Global Initiative for Chronic Obstructive Lung Disease 2025 REPORT

COPD: (GOLD) update 2025



Global Strategy for the Diagnosis, Management, and Prevention of Chronic Obstructive Pulmonary Disease



COPD: (GOLD) update 2025

Definitie

Wat is COPD?

Diagnose

Hoe stel ik de diagnose op een correcte manier?

Classificatie

Wat is al dat gedoe met 1, 2, 3, 4, A, B, C, D, E, ...?

Behandeling

Hoe starten en aanpassen van de behandeling?

Exacerbaties

Hoe behandel ik een exacerbatie?

Comorbiditeiten

Welke comorbiditeiten zijn relevant?





Definitie van COPD anno 2025



COPD is a heterogeneous lung condition characterized by chronic respiratory symptoms due to abnormalities of the <u>airways</u> (bronchitis, bronchiolitis) and/or <u>alveoli</u> (emphysema) that cause persistent, often progressive, airflow obstruction

Diagnose COPD



Clinical Indicators for Considering a Diagnosis of COPD

Figure 2.1

Consider the diagnosis of COPD, and perform spirometry, if any of these clinical indicators are present:

(these indicators are not diagnostic themselves, but the presence of multiple key indicators increases the probability of the presence of COPD; in any case, spirometry is required to establish a diagnosis of COPD)

Dyspnea that is	Progressive over time Worse with exercise Persistent
Recurrent wheeze	
Chronic cough	May be intermittent and may be non-productive
Recurrent lower respiratory tract infections	
History of risk factors	Tobacco smoke (including popular local preparations)
	Smoke from home cooking and heating fuels
	Occupational dusts, vapors, fumes, gases and other chemicals
	Host factors (e.g., genetic factors, developmental abnormalities, low birthweight, prematurity, childhood respiratory infections etc.)

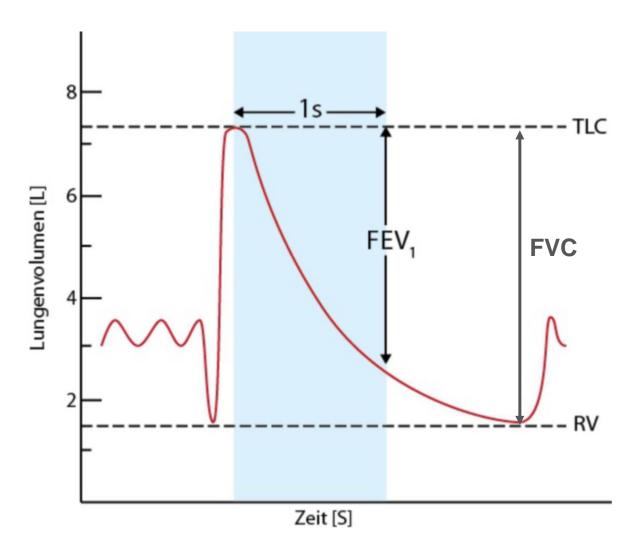
Differential Diagnosis of COPD

Figure 2.3

Diagnosis	Suggestive Features
COPD	Symptoms slowly progressive
	History of tobacco smoking or other risk factors
Asthma	Variable airflow obstruction
	Symptoms vary widely from day to day
	Symptoms worse at night/early morning
	Allergy, rhinitis, and/or eczema also present
	Often occurs in children
	Family history of asthma
Congestive heart failure	Chest X-ray shows dilated heart, pulmonary edema
	Pulmonary function tests indicate volume restriction, not airflow obstruction
Bronchiectasis	Large volumes of purulent sputum
	Commonly associated with bacterial infection
	Chest X-ray/HRCT shows bronchial dilation
Tuberculosis	Onset at all ages
	Chest X-ray shows lung infiltrate
	Microbiological confirmation
	High local prevalence of tuberculosis
Obliterative	Can occur in children
bronchiolitis	Seen after lung or bone marrow transplantation
	HRCT on expiration shows hypodense areas
Diffuse panbronchiolitis	Predominantly seen in patients of Asian descent
	Most patients are male and nonsmokers
	Almost all have chronic sinusitis
	Chest X-ray & HRCT show diffuse small centrilobular nodular opacities & hyperinflation

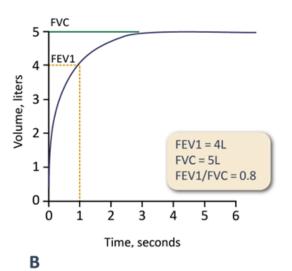
These features tend to be characteristic of the respective diseases, but are not mandatory. For example, a person who has never smoked may develop COPD (especially in LMICs where other risk factors may be more important than cigarette smoking).

