Impact and prevention of severe exacerbations of COPD: a review of the evidence

Abstract: Severe exacerbations of COPD, ie, those leading to hospitalization, have profound clinical implications for patients and significant economic consequences for society. The prevalence and burden of severe COPD exacerbations remain high, despite recognition of the importance of exacerbation prevention and the availability of new treatment options. Severe COPD exacerbations are associated with high mortality, have negative impact on quality of life, are linked to cardiovascular complications, and are a significant burden on the health-care system. This review identified risk factors that contribute to the development of severe exacerbations, treatment options (bronchodilators, antibiotics, corticosteroids [CSs], oxygen therapy, and ventilator support) to manage severe exacerbations, and strategies to prevent readmission to hospital. Risk factors that are amenable to change have been highlighted. A number of bronchodilators have demonstrated successful reduction in risk of severe exacerbations, including long-acting muscarinic antagonist or long-acting β₂-agonist mono- or combination therapies, in addition to vaccination, mucolytic and antibiotic therapy, and nonpharmacological interventions, such as pulmonary rehabilitation. Recognition of the importance of severe exacerbations is an essential step in improving outcomes for patients with COPD. Evidence-based approaches to prevent and manage severe exacerbations should be implemented as part of targeted strategies for disease management.

Keywords: severe COPD exacerbations, hospitalization, prevention, treatment, bronchodilators, long-acting muscarinic antagonist

Introduction

Exacerbations of COPD are important events in the course of the disease that have profound impact on patients’ health status, functional capacity, and lung function. The severity of exacerbations varies, and the clinical definition of a severe exacerbation is not always uniform; for practical reasons, clinical trials and epidemiological studies normally define severe exacerbations as those resulting in hospitalization. Severe exacerbations have a particularly significant clinical and socioeconomic impact. A recent database review of >73,000 patients found that fewer than half of patients hospitalized for an exacerbation survived for a further 5 years. Patients who survive severe exacerbations are likely to experience significantly impaired quality of life (QoL), are at increased risk of further exacerbations, and represent a major contributor to the overall health-care costs associated with COPD. Therefore, understanding the factors that lead to an exacerbation being so severe that treatment in hospital is needed, and optimization of patient management to reduce the risk of severe exacerbations, are crucial therapeutic goals for patients with stable COPD.

Currently, the prevalence and burden of severe exacerbations remain high, despite advances in COPD therapies and increased recognition of the importance
of preventing exacerbations. This manuscript reviews the risk factors, consequences, and treatment of severe COPD exacerbations, and highlights opportunities to improve the management and prevention of these serious events.

**What constitutes a severe exacerbation?**

An exacerbation is typically considered an acute episode characterized by worsening of the patient’s respiratory symptoms (ie, baseline dyspnea, cough, and/or sputum production) that is sufficient to require additional therapy. The American Thoracic Society (ATS)/European Respiratory Society (ERS) guidelines classify any exacerbation that can be managed at home (mild or moderate) as level I, while severe exacerbations are stratified further as those requiring hospitalization (level II) and those leading to respiratory failure (level III). For the purposes of this review, severe exacerbations refers to exacerbations requiring hospitalization, accepting that exacerbations requiring hospitalization may be quite varied in terms of the severity of respiratory symptoms and will also reflect the underlying severity of the patient’s lung impairment. In patients with very limited respiratory reserves, less severe exacerbation events may still require hospitalization for effective management. Additional factors that contribute to the decision to admit a patient to hospital include the presence of such comorbidities as heart failure, cardiac arrhythmia, or pneumonia, or the onset of other clinical signs, such as central cyanosis or peripheral edema. Patients may go into hospital as an acute emergency admission or be admitted only after outpatient management of an exacerbation has failed to produce an adequate response. Psychosocial factors may also be involved: the patient’s ability to cope at home, which may be a product of their age, mental status, and the level of support they have available to them, is an important consideration in deciding whether they should be treated for an exacerbation as an inpatient or outpatient. Finally, regional differences in local health-care practices and access to health-care services also influence the threshold that determines hospitalization for patients with COPD suffering an exacerbation of COPD.

**An exacerbation has a significant effect on clinical and patient-centered outcomes, including obstruction, dyspnea, and exercise capacity. Patients take a long time to recover from these effects, and may never return to their preexacerbation state.**

**Impact on mortality**

Hospitalization for COPD exacerbations is associated with a high mortality rate. In an EU COPD audit, 5% of patients admitted to hospital with a COPD exacerbation died while in hospital. Consistent with this finding, a recent database review of >73,000 patients with up to 17 years’ follow-up found that mortality peaked in the first week and remained high during the first 3 months following hospital admission. Fewer than half of patients hospitalized for an exacerbation were still alive after 5 years’ follow-up in this study. This is consistent with previous reports of >50% mortality at 5 years following hospitalization for acute exacerbations of COPD. Long-term posthospitalization mortality rates reported in the literature are quite variable: 12%–33% after 1 year, 26%–39% after 2 years, 39%–49% after 3 years, 45%–47% after 4 years. It has been suggested that survival following hospitalization for COPD is improving. However, mortality associated with COPD may be underestimated, as it is often cited as a contributory, rather than underlying, cause of death, or omitted from death certificates altogether if death is primarily attributed to comorbid conditions. The contribution of nonhospitalized exacerbations to mortality rates (ie, unreported exacerbations that should have warranted hospital treatment) has not been well quantified. Mortality risk increases with each new exacerbation: there is a fivefold greater risk of mortality after the tenth hospitalization compared with the first (Figure 1).

Identified risk factors for mortality associated with severe exacerbation include older age, male sex, prior hospitalizations, weight loss/low body-mass index, poor QoL, physiological parameters (eg, partial pressure of carbon dioxide [PaCO₂], pulmonary hypertension, lung cancer, cardiovascular comorbidity, and need for long-term oxygen therapy at discharge. Health-related quality of life

Health-related QoL deteriorates in patients experiencing exacerbations; the more frequent and severe the exacerbation, the more rapid and pronounced the decline. Severe exacerbations lead to impairment of the ability to perform usual activities and reduce work productivity. Activity limitation and reduced work productivity have been found to be strongly correlated with the total number of severe exacerbations in
the previous year.\textsuperscript{30} Assessment of health-related QoL varies considerably between studies, and some measures do not include all domains of health status (physiological functioning, symptoms, functional impairment, and QoL).\textsuperscript{32}

Cardiovascular sequelae

Cardiovascular comorbidities are common in patients with COPD, and particularly in those patients with more severe airflow limitation. A review of studies of patients hospitalized for COPD exacerbations found that up to 55% of patients had a history of cardiovascular disease.\textsuperscript{33} Exacerbations of COPD have a profound effect on patients’ cardiovascular status,\textsuperscript{33,34} and the increase in cardiac events after the first exacerbation is irrespective of the patient’s cardiac history.\textsuperscript{36} Markers of myocardial injury, including raised troponin, chest pain, and serial electrocardiogram changes, are commonly observed in patients admitted to hospital with exacerbations of COPD.\textsuperscript{33,34} Levels of N-terminal pro–brain natriuretic peptide and cardiac troponin T are elevated during acute exacerbations of COPD and thought to be a predictor of mortality among patients admitted to hospital.\textsuperscript{37} Arterial stiffness also rises acutely during COPD exacerbations, particularly when associated with airway infection.\textsuperscript{35} These changes may also be associated with an acute increase in the risk of myocardial infarction (MI): a UK observational database study found a 2.3-fold increase in the risk of MI 1–5 days postexacerbation,\textsuperscript{38} while analysis from the UPLIFT\textsuperscript{®} study showed 13-fold increased risk of MI in the 30 days postexacerbation compared with preexacerbation periods. A 17-fold increase in the occurrence of atrial fibrillation/flutter was also seen in the 30-day postexacerbation period.\textsuperscript{36}

Increased prevalence of pulmonary embolism has also been reported in patients with COPD exacerbations. Two meta-analyses have reported an overall incidence of pulmonary embolism of 16%\textsuperscript{39} and 20%,\textsuperscript{40} increasing to 25% in hospitalized patients, including those hospitalized specifically for a COPD exacerbation.\textsuperscript{30}

What is the epidemiology and economic impact of severe exacerbations?

