

Benralizumab Prevents Recurrent Exacerbations in Patients with Chronic Obstructive Pulmonary Disease: A Post Hoc Analysis

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Abstract: Exacerbations in chronic obstructive pulmonary disease (COPD), which tend to occur in clusters and increase with disease severity, come with high societal and economic burdens. Prevention and delay of recurrent exacerbations is an unmet and significant therapeutic need for patients with COPD. GALATHEA (NCT02138916) and TERRANOVA (NCT02155660) were trials assessing efficacy of benralizumab in patients with frequent COPD exacerbations despite treatment. Although these studies found that benralizumab given as an add-on treatment did not significantly reduce annual rates of COPD exacerbations after 56 weeks of treatment, in the following exploratory post hoc analysis of the GALATHEA and TERRANOVA trials we identified a potential responder population in which treatment with benralizumab prevents recurrent COPD exacerbations during 30- and 90-day periods following an initial exacerbation, a vulnerable period for an exacerbation to occur. This responder population was characterized by high blood eosinophil counts and frequent previous exacerbations despite optimized triple therapy. These results highlight the importance of targeted therapies for high-risk populations and merit further research into the benefits of biologic therapies for COPD exacerbations.

Keywords: COPD, recurrent exacerbations, benralizumab, eosinophil

Introduction

Preventing and delaying recurrent exacerbations is an unmet need and crucial therapeutic goal in chronic obstructive pulmonary disease (COPD). Studies have shown that COPD exacerbations often occur in clusters,^{1–3} with a US study showing that approximately 20% of patients hospitalized for acute COPD exacerbations are readmitted to hospital within 30 days,⁴ and a European study showing that over a third of patients are readmitted within 90 days.⁵ Benralizumab is a humanized afucosylated (IgG1) anti-interleukin-5 (IL-5) receptor α monoclonal antibody that has been shown to rapidly deplete eosinophils in blood, airway, and sputum by inducing apoptosis following recruitment of natural killer cells.^{6–8}

GALATHEA (NCT02138916) and TERRANOVA (NCT02155660) were 2 randomized, double-blind, placebo-controlled Phase 3 trials assessing the efficacy of benralizumab in patients experiencing frequent COPD exacerbations despite treatment.⁹ These trials showed that benralizumab added to inhaled maintenance treatments did not significantly reduce the annual COPD exacerbation rate after 56 weeks.⁹ However, a post hoc analysis identified a potential responder subgroup with distinct characteristics, including a history of ≥ 3 COPD exacerbations in the previous year despite receiving triple therapy (inhaled corticosteroid, a long-acting β_2 -agonist, and a long-acting muscarinic antagonist) and a baseline blood eosinophil (bEOS) count ≥ 300 cells/ μ L.¹⁰

Methods

In this exploratory post hoc analysis of the GALATHEA and TERRANOVA trials, we examined the effect of benralizumab 100 mg versus placebo on recurrent COPD exacerbations in the potential responder population (≥ 3 COPD exacerbations in the previous year while receiving triple therapy plus $\text{bEOS} \geq 300$ cells/ μL) who had ≥ 1 exacerbation following randomization despite treatment. In this subpopulation, we evaluated recurrent COPD exacerbations within 30 and 90 days following an initial moderate or severe exacerbation (a vulnerable period for another exacerbation to occur). Moderate COPD exacerbations were defined as increased respiratory symptoms leading to systemic corticosteroid and/or antibiotic use; severe exacerbations were defined as hospitalizations. Severe exacerbations resulting in death were excluded from these analyses. Statistical analyses were performed using a negative binomial model, including the following covariates: treatment, study (GALATHEA or TERRANOVA), geographic region, and the number of prior exacerbations.