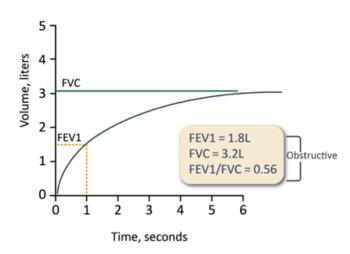
FEV1/FVC-ratio







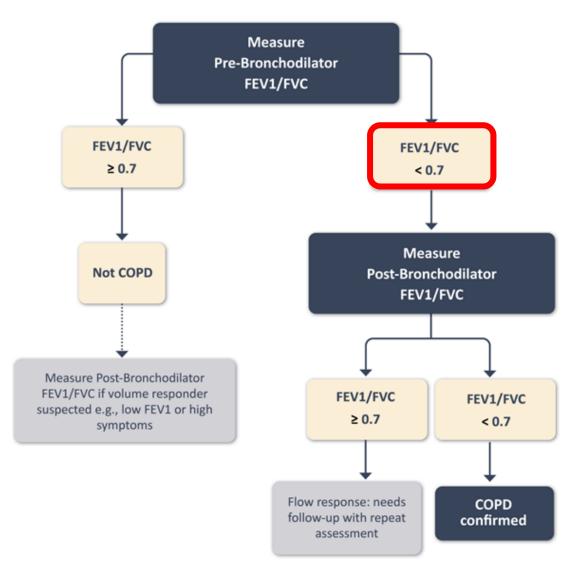








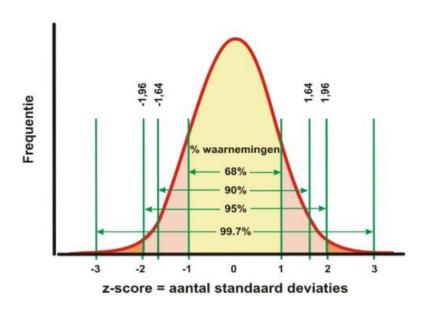
Spirometrische diagnose COPD



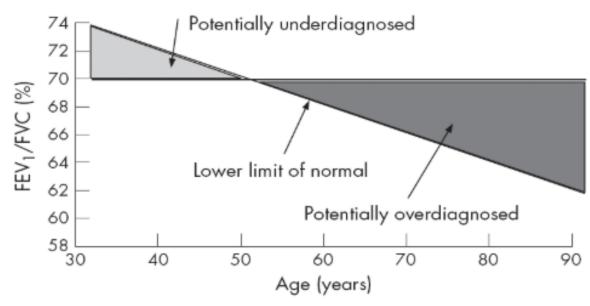




Spirometrische diagnose COPD

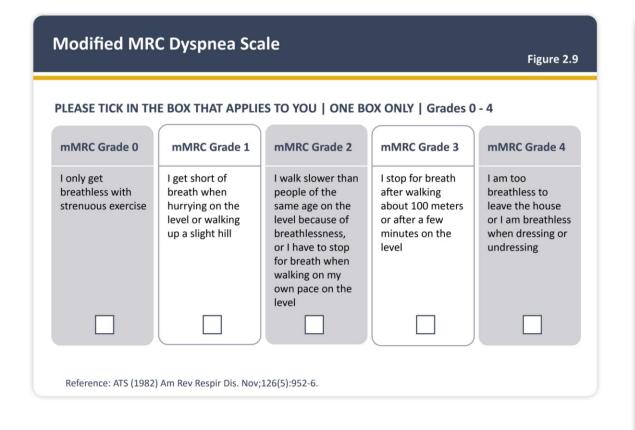


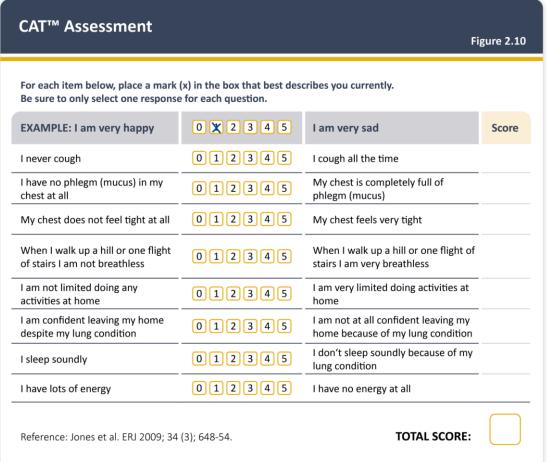
Datum onderzoek	19.12.24	Pred.	Gemeten	%Pred	Z-score
SPIROMETRIE/FLOW	-VOLUME				
FVC	L	3.53	2.89	82	-1.15
FEV 1	L	2.70	1.02	38	-3.43
FEV 1 % VC MAX	%	77	35	4	-4.23
PEF	L/s	7.25	3.13	43	-3.40
MFEF 75/25	L/s	2.12	0.26	12	-3.10
FEF 75	L/s	0.55	0.10	19	-2.67