Rates of hospitalization for exacerbations of COPD vary in different countries, which may reflect differences in the organization of health-care systems. A large international survey of more than 4,000 patients with COPD sampled from 12 countries in Europe, the US, and Asia (Continuing to Confront COPD International Patient Survey) found that overall, 15% of patients had been hospitalized for an exacerbation in the previous 12 months, with rates ranging 5%–25% across individual countries (Figure 2). In another large multinational survey of more than 1,000 patients in Germany, France, Italy, Spain, the UK, and the US (PERCEIVE), 21% of patients who reported a COPD exacerbation in the preceding year were hospitalized.\textsuperscript{42}

Severe COPD exacerbations are estimated to account for less than 10% of exacerbations,\textsuperscript{43,44} but have a disproportionate and significant socioeconomic impact, with hospitalizations estimated to account for approximately 60%–70% of health-care costs associated with COPD, depending on the region.\textsuperscript{45} Illustrative of the costs associated with COPD exacerbations, a recent health-economics analysis conducted in the US, based on a data set from 2006, identified a total of 1,254,703 hospitalizations for COPD exacerbations, with total inpatient costs of $11.9 billion over the year at a mean cost of $9,545 per hospitalized exacerbation.\textsuperscript{46} The substantial burden of severe COPD exacerbations is apparent when considered in the context of overall health-care resource utilization. In the UK, acute exacerbations account for 115,000 hospital admissions per year, representing one of the most common reasons for emergency admission to hospital,\textsuperscript{47} while in Spain it is estimated that 1%–2% of all emergency-service visits and 10% of all medical admissions are attributable to COPD exacerbations.\textsuperscript{18}

Despite trends toward an improvement in in-hospital mortality rates and a reduction in average length of hospital stay in the US,\textsuperscript{9,48} there has been no significant improvement in other indicators, including the number of hospital discharges,
emergency-department visits, and 30-day readmissions within the period 2001–2012,48 highlighting considerable room for improvement and opportunities for reduction in the costs associated with admissions for COPD exacerbations. Patients who have been admitted to hospital for a severe exacerbation of COPD are at substantial risk for rehospitalization,19 with up to a quarter of patients readmitted for COPD exacerbations within 1 year following discharge.9,19 Strategies to reduce readmission rates are thus a priority in efforts to reduce the burden of severe COPD exacerbations.

What are the risk factors associated with severe COPD exacerbations?

The majority of COPD exacerbations, including severe exacerbations, are precipitated by respiratory tract infections, either viral or bacterial, or by environmental agents, e.g., worsening air pollution.49 Numerous disease-related and demographic characteristics that modify patients’ risk of having an exacerbation have been identified.

Severity of airflow limitation is a key risk factor for exacerbation requiring hospitalization. In a retrospective study of ambulatory COPD patients from general practices, severity of forced expiratory volume in 1 second impairment was a significant predictor for increased risk of hospitalizations, as was the presence of significant comorbidities, such as diabetes or ischemic heart disease.49 The cross-sectional analysis of the PAC-COPD study found that patients with more severe respiratory symptoms, poorer QoL, worse lung function, and lower exercise capacity were at greater than threefold-higher risk of hospitalization for COPD than those with milder airflow limitation, with or without significant comorbidities (hazard ratio 3.28, \( P = 0.001 \)).51

A summary of variables that have been shown to modify the risk of severe exacerbations of COPD is shown in Table 1.4,6,18,19,26,50–78 These include lung function, symptoms (i.e., severity of dyspnea), QoL, exacerbation history, disease history and treatment, age and sex, body-mass index, smoking status, and comorbidities. Other influences on the risk of severe COPD exacerbations include the patient’s socioeconomic background, education level, and marital status.

Exacerbations may occur regardless of the degree of functional impairment, although it is generally agreed that the frequency of exacerbations increases with decreasing lung function.1 A major audit of COPD treatment and outcomes in Europe was conducted by the ERS between 2010 and 2011. This gathered a wealth of data (from more than 16,000
Poorer health literacy increases the likelihood of COPD-related hospitalizations and emergency department visits. Prior ER visits are a significant risk factor for exacerbation-related hospitalization. Low BMI is associated with hospitalization and readmission for COPD exacerbation. Underprescription of LTOT increases the risk of hospitalization, due to severe exacerbations.

Lower lung-function levels lead to higher rates of severe exacerbations and hospitalizations. Severe dyspnea and recurring exacerbations are related to high health-care resource utilization, including frequent emergency visits and hospital admissions. More than half (33%) of patients with MRC breathlessness scale scores 3–5 in the Global Hidden Depths of COPD survey had experienced an exacerbation requiring hospitalization in the previous year.

Worsening (higher) BODE index score is associated with increased risk of exacerbation and hospital admission for an exacerbation. Poor health status predicts increased hospitalizations. Frequency and proximity of previous exacerbations predicts subsequent exacerbations and hospitalizations. History of hospitalized exacerbations in the past 12 months is the strongest risk factor for future hospitalized exacerbation in ECLIPSE. Severe exacerbation risk increases threefold after the second exacerbation requiring hospitalization and 24-fold after the tenth exacerbation. Prior ER visits are a significant risk factor for exacerbation-related hospitalization.

Older age is a significant risk factor for hospitalization. Older age is associated with shorter time to first readmission. Patients with a longer history of COPD (>5 years) are approximately twice as likely to have frequent readmissions for COPD exacerbations. Presence of significant comorbidities (eg, diabetes mellitus, cardiac insufficiency, ischemic heart disease) is associated with increase in risk, specifically for severe exacerbations requiring hospitalization. Comorbidities associated with increased risk of hospitalization include cardiovascular disease, diabetes, asthma, psychiatric disorders (anxiety, depression), gastroesophageal reflux disease, and pulmonary hypertension. Eighty-two percent of patients admitted to hospital for exacerbations across the continent.

The median age of patients admitted to hospital was 72 years. Many patients requiring hospitalization due to an exacerbation had severe (Global Initiative for Chronic Obstructive Lung Disease [GOLD] grade 3, 39%) or very severe (GOLD grade 4, 22%) underlying COPD, but the proportion of patients with mild or moderate disease who experienced a severe exacerbation was not insignificant (GOLD grade 1, 15%; GOLD grade 2, 23%). Major comorbidities included other pulmonary diseases (21%), congestive heart failure (20%), and diabetes (20%).

