California URGENT CARE ASSOCIATION

2024 WESTERN REGIONAL URGENT CARE CONFERENCE

COPD in Urgent Care GOLD Guidelines CUCA - 2024

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Disclosures

INDUSTRY AFFILIATIONS Grifols Pharmaceutical - speaker, consultant AstraZeneca – advisory board, speaker Regeneron – advisory board Pfizer – speaker (Paxlovid)

CLINICAL RESEARCH

- 2017 Sub-I, Genetech Zenyatta Severe Asthma Study
- 2016 Sub-I, Biota Human Rhinovirus Study
- 2015 Sub-I, Sanofi Traverse Severe Asthma Study
- 2015 Sub-I, Sanofi Liberty Severe Asthma Study
- 2013 Study Coordinator: MediVector Influenza Study

Brian Bizik does not intend to discuss the use of any off-label use/unapproved use of drugs or devices with the exception of NON-APPROVED inhaler recommendations that are Guideline based but not yet FDA approved (asthma only)



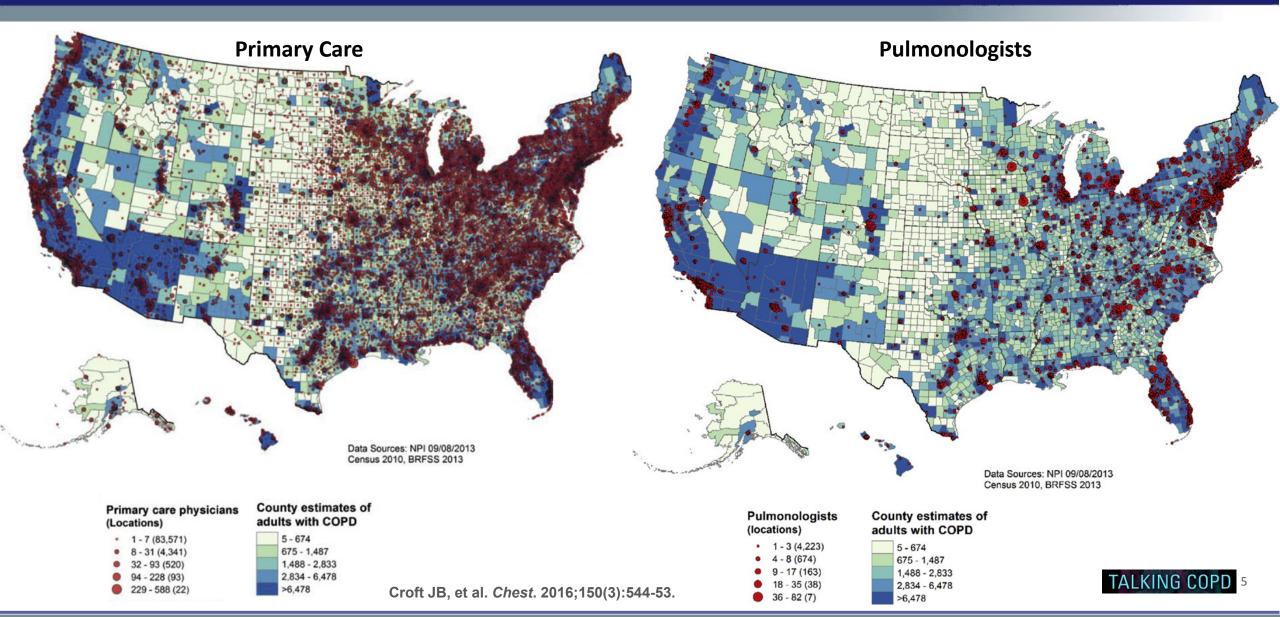
- Review medication classes, there are only three.
- Talk over the guidelines, what changes have there been and why does this matter for the UC?
- We need to GET patients better then take steps to KEEP them better. Next level care!



NEW PRODUCT SPOTLIGHT

Plan For Today

Primary vs Specialist Care – UCs is where many seek care

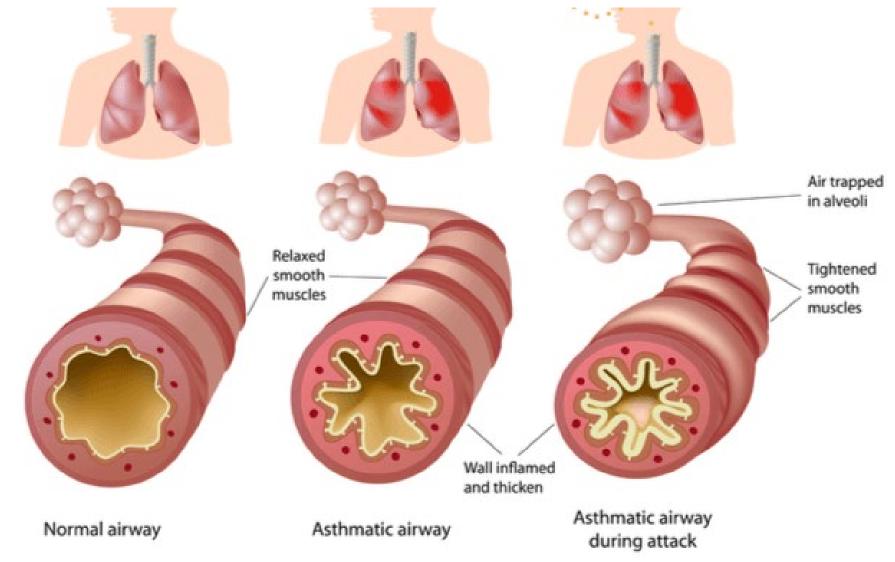


• Asthma – bronchoconstriction, airway inflammation, mucous production

• COPD – tissue destruction, chronic cough, due to exposure (tobacco) most of the time.

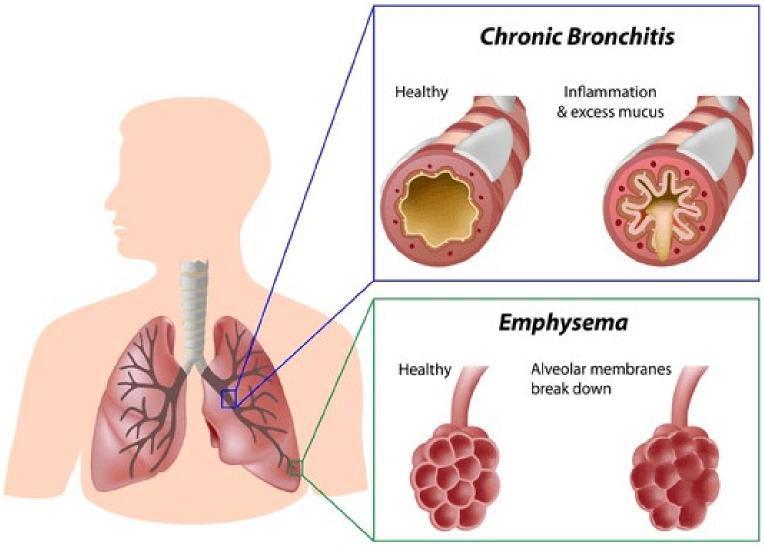


Asthma – bronchoconstriction, airway inflammation, mucous production



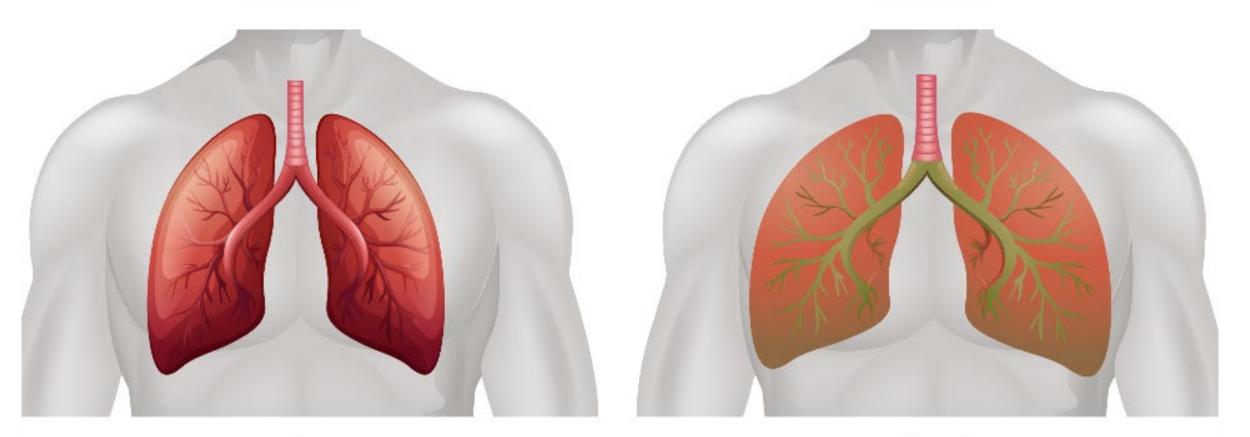
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COPD – Chronic (long term, you get this over time), Obstructive (elasticity is gone, things get floppy and weak, alveoli break down)