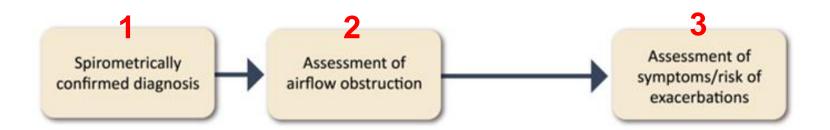
Kwantificatie van de ernst van de symptomen





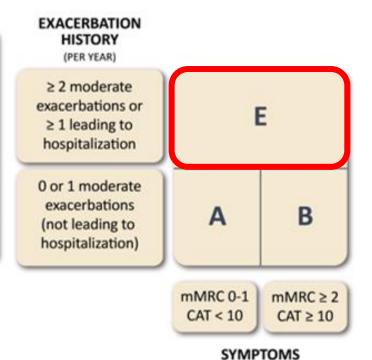


Classificatie van COPD



Post-bronchodilator FEV1/FVC < 0.7

GRADE	FEV1 (% predicted)
GOLD 1	≥ 80
GOLD 2	50-79
GOLD 3	30-49
GOLD 4	< 30



≥2 mod-severe exacerbations or ≥1 leading to hospital admission

O or 1 exacerbations (not leading to hospital admission)

C D

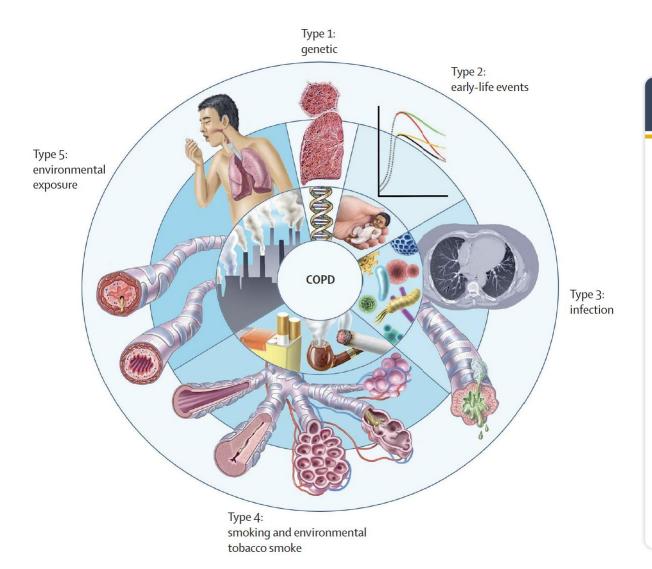
B

B

mMRC mMRC ≥2, 0-1, CAT CAT ≥10 <10

Etiotypes van COPD





PD	for CO	mes)	Etioty	Taxonomy	Proposed
г		(DES)	ELIOLY	Taxononing	Proposeu

Figure 1.2

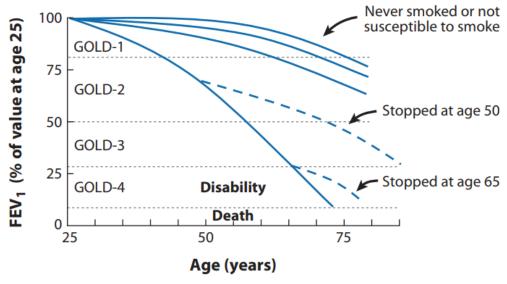
Classification	Description
Genetically determined COPD (COPD-G)	Alpha-1 antitrypsin deficiency (AATD) Other genetic variants with smaller effects acting in combination
COPD due to abnormal lung development (COPD-D)	Early life events, including premature birth and low birthweight, among others
Environmental COPD	
Cigarette smoking COPD (COPD-C)	 Exposure to tobacco smoke, including in utero or via passive smoking Vaping or e-cigarette use Cannabis
Biomass and pollution exposure COPD (COPD-P)	Exposure to household pollution, ambient air pollution, wildfire smoke, occupational hazards
COPD due to infections (COPD-I)	Childhood infections, tuberculosis-associated COPD, HIV-associated COPD
COPD & asthma (COPD-A)	Particularly childhood asthma
COPD of unknown cause (COPD-U)	

