


Clinically Important Deterioration Among Patients with Chronic Obstructive Pulmonary Disease (COPD) Treated with Nebulized Glycopyrrolate: A Post Hoc Analysis of Pooled Data from Two Randomized, Double-Blind, Placebo-Controlled Studies

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Edward M Kerwin¹
Lindsey Murray²
Xiaoli Niu³
Carole Dembek³

¹Clinical Research Institute of Southern Oregon, Medford, OR, USA; ²Evidera, Bethesda, MD, USA; ³Sunovion Pharmaceuticals Inc., Marlborough, MA, USA

Purpose: Using a composite endpoint, pooled data from two 12-week Phase III placebo-controlled trials (GOLDEN 3, NCT02347761; GOLDEN 4, NCT02347774) were analyzed to determine whether glycopyrrolate inhalation solution (25 mcg and 50 mcg) administered twice daily (BID) via the eFlow[®] Closed System nebulizer (GLY) reduced the risk of clinically important deterioration (CID) in patients with moderate-to-very-severe COPD.

Methods: CID was defined as ≥ 100 -mL decrease from baseline in post-bronchodilator trough forced expiratory volume in one second (FEV₁), or ≥ 4 -unit increase in baseline St. George's Respiratory Questionnaire (SGRQ) total score, or moderate/severe exacerbation. The relative treatment effect of GLY versus placebo on the odds of CID (any and by component endpoints) was expressed as the odds ratio (OR) and 95% confidence interval (CI). Subgroups categorized by age ($<65/\geq 65$ years), sex, smoking status (current/former), long-acting beta agonist (LABA) use, FEV₁ ($<50\%/ \geq 50\%$), and peak inspiratory flow rate (PIFR) (<60 L/min/ ≥ 60 L/min) were analyzed.

Results: Compared to placebo, GLY 25 mcg and 50 mcg BID over 12 weeks significantly reduced the risk of CID by 50% (OR: 0.50 [0.37–0.68]) and 40% (OR: 0.60 [0.44–0.80]), respectively. Subjects treated with GLY 25 mcg BID were 59% less likely to experience CID in FEV₁ (OR: 0.41 [0.27–0.62]) and 48% less likely to perceive CID in health status (OR: 0.52 [0.37–0.73]). Statistically significant reductions were also observed at the higher dose. The incidence of moderate/severe exacerbations was low and comparable among the cohorts. GLY 25 mcg BID was significantly more effective than placebo ($p < 0.05$) in preventing CID irrespective of age, smoking status, LABA use, COPD severity, or PIFR. Subjects < 65 years (OR 0.45 [0.29–0.68]) and those with PIFR < 60 L/min (OR 0.36 [0.20–0.67]) exhibited the largest benefit.

Conclusion: Nebulized GLY over 12 weeks significantly reduced the risk of CID and provided greater short-term stability in patients with moderate-to-very-severe COPD.

Keywords: nebulized glycopyrrolate, clinically important deterioration, composite measure

Correspondence: Xiaoli Niu
Sunovion Pharmaceuticals Inc., 84
Waterford Dr, Marlborough, MA 01752,
USA
Tel +1 508 357 7863
Email xiaoli.niu@sunovion.com

Introduction

Chronic obstructive pulmonary disease (COPD) is a common condition characterized by frequently progressive airflow limitation, persistent respiratory symptoms, and high

morbidity and mortality.¹ Without effective management, the burden of pervasive symptoms, particularly dyspnea, can increase the risk of exacerbations, impair quality of life, and contribute to substantial health care costs.^{2,3} Recommended by the Global Initiative for Chronic Obstructive Lung Disease (GOLD), inhaled long-acting beta-agonists (LABA) and long-acting muscarinic antagonists (LAMA) are central to the treatment strategy for alleviating symptoms, minimizing the frequency and severity of exacerbations, and improving health status and exercise tolerance.¹ Lonhala[®] Magnair[®] (glycopyrrolate, GLY) inhalation solution (Sunovion Pharmaceuticals Inc., Marlborough, MA, USA) is a LAMA approved in the US for the long-term maintenance treatment of airflow obstruction in patients with COPD.⁴ The licensed recommended dose is 25 mcg administered twice daily (BID) using the eFlow[®] Closed System (Magnair[®], PARI Pharma GmbH, Starnberg, Germany), a hand-held vibrating membrane nebulizer. The nebulizer is designed to deliver 1.0 mL of GLY within three minutes via normal breathing rather than a single-breath actuation.^{5,6} Two replicate, 12-week, randomized, placebo-controlled Phase III trials in the US, Glycopyrrolate for Obstructive Lung Disease via Electronic Nebulizer (GOLDEN) 3 and GOLDEN 4 (NCT02347761 and NCT02347774, respectively), evaluated the safety and efficacy of nebulized GLY 25 mcg BID and 50 mcg BID (not licensed for commercial use) in patients with moderate-to-very-severe COPD (Figure 1). In both trials, GLY administered for 12 weeks was well tolerated and yielded statistically significant improvements from baseline over placebo in the forced expiratory volume in one second (FEV₁), forced vital capacity (FVC), and the St. George Respiratory Questionnaire (SGRQ) total score.⁶

