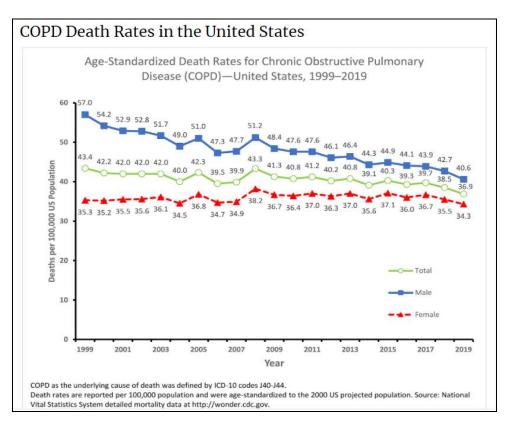


Diagnosis and Management of COPD Clinical Practice Guideline

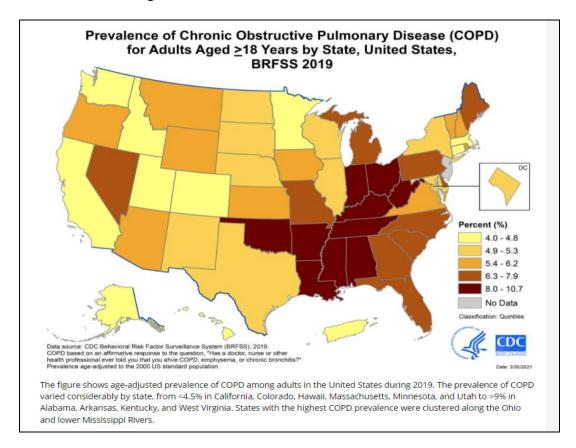
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"These guidelines are provided to assist physicians and other clinicians in making decisions regarding the care of their patients. They are not a substitute for individual judgment brought to each clinical situation by the patient's primary care provider-in collaboration with the patient. As with all clinical reference resources, they reflect the best understanding of the science of medicine at the time of publication but should be used with the clear understanding that continued research may result in new knowledge and recommendations".

According to the CDC, COPD affects more than 15 million Americans.¹ While death rates have been improving (2), COPD still accounts for more than 150,000 deaths each year.



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Prevalance in our region is between 4.9% and 6.2% of adults.²

The MedStar Health Ambulatory Best Practices Committee endorses and accepts the recommendations for care in *Global Strategy for the Diagnosis, Management and Prevention of COPD*, Global Initiative for Chronic Obstructive Lung Disease (GOLD) 2022 Report and the accompanying 2022 GOLD Pocket Guide.³ A complete copy of the documents can be downloaded at no charge for personal use at: <u>https://goldcopd.org/2022-gold-reports-2/</u>. Readers are encouraged to obtain copies of the entire document from the GOLD website. The excerpts included here are used with permission of GOLD, however, the copyright watermarks are left in place to remind the user that permission is required for uses beyond personal use.

Of note in this era of the continuing COVID-19 pandemic, the CDC recognizes chronic lung disease including COPD as a risk factor for severe disease and the most recent GOLD Guideline includes a new chapter devoted to COVID-19 and COPD. Information is evolving rapidly in this area and the reader is encouraged to seek the most up-to-date information from a variety of other sources to supplement this guideline.

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Below are the Key Points and key tables for each chapter of the GOLD guideline and are used with permission of the Global Initiative for Chronic Obstructive Lung Disease. The reader is referred to the complete document for expanded information and the references behind the key points.

KEY POINTS: Chapter 1: Definition and Overview

• Chronic Obstructive Pulmonary Disease (COPD) is a common preventable and treatable disease that is

characterized by persistent respiratory symptoms and airflow limitation that is due to airway and/or alveolar abnormalities usually caused by significant exposure to noxious particles or gases.

• The most common respiratory symptoms include dyspnea, cough and/or sputum production. These symptoms may be under-reported by patients

• The main risk factor for COPD is tobacco smoking but other environmental exposures such as biomass fuel exposure and air pollution may contribute. Besides exposures, host factors predispose individuals to develop COPD. These include genetic abnormalities, abnormal lung development and accelerated aging.

COPD may be punctuated by periods of acute worsening of respiratory symptoms, called exacerbations.

• In most patients, COPD is associated with significant concomitant chronic diseases, which increase its morbidity and mortality.

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| | FOR CONSIDERING A DIAGNOSIS OF COPD |
|---|--|
| These indicators are not diag | spirometry, if any of these indicators are present in an individual over age 40. nostic themselves, but the presence of multiple key indicators increases the OPD. Spirometry is required to establish a diagnosis of COPD. |
| Dyspnea that is: | Progressive over time. Characteristically worse with exercise. Persistent. |
| Chronic Cough: | May be intermittent and may be unproductive. Recurrent wheeze. |
| Chronic Sputum Production: | Any pattern of chronic sputum production may indicate COPD. |
| Recurrent Lower Respiratory | Tract Infections |
| History of Risk Factors: | Host factors (such as genetic factors, congenital/developmental abnormalities etc.). Tobacco smoke (including popular local preparations). Smoke from home cooking and heating fuels. Occupational dusts, vapors, fumes, gases and other chemicals. |
| Family History of COPD and/or Childhood Factors: | For example low birthweight, childhood respiratory infections etc. |
| TABLE 2.1 | |

KEY POINTS: Chapter 2: Diagnosis and Initial Assessment

• COPD should be considered in any patient who has dyspnea, chronic cough or sputum production, a history of recurrent lower respiratory tract infections and/or a history of exposure to risk factors for the disease.