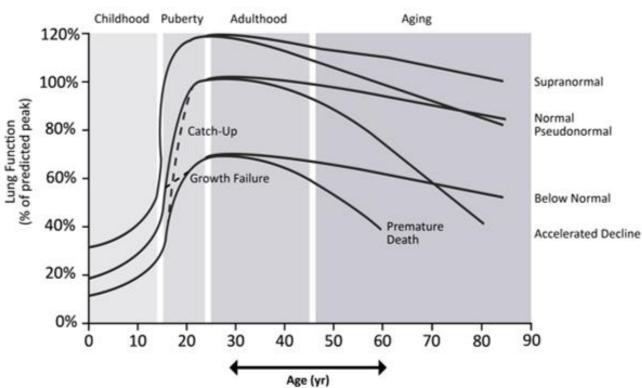
*Adapted from Celli et al. (2022) and Stolz et al. (2022)



Ziektetraject van COPD

Decline in FEV₁ with aging







Nieuwe terminologie

A: Early COPD

B: Mild COPD

C: Young COPD

D: Pre-COPD

E: PRISM

"near the beginning of a process"

"biological" first steps of the disease in an experimental setting

we propose that "mild" should not be used to identify "early" COPD and used only to describe the severity of airflow obstruction measured spirometrically

we propose to operationally consider "young COPD" in patients aged 20–50 years.

individuals (importantly, of any age) who have respiratory symptoms and/or other detectable structural and/or functional abnormalities, in the absence of airflow obstruction on forced spirometry.

preserved ratio impaired spirometry FEV1/FVC >0.7 + FEV1 <80%vv



Doelstellingen behandeling

- Relieve Symptoms
- Improve Exercise Tolerance
- Improve Health Status

REDUCE SYMPTOMS

AND

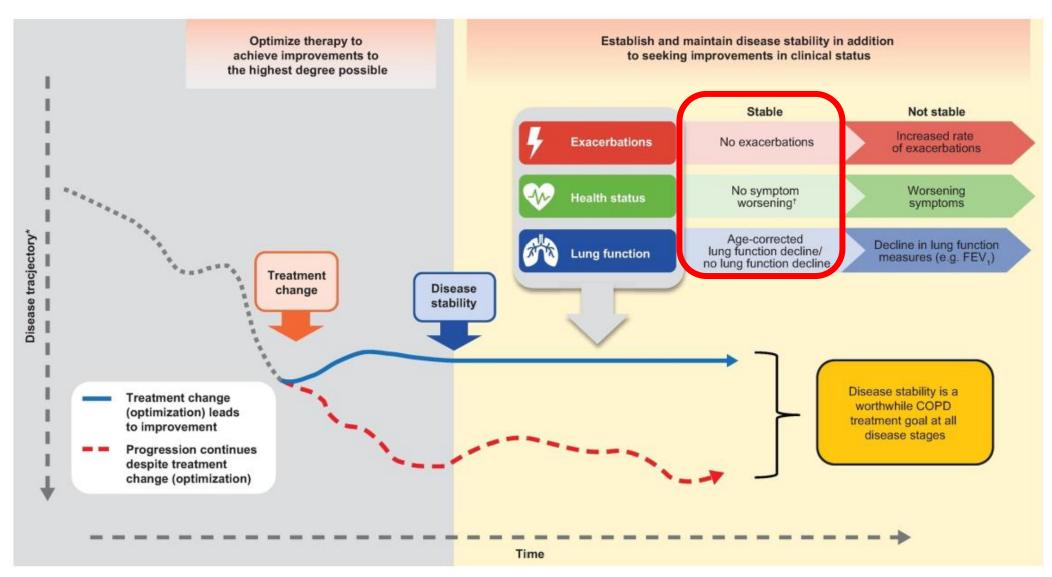
- Prevent Disease Progression
- Prevent and Treat Exacerbations
- Reduce Mortality



REDUCE RISK



Nieuw concept: ziektestabiliteit





Classificatie van COPD

Difficult COPD

(ICS/LABA/LAMA or LABA/LAMA during >12 months)

Systematic assessment

Difficult-to-treat COPD

Severe COPD

Poor control related to:

- Poor adeherence
- Suboptimal inhalation technique
- Exposure to cigarette smoke, ...
- Co-morbidities

Poor control despite:

- Good adherence
- Optimal inhalation technique
- Management of exposures
- Management of co-morbidities