Considering the progressive nature of COPD, it is essential to determine whether bronchodilator therapies can improve airway stability and prevent disease worsening. Exacerbations and rapid decline in lung function can signify poor long-term prognosis in COPD.^{7,8} Further, deterioration in health status over one year, as demonstrated by an increase in SGRQ score exceeding the within-patient clinically meaningful change threshold,⁹ has been associated with an increased risk of hospitalization and mortality.¹⁰ To complement efficacy assessments, a composite endpoint for clinically important deterioration (CID) in COPD has been used to provide a more comprehensive, multidimensional measure of treatment effectiveness. The CID captures changes in lung function, health status, and exacerbations that can reflect temporary fluctuations in disease state or more sustained worsening.^{11,12} A post hoc analysis of pooled data from the GOLDEN 3 and GOLDEN 4 trials was performed to examine whether nebulized GLY (25 mcg and 50 mcg) BID administered for 12 weeks decreased the risk of CID compared to placebo in patients with moderate-to-very-severe COPD.

Methods

Full methods and patient eligibility criteria for the GOLDEN-3 (NCT02347761) and GOLDEN-4 (NCT02347774) trials have been published elsewhere.⁶ In brief, the studies included male or female current or past smokers (≥ 10 pack-year) aged ≥ 40 years. Eligible patients had a clinical diagnosis of moderate-to-very-severe COPD by the GOLD classification and met the post-bronchodilator spirometry criteria at screening (ie, FEV₁ $\leq 80\%$ of predicted normal, FEV₁ > 0.7 L, and FEV₁/FVC ratio < 0.70).^{1,6} Patients with severe comorbidities or exacerbations

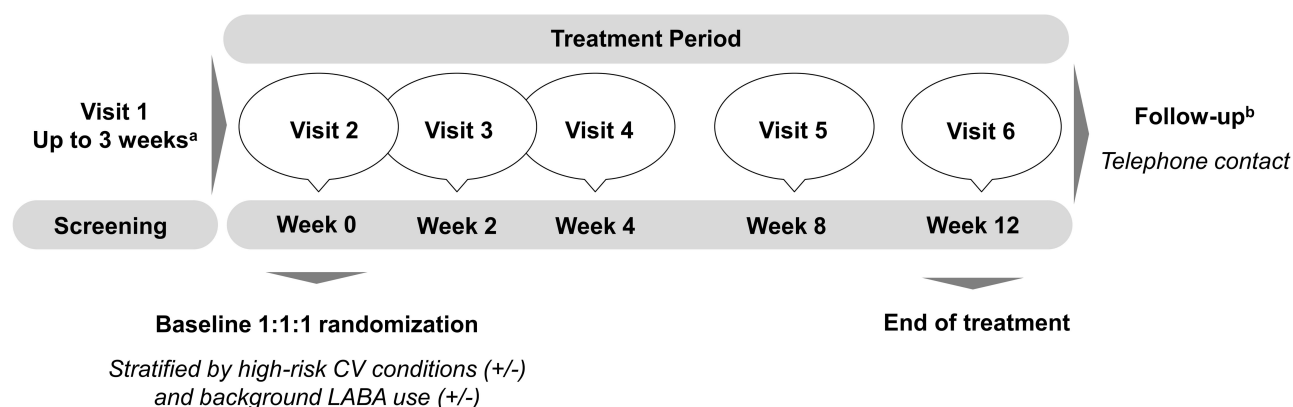


Figure 1 GOLDEN 3 and GOLDEN 4 study design.

Notes: ^aFlexible, between 1 and 7 days. ^bContacted by telephone 5 to 7 days after the last study visit.

Abbreviations: CV, cardiovascular; GOLDEN, Glycopyrrolate for Obstructive Lung Disease via Electronic Nebulizer; LABA, long-acting beta agonist.

requiring hospitalization or increased COPD treatments within six weeks of screening were excluded. Approximately 30% of subjects (limited by protocol) continued background LABA medications with or without inhaled corticosteroids (ICS). The study protocols were approved by the Quorum Review IRB North American (US and Canada) Board (Panel II) prior to patient enrollment, and were conducted in accordance with the protocols, International Council for Harmonization Good Clinical Practice guidelines and the Declaration of Helsinki. All subjects or their legal representatives provided written informed consent.⁶

The current study analyzed pooled data for efficacy assessments conducted at baseline and at the end of the study (week 12). Applying criteria described by Singh et al,¹³ subjects were considered to experience CID if any of the following events occurred: 1) a ≥ 100 -mL decrease from baseline in post-bronchodilator trough FEV₁; 2) a ≥ 4 -unit increase in the baseline SGRQ total score; or 3) a moderate or severe healthcare resource utilization (HCRU)-related exacerbation. An exacerbation was considered “moderate” if it required a clinic visit or treatment with antibiotics or oral corticosteroids with or without a clinic visit but no hospitalization, and “severe” if it necessitated an overnight stay in a hospital or emergency room. These component endpoints were based on the established within-patient clinically important change thresholds for FEV₁ and the SGRQ and an expert consensus panel definition for exacerbations.^{9,12,14–16}

Statistical Analysis

Statistical analyses were performed using SAS statistical software version 9.4 (SAS Institute Inc., Cary, NC). Analyses were conducted on the intent-to-treat population, defined as randomized patients who received at least one dose of the study treatment and had at least one post-dose pulmonary function assessment. Baseline patient demographics and clinical characteristics were summarized descriptively using the mean and standard deviation (SD) for continuous variables and the number of patients and proportion for categorical variables. The proportion of patients in each cohort experiencing CID and achieving the within-patient clinically meaningful change thresholds for each of the individual component endpoints was assessed at week 12.