Overall, 145 patients (benralizumab, $n = 73$; placebo, $n = 72$) were included in this post hoc analysis, representing approximately 6.5% of the relevant population from GALATHEA and TERRANOVA (benralizumab, 73/1114; placebo, 72/1118). Baseline demographics and characteristics of the subpopulation included in these analyses are shown in Table 1. The mean (SD) patient age was 65.3 (8.4) years, most patients were male ($n = 92$; 63.4%) and white ($n = 119$; 82.1%), and 30.3% were current smokers. The median bEOS count was 450.0 (range: 300–3860) cells/ μL , and the mean (SD) pre-bronchodilator forced expiratory volume in 1 second (FEV_1) was 37.4% (12.3) of predicted. Patient demographics and clinical characteristics were similar between groups, although there was a higher number of exacerbations in the previous year in the placebo group (4.3 [1.9]) compared to the benralizumab group (3.9 [1.3]). By design, the patients in this analysis had more exacerbations in the past 12 months (4.1 [1.7]) compared to the overall patient population in the GALATHEA and TERRANOVA trials.⁹

Table 1 Baseline Patient Demographics and Characteristics

	Benralizumab (n = 73)	Placebo (n = 72)	All Patients (n = 145)
Age, mean (SD), year	64.6 (8.1)	66.1 (8.7)	65.3 (8.4)
Sex, male, n (%)	41 (56.2)	51 (70.8)	92 (63.4)
Race, White, n (%)	59 (80.8)	60 (83.3)	119 (82.1)
Body mass index, mean (SD), kg/m^2	27.63 (6.5)	26.63 (4.9)	27.14 (5.8)
Nicotine use status			
Current, n (%)	25 (34.2)	19 (26.4)	44 (30.3)
Former, n (%)	48 (65.8)	53 (73.6)	101 (69.7)
Nicotine consumption, mean (SD), pack-year	44.6 (25.6)	47.1 (25.5)	45.9 (25.5)
On maintenance COPD medication (ICS/LABA/LAMA), n (%)	73 (100.0)	72 (100.0)	145 (100.0)
Baseline bEOS count, mean (SD), cells/mL	589.3 (468.8)	617.4 (460.2)	603.2 (463.2)
Baseline bEOS count, median (range), cells/mL	450 (300–3860)	445 (300–2830)	450 (300–3860)
Pre-bronchodilator FEV_1 , mean (SD), %PN	38.4 (12.4)	36.3 (12.1)	37.4 (12.3)
Post-bronchodilator FEV_1/FVC ratio, mean (SD), %	42 (10)	41 (11)	41 (11)
Exacerbations in the previous year, mean (SD), range	3.93 (1.3), 3–10	4.29 (1.9), 3–12	4.11 (1.6), 3–12
Atopic status by Phadiatop test, n (%)	25 (34.2)	25 (34.7)	50 (34.5)

Abbreviations: bEOS, blood eosinophils; COPD, chronic obstructive pulmonary disease; FEV_1 , forced expiratory volume in 1 second; FVC, forced vital capacity; ICS, inhaled corticosteroids; LABA, long-acting β_2 -agonist; LAMA, long-acting muscarinic antagonist; PN, predicted normal.