COPD – Big, floppy lungs. Flattened diaphragm. Harder to inhaler but MUCH hard to exhale, air is trapped, stale.

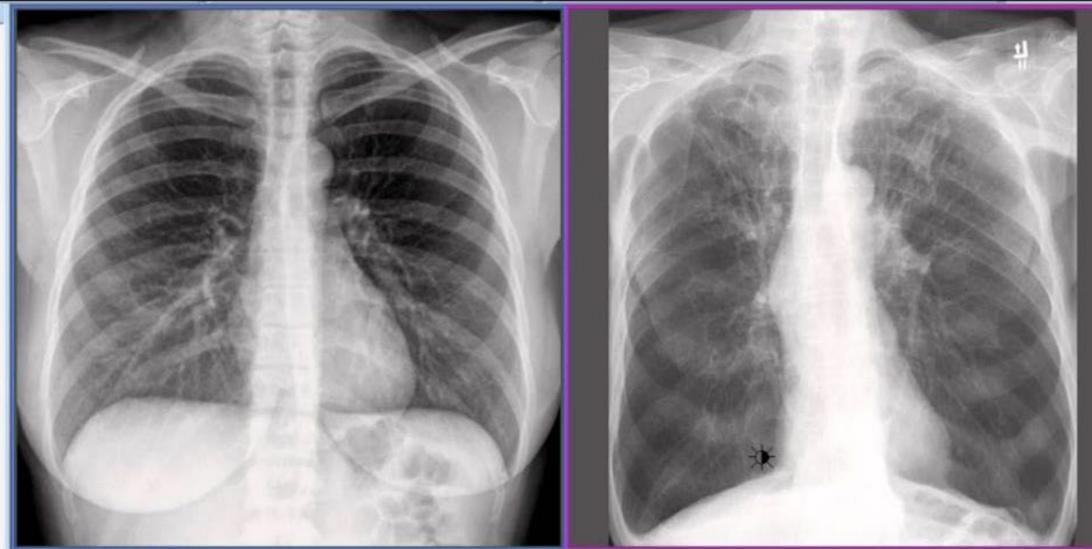


Normal Lungs

Hyperinflated Lungs

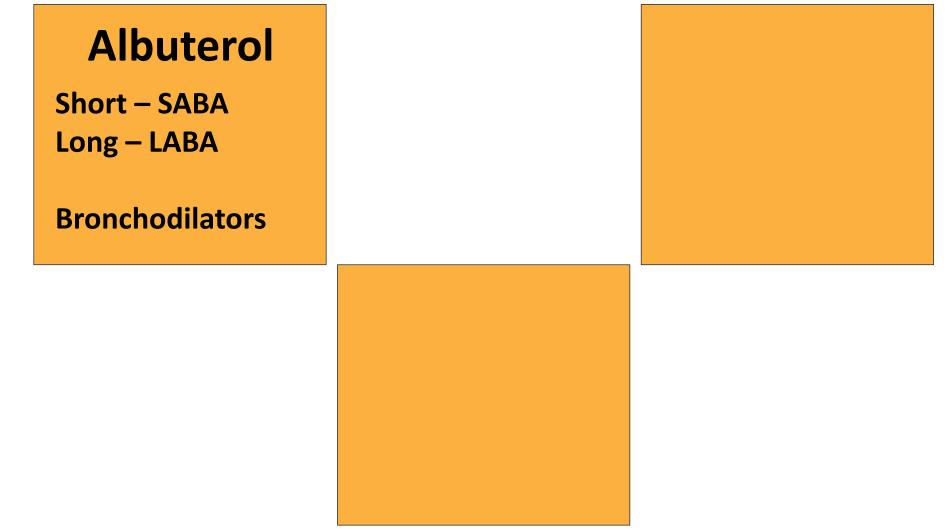


COPD – normal vs hyperinflated lungs





Respiratory medications: Three categories of medications

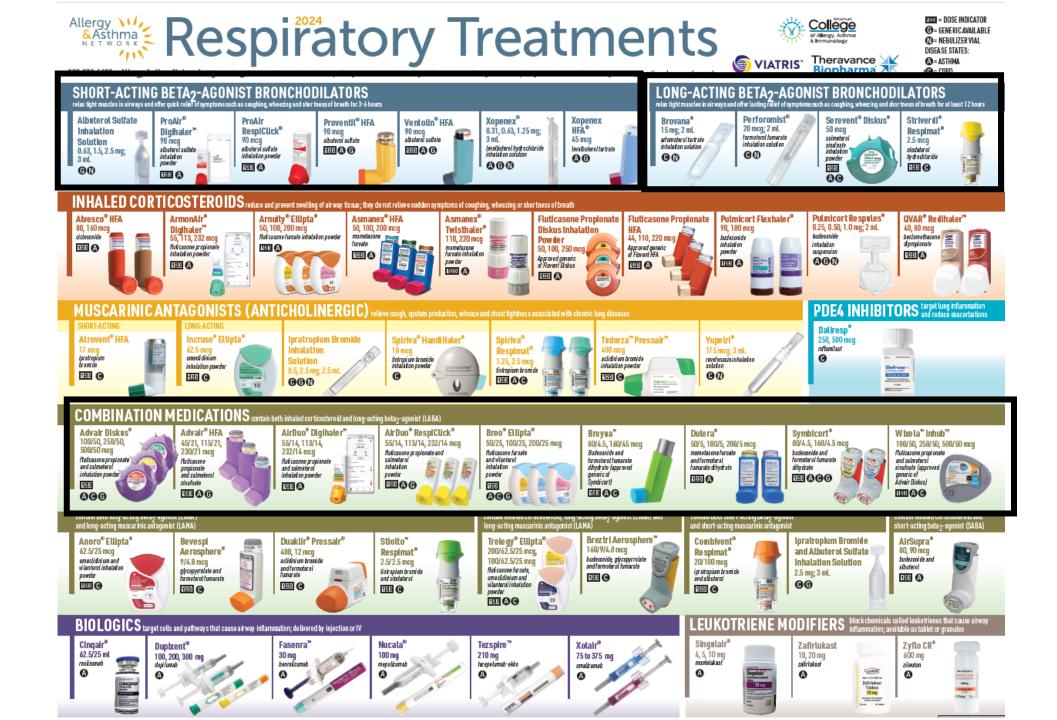




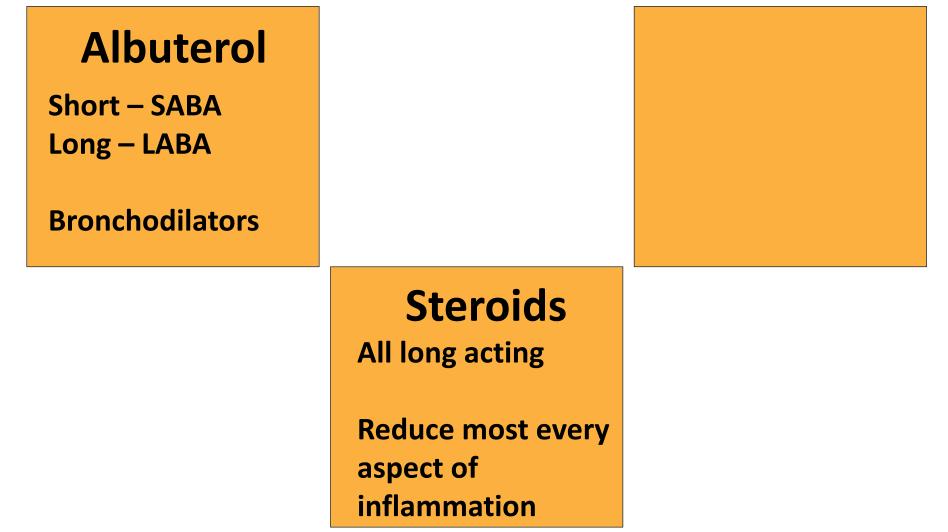
Albuterol – short acting bronchodilator, relaxes smooth muscle. Binds to beta receptors on smooth muscle, causing about a billion things to happen that drop the calcium in the cell and it relaxes.

Salmeterol/formoterol/vilanterol – Same thing as above but lasts 12 or 24 hours





Respiratory medications: Three categories of medications





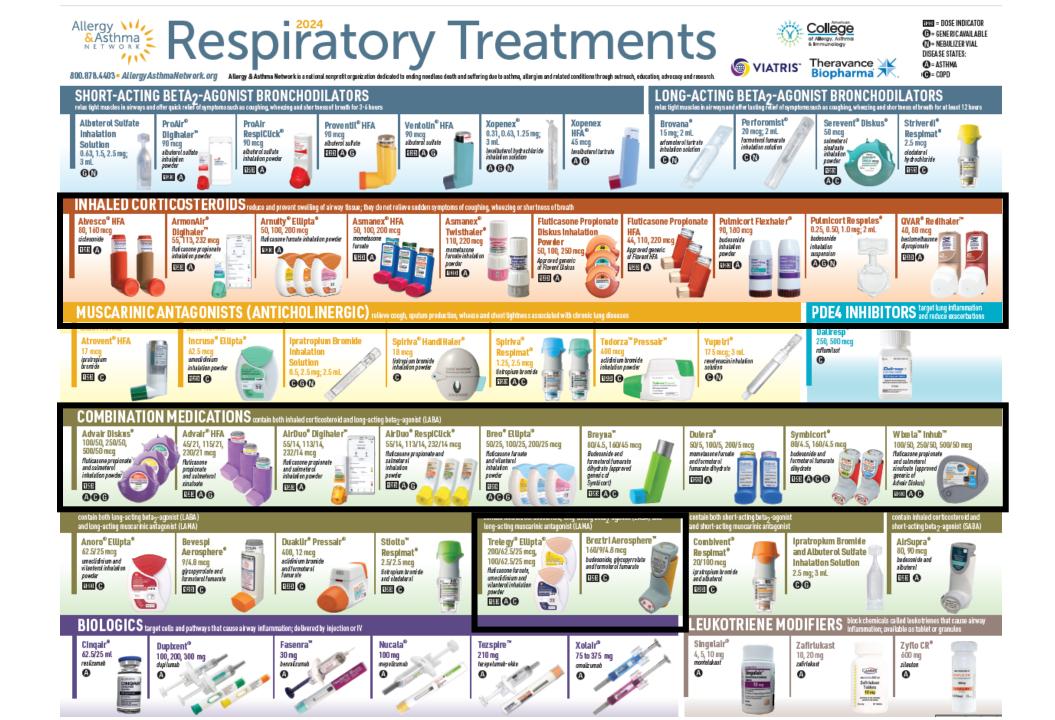