Separate logistic regression models were analyzed to compare the risk of experiencing CID (ie, any CID and by the individual component measures) between each GLY cohort and the placebo (reference) cohort. Subgroup analyses were performed to examine the risk of experiencing CID in the cohorts stratified by baseline age (<65 or ≥ 65

years), sex (male or female), smoking status (current or former), background LABA use (yes/no), COPD severity (FEV₁ $\geq 50\%$ or $<50\%$), and peak inspiratory flow rate (PIFR) (<60 or ≥ 60 L/min). Due to the small number of patients with GOLD 4 classification, aggregate data for the GOLD 3 and GOLD 4 subgroups were analyzed in the subgroup with FEV₁ $<50\%$. All models were adjusted for the presence (yes/no) of high cardiovascular risk conditions and background LABA use, baseline post-bronchodilator trough FEV₁ (continuous variable), baseline SGRQ total score (continuous variable), and history (yes/no) of moderate or severe HCRU-related exacerbations in the prior 12 months. Results were expressed as the adjusted odds ratio (OR) and 95% confidence interval (CI). An OR >1.0 (excluding “1.0” in the 95% CI) indicated an increased risk of CID in a GLY cohort versus placebo, whereas an OR <1.0 (excluding “1.0” in the 95% CI) signified a decreased risk of CID. A p-value <0.05 indicated statistical significance.

Results

Baseline Characteristics of the Pooled Study Cohorts

The pooled study population consisted of 1293 subjects randomized to GLY 25 mcg BID (N=431), GLY 50 mcg BID (N=432), or placebo BID (N=430) (Table 1). Baseline demographic and clinical characteristics, including the mean SGRQ total scores, were well matched across the cohorts. Subjects were predominantly male (54.7–57.2%), Caucasian (89.3–90.2%), and current smokers (50.7–55.7%). Nearly two-thirds (63.7–64.7%) had high cardiovascular risk conditions. Patients with moderate airflow obstruction (GOLD 2) comprised the majority (55.3–58.8%). More than one-third (35.6–36.6%) had severe COPD (GOLD 3), and 5.6–8.1% had very severe disease (GOLD 4). During the prior 12 months, 19.5% and 16.9% of patients who were randomized to GLY 25 mcg and 50 mcg BID, respectively, and 21.6% in the placebo cohort had experienced a COPD exacerbation. Utilization of background LABA was slightly higher than that of ICS (30.7–31.3% versus 28.2–29.5%, respectively).

Effects of Nebulized GLY on the Risk of CID at 12 Weeks

At week 12, lower proportions of patients treated with GLY 25 mcg and 50 mcg BID had experienced ≥ 1 qualifying CID event compared to those who had received

Table 1 Baseline Demographic and Clinical Characteristics of the Pooled GOLDEN 3 and GOLDEN 4 ITT Study Cohorts

Characteristics ^a	Placebo BID	GLY 25 mcg BID	GLY 50 mcg BID
Patients, n	430	431	432
Age, mean (SD)	63.7 (8.8)	63.3 (8.7)	62.6 (8.6)
Female, n (%)	195 (45.3)	189 (43.9)	185 (42.8)
Race, n (%)			
Caucasian	388 (90.2)	385 (89.3)	386 (89.4)
Black or African American	39 (9.1)	42 (9.7)	44 (10.2)
Smoking status, n (%)			
Current	218 (50.7)	240 (55.7)	226 (52.3)
Former	212 (49.3)	191 (44.3)	206 (47.7)
High cardiovascular risk: Yes, n (%) ^b	278 (64.7)	275 (63.8)	275 (63.7)
LABA use: Yes, n (%) ^c	132 (30.7)	135 (31.3)	135 (31.3)
ICS use: Yes, n (%)	127 (29.5)	126 (29.2)	122 (28.2)
Had a COPD exacerbation in the past 12 months: Yes, n (%) ^d	93 (21.6)	84 (19.5)	73 (16.9)
Post-bronchodilator FEV ₁ , L, mean (SD)	1.521 (0.5)	1.514 (0.5)	1.533 (0.5)
Post-bronchodilator FEV ₁ , n (%)			
<30% predicted (GOLD 4)	24 (5.6)	29 (6.7)	35 (8.1)
≥30% to <50% predicted (GOLD 3)	153 (35.6)	156 (36.2)	158 (36.6)
≥50% to <80% predicted (GOLD 2)	253 (58.8)	245 (56.8)	239 (55.3)
Total SGRQ score, mean (SD)	48.0 (17.5)	49.2 (18.7)	49.2 (18.5)
Completed study, n (%)	379 (88.1)	396 (91.9)	400 (92.6)

