</td>
<td>87</td>
<td>145</td>
</tr>
<tr>
<td>Jung et al5</td>
<td>39</td>
<td>223</td>
</tr>
<tr>
<td>Hanania et al6</td>
<td>25</td>
<td>173</td>
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<tr>
<td>Welte et al10</td>
<td>25</td>
<td>329</td>
</tr>
<tr>
<td>Overall</td>
<td></td>
<td></td>
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</table>

Estimate of heterogeneity between studies (SD = 0.51)

C

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<tr>
<th>Study</th>
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</tr>
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<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Aaron et al4</td>
<td>86</td>
<td>217</td>
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<tr>
<td>Cazzola et al6</td>
<td>186</td>
<td>66</td>
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<tr>
<td>Hanania et al6</td>
<td>201</td>
<td>287</td>
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<tr>
<td>Jung et al5</td>
<td>171</td>
<td>207</td>
</tr>
<tr>
<td>Hoshino and Ohtawa1</td>
<td>123</td>
<td>339</td>
</tr>
<tr>
<td>Welte et al10</td>
<td>64</td>
<td>198</td>
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<td>Overall</td>
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Estimate of heterogeneity between studies (SD = 0.93)

D

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<thead>
<tr>
<th>Study</th>
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</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Aaron et al4</td>
<td>-8.6</td>
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<td>12.8</td>
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<td>Overall</td>
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Estimate of heterogeneity studies (SD = 0.39)

Table 3 Summary of adverse events

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Number of studies</th>
<th>Total number of patients included</th>
<th>RR (95% CrI)</th>
<th>Posterior probability</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Total</td>
<td>Mean</td>
<td>SD</td>
<td></td>
</tr>
<tr>
<td>Total adverse events</td>
<td>5</td>
<td>800</td>
<td>813</td>
<td></td>
<td>1.12 (0.87–1.40) P (RR &gt; 1) = 0.84</td>
</tr>
<tr>
<td>Severe adverse events</td>
<td>6</td>
<td>1,031</td>
<td>1,051</td>
<td></td>
<td>0.62 (0.17–1.25) P (RR &lt; 1) = 0.93</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>4</td>
<td>829</td>
<td>856</td>
<td></td>
<td>1.07 (0.05–4.28) P (RR &gt; 1) = 0.27</td>
</tr>
<tr>
<td>Oral candidiasis</td>
<td>3</td>
<td>647</td>
<td>656</td>
<td></td>
<td>3.63 (0.46–12.82) P (RR &gt; 1) = 0.88</td>
</tr>
</tbody>
</table>

Abbreviations: TIO, tiotropium; RR, relative risk; CrI, credible interval; SD, standard deviation (between studies).
MCID. All of the trials investigating lung function reported significant improvements in the prebronchodilator FEV₁; however, the increase in FEV₁ was lower in magnitude than the MCID (100 mL) \( (P \ [MD > 100 \text{ mL}] = 0.002) \).\(^{24} \) Triple therapy also improved SGRQ scores; however, the magnitude of improvement was lower than the MCID \( (P \ [MD < -4.0 \text{ points} \ [\text{MCID}]) = 0.18) \). Thus, triple therapy can marginally improve the prebronchodilator FEV₁ and SGRQ scores of patients with COPD.

There are several possible explanations for the marginal difference in efficacy between triple therapy and TIO monotherapy. First, even though we used Bayesian methods, we were able to identify only a few trials, which could suggest the existence of bias. Second, the definition of acute exacerbation varied between trials. Third, TIO monotherapy is known to be efficacious; therefore, a ceiling effect may exist, and combining TIO with other inhaled drugs may not confer additional or synergistic effects. TIO is considered to be the most effective bronchodilator. TIO is superior to most LABAs in reducing the incidence of acute exacerbation,\(^{36,37} \) including indacaterol, a new LABA.\(^{38} \) TIO is also superior or at least equivalent to LABAs in improving the FEV₁ and SGRQ scores. Certain studies have reported that indacaterol is superior to TIO in improving the FEV₁; however, this finding was not confirmed in a large-scale trial or in the trial sponsored by the manufacturer of indacaterol.\(^{39} \) Furthermore, no studies using indacaterol were included in our systematic review because they lacked a triple therapy group. In fact, TIO monotherapy has been shown to be as effective as TIO + LABA dual therapy. Several RCTs\(^{40,41} \) and a meta-analysis\(^{41} \) showed no significant differences in the incidence of acute exacerbation in patients receiving TIO monotherapy and patients receiving a combination of TIO and LABAs. Although TIO + LABA dual therapy has been reported to be superior to TIO monotherapy in improving the FEV₁ and SGRQ scores,\(^{41} \) the MDs encountered in our analysis were lower than the MCID.