Corticosteroids bind to the glucocorticoid receptor and <u>mediate changes in gene expression</u> that lead to multiple downstream effects over hours to days.

Almost every inflammation mediator is reduced

Many actions, all with a central goal of reducing inflammation at the source

Most aspects of inflammation are affected





Respiratory medications: Three categories of medications

Albuterol Short – SABA Long – LABA Bronchodilators		SAMA/LAMA Short – SAMA Long – LAMA Anticholinergic and constriction prevention
	Steroids All long acting	
	Reduce most every aspect of inflammation	

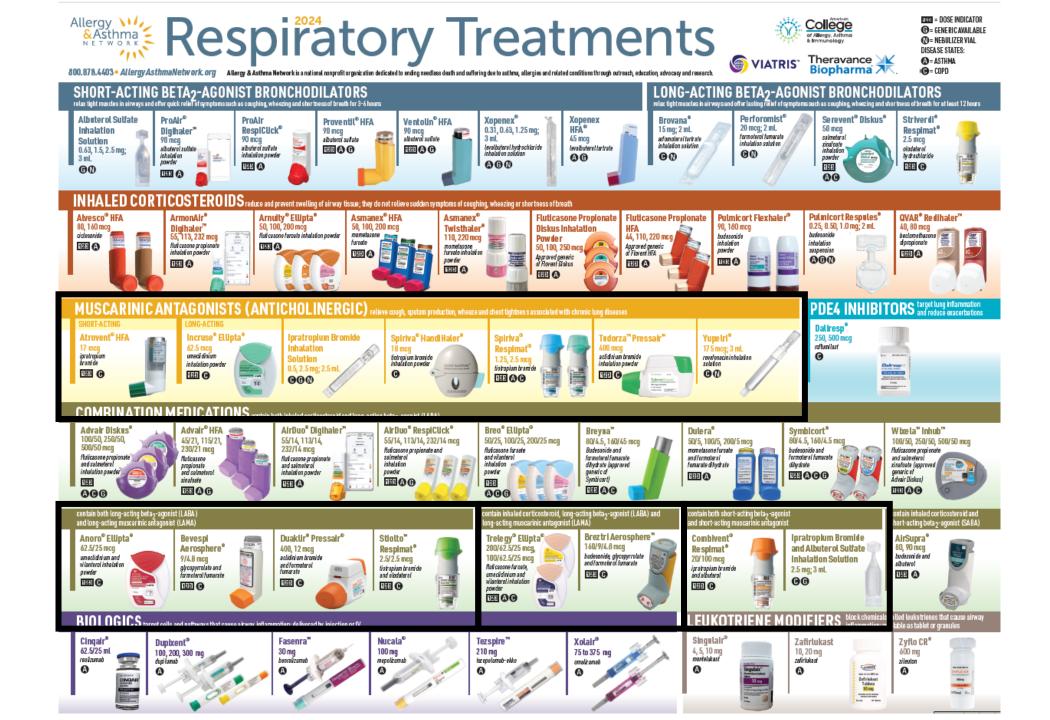


Medication Categories: SAMA/LAMA

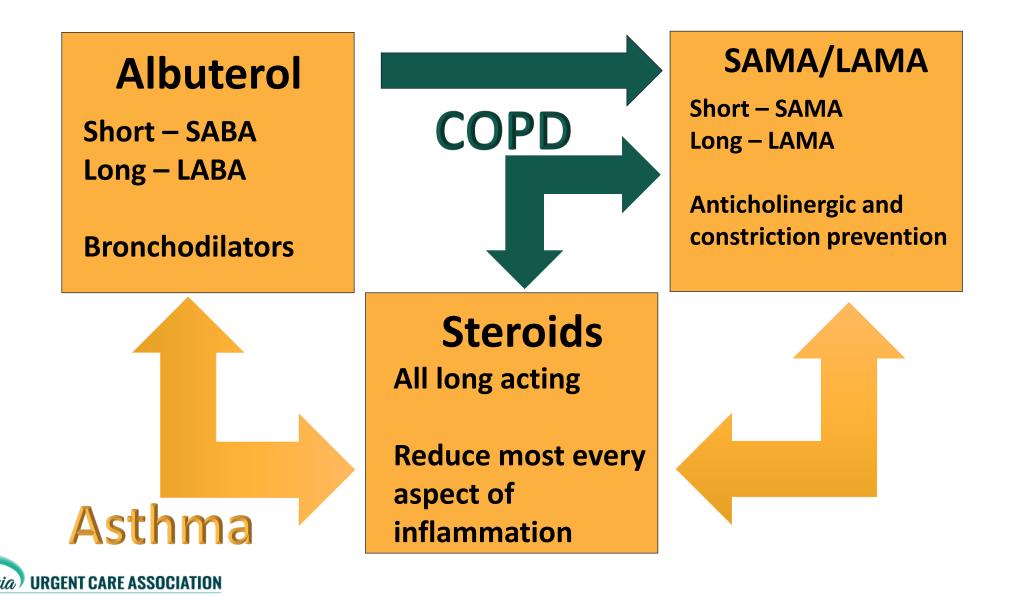
 Ipratropium bromide is our only short acting muscarinic, and there are several long acting