• Spirometry is required to make the diagnosis; the presence of a post-bronchodilator FEV1/FVC < 0.70 confirms the presence of persistent airflow limitation.

• The goals of COPD assessment are to determine the level of airflow limitation, the impact of disease on the patient's health status, and the risk of future events (such as exacerbations, hospital admissions, or death), in order to guide therapy.

• Concomitant chronic diseases occur frequently in COPD patients, including cardiovascular disease, skeletal muscle dysfunction, metabolic syndrome, osteoporosis, depression, anxiety, and lung cancer. These comorbidities should be actively sought and treated appropriately when present as they can influence mortality and hospitalizations independently.

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Key Indicators for Considering a Diagnosis of COPD

- Dyspnea persistent, progressive over time, worse with exercise
- Chronic cough intermittent and maybe unproductive; recurrent wheezing
- Chronic sputum production
- Recurrent lower respiratory tract infection
- Risk factors tobacco smoke, occupational exposures, indoor smoke, host factors such as genetics or congenital/developmental abnormalities

Differential Diagnosis

A major differential diagnosis is asthma. Sometimes a clear distinction from COPD is not possible, in which case management is similar to that of asthma. Other conditions to consider include congestive heart failure, bronchiectasis, tuberculosis, obliterative bronchiolitis and diffuse panbronchiolitis and others.

The WHO recommends that all patients with a diagnosis of COPD should be screened once for alpha-1 antitrypsin deficiency (AATD), especially in areas with high prevalence.

Assessment

COPD assessment must consider the following aspects of the disease:

- The presence and severity of the spirometric abnormality
- Current nature and magnitude of symptoms
- History of moderate and severe exacerbations and future risk
- Presence of comorbidities

The diagnosis should be confirmed with spirometry before treatmant plan is made. Spirometry should be performed post bronchodilator. Additional investigations may include chest Xray, lung volumes and carbon monoxide diffusion capacity (DLCO), oximetry, arterial blood gas measurement and exercise testing.

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| | | 14- | |
|------------------|------------------|--|--|
| | | DW LIMITATION SEVERITY BRONCHODILATOR FEV1) | |
| In patients with | FEV1/FVC < 0.70: | | |
| GOLD 1: | Mild | $M^{FEV_1 \ge 80\%}$ predicted | |
| GOLD 2: | Moderate | $50\% \leq FEV_1 < 80\%$ predicted | |
| GOLD 3: | Severe | $30\% \leq FEV_1 < 50\%$ predicted | |
| GOLD 4: | Very Severe | FEV ₁ < 30% predicted | |
| TABLE 2.4 | | | |

Assessment of symptoms can be measured using the modified MRC (mMRC) dyspnea scale.

| | D MRC DYSPNEA SCALE [®] | |
|--|--|--|
| PLEASE TICK IN THE BC | DX THAT APPLIES TO YOU ONE BOX ONLY Grades 0 - 4 | |
| mMRC Grade 0. | I only get breathless with strenuous exercise. | |
| mMRC Grade 1. | I get short of breath when hurrying on the level or walking up a slight hill. | |
| mMRC Grade 2. | I walk slower than people of the same age on the level because of breathlessness, or I have to stop for breath when walking on my own pace on the level. | |
| mMRC Grade 3. | I stop for breath after walking about 100 meters or after a few minutes on the level. | |
| mMRC Grade 4. | I am too breathless to leave the house or I am breathless when dressing or undressing. | |
| ^a Fletcher CM. BMJ 196 TABLE 2.5 | 50; 2: 1662. | |

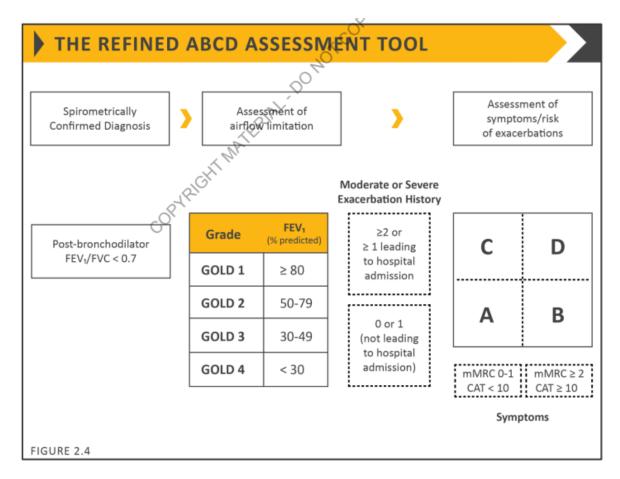
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Because patients may experience symptoms beyond dyspnea, a comprehensive assessment of symptoms using a tool such as the COPD Assessment Test CAT (below) is preferred.