A systematic review of 37 COPD-treatment studies, with mean forced expiratory volume in 1 second percentage predicted lung function of mostly 35–60%, estimated the annual frequency of severe exacerbations according to GOLD severity at 0.11 (GOLD grade 1, 95% CI 0.02–0.56), 0.16 (GOLD grade 2, 95% CI 0.07–0.33), 0.22 (GOLD grade 3, 95% CI 0.2–0.23), and 0.28 (GOLD grade 4, 95% CI 0.14–0.63). In the ECLIPSE study, exacerbations were more frequent and severe with increased severity of disease: the proportion of patients hospitalized for COPD exacerbations with GOLD grade 2–4 (moderate–very severe COPD) was 7%, 18%, and 33%, respectively.

When we used the GOLD 2017 “ABCD” assessment tool (in which the lettering scale reflects symptom burden and risk of exacerbation) to analyze patients included in the UPLIFT trial, a greater proportion of patients

<table>
<thead>
<tr>
<th>Table 1 Risk factors for severe exacerbations in COPD</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Proposed risk factors for severe exacerbations by category</strong></td>
</tr>
<tr>
<td><strong>Lung function</strong></td>
</tr>
<tr>
<td>• Lower lung-function levels lead to higher rates of severe exacerbations and hospitalizations.</td>
</tr>
<tr>
<td><strong>Dyspnea</strong></td>
</tr>
<tr>
<td>• Severe dyspnea and recurring exacerbations are related to high health-care resource utilization, including frequent emergency visits and hospital admissions.</td>
</tr>
<tr>
<td>• More than half (33%) of patients with MRC breathlessness scale scores 3–5 in the Global Hidden Depths of COPD survey had experienced an exacerbation requiring hospitalization in the previous year.</td>
</tr>
<tr>
<td><strong>BODE index</strong></td>
</tr>
<tr>
<td>• Worsening (higher) BODE index score is associated with increased risk of exacerbation and hospital admission for an exacerbation.</td>
</tr>
<tr>
<td><strong>Health status</strong></td>
</tr>
<tr>
<td>• Poor health status predicts increased hospitalizations.</td>
</tr>
<tr>
<td>• Impairment in health status and quality of life in COPD is a marker of risk for both frequent exacerbations and hospital admissions.</td>
</tr>
<tr>
<td><strong>Prior exacerbations, hospital visits, and hospital admissions</strong></td>
</tr>
<tr>
<td>• Frequency and proximity of previous exacerbations predicts subsequent exacerbations and hospitalizations.</td>
</tr>
<tr>
<td>• History of hospitalized exacerbations in the past 12 months is the strongest risk factor for future hospitalized exacerbation in ECLIPSE.</td>
</tr>
<tr>
<td>• Severe exacerbation risk increases threefold after the second exacerbation requiring hospitalization and 24-fold after the tenth exacerbation.</td>
</tr>
<tr>
<td>• Prior ER visits are a significant risk factor for exacerbation-related hospitalization.</td>
</tr>
<tr>
<td><strong>Age</strong></td>
</tr>
<tr>
<td>• Older age is a significant risk factor for hospitalization.</td>
</tr>
<tr>
<td>• Older age is associated with shorter time to first readmission.</td>
</tr>
<tr>
<td><strong>Duration of COPD</strong></td>
</tr>
<tr>
<td>• Patients with a longer history of COPD (&gt;5 years) are approximately twice as likely to have frequent readmissions for COPD exacerbations.</td>
</tr>
<tr>
<td><strong>Comorbidities</strong></td>
</tr>
<tr>
<td>• Presence of significant comorbidities (eg, diabetes mellitus, cardiac insufficiency, ischemic heart disease) is associated with increase in risk, specifically for severe exacerbations requiring hospitalization.</td>
</tr>
<tr>
<td>• Comorbidities associated with increased risk of hospitalization include cardiovascular disease, diabetes, asthma, psychiatric disorders (anxiety, depression), gastroesophageal reflux disease, and pulmonary hypertension.</td>
</tr>
<tr>
<td><strong>Blood-gas levels</strong></td>
</tr>
<tr>
<td>• Hypercapnia and poor arterial blood-oxygen saturation have been reported as being associated with hospitalization for a COPD exacerbation.</td>
</tr>
<tr>
<td><strong>BMI</strong></td>
</tr>
<tr>
<td>• Low BMI is associated with hospitalization and readmission for COPD exacerbation.</td>
</tr>
<tr>
<td><strong>Smoking status</strong></td>
</tr>
<tr>
<td>• Evidence regarding impact of smoking status on risk of hospitalization is mixed, with some studies suggesting current smoking was associated with lower risk of hospitalization, others finding no association, and others indicating that smoking cessation reduces risk.</td>
</tr>
<tr>
<td><strong>Use of LTOT</strong></td>
</tr>
<tr>
<td>• Patients on LTOT are at increased risk of hospitalization for exacerbation.</td>
</tr>
<tr>
<td>• Underprescription of LTOT increases the risk of hospitalization, due to severe exacerbations.</td>
</tr>
<tr>
<td><strong>Socioeconomic status</strong></td>
</tr>
<tr>
<td>• Poorer health literacy increases the likelihood of COPD-related hospitalizations and emergency department visits.</td>
</tr>
<tr>
<td>• Lowest education category and lowest income class are associated with a greater risk of hospitalization for exacerbations.</td>
</tr>
<tr>
<td>• Unmarried patients are at greater risk of hospital readmission than those with spousal support; patients requiring social work intervention are likely to have an increased length of stay for hospitalized COPD exacerbations.</td>
</tr>
</tbody>
</table>

**Abbreviations:** BMI, body-mass index; BODE, BMI, airway obstruction, dyspnea, and exercise capacity; ER, emergency room; LTOT, long-term oxygen therapy; MRC, Medical Research Council.
(35%) in GOLD group D with higher symptom burden or exacerbation risk had experienced a severe exacerbation compared with patients with lower symptom burden or exacerbation risk (GOLD group A, 14%; group B, 22%; group C, 21%). Adjusted rates of severe COPD exacerbations (per patient-year) were (mean [95% CI]): GOLD group A, 0.07 (0.05–0.09); group B, 0.13 (0.12–0.14); group C, 0.13 (0.09–0.19); group D 0.28 (0.26–0.31) (Boehringer Ingelheim, data on file, 2017).

What are the recommended treatment strategies for severe COPD exacerbations?