 These are anti-cholinergic medications that dry up secretions and help prevent constriction





Respiratory medications: Three categories of medications



QR Code for Inhaler Chart – English





QR Code for Inhaler Chart – Spanish

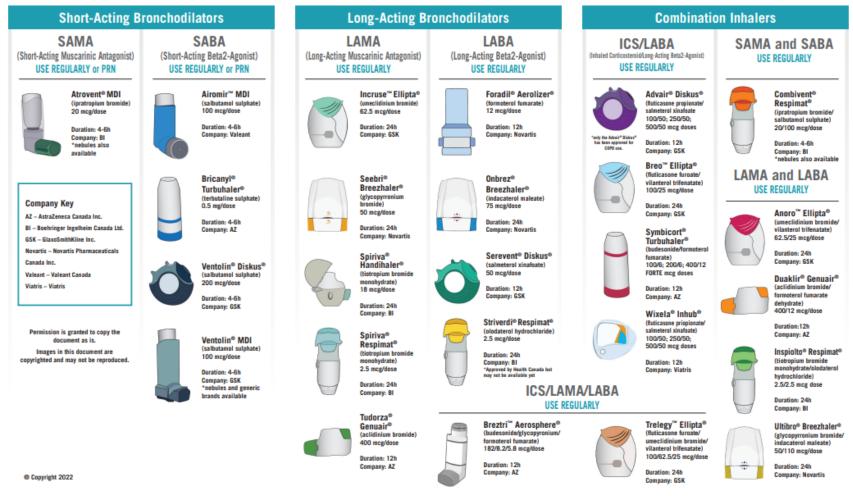




For Reference

RESPTREC® COPD MEDICATIONS & EDUCATOR COURSE

www.resptrec.org www.lungsask.ca



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GLOBAL INITIATIVE FOR CHRONIC OBSTRUCTIVE LUNG DISEASE (GOLD):



www.goldcopd.org

Global Initiative for Chronic Obstructive Lung Disease

2024 REPORT





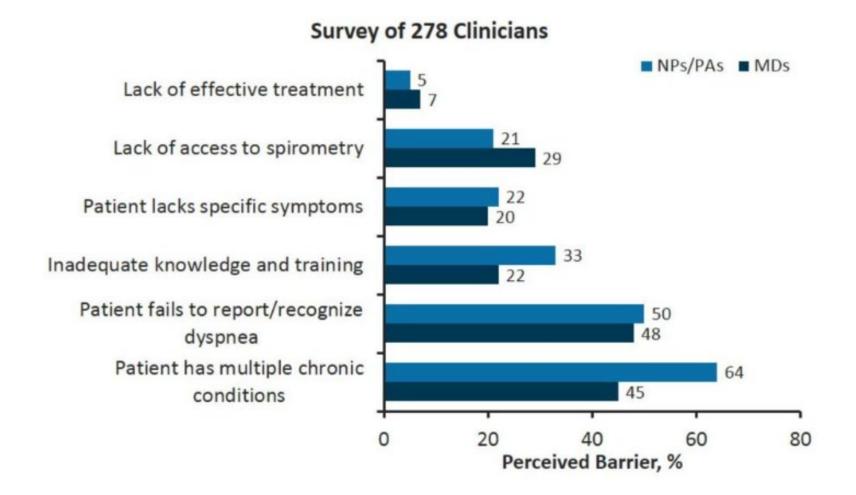
https://goldcopd.org/2024-gold-report/

COPD Defined

'A common preventable and treatable disease, is characterized by persistent airflow limitation that is usually progressive and associated with an enhanced chronic inflammatory response in the airways and the lung to noxious particles or gases. <u>Exacerbations and</u> <u>comorbidities contribute to the overall severity</u> in individual patients.'



Why is COPD so Underdiagnosed?



Yawn BP, et al. Int J Chron Obstruct Pulmon Dis. 2008;3:311-317.



Consider COPD in a patient with any of these characteristics

Symptom	Detail
Dyspnea that is:	 Progressive over time Characteristically worse with exercise Persistent
Chronic cough	May be intermittent and unproductiveRecurrent wheeze
Chronic sputum production	Any pattern of chronic sputum production may indicate COPD
Recurrent LRTIs	
History of risk factors	 Host factors (eg, genetic factors, congenital/developmental abnormalities) Tobacco smoke Smoke from home cooking and heating fuels Occupational dusts, vapors, fumes, gases and other chemicals
Family history of COPD and/or childhood factors	 Examples include: low birthweight, childhood respiratory infections, Hx of Alpha-1 Antitrypsin Deficiency or unexplained pulmonary disease

These indicators are not diagnostic themselves, but the presence of multiple key indicators increases the probability of a diagnosis of COPD. COPD, chronic obstructive pulmonary disease; GOLD, Global Initiative for Chronic Obstructive Lung Disease; LRTI, lower respiratory tract infection. 2023 GOLD Report. https://goldcopd.org/2023-gold-report-2/.



Diagnose COPD and then Staging

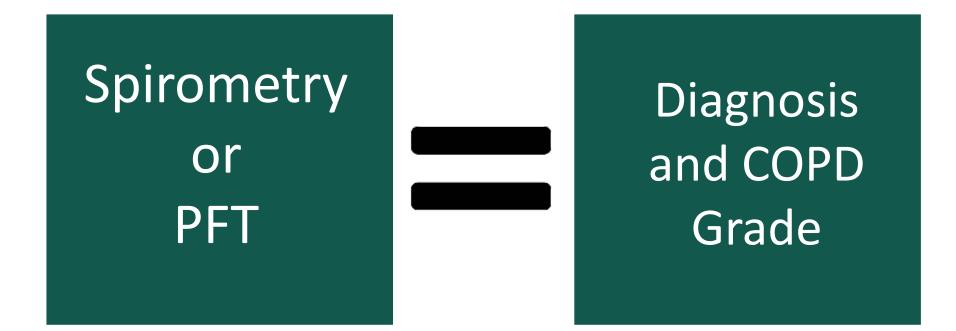
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		W LIMITATION SEVERITY SRONCHODILATOR FEV ₁)	
In patients with FE	V1/FVC < 0.70:		
GOLD 1:	Mild	$FEV_1 \ge 80\%$ predicted	
GOLD 2:	Moderate	$50\% \le FEV_1 < 80\%$ predicted	
GOLD 3:	Severe	$30\% \le FEV_1 < 50\%$ predicted	
GOLD 4:	Very Severe	$FEV_1 < 30\%$ predicted	