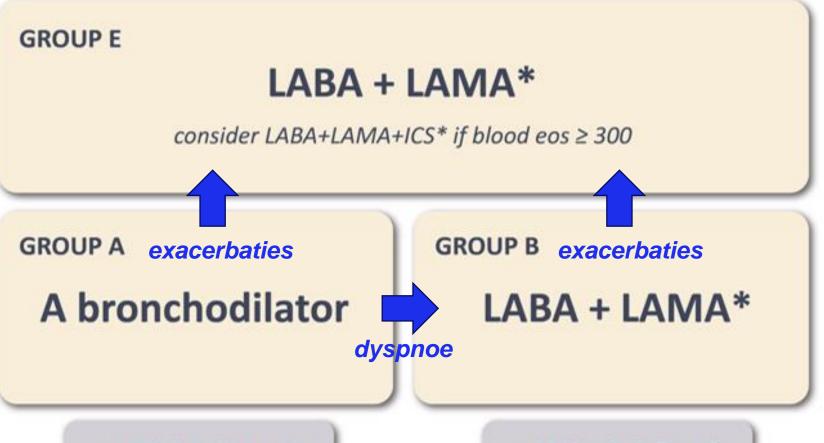




Chronische behandeling van COPD

≥ 2 moderate exacerbations or ≥ 1 leading to hospitalization

0 or 1 moderate exacerbations (not leading to hospital admission)

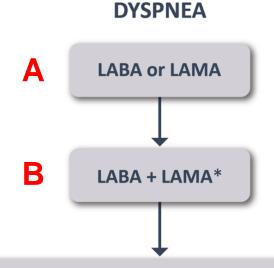


mMRC 0-1, CAT < 10

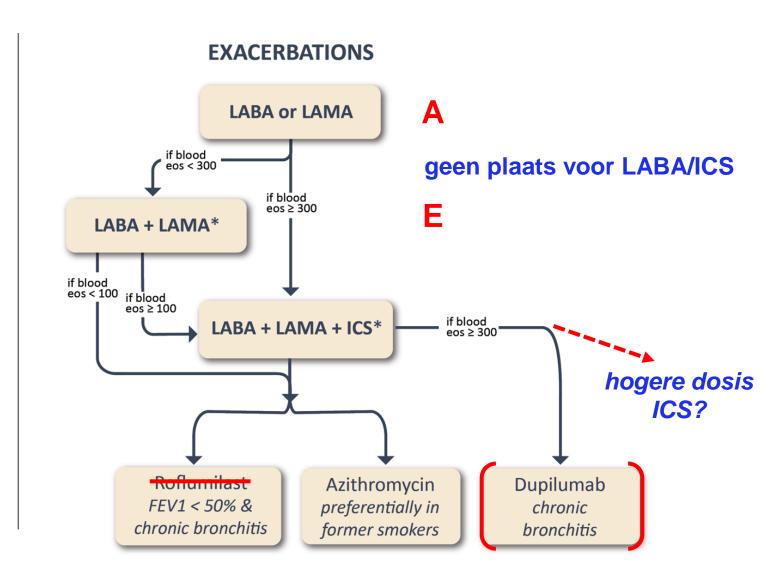
 $mMRC \ge 2$, $CAT \ge 10$



Step-up behandeling



- Consider switching inhaler device or molecules
- Implement or escalate non-pharmacological treatment(s)
- Consider adding ensifentrine
- Investigate (and treat) other causes of dyspnea







Factors to Consider when Initiating ICS Treatment

Figure 3.21

Factors to consider when adding ICS to long-acting bronchodilators:

(note the scenario is different when considering ICS withdrawal)

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

FAVORS USE

1 moderate exacerbation of COPD per year*

Blood eosinophils 100 to < 300 cells/μL

Repeated pneumonia events

Blood eosinophils < 100 cells/μL

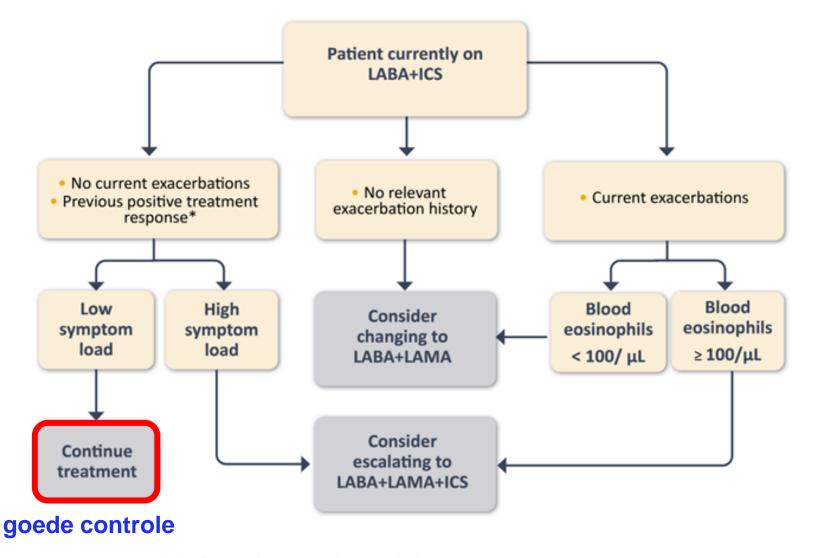
History of mycobacterial infection

"despite appropriate long-acting bronchodilator maintenance therapy (see Figures 3.7 & 3.18 for recommendations); *note that blood eosinophils should be seen as a continuum; quoted values represent approximate cut-points; eosinophil counts are likely to fluctuate.