Mortality statistics showing that the risk of death is highest in the first week following admission highlight the importance of prompt and effective intervention for patients admitted to hospital with severe exacerbations.4 Pharmacological treatment options for severe exacerbations include bronchodilators (short-acting β-agonists with or without short-acting anticholinergics), corticosteroids (CSs) (eg, prednisone), and antibiotics when the symptoms are suggestive of bacterial infection (eg, change in sputum characteristics).11,12 Full recommendations for the pharmacological treatment of exacerbations using bronchodilators can be found in published guidelines.11,12

Antibiotics

Based on available evidence that shows a reduced risk of short-term mortality and treatment failure, current GOLD recommendations are for antibiotic treatment in those patients whose exacerbation symptoms indicate likely bacterial infection, ie, increased sputum volume and purulence, as well as increased dyspnea.11 Antibiotics are also recommended for patients with exacerbations requiring mechanical ventilation, as this has been shown significantly to reduce mortality and the risk of secondary pneumonia.11

In this era of increasing antibiotic resistance, two key considerations are avoiding the use of unnecessary antibiotic treatment (ie, in patients who do not show an increase in purulent sputum)13 and ensuring that empiric treatment choices are selected based on knowledge of local bacteria-resistance patterns.11 Treatment selection might also consider how the microbiology may differ in patients with severe exacerbations versus patients with mild or moderate disease.79 The most common pathogens isolated from sputum and bronchoscopy samples are *Haemophilus influenzae*, *Streptococcus pneumoniae*, and *Moraxella catarrhalis*. However, in patients with more severe disease and/or exacerbations, Gram-negative pathogens, such as *Pseudomonas aeruginosa*, are common,79,80 and may prove more challenging to treat in light of increasing resistance in some regions. The role of procalcitonin as a potential biomarker for optimizing antibiotic-treatment strategy in COPD exacerbations (especially those induced by bacterial infection) is also being investigated; however, as yet its role is unproven.81,82 Further studies are required to guide optimal antibiotic treatment, based on investigation of biomarkers to identify patients expected to benefit, and to establish the optimal duration of treatment.83

Corticosteroids

In the treatment of acute exacerbations of COPD, systemic CSs have been shown to reduce length of stay in hospital, provide earlier improvement in lung function and symptoms, and reduce the risk of treatment failure or relapse.84 However, a US study has noted that critically ill patients admitted directly to an intensive care unit for the treatment of severe exacerbations often receive higher-than-recommended doses of CSs to combat inflammation in the lungs.85 A large cohort study of patients receiving low-dose (methylprednisolone ≤240 mg/day; ~6,000 patients) or high-dose (methylprednisolone >240 mg/day; ~11,000 patients) CS treatment found that outcomes were in fact worse for patients treated with high doses of CSs.86 Higher doses were on average associated with longer hospital stays, longer time in the intensive care unit, and corresponding higher costs, as well as more steroid-related adverse events, compared with patients on lower doses of CSs. No significant difference was observed between patients treated with oral or intravenous CSs in terms of treatment failure, hospital readmissions, or length of hospitalization; however, intravenous therapy might increase the risk of adverse events. Overall, available evidence does not recommend one method of administration over the other.86

Optimal dosing regimens for oral CSs in patients with severe exacerbations have not been studied extensively; however, prolonged CS treatment is not recommended, in consideration of the adverse effects and lack of evidence for benefits of extended treatment. A recent meta-analysis suggested that a 5-day course of oral CSs was likely to be sufficient to treat patients hospitalized with a COPD exacerbation, and outcomes were not inferior to those with longer (10- to 14-day) courses.87

Oxygen therapy

During exacerbations, patients with COPD may become significantly hypoxic.88 Severe exacerbations disrupt the normal ventilation:perfusion ratio in the lungs, resulting in systemic hypoxemia, with poorly oxygenated blood returned to the left atrium.89 In a Spanish cohort of 2,487 patients
attending the emergency department due to acute exacerbation of COPD, 50% had hypoxemia (arterial oxygen saturation \([\text{SatO}_2] <90\%) .^{99} \) Oxygen therapy is a key component in hospital treatment of patients with a severe exacerbation.\(^{11}\)

There are marked variations in the response of individual patients to oxygen, and oxygen therapy may lead to hypercapnia and acidosis.\(^{99} \) Titrated controlled oxygen treatment for acute exacerbations has been associated with less acidosis, a lower requirement for assisted ventilation, and reduced mortality compared with high-flow oxygen.\(^{91,92} \) GOLD recommends titrating supplemental oxygen therapy to achieve a target saturation of 88%–92%.\(^{11} \) To achieve optimal oxygenation without carbon dioxide retention and/or worsening acidosis, blood gases should be monitored frequently.\(^{11} \)

**Ventilatory support**
Several studies have characterized the physiological changes that occur during or immediately after acute exacerbations, and have shown that lung hyperinflation and worsening airflow obstruction are the critical factors that determine the course of respiratory symptoms.\(^{93-95} \) Ventilatory support for the management of respiratory failure can be provided by either noninvasive or invasive methods, dependent upon patient criteria. Since its introduction in the 1980s, the use of noninvasive ventilation (NIV) has significantly increased, and has been associated with a reduction in mortality of approximately 40% and significantly reduced risk of treatment failure, complications, and endotracheal intubation.\(^{96} \) NIV is indicated in patients with respiratory acidosis (arterial pH \( \leq 7.35 \)), severe dyspnea with clinical indicators of respiratory muscle fatigue, increased work of breathing, or both, and in patients on supplemental oxygen therapy who continue to experience persistent hypoxemia.\(^{11} \)

It has been observed that patients who fail NIV and then transition to invasive mechanical ventilation have a higher rate of mortality compared with patients who start directly with invasive ventilation.\(^{97} \) Therefore, it is important to evaluate further the factors that predict early failure of NIV: to date, studies have identified severe acidosis, tachycardia, high APACHE II score, and severe hypoxemia as risk factors for NIV failure in patients with COPD exacerbations.\(^{12,97-99} \) Further guidance on the use of ventilator support can be found in the GOLD report and ATS/ERS guidelines.\(^{11,12} \)

**Preventing severe COPD exacerbations**
The profound impact of severe COPD exacerbations on patients’ well-being and the substantial economic burden associated with hospitalization highlights the importance of risk reduction as a key component of COPD management. Strategies to reduce the risk of subsequent exacerbations following hospitalization might be considered secondary prevention in a similar way to the approach used in cardiology, using the index event as a critical point to review treatment and ensure optimal patient management to minimize the risk of future exacerbations.

**Risk-factor modification**

**Smoking cessation**
Smoking cessation has a substantial influence on the natural history of COPD, and is associated with a decrease in symptoms and improved health status. In a large observational cohort study of 23,971 US veterans, ex-smokers had a significantly reduced risk of COPD exacerbation compared with current smokers.\(^{100} \) The magnitude of reduction in risk was dependent on the duration of smoking cessation. Despite the evidence supporting the significant risk reductions that can be achieved through smoking cessation, a substantial proportion of patients continue to smoke. For example, the EU COPD audit data showed that approximately a third of patients admitted to hospital with severe exacerbations in Europe in 2010–2011 were current smokers.\(^{10} \) Reinforcing participation in smoking-cessation programs represents a significant opportunity to reduce exacerbation risk by targeting a modifiable risk factor, particularly at hospital discharge, when patients may be particularly motivated to stop smoking to avoid experiencing another severe exacerbation.