We found no clear evidence that triple therapy is significantly more efficacious in improving the health status of the patient and reducing future risks; however, triple therapy was also not associated with an increased incidence of adverse events. The total number of adverse events was nonsignificantly higher in the ICS/LABA + TIO group than in the TIO monotherapy group. However, significantly fewer SAEs occurred in the triple therapy group than in the TIO monotherapy group. Pneumonia, a critical possible adverse event in patients using ICS,\(^{31,42,43} \) was not found to occur more often in any particular group in this systematic review. Additionally, patients in the triple therapy group were not significantly more likely to experience oral candidiasis.

Our study has several strengths. First, the Bayesian methods we used are more appropriate for analyzing rare events and for assessing a small number of studies. Second, we could evaluate whether the effect of interest was greater than the MCID because Bayesian meta-analysis can provide a probability that the effect is larger (or smaller) than a specific value, which was overlooked in previous meta-analyses.\(^{12} \) We also acknowledge the limitations of our systematic review. First of all, studies have different study durations, which could lead to biased results. Inclusion of COPD patients with wide-range severity might also be a weak point, because mortality, lung function decline, and exacerbation can be affected by FEV₁ of patients.\(^{44-46} \) We initially intended to compare triple therapy and dual therapies; however, it was impossible to do so because no common outcomes were described in the available studies. We also attempted to include studies evaluating new ICSs, LABAs, and LAMAs; however, no studies compared these therapies with triple therapy. Additionally, we were not always able to obtain more detailed clinical information when contacting authors by email.

**Conclusion**

In conclusion, our systematic review using Bayesian meta-analysis showed that triple therapy with TIO and ICS/LABA was more efficacious than TIO monotherapy. However, the increase in efficacy was marginal, and the clinical relevance of the improvement was unclear. Further investigations evaluating new bronchodilators are needed.

**Acknowledgment**

Seoul National University Hospital Research Fund (2320130040 (2013-2423)).

**Author contributions**

CHL and MSK: conception and design of the study, selection of study, independent assessment of risk of bias, extraction, analysis, interpretation of the data, drafting of the manuscript, critical revision of the manuscript for important intellectual content, and final approval of the manuscript. HJK, EJJ, and EK: design of the study, analysis of data, interpretation of the data, critical revision of the manuscript for important intellectual content, and final approval of the manuscript.

**Disclosure**

The authors report no conflicts of interest in this work.
References


Supplementary materials

Details of the search strategy.

MEDLINE


2. ((((((((groups[tiab]) OR (trial[tiab]) OR (randomly[tiab]) OR (drug therapy[sh]) OR (placebo[tiab]) OR (randomized[tiab]) OR (controlled clinical trial[pt]) OR (randomized controlled trial[pt]))) NOT (animals[Mesh] NOT (humans[Mesh] AND animals[Mesh]))) 2865448