Adapted from & reproduced with permission of the © ERS 2019: European Respiratory Journal 52 (6) 1801219; DOI: 10.1183/13993003.01219-2018 Published 13 December 2018



Wat met patiënten op LABA/ICS?

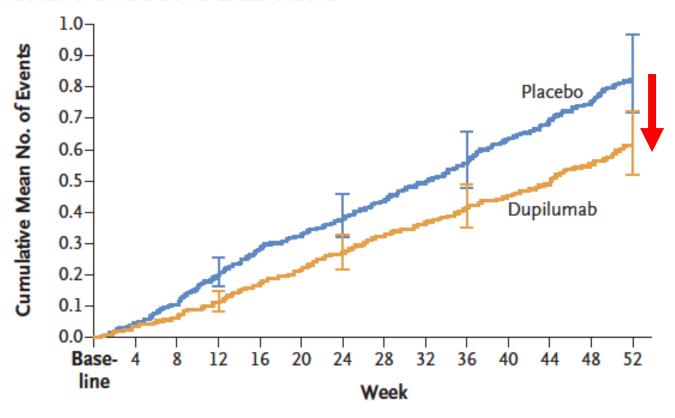


^{*}Patient previously had exacerbations and responded to LABA+ICS treatment



Dupilumab (anti IL-4 en IL-13)

A Moderate or Severe COPD Exacerbations



34% reductie in exacerbaties

No. at Risk
Placebo 465 464 458 453 453 448 430 415 403 394 384 368 351 303
Dupilumab 469 464 464 464 460 455 438 424 408 395 385 370 354 344

EOS ≥300/µI 2 matige of 1 ernstige AE







Andere farmacologische behandelingen

Other Pharmacological Treatments

Figure 3.23

Alpha-1 Antitrypsin Augmentation Therapy

 Intravenous augmentation therapy may slow down the progression of emphysoma (Evidence B)

Antitussives

• There is no conclusive evidence of a beneficial role of antitussives in people with COPD (Evidence C)

Vasodilators

 Vasodilators do not impreve outcomes and may worsen oxygenation (Evidence B)

Opioids

 Low-dose long acting oral and parenteral opioids may be considered for treating dyspnea in COPD patients with severe disease (Evidence B)

Pulmonary Hypertension Therapy

 Drugs approved for primary pulmonary hypertension are not recommended for patients we transfer hypertension secondary to COPD (Evidence B)



Niet-farmacologische behandeling

Patient Group	Essential	Recommended	Depending on Local Guidelines
A	Smoking cessation (can include pharmacological treatment)	Physical activity	Influenza vaccination COVID-19 vaccinations Pneumococcal vaccination Pertussis vaccination Shingles vaccination RSV vaccination
B and E	Smoking cessation (can include pharmacological treatment) Pulmonary rehabilitation	Physical activity	Influenza vaccination COVID-19 vaccinations Pneumococcal vaccination Pertussis vaccination Shingles vaccination RSV vaccination



Niet-farmacologische behandeling

Vaccination for Stable COPD

Figure 3.6

People with COPD should receive all recommended vaccinations in line with the relevant local guidelines:

- Yearly influenza vaccination (Evidence B)
- SARS-CoV-2 (COVID-19) vaccination based on WHO and CDC updated recommendations (Evidence B)
- Either one dose of 21-valent pneumococcal conjugate vaccine (PCV21) or one dose PCV20, as recommended by the CDC (Evidence B). Pneumococcal vaccination has been shown to reduce the incidence of community-acquired pneumonia and exacerbations for people with COPD (Evidence B)
- Respiratory syncytial virus (RSV) vaccination for individuals aged ≥ 60 years and/or with chronic heart or lung disease, as recommended by the CDC (Evidence A)
- Tdap (dTaP/dTPa) vaccination to protect against pertussis (whooping cough) for people with COPD that
 were not vaccinated in adolescence, as recommended by the CDC (Evidence B)
- Zoster vaccine to protect against shingles for people with COPD aged > 50 years, as recommended by the CDC (Evidence B)