| CAT™ ASSESSM | ENT | |
|---|---|-------|
| For each item below, place a mark Be sure to only select one respons | (x) in the box that best describes you currently. e for each question. | |
| EXAMPLE: I am very happy | 0 🗶 2 3 4 5 I am very sad | SCORE |
| l never cough | 012345 I cough all the time | |
| l have no phlegm (mucus) in my chest at all | 0 1 2 3 4 5 My chest is completely full of phlegm (mucus) | |
| My chest does not feel tight at all | 0 1 2 3 4 5 My chest feels very tight | |
| When I walk up a hill or one flight of stairs I am not breathless | 0 1 2 3 4 5 When I walk up a hill or one flight of stairs I am very breathless | |
| I am not limited doing any activities at home | 0 1 2 3 4 5 I am very limited doing activities at home | |
| I am confident leaving my home despite my lung condition | 0 1 2 3 4 home because of my lung condition | |
| l sleep soundly | 0 1 2 3 4 5 I don't sleep soundly because of my lung condition | |
| I have lots of energy | 0 1 2 3 4 5 I have no energy at all | |
| Reference: Jones et al. ERJ 2009 3 FIGURE 2.3 | 34 (3); 648-54. TOTAL SCORE | |

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Combining spirometry, symptomatic assessment, and risk of exacerbation in the refined ABCD assessment tool may facilitate consideration of individual therapies for a specific patient.



KEY POINTS: Chapter 3: Evidence Supporting Prevention and Maintenance Therapy

- Smoking cessation is key. Pharmacotherapy and nicotine replacement reliably increase long-term smoking abstinence rates. Legislative smoking bans and counselling, delivered by healthcare professionals, improve quit rates.
- The effectiveness and safety of e-cigarettes as a smoking cessation aid is uncertain at present.
- Pharmacologic therapy can reduce COPD symptoms, reduce the frequency and severity of exacerbations, and improve health status and exercise tolerance. Recent data suggests beneficial effects on mortality.
- Each pharmacologic treatment regimen should be individualized and guided by the severity of symptoms, risk of exacerbations, side-effects, co-morbidities, drug availability

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and cost, and the patient's response, preference and ability to use various drug delivery devices.

- Inhaler technique needs to be assessed regularly.
- COVID-19 vaccines are highly effective against SARS-CoV-2 infection and people with COPD should have the COVID-19 vaccination in line with national recommendations.
- Influenza vaccination decreases the incidence of lower respiratory tract infections.
- Pneumococcal vaccination decreases lower respiratory tract infections.
- CDC recommends the Tdap vaccination (dTaP/dTPa) in COPD patients to protect against pertussis, tetanus and diphtheria, in those who were not vaccinated in adolescence and Zoster vaccine to protect against shingles for adults with COPD aged <u>></u> 50 years.
- Pulmonary rehabilitation with its core components, including exercise training combined with disease-specific education, improves exercise capacity, symptoms and quality of life across all grades of COPD severity.
- In patients with severe resting chronic hypoxemia, long-term oxygen therapy improves survival.
- In patients with stable COPD and resting or exercise-induced moderate desaturation, long-term oxygen treatment should not be prescribed routinely. However, individual patient factors must be considered when evaluating the patient's need for supplemental oxygen.
- In patients with severe chronic hypercapnia and a history of hospitalization for acute respiratory failure, long-term non-invasive ventilation may decrease mortality and prevent re-hospitalization.
- In select patients with advanced emphysema refractory to optimized medical care, surgical or bronchoscopic interventional treatments may be beneficial.
- Palliative approaches are effective in controlling symptoms in advanced COPD.

Smoking Cessation should be encouraged. Counseling, nicotine replacements and other pharmacological support should be provided.

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Vaccinations for Stable COPD

| VACCINATION FOR STABLE COPD |
|--|
| • Influenza vaccination reduces serious illness and death in COPD patients (Evidence B). |
| • The WHO and CDC recommend SARS-Cov-2 (COVID-19) vaccination for people with COPD (Evidence B). |
| The 23-valent pneumococcal polysaccharide vaccine (PPSV23) has been shown to reduce the incidence of community-acquired pneumonia in COPD patients aged < 65 years with an OEV₁ < 40% predicted and in those with comorbidities (Evidence B). |
| In the general population of adults ≥ 65 years the 13-valent conjugated pneumococcal vaccine (PCV13) has demonstrated significant efficacy in reducing bacteremia & serious invasive pneumococcal disease (Evidence B). |
| The CDC recommends Tdap (dTaP/dTPa) vaccination to protect against pertussis (whooping cough) for adults with COPD who were not vaccinated in adolescence (Evidence B) and Zoster vaccine to protect against shingles for adults with COPD aged ≥ 50 years (Evidence B). |
| TABLE 3.2 |
| |