This is comparing the patient to themselves – how much of their own air can they get out in 1 second This is comparing the patient to a peer based on height, weight, age, gender and ethnicity

https://goldcopd.org/2024-gold-report/

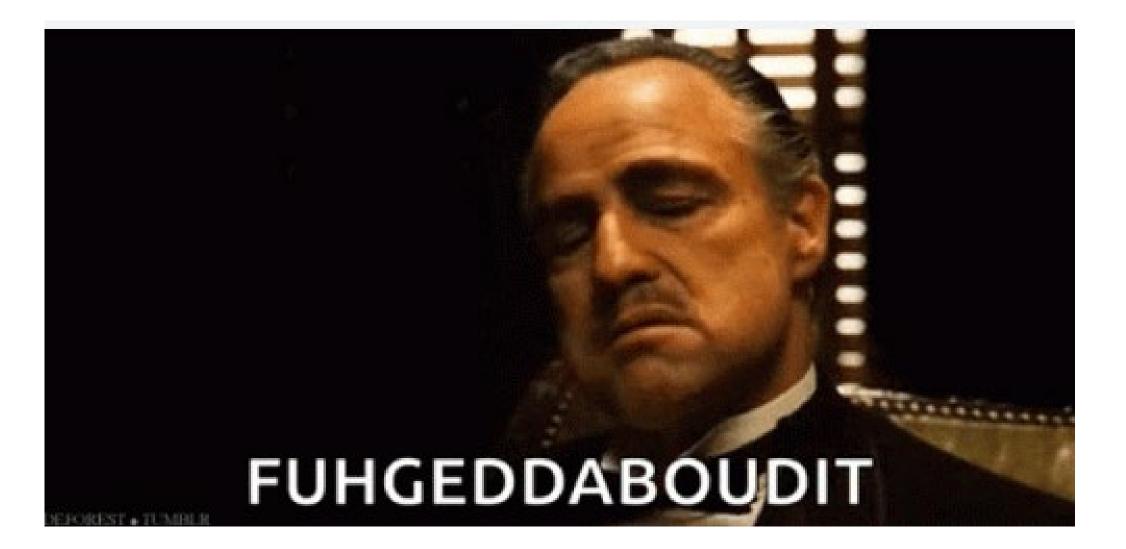
COPD Diagnosis and Treatment



So do this once, then, the good news . . .



COPD Diagnosis and Treatment



COPD Diagnosis and treatment

 Diagnosis is based on how much air can you exhale in 1 second (FEV1/FVC)

• Staging is based on comparing you to a peer

- TREATMENT is based on two factors what GOLD calls "Treatable Traits"
- The traits are exacerbation and symptoms



Review

- COPD is widespread and largely underdiagnosed, flares are frequent
- Most are tobacco related but there are others
- In Urgent Care consider this even if they don't have a diagnosis, most wont.
- You need spirometry to get the diagnosis and stage of COPD
- But the stage DOES NOT equal quality of life, life expectancy and does not affect treatment decisions.
- Now you want to know –
- How are you?
- How often are you sick?



For Reference

CAT[™] ASSESSMENT

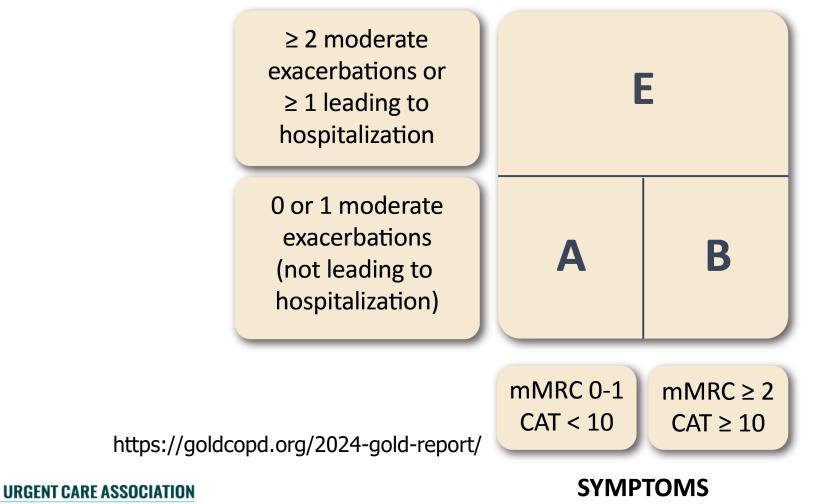
For each item below, place a mark (x) in the box that best describes you currently. Be sure to only select one response for each question.

EXAMPLE: I am very happy	0 2 3 4 5	I am very sad	SCORE
l never cough	012345	I cough all the time	
I have no phlegm (mucus) in my chest at all	012345	My chest is completely full of phlegm (mucus)	
My chest does not feel tight at all	012345	My chest feels very tight	
When I walk up a hill or one flight of stairs I am not breathless	012345	When I walk up a hill or one flight of stairs I am very breathless	
I am not limited doing any activities at home	012345	I am very limited doing activities at home	
I am confident leaving my home despite my lung condition	012345	I am not at all confident leaving my home because of my lung condition	
I sleep soundly	012345	l don't sleep soundly because of my lung condition	
I have lots of energy	012345	I have no energy at all	
			\bigcap
Reference: Jones et al. ERJ 2009; 34 (3); 648-54.			



GOLD 2024 Treatment Guidelines

EXACERBATION HISTORY



GOLD 2024 Treatment Guidelines

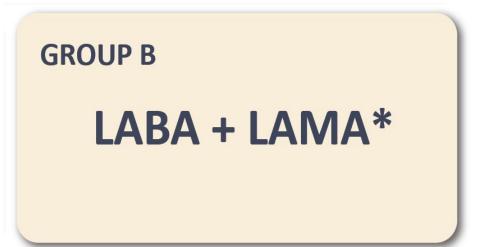
0 or 1 moderate exacerbations (not leading to hospital admission)

GROUP A A bronchodilator

mMRC 0-1, CAT < 10

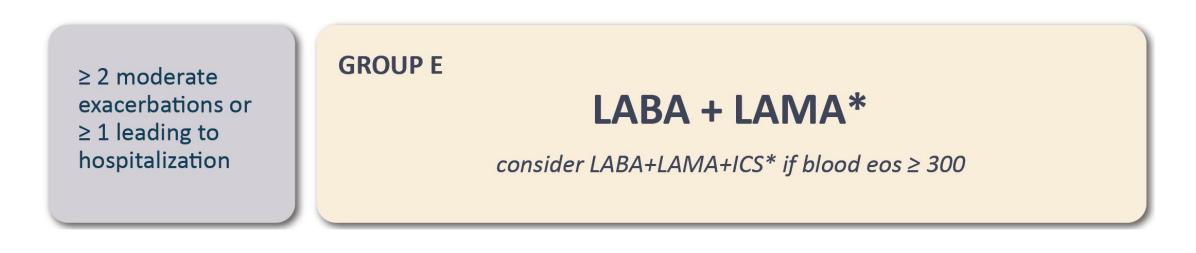


0 or 1 moderate exacerbations (not leading to hospital admission)