OR “Muscarinic Antagonists/therapeutic use”[Mesh]
OR “Muscarinic Antagonists/toxicity”[Mesh]) OR “tiotropium”[Supplementary Concept]) OR (((“Glycopyrronium Bromide”[tiab] OR “aclidinium bromide”[tiab]))
OR (((“aclidinium bromide”[Supplementary Concept]) OR “Glycopyrrolate”[Mesh])) 17656
7. 1 AND ((3 AND 4) OR 5) AND 6) 1324
8. 7 AND 2 1123
EMBASE
1. ‘momentasone’:ab, ti OR ‘ics’:ab, ti OR ‘inhaled
corticosteroid’:ab, ti OR ‘budesonide’:ab, ti OR ‘glucocorticoids’:ab, ti OR ‘adrenal cortex hormones’:ab,
ti OR ‘corticosteroids’:ab, ti OR ‘pulmicort’:ab, ti OR ‘horacort’:ab, ti OR ‘rhinocort’:ab, ti OR ‘volon’:ab,
ti OR ‘aristocort’:ab, ti OR ‘glucocorticoid’:ab, ti OR ‘flunisolide’:ab, ti, OR ‘tiotropium’:ab, ti
2. ‘glucocorticoid’/exp OR ‘corticosteroid’/exp OR ‘steroid’/exp OR ‘pregnane derivative’/exp OR ‘androstan
de derivative’/exp NOT (‘hydroxycorticosteroid’/exp OR ‘mineralocorticoid’/exp) 1254293
3. 1 OR 2 1280780
4. ‘onbrez’/exp OR ‘onbrez’:ab, ti 607
5. ‘beta 2 adrenergic receptor stimulating agent’/exp
OR ‘beta 2 adrenergic receptor stimulating agent’:ab, ti 10767
6. ‘indacaterol’/exp OR ‘indacaterol’:ab, ti 317
7. 4-6/OR 11198
8. 3 AND 7 7917
9. (‘corticosteroid’/exp OR ‘antiasthmatic agent’/exp)
AND (‘beta 2 adrenergic receptor stimulating agent’/exp OR ‘bronchodilating agent’/exp) AND ‘drug combina
tion’/exp 1879
10. ‘fluticasone propionate plus salmeterol’/exp OR ‘fluti
casone propionate plus salmeterol xinafoate’/exp OR ‘seretide’:ab, ti OR ‘androstane derivative’/exp OR
‘budesonide plus formoterol’/exp OR ‘budesonide plus formoterol fumarate’/exp OR ‘symbicort’:ab, ti
OR ‘fluticasone/salmeterol’:ab, ti OR ‘budesonide/ formoterol’:ab, ti 7345
11. ‘fluticasone furoate plus vilanterol’/exp OR ‘fluticasone
propionate plus salmeterol’/exp OR ‘budesonide plus formoterol’/exp 2984
12. 9–11/OR 8942
13. 8 OR 12 15889
14. ‘cholinergic receptor blocking agent’/exp OR ‘scopol
amine derivative’/exp OR ‘muscarinic receptor blocking
agent’/exp OR ‘tiotropium bromide’/exp 150859
15. ‘spiriva’:ab, ti OR ‘tiotropium’:ab, ti OR ‘lama’:ab, ti
OR ‘long-acting muscarinic antagonist’:ab, ti 2268
16. ‘aclidinium bromide’/exp OR ‘glycopyrronium bro
mide’/exp 4527
17. ‘aclidinium bromide’:ab, ti OR ‘glycopyrronium
bromide’:ab, ti 177
18. 14–17/OR 151653
19. 13 AND 18 3997
20. ‘emphysema’/exp OR ‘emphysema’ OR ‘emphysema’:ab,
ti OR ‘chronic bronchitis’/exp OR ‘chronic bronchitis’
OR ‘chronic bronchitis’:ab, ti OR ‘obstructive lung
disease’/exp OR ‘obstructive lung disease’ OR ‘obstruc
tive lung disease’:ab, ti OR ‘obstructive pulmonary
disease’/exp OR ‘obstructive pulmonary disease’ OR
‘obstructive pulmonary disease’:ab, ti OR ‘obstruc
tive lung diseases’:ab, ti OR ‘chronic obstructive lung
disease’/exp OR ‘chronic obstructive lung disease’
OR ‘chronic obstructive lung disease’:ab, ti 125588
21. 19 AND 20 2073
22. ‘crossover procedure’/exp OR ‘crossover procedure’
OR ‘double blind procedure’/exp OR ‘double blind
procedure’ OR ‘randomized controlled trial’/exp OR ‘randomized controlled trial’ OR ‘single blind proce
dure’/exp OR ‘single blind procedure’ OR random* OR factorial* OR crossover* OR ‘cross over’ OR ‘cros
sover’ OR placebo* OR (doub* AND blind*) OR (singl*
AND blind*) OR assign* OR allocat* OR volunteer*
1591223
23. 21 AND 22 786

**COCHRANE**

1. ICS:ti, ab, kw (Word variations have been searched) 694
3. (inhal*) and (Corticosteroid* or cortico-steroid* or beclomethasone or beclazone or becotide or becloforte or budesonide or pulmicort* or fluticasone or flixotide or qvar or filair or aerobec or asmabec or becodisk* or triamcinolone or mometasone or flunisolide):ti, ab, kw (Word variations have been searched) 5191
5. 1–4/or 11759
6. onbrez or indacaterol:ti, ab, kw (Word variations have been searched) 141
7. adrenergic beta-2 receptor agonists 87
8. #6 or #7 277
9. (fluticasone):ti, ab, kw and (salmeterol):ti, ab, kw 965
10. (formoterol):ti, ab, kw and (budesonide):ti, ab, kw 643
11. (seretide):ti, ab, kw or (symbicort):ti, ab, kw or (fluticasone salmeterol):ti, ab, kw or (budesonide formoterol): ti, ab, kw 1557
12. MeSH descriptor: [Drug Combinations] explode all trees 9258
13. fluticasone furoate plus vilanterol:ti, ab, kw (Word variations have been searched) 1
14. MeSH descriptor Cholinergic Antagonists explode all trees 879
15. MeSH descriptor Scopolamine Derivatives explode all trees 868
16. MeSH descriptor Muscarinic Antagonists explode all trees 575
17. (spiriva):ab, ti OR (tiotropium):ab, ti OR (lama):ab, ti OR (long-acting muscarinic antagonist):ab, ti 629
18. “glycopyrronium bromide” or “aclidinium bromide”:
ti, ab, kw (Word variations have been searched) 239
19. MeSH descriptor Emphysema explode all trees 97
20. emphysema:ti, ab, kw OR chronic bronchitis:ti, ab, kw OR chronic obstructive lung disease:ti, ab, kw OR obstructive lung disease:ti, ab, kw OR obstructive pulmonary disease:ti, ab, kw 813
21. COPD:ti, ab, kw 6441
22. MeSH descriptor: [Pulmonary Disease, Chronic Obstructive] explode all trees 2349
23. (#19 OR #20 OR #21 OR #22) AND ((#5 AND #8) OR (#9 OR #10 OR #11 OR #12 OR #13)) AND (#14 OR #15 OR #16 OR #17 OR #18) 84