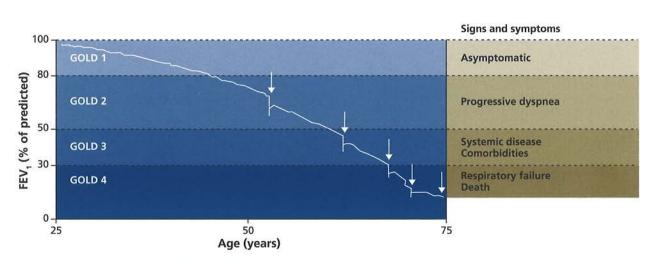
Exacerbations May Negatively Impact COPD Patients

Exacerbations are a significant clinical component of COPD, and as the disease progresses, exacerbations may become more frequent¹⁻³

Exacerbations are important events in the disease course of COPD for many reasons, including4:

- Negative effect on a patient's quality of life
- Physical, social, and emotional impairments
- Effects on symptoms and lung function
- Accelerated rate of decline in lung function
- Association with significant mortality, particularly in exacerbations that require hospitalizations

The Role of Exacerbations in Accelerating Lung Function Decline5,



Exacerbations, indicated by white arrows, punctuate and accelerate lung function decline.

* Figure represents the physiology of exacerbations by severity of airflow limitation in a hypothetical habitual smoker with COPD. The impact of COPD exacerbations on FEV, decline remains uncertain.

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Even when an exacerbation resolves, the effects may linger,4 and irreversible impairment may occur^{6,7}

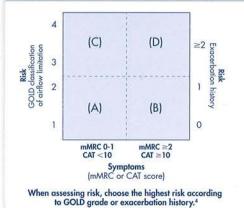
- Higher exacerbation rates are associated with greater decline of FEV, and worsening of health status^{4,8}
- Patients with frequent exacerbations will continue to have exacerbations often³

Early detection and reducing the risk of exacerbations are vital to reducing the burden of COPD⁴

COPD=chronic obstructive pulmonary disease; FEV,=forced expiratory volume in the first second; GOLD=Global Initiative for Chronic Obstructive Lung Disease.

GOLD Guidelines Recommend Maintenance Therapy to Reduce COPD Exacerbation Risk

COPD Assessment Utilizes Reported Symptoms, Airflow Limitation, and Exacerbation History⁴



Patient Group	Characteristics	Spirometric Classification	Exacerbations per Year	mMRC	CAT
Α	Low risk, less symptoms	GOLD 1-2	≤l	0-1	<10
В	Low risk, more symptoms	GOLD 1-2	≤1	≥2	≥10
С	High risk, less symptoms	GOLD 3-4	≥2	0-1	<10
D	High risk, more symptoms	GOLD 3-4	≥2	≥2	≥10

Exacerbation risk can be reduced with appropriate pharmacotherapy^{3,4}

Initial Pharmacologic Management of COPD4,

Patient Group	First Choice	Second Choice	Alternative Choice†	
A Low risk, less symptoms	SA anticholinergic prn or SA B ₂ -agonist prn	LA anticholinergic or LA β_2 -agonist <i>or</i> SA β_2 -agonist + SA anticholinergic	Theophylline	
B Low risk, more symptoms	LA anticholinergic <i>or</i> LA B ₂ -agonist	LA anticholinergic and LA \upbeta_2 -agonist SA \upbeta_2 -agonist and/or SA anticholinergic Theophylline		
C High risk, less symptoms	ICS + LA β_2 -agonist or LA anticholinergic	LA anticholinergic and LA $\ensuremath{\beta_2}\xspace$ -agonist	PDE-4 inhibitor SA B ₂ -agonist <i>and/or</i> SA anticholinergic Theophylline	
D High risk, more symptoms	ISC+ LA B ₂ -agonist <i>or</i> LA anticholinergic	ICS and LA anticholinergic or ISC + LA B_2 -agonist and LA anticholinergic or ISC + LA B_2 -agonist and PDE-4 inhibitor or LA anticholinergic and LA B_2 -agonist or LA anticholinergic and PDE-4 inhibitor	Carbocysteine SA B ₂ -agonist <i>and/or</i> SA anticholinergic Theophylline	

^{*}Medications in each box are mentioned in alphabetical order and therefore not necessarily in order of preference; ¹Medications in this column can be used alone or in combination with other options in the first and second columns.

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CAT=COPD Assessment Test"; ICS=inhaled corticosteroid; LA=long-acting; mMRC=modified Medical Research Council; PDE-4=phosphodiesterase-4; prn=when necessary; SA=short-acting.

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