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Pharmacological Therapy for Stable COPD

| BRONCHODILATORS IN STABLE COPD |
|---|
| Inhaled bronchodilators in COPD are central to symptom management and commonly given on a regular basis to prevent or reduce symptoms (Evidence A). |
| Regular and as-needed use of SABA or SAMA improves FEV₁ and symptoms (Evidence A). |
| Combinations of SABA and SAMA are superior compared to either medication alone in improving FEV₁ and symptoms (Evidence A). |
| LABAs and LAMAs significantly improve lung function, dyspnea, health status, and reduce exacerbation rates (Evidence A). |
| LAMAs have a greater effect on exacerbation reduction compared with LABAs (Evidence A) and decrease hospitalizations (Evidence B). |
| Combination treatment with a LABA and LAMA increases FEV₁ and reduces symptoms compared to monotherapy (Evidence A). |
| Combination treatment with a LABA/LAMA reduces exacerbations compared to monotherapy (Evidence B). |
| Tiotropium improves the effectiveness of pulmonary rehabilitation in increasing exercise performance (Evidence B). |
| Theophylline exerts a small bronchodilator effect in stable COPD (Evidence A) and that is associated with modest symptomatic benefits (Evidence B). |
| TABLE 3.4 |

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ANTI-INFLAMMATORY THERAPY IN STABLE COPD

INHALED CORTICOSTEROIDS

- An ICS combined with a LABA is more effective than the individual components in improving lung function and health status and reducing exacerbations in patients with exacerbations and moderate to very severe COPD (Evidence A).
- Regular treatment with ICS increases the risk of pneumonia especially in those with severe disease (Evidence A).
- Triple inhaled therapy of LABA/LAMA/ICS improves lung function, symptoms and health status, and reduces exacerbations, compared to LABA/ICS, LABA/LAMA or LAMA monotherapy (Evidence A). Recent data suggest a beneficial effect versus fixed-dose LABA/LAMA combinations on mortality in symptomatic COPD patients with a history of frequent and/or severe exacerbations.

ORAL GLUCOCORTICOIDS

 Long-term use of oral glucocorticoids has numerous side effects (Evidence A) with no evidence of benefits (Evidence C).

PDE4 INHIBITORS

- In patients with chronic bronchitis, severe to very severe COPD and a history of exacerbations:
 - » A PDE4 inhibitor improves lung function and reduces modera@and severe exacerbations (Evidence A).
 - » A PDE4 inhibitor improves lung function and decreases exacerbations in patients who are on fixed-dose LABA/ICS combinations (Evidence A).

ANTIBIOTICS

- . Long-term azithromycin and erythromycin therapy reduces exacerbations over one year (Evidence A).
- Treatment with azithromycin is associated with an increased incidence of bacterial resistance (Evidence A) and hearing test impairments (Evidence B).

MUCOREGULATORS AND ANTIOXIDANT AGENTS

 Regular treatment with mucolytics such as erdosteine, carbocysteine and NAC reduces the risk of exacerbations in select populations (Evidence B).

OTHER ANTI-INFLAMMATORY AGENTS

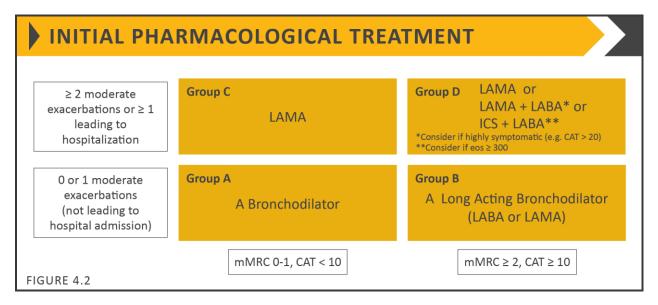
- Simvastatin does not prevent exacerbations in COPD patients at increased risk of exacerbations and without
 indications for statin therapy (Evidence A). However, observational studies suggest that statins may have positive
 effects on some outcomes in patients with COPD who receive them for cardiovascular and metabolic indications
 (Evidence C).
- Leukotriene modifiers have not been tested adequately in COPD patients.

TABLE 3.5

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KEY POINTS: Chapter 4: Management of Stable COPD

- The management strategy for stable COPD should be predominantly based on the individualized assessment of symptoms and future risk of exacerbations.
- All individuals who smoke should be strongly encouraged and supported to quit.
- The main treatment goals are reduction of symptoms and future risk of exacerbations.
- Management strategies include pharmacologic and non-pharmacologic interventions.



KEY POINTS FOR THE USE OF BRONCHODILATORS

- LABAs and LAMAs are preferred over short-acting agents except for patients with only occasional dyspnea (Evidence A), and for immediate relief of symptoms in patients already on long-acting bronchodilators for maintenance therapy.
- Patients may be started on single long-acting bronchodilator therapy or dual long-acting bronchodilator therapy. In patients with persistent dyspnea on one bronchodilator treatment should be escalated to two **(Evidence A)**.
- Inhaled bronchodilators are recommended over oral bronchodilators (Evidence A).
- Theophylline is not recommended unless other long-term treatment bronchodilators are unavailable or unaffordable (Evidence B).