Notes: ^aBaseline data were collected in Week 0 (Visit 2). ^bDefined as a clinical diagnosis of one or more of the following conditions: hypertension, ischemic heart disease, cerebrovascular disease, peripheral arterial disease, clinically significant arrhythmia, or heart failure. ^cLimited by protocol to approximately 30%. ^dCOPD exacerbation or worsening, defined as an increase in clinician-assessed COPD symptoms (eg, dyspnea, cough, sputum volume, or sputum purulence) during ≥2 consecutive days, and classified as mild (self-managed, with increased short-acting bronchodilator and/or ICS use), moderate (treated with antibiotics and/or systemic corticosteroids) or severe (requiring hospitalization). Any event occurring within 30 days of a prior event was considered a continuation of the index event.

Abbreviations: BID, twice daily; COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in one second; GLY, glycopyrrolate; GOLD, Global Initiative for Chronic Obstructive Lung Disease; GOLDEN, Glycopyrrolate for Obstructive Lung Disease via Electronic Nebulizer; ICS, inhaled corticosteroids; ITT, intent-to-treat; LABA, long-acting beta-agonist; SD, standard deviation; SGRQ, St. George's Respiratory Questionnaire.

placebo (34.2% and 37.8%, respectively, versus 51.2%) (Table 2). Compared to placebo, treatment with GLY reduced the risk of any CID by 50% (OR: 0.5; 95% CI: 0.37–0.68) when administered at 25 mcg BID, and by 40% (OR: 0.60; 95% CI: 0.44–0.80) at 50 mcg BID (both, $p < 0.05$) (Figure 2). Examining the CID component endpoints, patients treated with GLY BID, regardless of study dose, demonstrated sustained bronchodilator effects and maintained self-reported health status with lower proportions experiencing deterioration in trough FEV₁ and SGRQ total score versus placebo. The risk of a ≥100-mL decline in trough FEV₁ from baseline was reduced significantly by 59% (OR: 0.41; 95% CI: 0.27–0.62) with GLY 25 mcg BID, and by 52% (OR: 0.48; 95% CI: 0.32–0.72) with GLY 50 mcg BID (Figure 2), while the risk of a ≥4-unit increase (worsening) in the total SGRQ score was significantly decreased by 48% (OR: 0.52; 95% CI: 0.37–0.73) and 37% (OR: 0.67; 95% CI: 0.48–0.93), respectively (all, $p < 0.05$). During the 12 weeks, less than

10% of patients in each cohort had experienced an HCRU-related moderate or severe exacerbation, resulting in no significant differences in the risk of this component endpoint between GLY BID and placebo (Table 2).

Subgroup Analysis

In the subgroup analysis, the risks of CID were significantly lower ($p < 0.05$) for GLY 25 mcg BID than placebo irrespective of age group, smoking status, background LABA use, COPD severity, or PIFR (Figure 3). Patients aged <65 years (OR 0.45, 95% CI 0.29, 0.68) and those with a PIFR <60 L/min (OR 0.36, 95% CI 0.20, 0.67) exhibited the most pronounced reductions in the risk of CID. Treatment with GLY 50 mcg BID also significantly decreased the risk of CID in most subgroups (Figure 4), except for the subgroup aged ≥65 years and females. The risk of CID in these subgroups was numerically lower but not significantly different from placebo. For both GLY regimens, patients <65 years, males, current smokers, and patients with PIFR <60 min/L experienced larger

Table 2 Proportions of Subjects Meeting Criteria for CID at Treatment Week 12

CID Criteria	Placebo BID		GLY			
	CID n/Total N	CID, %	25 mcg BID		50 mcg BID	
			CID n/Total N	CID, %	CID n/Total N	CID, %
Any CID endpoint	200/391	51.2%	136/398	34.2%	152/402	37.8%
≥100-mL decrease in baseline post-bronchodilator trough FEV ₁	88/380	23.2%	48/393	12.2%	56/399	14.0%
≥4-unit increase in SGRQ total score	115/357	32.2%	72/368	19.6%	92/373	24.7%
HCRU-related moderate or severe exacerbation ^a	40/430	9.3%	33/431	7.7%	39/432	9.0%

Notes: ^aDefined as an increase in clinician-assessed COPD symptoms (eg, dyspnea, cough, sputum volume, or sputum purulence) during at least two consecutive days, and classified as “moderate” (treated with clinic visit with or without antibiotics or oral corticosteroids but no hospitalization) or “severe” (requiring overnight stay in a hospital or emergency room).