**Vaccinations**
Vaccination against influenza is a highly cost-effective intervention for exacerbation reduction,\(^{101} \) and is recommended for all patients.\(^{11} \) In a retrospective cohort study of 1,323 patients with COPD, vaccination against seasonal influenza significantly reduced hospitalization due to exacerbation versus patients who were not vaccinated.\(^{102} \) Protection may be enhanced by the addition of pneumococcal vaccine (although additive effects of influenza and pneumococcal vaccinations were only observed during the first year after vaccination).\(^{103} \)

**Pulmonary rehabilitation**
Pulmonary rehabilitation may be offered after a severe exacerbation with the aim of restoring preexacerbation functional status, resuming physical activities in daily life, improving QoL, and reducing the risk of further exacerbations.\(^{11,101} \) The reported benefits of pulmonary rehabilitation on exacerbations of COPD have been mixed, but the balance of evidence supports its use.\(^{104,105} \) In a meta-analysis of nine trials in
patients who had been hospitalized for an exacerbation, pulmonary rehabilitation significantly reduced future hospital admissions and mortality and improved QoL compared with usual community care. Pulmonary rehabilitation may reduce future exacerbations by targeting modifiable risk factors for readmission, such as physical inactivity, reduced exercise capacity, impaired physical function, central desensitization to dyspnea, anxiety, and depression. Although the benefits of pulmonary rehabilitation are increasingly recognized and it is recommended in treatment guidelines for the prevention of acute exacerbations, availability of pulmonary rehabilitation programs is often limited. For example, among hospitals surveyed in the EU COPD audit, only half offered a pulmonary rehabilitation program to patients discharged following a severe exacerbation (0%-90% across countries). Identifying barriers to availability and uptake of pulmonary rehabilitation would be a useful step toward improving prevention strategies.

**Self-management intervention programs**

Self-management intervention programs improve outcomes for many chronic diseases. A disease-management plan for COPD should include an action plan for exacerbation prevention, designed in partnership with the physician and the patient, and taking into account the patient’s experience of an acute (severe) exacerbation. Evidence for benefits of self-management plans for the prevention of COPD exacerbations is mixed. In one study, a self-management program for patients with COPD (comprising education on the disease, coughing and breathing techniques, energy conservation during daily activities, relaxation exercises, symptom prevention and control, an acute-exacerbation plan of action, and lifestyle advice, including nutrition and exercise) led to a 40% reduction in hospital admissions for exacerbations compared with usual care over 12 months. Emergency visits were also reduced by 41% and unscheduled physician visits by 59%. Extended follow-up for 2 years demonstrated sustained benefits. A similar benefit was observed using a simplified disease-management program (a single education session, action plan for self-management of exacerbations, monthly follow-up calls). However, another randomized trial of patients with COPD found that a home education and management program did not result in a reduction in admissions for exacerbations, and in fact showed an unexpected increase in mortality. A Cochrane review suggested that overall, the balance of evidence supports the use of self-management programs to reduce the probability of respiratory-related hospitalizations, and the American College of Chest Physicians and Canadian Thoracic Society and ATS/ERS guidelines advocate their use.

**Pharmacological strategies**

**Bronchodilators**

Pharmacological agents currently available as maintenance treatment for COPD have varying efficacy for the prevention of COPD exacerbations. Long-term bronchodilator treatment with a long-acting muscarinic antagonist (LAMA), long-acting β2-agonist (LABA), or a combination of both have been shown to reduce the risk of severe exacerbations significantly. Table 2 provides a summary of large-scale randomized trials comparing the efficacy of inhaled maintenance COPD therapies in reducing exacerbation risk. The evidence base for LAMAs appears to be strongest, with a recent systematic review demonstrating that tiotropium is beneficial in reducing exacerbation risk versus placebo or other maintenance treatments, with longer time to first exacerbation event and fewer exacerbations (including hospitalizations) than either placebo or most active-comparator treatments. A fixed-dose LAMA–LABA combination of glycopyrronium and indacaterol successfully reduced moderate–severe exacerbations in patients of GOLD grade 3 or 4 compared with glycopyrronium alone, but this was not significant for severe exacerbations only. The FLAME study found that the combination of indacaterol–glycopyrronium significantly prolonged time to first severe exacerbation compared with salmeterol–fluticasone, although the annual rate of severe exacerbations was not significantly different between the two treatment groups; however, it has to be noted that the FLAME study was not powered to analyze differences in severe exacerbations between groups.

Despite the wealth of evidence to support tiotropium and other LAMAs in reducing the risk of severe exacerbations, data from the EU COPD audit indicate that they may be underutilized. Fewer than half of patients were on LAMA treatment prior to hospitalization, even though this patient population had been hospitalized once on average in the past year. Discharge following an exacerbation presents a pivotal opportunity to ensure that maintenance treatment is optimized, in line with goal-directed treatment guidelines to reduce the risk of disease progression, exacerbations, and mortality. Medication review at discharge, and at the same time also ensuring correct inhaler technique, represents a valuable opportunity for improvements in pharmacological approaches to reducing exacerbation risk. Adhering to COPD-treatment guideline recommendations is likely to...
Table 2: Randomized, controlled clinical trials assessing the effectiveness of inhaled long-acting bronchodilators, ICS, or combination therapy for reducing the risk of severe (hospitalized) exacerbations of COPD

<table>
<thead>
<tr>
<th>Treatment comparison</th>
<th>Trial</th>
<th>Patients, N; study length</th>
<th>Outcomes</th>
<th>Results</th>
<th>Description</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Monotherapy versus placebo or monotherapy</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tiotropium Respimat® versus placebo</td>
<td>Bateman et al(^{114})</td>
<td>3,991; 48 weeks</td>
<td>Time to first severe exacerbation</td>
<td>Placebo &lt; tiotropium</td>
<td>HR 0.73, 95% CI 0.59–0.9</td>
<td>&lt;0.005</td>
</tr>
<tr>
<td></td>
<td>Bateman et al(^{115}) (two duplicate studies)</td>
<td>1,990; 1 year</td>
<td>Risk of severe exacerbation</td>
<td>Tiotropium ↓ placebo</td>
<td>RR 0.81, 95% CI 0.7–0.93</td>
<td>&lt;0.005</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Annual incidence of severe exacerbations</td>
<td>Tiotropium (Respimat® 5 µg) ↓ placebo</td>
<td>0.12 and 0.2 hospitalizations per patient-year</td>
<td>Not reported</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Patients with ≥1 hospitalization</td>
<td>Tiotropium (Respimat® 5 µg) = placebo*</td>
<td>5.8% versus 6.7% for placebo</td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Rate of severe exacerbations</td>
<td>Tiotropium ↓ placebo, by 47%</td>
<td>0.086 versus 0.161</td>
<td>0.019</td>
</tr>
<tr>
<td></td>
<td>Casaburi et al(^{116}) (two duplicate studies)</td>
<td>921; 1 year</td>
<td>Patients with ≥1 hospitalization</td>
<td>Tiotropium (Respimat® 5 µg) ↓ placebo, by 41%</td>
<td>0.12 and 0.2 hospitalizations per patient-year</td>
<td>0.05</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Days in hospital for exacerbation</td>
<td>Tiotropium ↓ placebo, by 50%</td>
<td>0.6 days per patient-year versus 1.2 days/patient-year</td>
<td>0.023</td>
</tr>
<tr>
<td></td>
<td>Chan et al(^{117})</td>
<td>913; 48 weeks</td>
<td>Number of severe exacerbations</td>
<td>Tiotropium = placebo</td>
<td>0.13 versus 0.15</td>
<td>0.557</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Number of days in hospital</td>
<td>Tiotropium = placebo</td>
<td>1.14 versus 1.16 days/patient-year</td>
<td>0.775</td>
</tr>
<tr>
<td></td>
<td>Niewoehner et al(^{118})</td>
<td>1,829; 6 months</td>
<td>Time to first severe exacerbation</td>
<td>Placebo = tiotropium(^{11})</td>
<td>HR 0.73, 95% CI 0.53–1.01</td>
<td>0.055</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Patients with ≥1 hospitalization</td>
<td>Tiotropium = placebo(^{11})</td>
<td>7% versus 9.5%, OR 0.72, 95% CI 0.51–1.01</td>
<td>0.056</td>
</tr>
<tr>
<td></td>
<td>UPLIFT(^{119})</td>
<td>5,993; 4 years</td>
<td>Time to first hospitalized exacerbation</td>
<td>Placebo &lt; tiotropium</td>
<td>HR 0.86, 95% CI 0.78–0.95</td>
<td>0.002</td>
</tr>
<tr>
<td><strong>Glycopyrronium versus placebo</strong></td>
<td>GLOW(^{120})</td>
<td>822; 26 weeks</td>
<td>Risk of severe exacerbation</td>
<td>Glycopyrronium &lt; placebo</td>
<td>HR 0.35, 95% CI 0.14–0.857</td>
<td>0.022</td>
</tr>
<tr>
<td><strong>Tiotropium HandHaler® versus glycopyrronium study also included an indacaterol-glycopyrronium arm</strong></td>
<td>SPARK (Wedzicha et al(^{120}))</td>
<td>2,224; 64 weeks</td>
<td>Annual rate of severe exacerbations</td>
<td>Tiotropium ↓ glycopyrronium</td>
<td>0.08 versus 0.12, RR 1.43, 95% CI 1.05–1.97</td>
<td>0.025</td>
</tr>
<tr>
<td></td>
<td>Brusasco et al(^{121}) (two duplicate studies)</td>
<td>1,207; 6 months</td>
<td>Time to first hospitalized exacerbation</td>
<td>Not reported</td>
<td>Not applicable</td>
<td>Not reported</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Hospital admissions</td>
<td>Tiotropium = salmeterol or placebo(^{1})</td>
<td>0.1 versus 0.17 or 0.15</td>
<td>NS</td>
</tr>
<tr>
<td></td>
<td>INVIGORATE(^{122})</td>
<td>3,444; 52 weeks</td>
<td>Annual rate of severe (hospitalized) exacerbations</td>
<td>Tiotropium ↓ indacaterol</td>
<td>0.07 versus 0.1, RR 1.36, 95% CI 1.03–1.79</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td>POET-COPD(^{123})</td>
<td>7,376; 1 year</td>
<td>Risk of severe exacerbations</td>
<td>Tiotropium &lt; salmeterol, by 28%</td>
<td>HR 0.72, 95% CI 0.61–0.85</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>TIOSPIR(^{124})</td>
<td>17,135; 2.3 years</td>
<td>Risk of severe exacerbation</td>
<td>Tiotropium (Respimat®) = tiotropium (HandHaler®)</td>
<td>HR 1.02, 95% CI 0.93–1.13</td>
<td>0.64</td>
</tr>
</tbody>
</table>