TABLE 4.5

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KEY POINTS FOR THE USE OF ANTI-INFLAMMATORY AGENTS

- Long-term monotherapy with ICS is not recommended (Evidence A).
- Long-term treatment with ICS may be considered in association with LABAs for patients with a history of exacerbations despite appropriate treatment with long-acting bronchodilators (Evidence A).
- Long-term therapy with oral corticosteroids is not recommended (Evidence A).
- In patients with severe to very severe airflow limitation, chronic bronchitis and exacerbations the addition of a PDE4 inhibitor to a treatment with long acting bronchodilators with/without ICS can be considered (Evidence B).
- Preferentially, but not only in former smokers with exacerbations despite appropriate therapy, macrolides, in particular azithromycin, can be considered (Evidence B).
- Statin therapy is not recommended for prevention of exacerbations (Evidence A).
- J.H. • Antioxidant mucolytics are recommended only in selected patients (Evidence A). STRIB

TABLE 4.6

| Factors to consider when initiating IC (note the scenario is different when c | S treatment in combination with one or onsidering ICS withdrawal): | two long-acting bronchodilators |
|--|--|---|
| · STRONG SUPPORT · | · CONSIDER USE · | · AGAINST USE · |
| History of hospitalization(s) for exacerbations of COPD# ≥ 2 moderate exacerbations of COPD per year# Blood eosinophils ≥ 300 cells/µL History of, or concomitant, asthma | 1 moderate exacerbation of COPD per year# Blood eosinophils ≥ 100 to < 300 cells/µL | Repeated pneumonia events Blood eosinophils <100 cells/μL History of mycobacterial infection |
| *note that blood eosinophils should be s eosinophil counts are likely to fluctuate. | odilator maintenance therapy (see Table 3.4 eeen as a continuum; quoted values represe RS 2019: <i>European Respiratory Journal 52 (</i> blished 13 December 2018 | ent approximate cut-points; |

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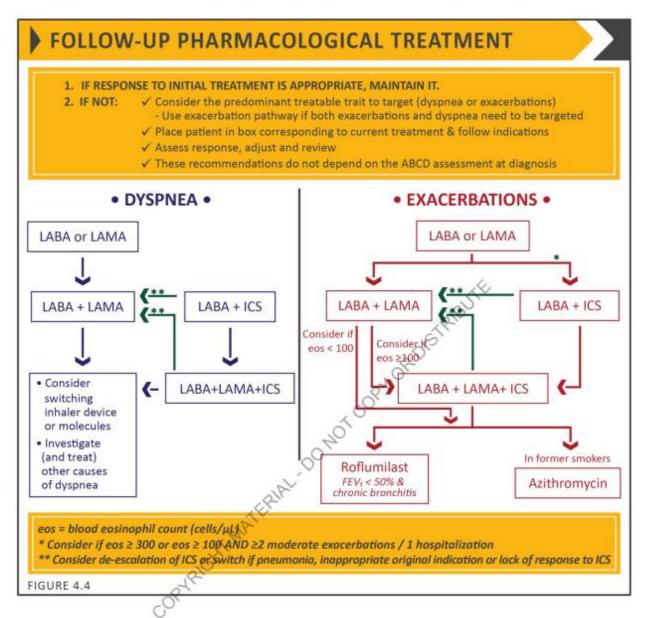
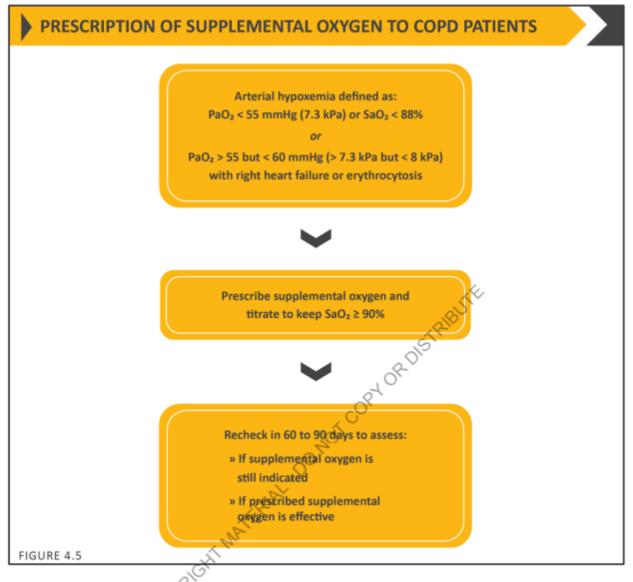


Figure 4.4 suggests escalation and de-escalation strategies based on available efficacy as well as safety data. The response to treatment escalation should always be reviewed, and de-escalation should be considered if there is a lack of clinical benefit and/or side effects occur. De-escalation may also be considered in COPD patients receiving treatment who return with resolution of some symptoms that subsequently may require less therapy. Patients, in whom treatment modification is considered, in particular de-escalation, should be undertaken under close medical supervision. We are fully aware that treatment escalation has not been systematically tested; trials of de-escalation are also limited and only include ICS.