Abbreviations: BID, twice daily; CID, clinically important deterioration; FEV₁, forced expiratory volume in one second; GLY, glycopyrrolate; HCRU, healthcare resource utilization; LABA, long-acting beta-agonist; SGRQ, St. George’s Respiratory Questionnaire.

reductions in the risk of CID than the respective alternate subgroups. Treatment with the GLY 25 mcg BID had a greater impact on reducing CID in the subgroup with baseline FEV₁ ≥50% predicted (versus FEV₁ <50%) and was

comparably effective in the subgroups with LABA and without LABA use. In contrast, larger effect sizes were observed in the 50 mcg BID subgroups with FEV₁ <50% (versus FEV₁ ≥50%) and no LABA use (versus LABA use).

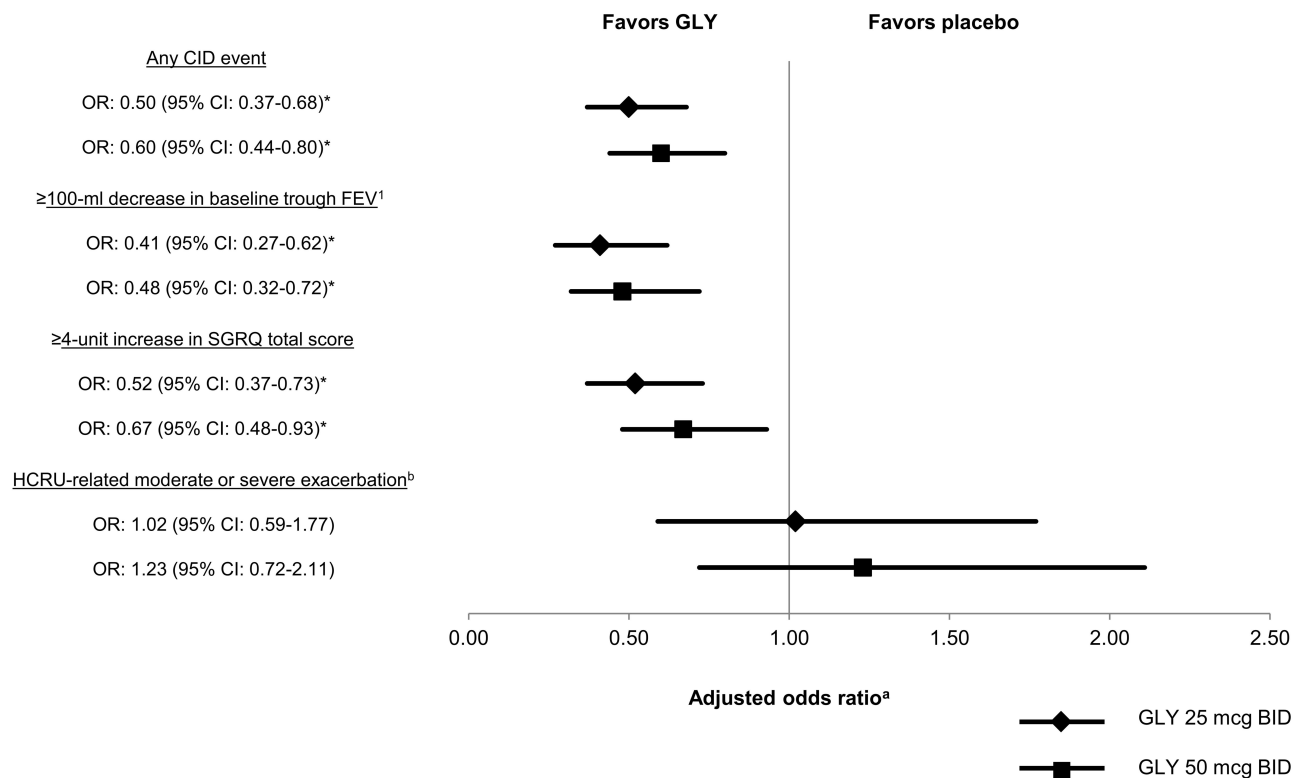


Figure 2 Risk (adjusted odds ratio)^a of CID at week 12 of treatment with GLY (25 mcg BID and 50 mcg BID) compared to placebo.

Notes: ^aOdds ratio (GLY/placebo), adjusted for high cardiovascular risk, LABA use, baseline FEV₁, baseline SGRQ total score, and 12-month prior history of exacerbations.

^bDefined as an increase in clinician-assessed COPD symptoms (eg, dyspnea, cough, sputum volume, or sputum purulence) during at least two consecutive days, and classified as “moderate” (treated with clinic visit, but no hospitalization, with or without antibiotics or oral corticosteroids) or “severe” (requiring overnight stay in a hospital or emergency room). *p<0.05.

Abbreviations: BID, twice daily; CI, confidence interval; CID, clinically important deterioration; FEV₁, forced expiratory volume in one second; GLY, glycopyrrolate; HCRU, healthcare resource utilization; LABA, long-acting beta-agonist; OR, odds ratio; SGRQ, St. George’s Respiratory Questionnaire.