(Continued)
Table 2 (Continued)

<table>
<thead>
<tr>
<th>Treatment comparison</th>
<th>Trial</th>
<th>Patients, N; study length</th>
<th>Outcomes Measures</th>
<th>Results</th>
<th>Description</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Combination therapy versus monotherapy or placebo</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mometasone–formoterol versus mometasone, formoterol, and placebo</td>
<td>Doherty et al125</td>
<td>1,201; 1 year (26-week treatment, 26-week safety extension)</td>
<td>Incidence of severe exacerbations</td>
<td>Mometasone + formoterol 400+10 µg BID, mometasone + formoterol 200+10 µg BID, mometasone and formoterol monotherapies BID ↓ placebo BID</td>
<td>4.4%, 1.7%, 2.4%, 2.1%, and 5.1%, respectively</td>
<td>Not reported</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Incidence of moderate or severe first exacerbations</td>
<td>Mometasone + formoterol 400+10 µg BID, mometasone + formoterol 200+10 µg BID ↓ placebo</td>
<td>15.4% and 12.8%, respectively, versus 24.6%</td>
<td>≤0.006</td>
</tr>
<tr>
<td>Umeclidinium–vilanterol versus umeclidinium and placebo</td>
<td>Donohue et al126</td>
<td>563; 1 year</td>
<td>Patients with ≥1 hospitalization</td>
<td>Umeclidinium–vilanterol and umeclidinium ↓ placebo</td>
<td>6% and 7% versus 12%</td>
<td>Not reported</td>
</tr>
<tr>
<td>Tiotropium HandiHaler® versus fluticasone–salmeterol</td>
<td>INSPIRE127</td>
<td>1,323; 2 years</td>
<td>Incidence of exacerbations requiring hospitalizations</td>
<td>Tiotropium = fluticasone–salmeterol</td>
<td>13% versus 16%</td>
<td>0.085</td>
</tr>
<tr>
<td>Fluticasone–vilanterol versus vilanterol</td>
<td>NCT01009463/NCT01017952128</td>
<td>3,255; 1 year</td>
<td>Annual rate of severe exacerbations (pooled analysis of study 1 and study 2)</td>
<td>Fluticasone + vilanterol 50+25 µg = vilanterol 25 µg</td>
<td>0.08 versus 0.1, RR 0.8, 95% CI 0.6–1.2</td>
<td>0.3133</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Fluticasone + vilanterol 100+25 µg = vilanterol 25 µg</td>
<td>0.09 versus 0.1, RR 0.9, 95% CI 0.6–1.4</td>
<td>0.6948</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Fluticasone + vilanterol 200+25 µg = vilanterol 25 µg</td>
<td>0.08 versus 0.1, RR 0.8, 95% CI 0.5–1.2</td>
<td>0.2802</td>
</tr>
<tr>
<td>Budesonide–formoterol versus formoterol</td>
<td>Sharafkhaneh et al129</td>
<td>1,219; 1 year</td>
<td>Number of severe exacerbations per patient-treatment year</td>
<td>Budesonide + formoterol 320+9 µg BID = formoterol¹</td>
<td>0.106 versus 0.14, treatment ratio 0.732, 95% CI 0.52–1.03</td>
<td>0.07</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Budesonide + formoterol 160+9 µg BID = formoterol¹</td>
<td>0.127 versus 0.14, treatment ratio 0.878, 95% CI 0.64–1.22</td>
<td>0.433</td>
</tr>
<tr>
<td>Tiotropium HandiHaler® versus indacaterol–glycopyrronium study also included a glycopyrronium arm</td>
<td>SPARK (Wedzicha et al)130</td>
<td>2,224; 64 weeks</td>
<td>Annual rate of severe exacerbations</td>
<td>Tiotropium = indacaterol–glycopyrronium</td>
<td>0.08 versus 0.09, RR 1.16, 95% CI 0.84–1.61</td>
<td>0.36</td>
</tr>
<tr>
<td>Mometasone–formoterol versus mometasone, formoterol, and placebo</td>
<td>Tashkin et al131</td>
<td>2,251; 26 weeks</td>
<td>Incidence of severe exacerbations</td>
<td>Mometasone + formoterol 200+10 µg BID, mometasone + formoterol 400+10 µg BID, mometasone and formoterol monotherapies BID ↓ placebo</td>
<td>1.6%, 3.4%, 2.4%, and 2.4%, versus 4.2%, respectively</td>
<td>Not reported</td>
</tr>
<tr>
<td>Salmeterol–fluticasone versus placebo, salmeterol alone, and fluticasone alone</td>
<td>TORCH132</td>
<td>6,112; 3 years</td>
<td>Annual rate of severe exacerbation</td>
<td>Salmeterol–fluticasone ↓ placebo</td>
<td>0.16 versus 0.19, RR 0.83, 95% CI 0.71–0.98</td>
<td>0.03</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Salmeterol–fluticasone = salmeterol</td>
<td>0.16 versus 0.16, RR 1.02, 95% CI 0.87–1.2</td>
<td>0.79</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Salmeterol–fluticasone = fluticasone</td>
<td>0.16 versus 0.17, RR 0.95, 95% CI 0.82–1.12</td>
<td>0.56</td>
</tr>
</tbody>
</table>
Combination therapy versus other combination therapy