Oxygen Therapy and Ventilatory Support in Stable COPD

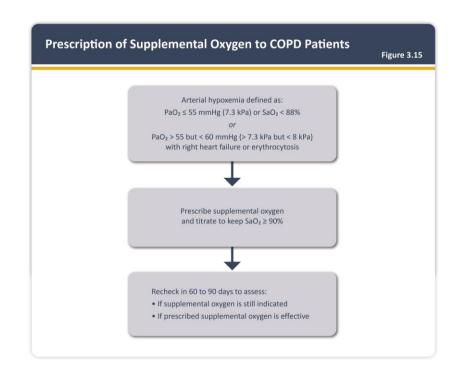
Figure 3.14

Oxygen Therapy

- The long-term administration of oxygen increases survival in patients with severe chronic resting arterial hypoxemia (Evidence A)
- In patients with stable COPD and moderate resting or exercise-induced arterial desaturation, prescription of long-term oxygen does not lengthen time to death or first hospitalization or provide sustained benefit in health status, lung function and 6-minute walk distance (Evidence A)
- Resting oxygenation at sea level does not exclude the development of severe hypoxemia when traveling by air (Evidence C)

Ventilatory Support

- NPPV may improve hospitalization-free survival in selected patients after recent hospitalization, particularly in those with pronounced daytime persistent hypercapnia (PaCO₃ > 53 mmHg) (Evidence B)
- In patients with severe chronic hypercapnia and a history of hospitalization for acute respiratory failure, long-term noninvasive ventilation may be considered (Evidence B)







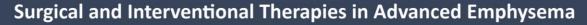
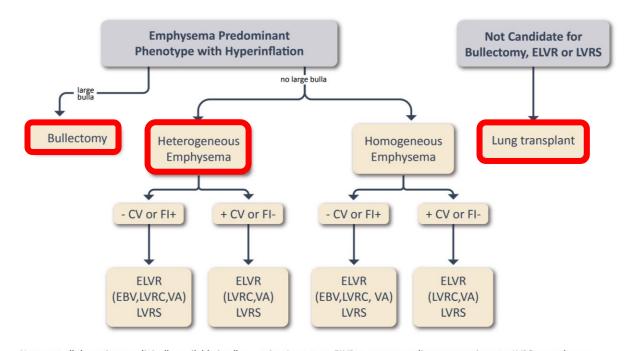


Figure 3.26

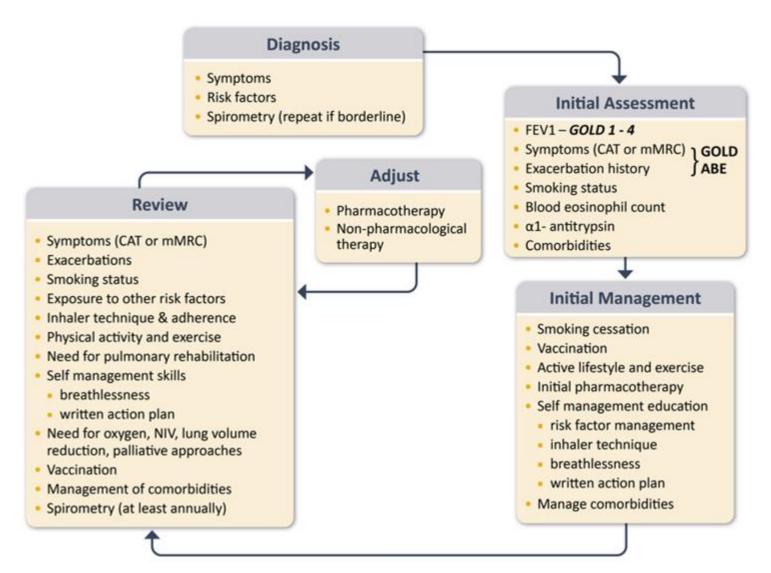


Note: not all therapies are clinically available in all countries. Long term ELVR outcomes or direct comparisons to LVRS are unknown.

Definition of abbreviations: CV, collateral ventilation measure by Chartis; FI + fissure integrity > 90% by HRCT; FI-, fissure integrity < 90% by HRCT; ELVR, Endoscopic Lung Volume Reduction, EBV, Endobronchial Valve; VA, Vapor Ablation; LVRC, Lung Volume Reduction Coil; LVRS, Lung Volume Reduction Surgery. Modified from Vogelmeier, AJRCCM, 2017.