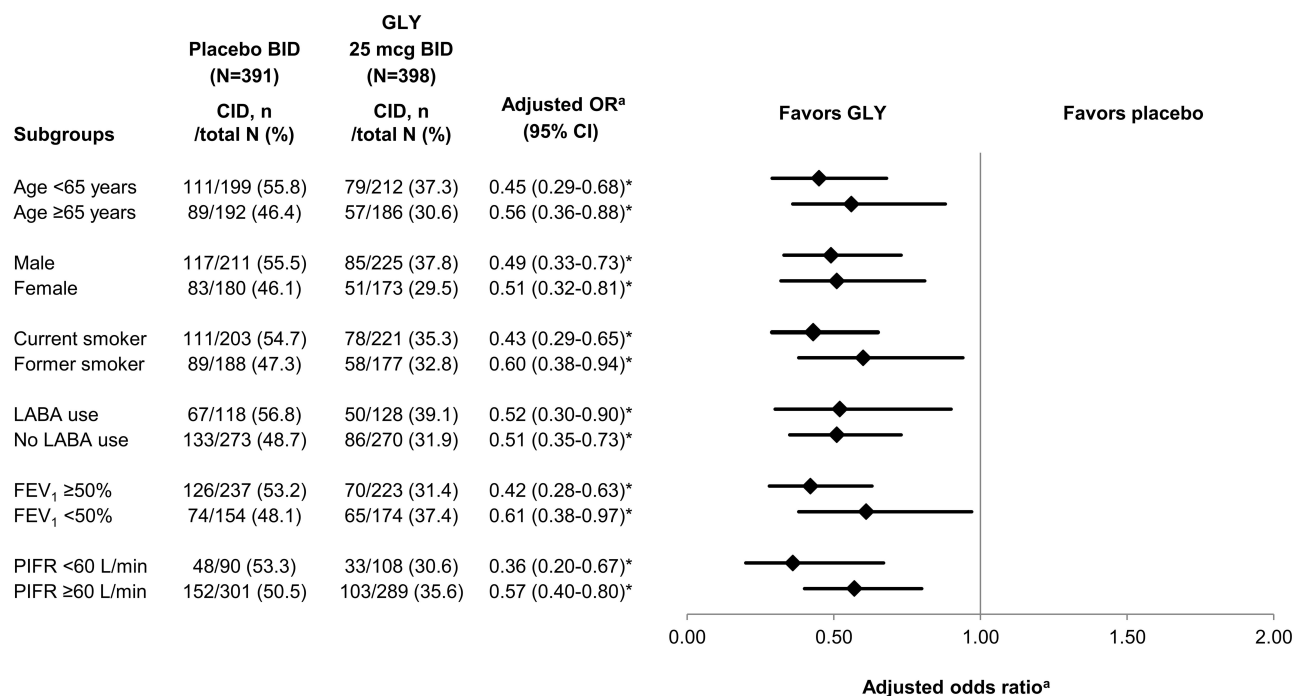


Figure 3 Subgroup analysis of the risk (adjusted odds ratio)^a of CID at week 12 of treatment with GLY 25 mcg BID compared to placebo.
Notes: ^aOdds ratio (GLY/placebo), adjusted for high cardiovascular risk, LABA use, baseline FEV₁, baseline SGRQ total score, and 12-month prior history of exacerbations. *p<0.05.

Abbreviations: BID, twice daily; CI, confidence interval; CID, clinically important deterioration; FEV₁, forced expiratory volume in one second; GOLD, Global Initiative for Chronic Obstructive Lung Disease; GLY, glycopyrrolate; LABA, long-acting beta-agonist; OR, odds ratio; PIFR, peak inspiratory flow rate.