mMRC \geq 2, CAT \geq 10

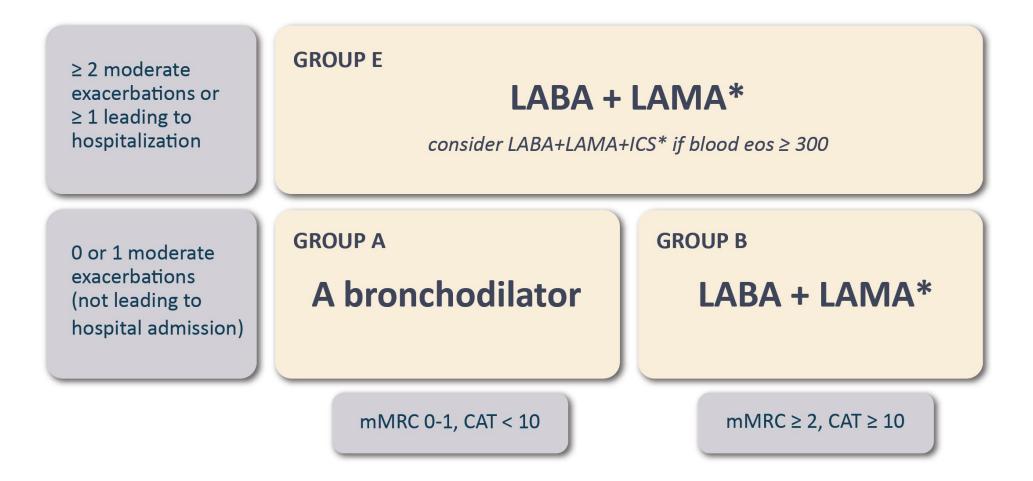




mMRC 0-1, CAT < 10

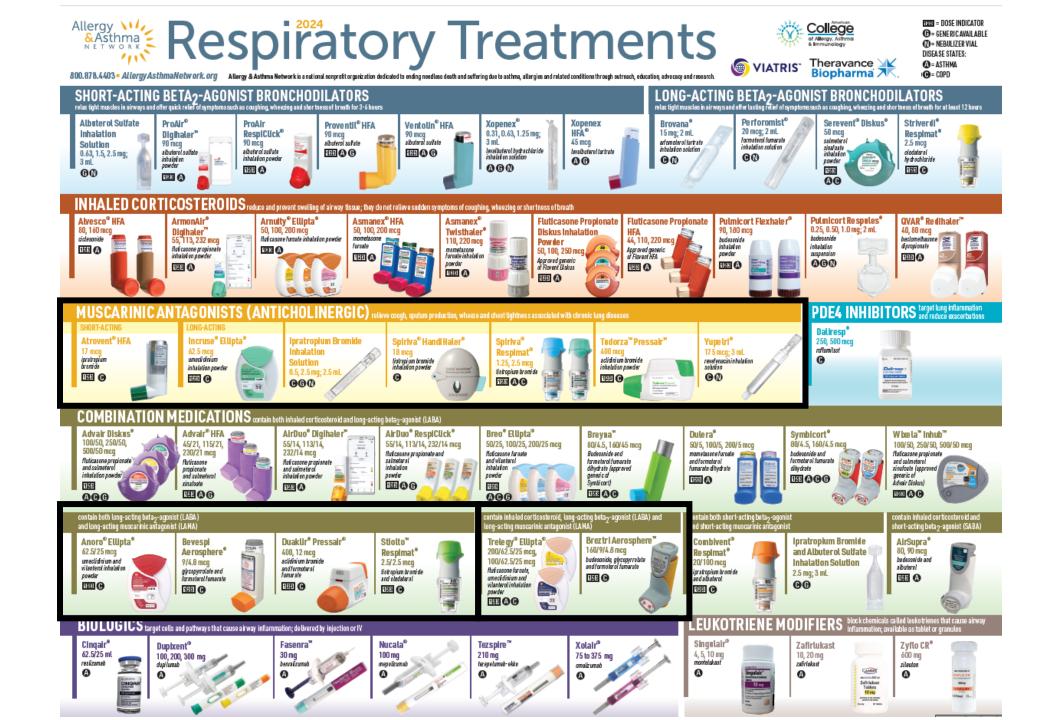
mMRC ≥ 2, CAT ≥ 10





*single inhaler therapy may be more convenient and effective than multiple inhalers





Why not Inhaled Steroids?

If not needed don't use them!

Increased risk of all URIs and increased risk of pneumonia and exacerbations

Fluticasone causes the most URIs

GROUP E

LABA + LAMA*

consider LABA+LAMA+ICS* if blood eos ≥ 300

Inhaled corticosteroids and risk of pneumonia in patients with chronic obstructive pulmonary disease: A meta-analysis of randomized controlled trials

Mingjin Yang ¹, Yuejun Du ¹, Hong Chen ¹, Depeng Jiang ², Zhibo Xu ³

Affiliations + expand PMID: 31629940 DOI: 10.1016/j.intimp.2019.105950

Abstract

Objective: Inhaled corticosteroids (ICS) are generally used to treat patients with chronic obstructive pulmonary disease (COPD) who suffer from repeated exacerbations. Recently, it was reported that ICS treatment increased the risk of pneumonia in COPD patients. But it is controversial. The objective of this paper is to clarify the associations between ICS treatment and the risk of pneumonia in COPD patients.

Methods: PubMed, Cochrane Library, Clinical Trials.gov, and Embase were searched from February 2019 to June 2019. Randomized clinical trials (RCTs) were incorporated that compared ICS with non-ICS treatment on the risk of pneumonia in COPD patients. Meta-analyses were conducted by the Peto and Mantel-Haenszel approaches with corresponding 95% CIs.

Results: Twenty-five trials (N = 49,982 subjects) were included. Pooled results demonstrated a significantly increased risk of pneumonia with ICS use in COPD patients (RR, 1.59, 95% CI, 1.33-1.90; $I^2 = 51\%$). ICS treatment also increased the risk of severe pneumonia (RR, 2.17, 95% CI, 1.47-3.22; $I^2 = 29\%$). The results of subgroup analysis based on doses of ICS were consistent with the above. However, subgroup analyses based on types of ICS revealed that fluticasone therapy was associated with an increased risk of pneumonia but not budesonide. In addition, medium- and low-doses of budesonide treatment also did not increase the risk of pneumonia.

Conclusions: Use of ICS increases the risk of pneumonia in patients with COPD. The above is prominent for fluticasone-containing ICSs but not for budesonide-containing ICSs.



Should Inhaled Steroids be used?

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

(note the scenario is different when considering ICS withdrawal)

STRONGLY FAVORS 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

FAVORS USE	1 moderate exacerbation of COPD per year [#]	
	Blood eosinophils 100 to < 300 cells/µL	

	Repeated pneumonia events
AGAINST USE	Blood eosinophils < 100 cells/µL
	History of mycobacterial infection

[#]despite appropriate long-acting bronchodilator maintenance therapy (see Table 3.4 and Figure 4.3 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

Review

- Outpatient therapy for COPD is determined by the number and frequency of exacerbations and overall symptom assessment. How sick? How are you?
- Most will do well on a LAMA/LABA. That is your go-to most of the time
- Inhaled steroids have significant s/e and concerns (2X risk of all URI and pneumonia), the benefits outweigh the negatives for patients with asthma and/or those with eosinophils over 300 and in general, for those with severe disease and controlling inflammation is one of the final steps – for all of these a triple inhaler is reasonable.
- UC providers have a SIGNIFICANT role in helping COPD patients get on the correct inhaler, helping them get set up to STAY WELL after we help them GET WELL!
- What about this getting them well. the exacerbation!



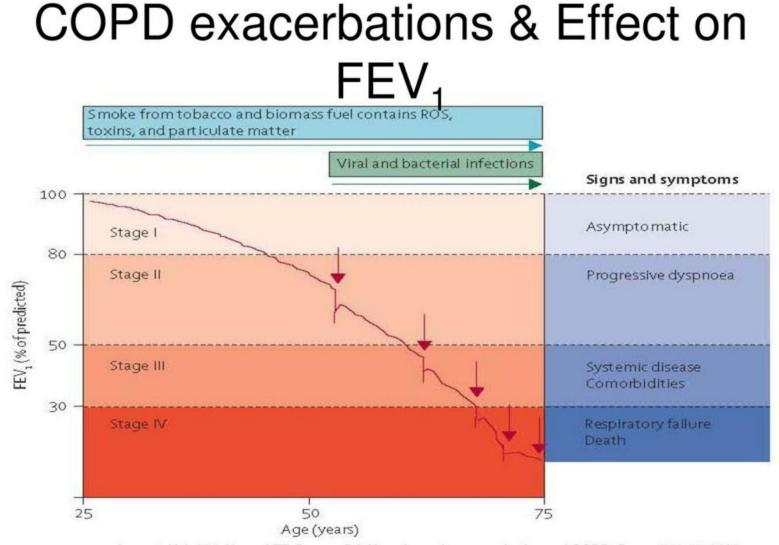
GOLD #1 Goal is Exacerbation Treatment and Prevention

 Exacerbations are not "bumps" in the road like they are for asthma

 Moderate to severe exacerbations are life altering, patients never recover fully.

 An exacerbation is an acute change in a patient's baseline dyspnea, cough, or sputum that is beyond normal variability, and that is sufficient to warrant a change in therapy.