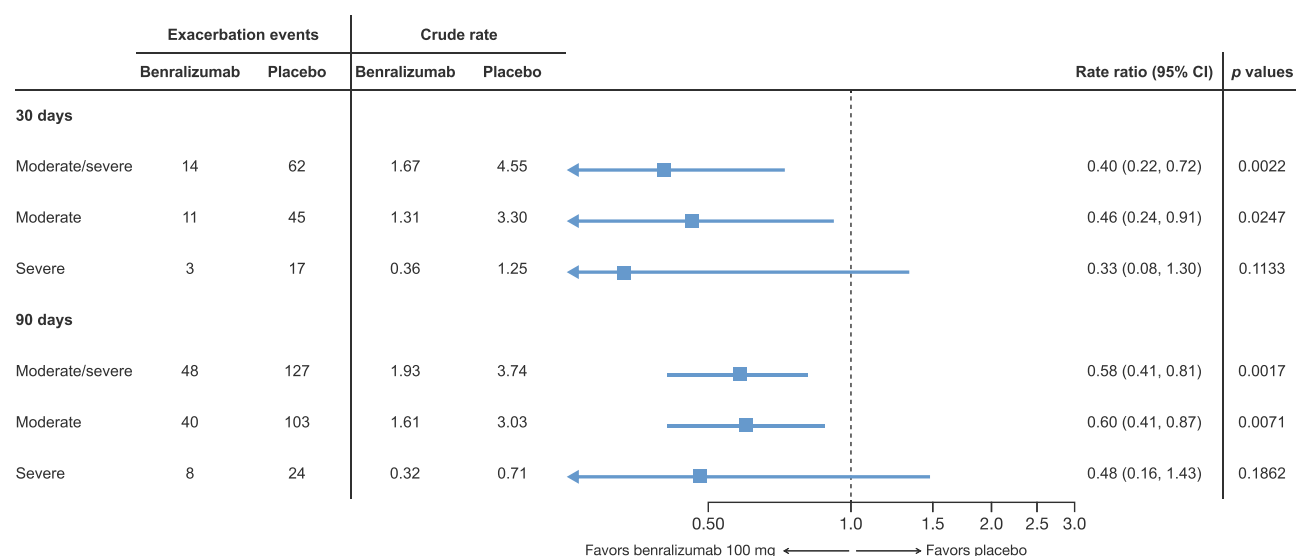


Figure 1 Risk of recurrent COPD exacerbations within 30^a and 90^b days.

Notes: ^aTotal follow-up time: benralizumab 100 mg, 8.4 years; placebo, 13.6 years. ^bTotal follow-up time: benralizumab 100 mg, 24.9 years; placebo, 34.0 years.

Abbreviations: CI, confidence interval; COPD, chronic obstructive pulmonary disease.

Results

Within 30 days of an initial exacerbation, 14 moderate or severe exacerbations occurred in the benralizumab group versus 62 in the placebo group. Benralizumab reduced the risk of recurrent moderate or severe COPD exacerbations within 30 days by 60% compared with placebo (rate ratio [RR], 0.40; 95% CI: 0.22–0.72; $p=0.0022$) (Figure 1).

Within 90 days of an initial exacerbation, 48 moderate or severe exacerbations occurred in the benralizumab group versus 127 in the placebo group. Benralizumab reduced the risk for recurrent moderate or severe COPD exacerbations within 90 days by 42% compared to placebo (RR, 0.58; 95% CI: 0.41–0.81; $p=0.0017$) (Figure 1). Additionally, fewer recurrent severe exacerbations occurred in the benralizumab group compared to placebo after 30 days (benralizumab = 3, placebo = 17; RR, 0.33; 95% CI: 0.08–1.30; $p=0.1133$) and 90 days (benralizumab = 8, placebo = 24; RR, 0.48; 95% CI: 0.16–1.43; $p=0.1862$), although the differences were not statistically significant.

Discussion

We acknowledge that a potential limitation of this post hoc analysis is the risk of a type I error. Yet, the effects of benralizumab on this patient population appear clinically significant and merit further investigation; indeed, the ongoing RESOLUTE (NCT04053634) phase 3 trial is designed to evaluate the efficacy of benralizumab specifically in this patient population. In addition, the ABRA trial (NCT04098718) will evaluate bEOS-guided benralizumab therapy in COPD patients with eosinophilic exacerbations in the acute setting. The results from these studies will examine the potential benefits of benralizumab therapy in COPD patients with an eosinophilic phenotype.

Conclusion

In conclusion, the results of this post hoc analysis support the hypothesis that benralizumab reduces the risk for recurrent COPD exacerbations within the vulnerable 30-to-90-day period in the previously identified responder population: those with high bEOS counts and frequent previous exacerbations despite optimized triple therapy. This is a promising finding, as recurrent COPD exacerbations carry serious risks for patients, including hospital readmittance 30 and 90 days following an event,^{11,12} treatment failure,¹² inpatient mortality of 10%, and up to 50% all-cause mortality 2 years after an event.¹² The results of this post hoc analysis highlight the importance of targeted interventions to reduce the risk for recurrent COPD exacerbations in high-risk patients, as well as the beneficial impact of biologic therapy on COPD exacerbations overall.

Data Sharing Statement

Data underlying the findings described in this manuscript may be obtained in accordance with AstraZeneca's data-sharing policy described at <https://astrazenecagrouptrials.pharmacm.com/ST/Submission/Disclosure>.

Ethics Approval and Consent to Participate

Ethical review and approval was not required for this post hoc analysis of data from previously completed trials. As described in the previous primary publication of that study, independent ethics committees of the trial centers or central institutional review boards approved the trial protocols for the GALATHEA and TERRANOVA trials.⁹ Both trials were conducted in accordance with the principles of the Declaration of Helsinki; all patients provided written informed consent.

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Author Contributions

All authors contributed to data analysis, drafting or revising the article, have agreed on the journal to which the article will be submitted, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

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