<table>
<thead>
<tr>
<th>Therapy</th>
<th>Study</th>
<th>Duration</th>
<th>Annual rate of severe exacerbations</th>
<th>Rate of severe exacerbations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td>FLAME (Wedzicha et al)</td>
<td>3.362; 1 year</td>
<td>0.046</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td>QUANTIFY (Buhl et al)</td>
<td>934; 26 weeks</td>
<td>2.1%</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td></td>
<td></td>
<td>0.15 versus 0.17, RR 0.87, 0.23</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td></td>
<td></td>
<td>95% CI 0.69–1.09</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td></td>
<td></td>
<td>HR 0.81, 95% CI 0.66–1</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td></td>
<td></td>
<td>2.1% versus 2.4%, RR 0.88, 0.759</td>
<td></td>
</tr>
<tr>
<td>Indacaterol–glycopyrronium</td>
<td></td>
<td></td>
<td>95% CI 0.38–2.01</td>
<td></td>
</tr>
</tbody>
</table>

Notes: Large numerical differences noted between comparators. Table includes randomized studies examining the outcome of severe (hospitalized) exacerbations in >500 patients with COPD; for HRs showing time to first severe exacerbation, < and > indicate shorter and longer time to event, or lower and higher risk of severe event versus comparator, respectively. Note: For POeT-COPD 0.05 is deemed statistically significant; \( \downarrow \) indicates a lower or RR, rate or incidence of severe exacerbations versus comparator; \( \uparrow \) indicates no statistically significant difference between comparators; NS indicates no statistically significant difference between comparators, as stated in the publication text (ie, no P-values provided).

Abbreviations: BID, bis in die (twice daily); HR, hazard ratio; ICS, inhaled corticosteroid; OR, odds ratio; RR, rate ratio.

Phosphodiesterase-4 inhibitors

Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides.

Roflumilast is an oral phosphodiesterase-4 inhibitor, indicated as a maintenance treatment for severe COPD associated with chronic bronchitis in adult patients with a history of frequent exacerbations (as add-on to bronchodilator treatment).

The 1-year REACT study showed that in symptomatic patients with severe COPD and a history of frequent exacerbations (more than three exacerbations in the previous year) and/or history of hospitalization, roflumilast significantly reduced the risk of exacerbations, 

ROSE study showed that roflumilast reduced the rate of moderate–severe exacerbations (reduced risk of 18% versus 0.88, but not for severe exacerbations only).

Large numerical differences noted between comparators. Table includes randomized studies examining the outcome of severe (hospitalized) exacerbations in >500 patients with COPD; for HRs showing time to first severe exacerbation, < and > indicate shorter and longer time to event, or lower and higher risk of severe event versus comparator, respectively. Note: For POeT-COPD 0.05 is deemed statistically significant; \( \downarrow \) indicates a lower or RR, rate or incidence of severe exacerbations versus comparator; \( \uparrow \) indicates no statistically significant difference between comparators; NS indicates no statistically significant difference between comparators, as stated in the publication text (ie, no P-values provided).

Abbreviations: BID, bis in die (twice daily); HR, hazard ratio; ICS, inhaled corticosteroid; OR, odds ratio; RR, rate ratio.

Phosphodiesterase-4 inhibitors

Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides. Phosphodiesterase-4 inhibitors do not have direct bronchodilator activity, but act as anti-inflammatory agents by inhibiting the breakdown of intracellular cyclic nucleotides.

Roflumilast is an oral phosphodiesterase-4 inhibitor, indicated as a maintenance treatment for severe COPD associated with chronic bronchitis in adult patients with a history of frequent exacerbations (as add-on to bronchodilator treatment).

The 1-year REACT study showed that in symptomatic patients with severe COPD and a history of frequent exacerbations (more than three exacerbations in the previous year) and/or history of hospitalization, roflumilast significantly reduced the risk of exacerbations.

Lack of significant benefits of ICS–LABA treatment over LABA monotherapy was demonstrated in a systematic review of 18 randomized controlled trials, which failed to show a significant decrease in the number of severe exacerbations.

Due to the possible adverse events of pneumonia and fractures resulting from inhaled CS (ICS) treatment, current guidelines recommend ICS–LABA combination therapy only in patients with severe airflow limitation at minimum and often with severe exacerbations that are not controlled adequately with long-acting bronchodilators alone.

The overall result of significant benefits of ICS–LABA treatment over LABA monotherapy was demonstrated in a systematic review of 18 randomized controlled trials, which failed to show a significant decrease in the number of severe exacerbations.

Due to the possible adverse events of pneumonia and fractures resulting from inhaled CS (ICS) treatment, current guidelines recommend ICS–LABA combination therapy only in patients with severe airflow limitation at minimum and often with severe exacerbations that are not controlled adequately with long-acting bronchodilators alone.
patients with severe COPD, and evidence from these trials suggests that the benefits of roflumilast are demonstrated most consistently in patients with severe–very severe COPD associated with chronic bronchitis and a history of frequent exacerbations. Concerns have been raised regarding the safety profile of roflumilast, due to higher incidences of adverse events compared to other COPD medications; however, a benefit–harm analysis of the use of roflumilast found a net benefit in patients with a high risk of severe exacerbations.

**Mucolytics**

Mucolytic agents (specifically N-acetylcysteine [NAC]) may have a beneficial role in exacerbation reduction, with studies indicative of a reduction in exacerbation frequency. Although a Cochrane systematic review concluded that the small reduction in overall acute exacerbations achieved with mucolytics may have been overestimated, some studies suggest that higher doses of mucolytics (NAC 600 mg twice daily) may be particularly effective in patients with stable COPD, moderate–severe COPD, or at high risk of exacerbations. However, these studies were not specifically powered for the analysis of severe exacerbations. The current GOLD report concludes that regular use of mucolytic agents, such as NAC and carbocysteine, reduce the risk of exacerbation when used in appropriate patients.

**Antibiotic prophylaxis**

Several studies suggest that the use of certain types of antibiotics may have a role in preventing exacerbations in certain groups of patients for whom the benefits outweigh the risks. Macrolides, which possess both antibacterial and anti-inflammatory effects, and fluoroquinolones are the classes of antibiotics studied most extensively for use in stable COPD, and prophylactic use over 1 year can reduce the rate of hospitalizations for COPD exacerbations compared with placebo. While a meta-analysis of nine studies showed that macrolides are effective in reducing the overall rate of COPD exacerbations when administered for 6–12 months, the analysis was not able to demonstrate an effect on hospitalization rates, due to the low number of hospitalizations in the studies, which were not powered to detect such differences. Prophylactic antibiotic therapy should be applied cautiously, as the evidence remains limited, and further studies will be required to determine the optimal treatment regimen and duration with prophylactic use of antibiotics, and in what patients, to maintain a positive benefit–risk balance.