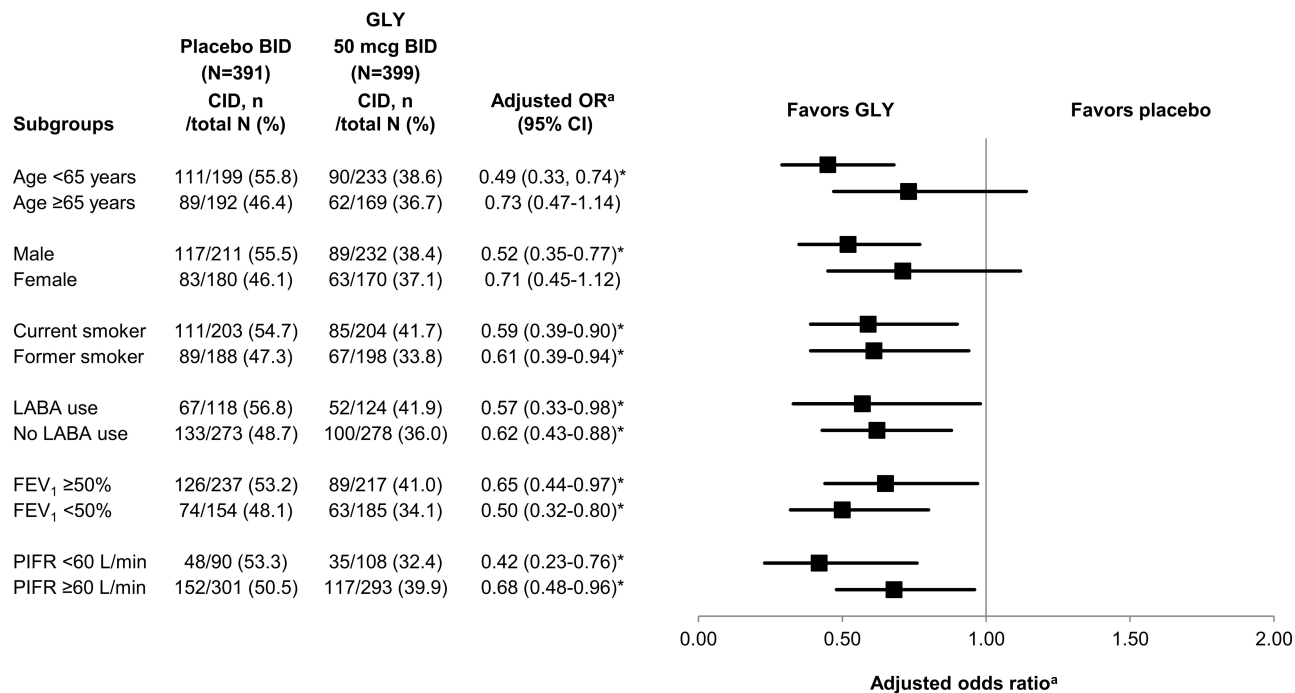


Figure 4 Subgroup analysis of the risk (adjusted odds ratio)^a of CID at week 12 of treatment with GLY 50 mcg BID compared to placebo.
Notes: ^aOdds ratio (GLY/placebo), adjusted for high cardiovascular risk, LABA use, baseline FEV₁, baseline SGRQ total score, and 12-month prior history of exacerbations. *p<0.05.

Abbreviations: BID, twice daily; CI, confidence interval; CID, clinically important deterioration; FEV₁, forced expiratory volume in one second; GOLD, Global Initiative for Chronic Obstructive Lung Disease; GLY, glycopyrrolate; LABA, long-acting beta-agonist; OR, odds ratio; PIFR, peak inspiratory flow rate.

Discussion

Quantifying the effectiveness of treatments in preventing worsening of COPD is essential given that minimizing the risk of progression is a primary goal of managing patients with stable disease.¹ To provide a more comprehensive assessment in COPD, CID has emerged as an important composite endpoint that captures the multidimensional effects of treatment on key measures related to lung function, health status, and exacerbations. Each CID endpoint represents a clinically relevant event in COPD that may portend worse outcomes in the absence of effective intervention. Progressive decline in lung function is accelerated by exacerbations.⁷ COPD progression, however, is not restricted to deterioration in FEV₁ and can manifest in worsening of patient-reported outcomes such as health status and quality of life.^{17,18} Further, managing CID is critical as published evidence suggests that patients experiencing CID during the early months of bronchodilator therapy may experience poorer medium to long-term outcomes. A comparative study of indacaterol/glycopyrronium and salmeterol/fluticasone observed that CID during the first 12 treatment weeks was associated with significantly earlier and higher rates of moderate-to-severe exacerbations between 12 and 52 weeks (both, $p < 0.00001$).¹⁵ A post hoc analysis of data from interventional and observational studies of fluticasone propionate/salmeterol found that at three years, the respective risk of CID in each component endpoint was significantly increased when CID in the endpoint occurred within 24 weeks. In contrast, no CID was significantly associated with higher quality of life scores and lower healthcare costs (all, $p < 0.001$).^{11,19}

In this post hoc analysis, patients with moderate-to-severe COPD treated with GLY over 12 weeks were significantly less likely to experience CID compared to placebo with those treated with 25 mcg BID, the licensed regimen in the US, attaining a 50%-reduction in risk. Results of the subgroup analyses were largely consistent with the overall findings. At both GLY doses, reduction in the risk of CID was most pronounced for patients who had low PIFR (< 60 L/min) at baseline, a finding which may be related to use of the nebulizer. Several studies have shown that individuals with poor inspiratory effort, such as the elderly and some patients with COPD, are unable to overcome the internal resistance of dry powder inhalers and therefore, have difficulty generating sufficient inspiratory flow for effective drug delivery.^{20–23} In these cases, the nebulizer's vibrating membrane technology may offer an

alternative mode as drug deposition is targeted to the lungs via regular tidal breathing.²⁴ This nebulization method achieves a high percentage of aerosolized particles within the respiratory range, and does not require strong PIFR for device actuation or careful hand-breath coordination to achieve an effective dose. For these reasons, older patients with concomitant health conditions may likely also benefit from nebulized GLY. The comparative effectiveness of different inhalation devices in patients with COPD and poor inspiratory effort has not been established, and the choice of inhalation devices depends ultimately on a multitude of factors.²⁵ However, a prospective, observational study in the US found that nearly one-third of patients hospitalized for a COPD exacerbation had a PIFR < 60 min/L at discharge.²³ It is possible that certain patients under such circumstances may benefit from inhalation devices that can enhance medication dispersal to the lungs.