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Ventilatory support

NIV is occasionally used in patients with stable very severe COPD. NIV may be considered of some use in a selected group of patients, particularly in those with pronounced daytime hypercapnia and recent hospitalization, although systematic review is unable to support or refute this.⁽⁹²⁾ However, in patients with both COPD and obstructive sleep apnea there are clear indications for continuous positive airway pressure (CPAP).⁽⁹³⁾

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KEY POINTS: Chapter 5: Management of Exacerbations

- An exacerbation of COPD is defined as an acute worsening of respiratory symptoms that results in additional therapy.
- As the symptoms are not specific to COPD relevant differential diagnoses should be considered.
- Exacerbations of COPD can be precipitated by several factors. The most common causes are respiratory tract infections.
- The goal for treatment of COPD exacerbations is to minimize the negative impact of the current exacerbation and to prevent subsequent events.
- Short-acting inhaled beta2-agonists with or without short-acting anticholinergics are recommended as the initial bronchodilators to treat an acute exacerbation.
- Maintenance therapy with long-acting bronchodilators should be initiated as soon as possible before hospital discharge.
- Systemic corticosteroids can improve lung function (FEV1), oxygenation and shorten recovery time and hospitalization duration. Duration of therapy should not be more than 5-7 days.
- Antibiotics, when indicated, can shorten recovery time, reduce the risk of early relapse, treatment failure, and hospitalization duration. Duration of therapy should be 5-7 days.
- Methylxanthines are not recommended due to increased side effect profiles.
- Non-invasive mechanical ventilation should be the first mode of ventilation used in COPD patients with acute respiratory failure who have no absolute contraindication because it improves gas exchange, reduces work of breathing and the need for intubation, decreases hospitalization duration and improves survival.
- Following an exacerbation, appropriate measures for exacerbation prevention should be initiated.

POTENTIAL INDICATIONS FOR HOSPITALIZATION ASSESSMENT*

- Severe symptoms such as sudden worsening of resting dyspnea, high respiratory rate, decreased oxygen saturation, confusion, drowsiness.
- Acute respiratory failure.
- Onset of new physical signs (e.g., cyanosis, peripheral edema).
- Failure of an exacerbation to respond to initial medical management.
- Presence of serious comorbidities (e.g., heart failure, newly occurring arrhythmias, etc.).
- Insufficient home support.

*Local resources need to be considered.

TABLE 5.2

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MANAGEMENT OF SEVERE BUT NOT LIFE-THREATENING EXACERBATIONS*

• Assess severity of symptoms, blood gases, chest radiograph.

- S • Administer supplemental oxygen therapy, obtain serial arterial blood gas, venous blood gas and pulse oximetry measurements.
- Bronchodilators:
 - » Increase doses and/or frequency of short-acting bronch dilators.
 - » Combine short-acting beta 2-agonists and anticholinergics.
 - » Consider use of long-active bronchodilators when patient becomes stable.
 - » Use spacers or air-driven nebulizers when appropriate.
- Consider oral corticosteroids.
- Consider antibiotics (oral) when signs of bacterial infection are present.
- Consider noninvasive mechanical ventilation (NIV).
- At all times:
 - » Monitor fluid balance.
 - » Consider subcutaneous heparin or low molecular weight heparin for thromboembolism prophylaxis.
 - » Identify and treat associated conditions (e.g., heart failure, arrhythmias, pulmonary embolism etc.).

*Local resources need to be considered.

TABLE 5.3

KEY POINTS FOR THE MANAGEMENT OF EXACERBATIONS

- Short-acting inhaled beta2-agonists, with or without short-acting anticholinergics, are recommended as the initial bronchodilators to treat an acute exacerbation (Evidence C).
- Systemic corticosteroids can improve lung function (FEV1), oxygenation and shorten recovery time and hospitalization duration. Duration of therapy should not be more than 5-7 days (Evidence A).
- Antibiotics, when indicated, can shorten recovery time, reduce the risk of early relapse, treatment failure, and hospitalization duration. Duration of therapy should be 5-7 days (Evidence B).
- Methylxanthines are not recommended due to increased side effect profiles (Evidence B).
- Non-invasive mechanical ventilation should be the first mode of ventilation used in COPD patients with acute respiratory failure who have no absolute contraindication because it improves gas exchange, reduces work of breathing and the need for intubation, decreases hospitalization duration and improves survival (Evidence A).

TABLE 5.4

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KEY POINTS: Chapter 6: COPD and Co-morbidities

• COPD often coexists with other diseases (co-morbidities) that may have a significant impact on disease course.

• In general, the presence of co-morbidities should not alter COPD treatment and co-morbidities should be treated per usual standards regardless of the presence of COPD.

- Lung cancer is frequently seen in patients with COPD and is a main cause of death.
 - Annual low-dose CT (LDCT) is recommended for lung cancer screening in patients with COPD due to smoking according to recommendations for the general population
 - Annual LDCT is not recommended for lung cancer screening in patients with COPD not due to smoking due to insufficient data to establish benefit over harm
- Cardiovascular diseases are common and important co-morbidities in COPD.

• Osteoporosis and depression/anxiety are frequent, important co-morbidities in COPD, are often under-diagnosed, and are associated with poor health status and prognosis.