**Vitamin D**

It has been suggested that vitamin D supplements may help to prevent COPD exacerbations. The ViDiCO study demonstrated that vitamin D supplementation reduced the risk of moderate and severe exacerbations in patients with COPD; however, findings have been conflicting and the benefits may only be evident in patients with severe vitamin D deficiency. The PRECOVID study will be the first randomized trial to evaluate the effects of vitamin D supplementation on exacerbation prevention (primary outcome; rates of hospitalization will be a secondary outcome), and will recruit vitamin D-deficient patients who have recently experienced an exacerbation. The results of the study will be beneficial for establishing guidelines for vitamin D supplementation with regard to reducing exacerbation rates.

**β-Blockers**

β-blocker usage is low in patients with COPD and cardiovascular comorbidities, as blockade of β-adrenergic receptors might induce bronchoconstriction and lead to bronchospasm and worsening of lung function. This is based largely on anecdotal evidence; however, review of the evidence base has indicated that cardioselective β-blockers (ie, those with greater affinity for β1-over β2-receptors) reduce overall mortality in patients with such conditions as coronary artery disease or MI and coexisting mild–moderate COPD, without producing clinically significant adverse respiratory effects, and should not be withheld. One observational cohort study (COPDGene®), which followed approximately 3,500 patients for a median of 2.1 years, indicated that the use of β-blockers in COPD patients with cardiovascular comorbidities was associated with a significant reduction in exacerbation rates, including severe exacerbations. Effects were most pronounced in patients with severe COPD (GOLD grade 3 and 4 on home oxygen therapy). Randomized trials may be warranted to confirm whether this observation represents beneficial effects of β-blockers in COPD and explore the mechanisms underlying potential protective effects.

**Readmissions**

Patients who have been admitted to hospital for a severe exacerbation of COPD are at substantial risk of rehospitalization. In the EU COPD audit, 35% of all patients admitted to hospital with COPD exacerbations were readmitted within 90 days, and approximately three-quarters of readmissions were recorded as COPD-related. Readmission is a poor outcome for patients, and presents a significant strain on resources. In some cases, readmission is the result of a poor discharge, while in others it reflects the severity of the
underlying COPD or the presence of comorbidities. Despite best efforts, many readmissions may not be preventable. The US National COPD Readmissions Summit, as well as a recent systematic review, found that the evidence base to support specific interventions that would significantly reduce 30-day readmission rates in patients hospitalized for COPD exacerbation was inadequate.

Nevertheless, readmission rates are of increasing significance to payers in both the US and the UK, where financial penalties have been imposed on hospitals with higher-than-expected readmission rates, as part of an incentive to promote optimized, value-driven care packages for patients discharged following a COPD exacerbation. From a clinical perspective, it is important to monitor whether these approaches have unintended negative consequences, such as an increase in out-of-hospital mortality resulting from the disincentive to readmit patients who might genuinely benefit from further hospital treatment.

Since a substantial proportion of readmissions are not due to COPD itself, but relate to comorbidities, effective management of these conditions and COPD is important in attempts to reduce readmission rates. Ensuring patients are on optimal maintenance treatment known to reduce exacerbation rates (as discussed earlier) at the time of discharge is essential to reduce the risk of further exacerbations and readmission.

Improving overall disease-management strategies
Several studies have revealed the frequency of clinical practices that are nonadherent to treatment guidelines, showing a disparity in treatment choices, overuse of CSs and antibiotics, and underuse of pulmonary rehabilitation. In a study conducted in the US, review of medical records indicated that patients with COPD received only 58% of the care recommended by guidelines. A survey of respiratory specialists in Canada identified important care gaps regarding interventions to prevent future exacerbations, eg, suboptimal vaccinations, care plans, and patient education (39% of patients had never been taught how to recognize the signs and symptoms of an exacerbation). In the Continuing to Confront COPD International Physician Survey (12 countries), only 57% of primary-care physicians and 58% of respiratory specialists provided treatment options concordant with the GOLD report for high-risk (GOLD grade 4) COPD. The EU COPD audit showed that COPD treatment received before admission for COPD exacerbation, inpatient treatment, use of mechanical ventilation, and discharge medications (including oxygen) varied widely across Europe. In addition, the organization of COPD care and hospital facilities was shown to vary considerably between countries.

Adherence to therapy is an important factor in successfully preventing exacerbations. Poor adherence to COPD medication detrimentally affects long-term disease outcomes. Among inhaled bronchodilators prescribed as maintenance therapy for COPD, adherence varies according to the type of medication, with higher rates of adherence observed with LAMAs (tiotropium in the study quoted) and relatively low adherence to ICSs or ICS–LABA combinations.

Conclusion
Severe exacerbations are an important cause of morbidity and mortality, and have substantial economic consequences. A large number of potential risk factors are associated with severe exacerbations, many of which are easily identifiable and modifiable with careful patient management. Previous severe exacerbations and hospital readmission are significant risk factors for mortality; therefore, targeted interventions aimed at preventing or reducing severe exacerbations should be a priority for improving patients' prognoses. Strategies to reduce hospital readmissions for COPD and related comorbidities are also required to reduce the economic burden associated with repeated hospitalizations.

Recent observational studies have reported lower rates of in-hospital mortality, reduced length of hospitalization, and fewer subsequent exacerbations over the past decade, reflecting improvements in care and greater awareness of the impact of COPD exacerbations. Nonetheless, effective prevention of severe exacerbations remains a significant unmet need in COPD. Both pharmacological and nonpharmacological interventions have a role as part of a holistic package to minimize exacerbation risk. Smoking cessation is a key intervention, and improved access to/uptake of pulmonary rehabilitation may be beneficial. Large randomized clinical trials have demonstrated the benefit of long-acting bronchodilators, such as LAMAs or LABAs, either as monotherapies or in combination, in reducing the risk of severe exacerbations in patients with COPD at all stages of severity. Overall, bronchodilators appear to be at least as effective as LABA–ICS therapy in preventing severe exacerbations of COPD. Antibiotics also have a role in selected patients, and influenza and pneumococcal vaccination can help prevent exacerbations. In conclusion, multiple opportunities to improve management of COPD and reduce risk of severe exacerbations are available, and should be considered in all patients with COPD, with particular attention paid to those who have already had a severe exacerbation.
Acknowledgments

Editorial and writing support was provided by Andree Rose, Jennifer Fuchs, and Deepi Sharda at PAREXEL, funded by Boehringer Ingelheim.

Disclosure

DMGH has received personal fees from Almirall, Boehringer Ingelheim, GlaxoSmithKline, InterMune, Pfizer, and Novartis, and non-financial support from Boehringer Ingelheim and Novartis. MM has received speaker fees from Almirall, Boehringer Ingelheim, Pfizer, AstraZeneca, Chiesi, Esteve, GlaxoSmithKline, Menarini, Novartis, Talecris-Grifols, Takeda-Nycomed, and Novartis, and consulting fees from Almirall, Boehringer Ingelheim, Pfizer, GlaxoSmithKline, Gebro Pharma, MedImmune, Novartis, Talecris-Grifols and Takeda-Nycomed. NM is an employee of Boehringer Ingelheim. BC has received fees for consulting from AstraZeneca, Boehringer Ingelheim, GlaxoSmithKline, MedImmune and Novartis.

References