Behandeling van exacerbaties



Figure 4.5

Assess severity of symptoms, blood gases, chest radiograph

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 bronchodilators
- Combine short-acting beta 2-agonists and anticholinergies
- · Consider use of long-acting 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:

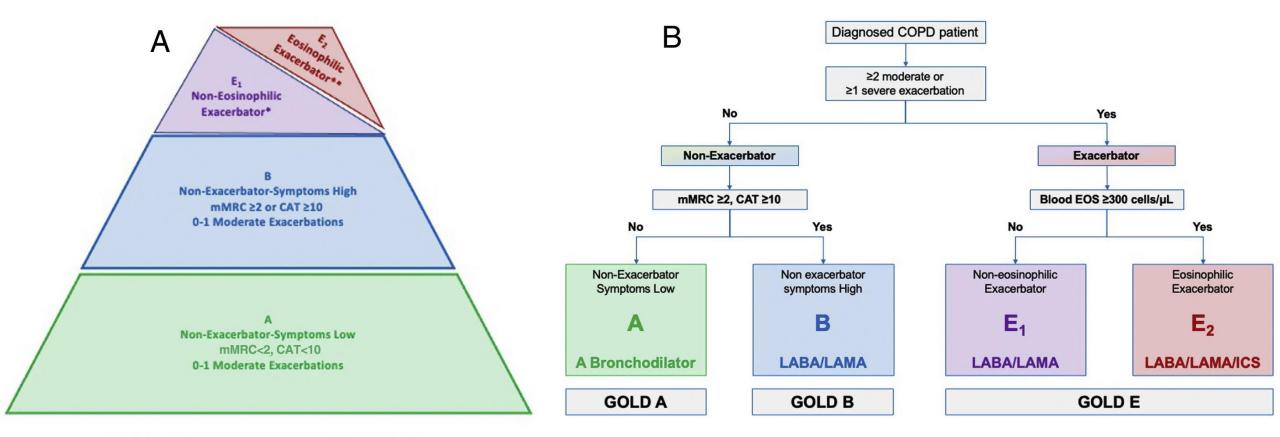
- Monitor fluid balance
- · Consider subcutaneous heparin or low molecular weight heparin for thromboembolism prophylaxis
- · Identify and treat associated conditions (e.g., heart failure, arrhythmias, pulmonary embolism etc.)



^{*}Local resources need to be considered



Eosinofiele en niet-eosinofiele exacerbaties



ABE_{1,2} Initial Patient Classification

Preventie van exacerbaties

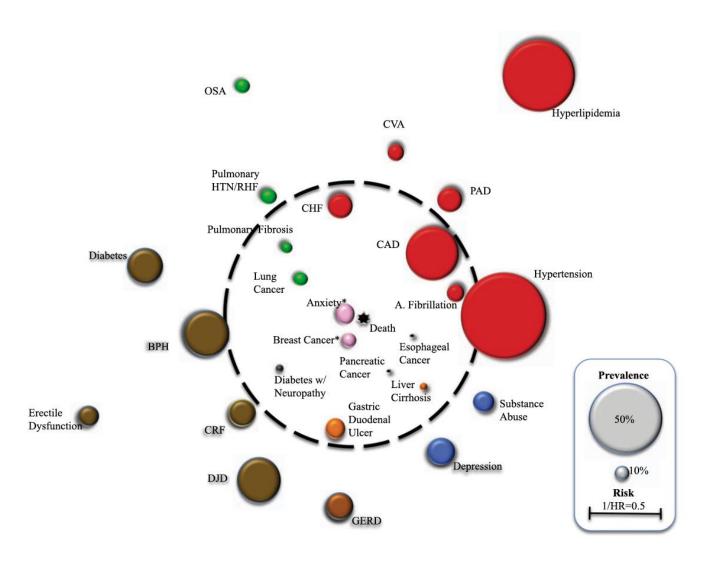
Interventions that Reduce the Frequency of COPD Exacerbations Figure 4.11

Intervention Class	Intervention
Bronchodilators	LABAs LAMAs LABA + LAMA
Corticosteroid-containing regimens	LABA + ICS LABA + LAMA + ICS
Anti-inflammatory (non-steroid)	Roflumilast Dupilumab
Anti-infectives	Vaccines Long Term Macrolides
Mucoregulators	N-acetylcysteine Carbocysteine Erdosteine
Various others	Smoking Cessation Rehabilitation Lung Volume Reduction Vitamin D Shielding measures (e.g., mask wearing, minimizing social contact, frequent hand washing)





Mortaliteit als gevolg van 12 comorbiditeiten





Milieuaspecten van inhalatiemedicatie

CO₂ uitstoot dosisaerosol vs poederinhalator

1 dosisaerosol ICS, LABA of LAMA =
 19kg CO₂e =
 100km met een benzine auto

100 km met benzine auto (19 kg CO₂e*)



1 dosisaerosol ICS, LABA of LAMA (19 kg CO₂e)



1 poederinhalator =1kg CO₂e

1 poederinhalator of softmistinhaler (<1 kg CO2e)