Lancet, Vol. 374, Hansel TT, Barnes PJ, New drugs for exacerbations of COPD, Pages 744-55, 2009

COPD Exacerbation Definition

An exacerbation of chronic obstructive pulmonary disease (ECOPD) is defined as an event characterized by increased dyspnea and/or cough and sputum that worsens in < 14 days which may be accompanied by tachypnea and/or tachycardia and is often associated with increased local and systemic inflammation caused by infection, pollution, or other insult to the airways.⁽³⁰⁴⁾



https://goldcopd.org/2024-gold-report/

COPD Exacerbation Definition

Practically – Patients present with an increase in medication use, an increase in cough, sputum increase and/or change in color. And the key is that they came in to see you.

"My inhalers don't work"

"I can't breathe"

"Chest is tight"

"Using my nebulizer more"

All of these strongly support that an intervention is needed.



Considerations for the Exacerbation Patient

	Pneumonia
	Chest radiograph
	Pulmonary embolism
Most frequent	 Clinical probability assessment (Hemoptysis, surgery, fracture, history of cancer, DVT) D-dimer
	 CT angiography for pulmonary embolism
	Heart failure
	 Chest radiograph NT Pro-Brain Natriuretic Peptide (Pro-BNP) and BNP Echocardiography
	Pneumothorax, pleural effusion
Less frequent	 Chest radiograph Thoracic ultrasound
	Myocardial infarction and/or cardic arrhythmias (atrial fibrillation/flutter)
	 Electrocardiography Troponin



COPD Exacerbation Diagnosis

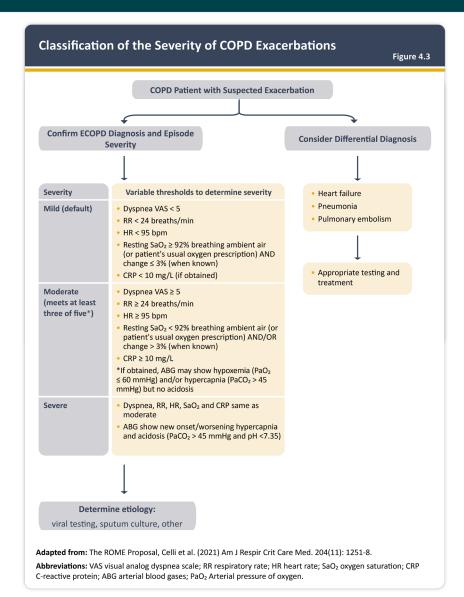
1.	Complete a thorough clinical assessment for evidence of COPD and potential respiratory and non-respiratory concomitant diseases, including consideration of alternative causes for the patient's symptoms and signs: primarily pneumonia, heart failure, and pulmonary embolism.
2.	 Assess: a. Symptoms, severity of dyspnea that can be determined by using a VAS, and documentation of the presence of cough. b. Signs (tachypnea, tachycardia), sputum volume and color, and respiratory distress (accessory muscle use).
3.	Evaluate severity by using appropriate additional investigations such as pulse oximetry, laboratory assessment, CRP, arterial blood gases.
4.	Establish the cause of the event (viral, bacterial, environmental, other).

Abbreviations: COPD = chronic obstructive pulmonary disease; CRP = C-reactive protein; VAS = visual analog scale.



https://goldcopd.org/2024-gold-report/

Exacerbation Severity Assessment





Exacerbation Severity Assessment

Severity	Variable thresholds to determine severity
Mild (default)	 Dyspnea VAS < 5 RR < 24 breaths/min HR < 95 bpm Resting SaO₂ ≥ 92% breathing ambient air (or patient's usual oxygen prescription) AND change ≤ 3% (when known) CRP < 10 mg/L (if obtained)
Moderate (meets at least three of five*)	 Dyspnea VAS ≥ 5 RR ≥ 24 breaths/min HR ≥ 95 bpm Resting SaO₂ < 92% breathing ambient air (or patient's usual oxygen prescription) AND/OR change > 3% (when known) CRP ≥ 10 mg/L *If obtained, ABG may show hypoxemia (PaO₂ ≤ 60 mmHg) and/or hypercapnia (PaCO₂ > 45 mmHg) but no acidosis
Severe	 Dyspnea, RR, HR, SaO₂ and CRP same as moderate ABG show new onset/worsening hypercapnia and acidosis (PaCO₂ > 45 mmHg and pH <7.35)



Considerations for Hospitalization.

- 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



- 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 normally be more than 5 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 normally be 5 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)



Why Nebulizers?

Respiratory Medicine 161 (2020) 105857



Contents lists available at ScienceDirect

Respiratory Medicine

journal homepage: http://www.elsevier.com/locate/rmed

Review article

The role of inspiratory flow in selection and use of inhaled therapy for patients with chronic obstructive pulmonary disease



spiratory MEDICINE

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ARTICLE INFO

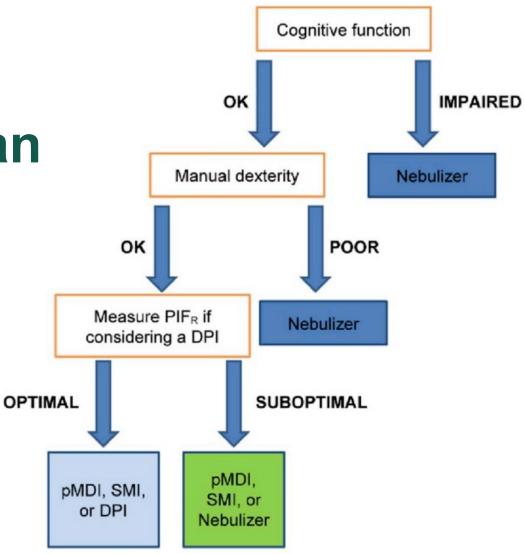
ABSTRACT

Keywords: Chronic obstructive pulmonary disease Hand-held inhalers Inhalation technique Inspiratory flow Peak inspiratory flow Inhalation therapy is the mainstay of chronic obstructive pulmonary disease management, and inhaler selection can have a profound impact on drug delivery and medication adherence, as well as on treatment outcomes. Although multiple delivery systems, such as pressurized metered-dose inhalers, dry powder inhalers, slow-mist inhalers, and nebulizers, are available, clinical benefits achieved by patients rely on effective delivery of the inhaled medication to the airways. Among several factors influencing drug deposition, inspiratory flow is one of the most important. Inspiratory flow impacts drug delivery and subsequent clinical efficacy, making it necessary to adequately train patients to ensure correct inhaler use. Peak inspiratory flow is the maximal airflow generated during a forced inspiratory maneuver. Health care professionals need to select the appropriate delivery system after carefully considering patient characteristics, including lung function, optimal inspiratory flow, manual dexterity, and cognitive function. Herein, the role of inspiratory flow in the selection and use of inhaled therapy in patients with COPD is reviewed.



Why Nebulizers?

Do what you can in the UC, but if you can ask about this you may do more to help them than any medication





Why Nebulizers?

- Measure this with an In-Check Device
- Can also see if they can "make noise" with their inhaler*
- Can they hold a Post-it note to their lips**?



• Do they feel nebulized medication is sig better?

*Barjaktarevic IZ, Milstone AP. Nebulized Therapies in COPD: Past, Present, and the Future. Int J Chron Obstruct Pulmon Dis. 2020 Jul 12;15:1665-1677. doi: 10.2147/COPD.S252435. PMID: 32764912; PMCID: PMC7367939. ** No data to support this whatsoever, but I like it.



For Reference about Azithromycin

Review > Rev Med Virol. 2021 Mar;31(2):e2163. doi: 10.1002/rmv.2163. Epub 2020 Sep 23.