• Gastroesophageal reflux (GERD) is associated with an increased risk of exacerbations and poorer health status.

• When COPD is part of a multimorbidity care plan, attention should be directed to ensure simplicity of treatment and to minimize polypharmacy.

See the Guideline for recommendations on treating the following comorbidities with COPD

| • | Heart failure | • | Osteoporosis |
|---|-----------------------------|---|---------------------------------|
| • | Ischemic heart disease | • | Anxiety and depression |
| • | Arrhythmias | • | Metabolic syndrome and diabetes |
| • | Peripheral vascular disease | • | Gastroesophageal reflux |
| • | Hypertension | • | Bronchiectasis |
| • | Lung cancer | • | Obstruct sleep apnea |
| • | Cognitive impairment | | |

KEY POINTS: Chapter 7: COVID-19 and COPD

- Patients with COPD presenting with new or worsening respiratory symptoms, fever, and/or any other symptoms that could be COVID-19 related, even if these are mild, should be tested for possible infection with SARS-CoV-2.
- Patients should keep taking their oral land inhaled respiratory medication for COPD as directed as there is no evidence that COPD medication should be changed during this COID-19 pandemic.

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| Effective 2/2012, 2/2014, 2/2016, 2/2018, | Date: 2/2022 | 2/2024 |
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- During periods of high prevalence of COVID-19 in the community, spirometry should be restricted to patients requiring urgent or essential tests for the diagnosis of COPD, and/or to assess lung function status for interventional procedures or surgery.
- Physical distancing and shielding, or sheltering-in-place, should not lead to social isolation and inactivity. Patients should stay in contact with their friends and families by telecommunications and continue to keep active. They should also ensure they have enough medication.
- Patients should be encouraged to use reputable resources for medical information regarding COVID-19 and its management.
- Guidance for remote (phone/virtual/online) COPD patient follow-up and printable checklist are provided [and can be found in the Guideline document].

| KEY POINTS FOR THE MANAGEMENT OF PATIENTS WITH COPD AND SUSPECTED OR PROVEN COVID-19 |
|---|
| SARS-CoV-2 TESTING |
| Swab/Saliva PCR if new or worsening respiratory symptoms, fever, and/or any other symptoms that could be COVID related |
| OTHER INVESTIGATIONS |
| Avoid spirometry unless essential Consider CT for COVID pneumonia and to exclude other diagnoses e.g. PE Avoid bronchoscopy unless essential Assess for co-infection |
| COPD PHARMACOTHERAPY |
| Ensure adequate supplies of medication Continue maintenance therapy unchanged including ICS Use antibiotics and oral steroids in line with recommendations for exacerbations Avoid nebulization when possible |
| COPD NON-PHARMACOLOGICAL THERAPY |
| Maintain physical activity as able |
| PROTECTIVE STRATEGIES |
| Follow basic infection control measures Maintain physical distancing Wear a face covering |
| COVID-19 THERAPY |
| Use systemic steroids and remdesivir as recommended for patients with COVID-19 Use HFNT or NIV for respiratory failure if possible Use invasive mechanical ventilation if HFNK or NIV fails Post COVID-19 rehabilitation Ensure appropriate post COVID-19 follow-up |
| |

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|---|-----------------------------------|-----------------------------|
| Effective 2/2012, 2/2014, 2/2016, 2/2018, | Date: 2/2022 | 2/2024 |
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Appendix: Medications for COPD

- a. May differ from product labeling.
- b. Wholesale cost for 30-day supply of highest strength, of generic if available unless otherwise specified. For short-acting agents, cost is for 200 inhalations.
- c. Only long-acting inhalers specifically approved for COPD are included.

| Inhalers for COPD | | | | |
|--|--|--------------------------------|---|--|
| Medication | Dosing Frequency _a | Costb | Adverse Reactions of Note | |
| Shor | t-acting Bronchodila | itors | | |
| | Beta-2 agonists | 1 | | |
| Albuterol (Salbutamol) (ProAir HFA, ProAir Digihaler, ProAir RespiClick, Proventil HFA, Ventolin HFA) | 2 inhalations every 4-6 hours as needed | \$78 | Tachycardia, excitement/nervousness, tremors, paradoxical bronchospasm | |
| Levalbuterol (<i>Xopenex HFA</i>) | 2 inhalations every 4-6 hours as needed | \$74 | Headache, rhinitis | |
| | Anticholinergic | | | |
| Ipratropium (<i>Atrovent HFA</i>) | 2 inhalations every 6 hours | Atrovent HFA \$513 | Bronchitis | |
| Combination Beta-2 agonist/Anticholinergic | | | | |
| Albuterol/ipratropium | 1 inhalation four times daily; | | | |
| (Combivent Respimat) | May take up to 2 additional inhalations daily as needed | Combivent Respimat \$532 | | |