Based on the ORs, treatment response to GLY appeared larger in the subgroup aged < 65 years than in the subgroup ≥ 65 years. This was likely due to age-related differences in airflow capacity. Compared to older patients, younger patients generally have larger airflow capacity with FEV₁ values averaging 2.0 liters.²⁶ A 10% improvement in FEV₁ in these younger patients could, therefore, represent an increase of 200 mL. FEV₁ usually diminishes with normal aging and declines at accelerated rates up to -60 mL per year with cigarette smoking and COPD. As a result, airflow capacity is reduced in older patients (age ≥ 65) and may be as low as 1.0 liter.²⁶ A 10% improvement in these patients could result in an increase of 100 mL. Therefore, a bronchodilator such as GLY would be expected to have a relatively larger positive FEV₁ impact > 100 mL in younger patients compared to older patients with COPD. Similarly, since younger patients likely experience greater net on-treatment milliliter improvements from baseline with nebulized GLY, they may be less likely to encounter net 100-mL declines meeting the FEV₁ criteria for CID.

Examining the individual CID endpoints, patients treated with GLY were significantly more likely to maintain FEV₁ and health status, but experienced no improvement in the risk of HCRU-related moderate or severe exacerbations over placebo. The latter may have been related to the low frequency of exacerbations during the study period and prior 12 months. Across all cohorts, deterioration in the total SGRQ score at week 12 was the strongest driver of CID (19.6–32.2% of patients). A prior analysis of the pooled data from the GOLDEN 3 and GOLDEN 4 trials showed that

GLY (25 mcg and 50 mcg BID) led to statistically significant improvements from baseline in the SGRQ total and component (Symptoms, Impacts, and Activity) scores compared to placebo.^{6,27} Change from baseline SGRQ total score at week 12 was measured as the least squares mean and compared between GLY and placebo using analysis of variance. Applying the minimal clinically important difference criteria for SGRQ response (≥ 4 decrease in total score), the odds of improvement at week 12 were also significantly greater with both GLY 25 mcg (OR 1.68; 95% CI 1.24, 2.28) and 50 mcg (OR 1.36; 95% CI 1.00, 1.85) compared with placebo.²⁷ In the present study, we identified a subset of patients who experienced CID in health status (based on the total SGRQ score). This finding highlights the potential utility of CID assessment in identifying “at-risk” individuals who may be otherwise obscured in assessments of SGRQ response, and who might benefit from modifications in management.

Findings from this post hoc analysis are noteworthy as the data were pooled from two Phase III double-blind, placebo-controlled studies with robust replicate designs. Examining CID by the component endpoints also provided greater insight into how worsening in specific factors associated with COPD progression, ie, pulmonary function and health status, might be individually impacted during GLY treatment. Further, the analysis adds to the evidence supporting the utility of the composite CID endpoint for identifying therapies that prevent disease worsening, an important goal of management, particularly for patients who fail to achieve clinically meaningful responses to treatment.^{1,28}

Study limitations include the 12-week study period of the GOLDEN 3 and GOLDEN 4 trials which precluded exploration of sustained CID. FEV₁ and SGRQ were assessed only at baseline and at 12 weeks, the incidence of moderate or severe HCRU-related exacerbations was low, and fluctuations in disease state were not captured. Nonetheless, published evidence supports the merits of assessing and preventing early, short-term CID. Further, generalizability of the findings to patients with COPD managed in the real world is limited as the outcomes reflect those of the population selected for the Phase III trials. Both GOLDEN 3 and GOLDEN 4, however, were prospectively designed to enroll subjects representative of the general COPD population and included proportions of subjects who had very severe disease (post-bronchodilator FEV₁% predicted <30%), continuing background LABA use with or without ICS, and high cardiovascular risk factors.⁶

Conclusion

Nebulized GLY (25 mcg and 50 mcg BID) administered over 12 weeks significantly reduced the risk of CID over placebo and was effective in preventing clinically meaningful worsening in FEV₁ and patient-reported health status. These findings contribute to the evidence supporting the use of nebulized GLY BID as a maintenance treatment for patients with moderate-to-very-severe COPD. Assessment over a longer period may confirm a sustained effect in stabilizing disease.

Data Sharing Statement

Sunovion Pharmaceuticals Inc. is part of a clinical trial data sharing consortium that facilitates access for qualified researchers to selected anonymized clinical trial data. For up-to-date information on data availability, please visit <https://www.clinicalstudydatarequest.com/Study-Sponsors.aspx> and click on “Sunovion”.

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Disclosure

LM was employed by Evidera at the time this work was conducted, which provides consulting and other research services to pharmaceutical, device, government and non-government organizations. As an Evidera employee, she worked with a variety of companies and organizations and was expressly prohibited from receiving any payment or honoraria directly from these organizations for services rendered. EMK has participated in consulting, advisory boards, speaker panels, or received travel reimbursement from Amphastar, AstraZeneca, Boehringer Ingelheim, Chiesi, Cipla, GlaxoSmithKline plc, Mylan, Novartis, Oriel, Pearl, Sunovion, Teva, and Theravance.

He has conducted multicenter clinical research trials for approximately 40 pharmaceutical companies. CD and XN are employees of Sunovion. The authors report no other conflicts of interest in this work.

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