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|---|-----------------------------------|-----------------------------|
| Effective 2/2012, 2/2014, 2/2016, 2/2018, | Date: 2/2022 | 2/2024 |
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| Long-Acting Beta-2 agonists (LABAs) | | | |
|---|---|---|--|
| Olodaterol (Striverdi Respimat) | 2 inhalations once daily | Striverdi Respimat \$280 | Nasopharyngitis |
| Salmeterol (Serevent Diskus) | One inhalation twice daily | Serevent Diskus \$493 | Headache |
| Long-Acting | Antimuscarinc agen | ts (LAMAs) | |
| Aclidinium (Tudorza Pressair) | 1 inhalation twice daily | Tudorza Pressair \$343 | |
| Glycopyrrolate (Seebri Neohaler) | 1 capsule inhaled twice daily | \$473 | |
| Tiotropium (Spiriva HandiHaler, Spiriva Respimat) | HandiHaler: 1 capsule inhaled once daily; each capsule should be inhaled twice <i>Respimat</i> : 2 inhalations once | Spiriva HandiHaler \$540 Respimat \$572 | Xerostomia, pharyngitis, upper respiratory tract infections |
| Umeclidinium (Incruse Ellipta) | daily 1 inhalation once daily | \$386 | |

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|---|-----------------------------------|-----------------------------|
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| Combination LABA/LAMA | | | | |
|---|------------------------------|------------------------------------|--|--|
| Olodaterol/Tiotropium (Stiolto Respimat) | 2 inhalations once daily | Stiolto Respimat \$90 | Nasopharyngitis | |
| Vilanterol/Umeclidinium (Anoro Ellipta) | 1 inhalation once daily | Anoro Ellipta \$269 | | |
| Formoterol/Aclidinium (Duaklir Pressair) | 1 inhalation twice daily | Duaklir Pressair \$1194 | | |
| Formoterol/Glycopyrrolate (Bevespi Aerosphere) | 2 inhalations twice daily | Bevespi Aerosphere \$261-474 | | |
| Combin | ation LABA/ Cortico | osteroid | | |
| Formoterol/Budesonide (Symbicort) | 2 inhalations twice daily | \$352-403 | Headache, nasopharyngitis, upper respiratory tract infections | |
| Salmeterol/Fluticasone propionate (Advair Diskus, Advair HFA, AirDuo Digihaler, AirDuo RespiClick, Wixela Inhub) | 1 inhalation twice daily | \$120 | Headache, upper respiratory tract infections | |
| Vilanterol/Fluticasone furoate (<i>Breo Ellipta</i>) | 1 inhalation once daily | \$369 | | |

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|---|-----------------------------------|-----------------------------|
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| Combination LABA/Corticosteroid/Anticholinergic | | | | |
|--|---|---|--|--|
| Fluticasone/Umeclidium/ Vilanterol (<i>Trelegy Ellipta</i>) | 1 inhalation once daily | \$361 | Pharyngitis | |
| | Other Agents | | | |
| | Methylxanthines | | | |
| Theophylline | Variable depending on formulation Toxicity is dose related; drug levels should be monitored | \$362 dollars (generic twice daily product) | Tachycardia, headache, tremor, nausea | |
| Sy | stemic Corticosteroi | ds | | |
| Prednisone | Once daily by mouth (usually, 40mg once daily for 5 days) | \$0.50 per day – therapy duration varies) | Increased appetite, fluid retention, electrolyte changes, weight gain, hypertension | |
| Methylprednisolone | 60-125 mg 1-4 times daily IV followed by oral therapy (prednisone preferred for oral route) | \$4 -\$8 per day – therapy duration varies | Hypertension, fluid retention, electrolyte changes | |
| Phosphodiesterase-4 Inhibitor | | | | |
| Roflumilast (<i>Daliresp</i>) | 250 mcg once daily by mouth for 4 weeks followed by 500mcg once daily | \$488 | Weight loss, diarrhea | |

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|---|-----------------------------------|-----------------------------|
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Patient Education:

https://www.cdc.gov/copd/infographics/copd-awareness.html

https://www.cdc.gov/copd/basics-about.html

https://www.uptodate.com/contents/chronic-obstructive-pulmonary-disease-copd-thebasics?search=COPD&source=search_result&selectedTitle=1~150&usage_type=default &display_rank=1

https://www.uptodate.com/contents/medicines-for-chronic-obstructive-pulmonarydisease-copd-the-basics?search=COPD&topicRef=4649&source=see_link

Patient education videos (in English and in Spanish) on proper use of metered dose inhalers with and without spacers can be found at: <u>https://www.cdc.gov/asthma/inhaler_video/default.htm</u>

References:

- 1) <u>https://www.cdc.gov/copd/data.html</u>
- 2) <u>https://www.cdc.gov/dotw/copd/index.html#:~:text=COPD%20affects%20more%20than %2015,smoke%20and%20other%20air%20pollutants.</u>
- 3) From the *Global Strategy for the Diagnosis, Management and Prevention of COPD*, Global Initiative for Chronic Obstructive Lung Disease (GOLD) 2022. Available from: <u>http://www.goldcopd.org/</u>.
- 4) Drugs for COPD. Med Lett Drugs Ther. 2017 Apr 10;59(1518):57-62
- 5) Drug Information Articles on <u>https://www.uptodate.com/contents/table-of-contents/drug-information/general-drug-information</u> (accessed February 16, 2022)

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