Azithromycin in viral infections

Madeleine E Oliver ¹, Timothy S C Hinks ²

Affiliations + expand PMID: 32969125 PMCID: PMC7536932 DOI: 10.1002/rmv.2163

Abstract

Azithromycin (AZM) is a synthetic macrolide antibiotic effective against a broad range of bacterial and mycobacterial infections. Due to an additional range of anti-viral and anti-inflammatory properties, it has been given to patients with the coronaviruses SARS-CoV or MERS-CoV. It is now being investigated as a potential candidate treatment for SARS-CoV-2 having been identified as a candidate therapeutic for this virus by both in vitro and in silico drug screens. To date there are no randomised trial data on its use in any novel coronavirus infection, although a large number of trials are currently in progress. In this review, we summarise data from in vitro, murine and human clinical studies on the anti-viral and anti-inflammatory properties of macrolides, particularly AZM. AZM reduces in vitro replication of several classes of viruses including rhinovirus, influenza A, Zika virus, Ebola, enteroviruses and coronaviruses, via several mechanisms. AZM enhances expression of anti-viral pattern recognition receptors and induction of anti-viral type I and III interferon responses. Of relevance to severe coronavirus-19 disease (COVID-19), which is characterised by an over-exuberant innate inflammatory response, AZM also has anti-inflammatory properties including suppression of IL-1beta, IL-2, TNF and GM-CSF. AZM inhibits T cells by inhibiting calcineurin signalling, mammalian target of rapamycin activity and NFKB activation. AZM particularly targets granulocytes where it concentrates markedly in lysosomes, particularly affecting accumulation, adhesion, degranulation and apoptosis of neutrophils. Given its proven safety, affordability and global availability, tempered by significant concerns about antimicrobial stewardship, there is an urgent mandate to perform welldesigned and conducted randomised clinical trials.

Keywords: COVID-19; SARS-CoV-2; azithromycin; coronavirus; macrolide; mechanism; review; virus.



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- Inhalers/Nebulizers Make sure they have them, review use. SVN if any doubt about ability. Duoneb best SVN, Combivent best rescue inhaler. Have them dose based on TIME, not symptoms, till the steroids kick in, usually 24 hours. Q4H while awake, at noc if they get up.
- Limit? None. Do what is needed to break through. But if >3 and not getting better then come in. Don't have to do the entre SVN.



- PO Prednisone (cheaper) 40mg for 3 days, 20mg for 2 days. Take QD, dose in the AM if possible. Warn of s/e, glucose.
- PO Azithromycin for MOST. 500mg QD x 1 day then 250 mg QD x 4 days.
- Both are QD for 5 days, compliance is high, cost low.
- May do a patient-controlled taper 40mg QD till they feel about half better then 20mg QD till close to baseline. No data on this.
- Tessalon? Sure, what does this do anyway?
- Mucinex DM



- If you can, talk with them about cost/refills. Help them by refilling chronic COPD medications if you are comfortable doing so.
- AstraZeneca has a cap of \$35 on inhalers for non-insured patients
- Keep an inhaler and a soft-mist inhaler with you at work, wash the actuator part and let them show you how they do it. More than half are using it wrong. PROMISE.
- GET them better, then help them STAY better. Be a Superstar!





For Reference All COPD Medications – including nebulized

Commonly Used Maintenance Medications in COPD*

Table 3.3

			DELIVERY OPTIONS		
Generic Drug Name	Inhaler Type	Nebulizer	Oral	Injection	Duration of Action
BETA ₂ -Agonists					
Short-acting (SABA)					
Fenoterol	MDI	1	pill, syrup		4-6 hours
Levalbuterol	MDI	1			6-8 hours
Salbutamol (albuterol)	MDI & DPI	1	pill, syrup, extended	1	4-6 hours
			release tablet		12 hours (ext. releas
Terbutaline	DPI		pill	1	4-6 hours
Long-acting (LABA)					
Arformoterol		1			12 hours
Formoterol	DPI	1			12 hours
Indacaterol	DPI				24 hours
Olodaterol	SMI				24 hours
Salmeterol	MDI & DPI				12 hours
Anticholinergics					
Short-acting (SAMA)					
Ipratropium bromide	MDI	1			6-8 hours
Oxitropium bromide	MDI				7-9 hours
Long-acting (LAMA)					
Aclidinium bromide	DPI,				MDI 12 hours
Glycopyrronium bromide	DPI		solution	1	12-24 hours
Tiotropium	DPI, SMI, MDI				24 hours
Umeclidinium	DPI				24 hours
Glycopyrrolate		1			12 hours
Revefenacin		1			24 hours
Combination Short-Acting Beta ₂ -Agonist P	lus Anticholiner	ic in One De	vice (SABA+SAMA)		
Fenoterol/ipratropium	SMI	1			6-8 hours
Salbutamol/ipratropium	SMI, MDI	1			6-8 hours
Combination Long-Acting Beta ₂ -Agonist Pl	us Anticholinerg	ic in One De	vice (LABA+LAMA)		
Formoterol/aclidinium	DPI		, í l		12 hours
Formoterol/glycopyrronium	MDI				12 hours
Indacaterol/glycopyrronium	DPI				12-24 hours
Vilanterol/umeclidinium	DPI				24 hours
Olodaterol/tiotropium	SMI				24 hours
Methylxanthines					
Aminophylline			solution	1	Variable, up to 24 ho
Theophylline (SR)			pill	1	Variable, up to 24 ho
Combination of Long-Acting Beta ₂ -Agonist	Plus Corticoster	oid in One D		•	
Formoterol/beclometasone	MDI, DPI		ence (LADATICS)		12 hours
Formoterol/budesonide	MDI, DPI				12 hours
Formoterol/mometasone	MDI, DFT				12 hours
Salmeterol/fluticasone propionate	MDI, DPI				12 hours
Vilanterol/fluticasone furoate	DPI				24 hours
Triple Combination in One Device (LABA+I					24110013
Fluticasone/umeclidinium/vilanterol	DPI				24 hours
Beclometasone/formoterol/glycopyrronium	MDI, DPI				12 hours
Budesonide/formoterol/glycopyrrolate	MDI, DPI				12 hours
Phosphodiesterase-4 Inhibitors					12 110013
			pill		24 hours
Roflumilast			piii		24 110015
Mucolytic Agents			n ill		12 hours
Erdosteine			pill		12 hours
Carbocysteine [†] N-acetylcysteine [†]			pill		
		1	pill		1

*Not all formulations are available in all countries. In some countries other formulations and dosages may be available. †Dosing regimens are under discussion MDI = metered dose inhaler; DPI = dry powder inhaler; SMI = soft mist inhaler. Note that glycopyrrolate & glycopyrronium are the same compound.



For Reference Smoking Cessation

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Practice Guidelines

Medications for Smoking Cessation: Guidelines from the American Thoracic Society

Reference

Am Fam Physician. 2021 Mar 15;103(6):380-381.

Author disclosure: No relevant financial affiliations.

Key Points for Practice

• Varenicline is more effective than nicotine patches and bupropion with similar or fewer adverse events, even with comorbid psychiatric or substance abuse conditions.

• Combining varenicline with nicotine patches appears to be more effective than using varenicline alone based on limited evidence.

• For people who smoke and are not ready to quit, prescribing varenicline increases six-month abstinence with an NNT of 6 compared with waiting for readiness.

• Extending treatment beyond 12 weeks increases abstinence, with an NNT of 19 compared with shorter treatment durations.



From the AFP Editors

For Reference COPD Action Plan

https://www.lung.org/get media/c7657648-a30f-4465-af92fc762411922e/copdaction-plan.pdf.pdf

American Lung Association.

My COPD Action Plan

Patients and healthcare providers should complete this action plan together. This plan should be discussed at each visit and updated as needed.

The green, yellow and red zones show symptoms of COPD. The list of symptoms is not complete. You may experience other symptoms. In the "Actions" column, your healthcare provider will recommend actions for you to take. Your healthcare provider may write down other actions in addition to those listed here.

Green Zone: I am doing well today	Actions
Usual activity and exercise level Usual amounts of cough and phlegm/mucus Sleep well at night Appetite is good	Take daily medicines Use oxygen as prescribed Continue regular exercise/diet plan Avoid tobacco product use and other inhaled irritants
Yellow Zone: I am having a bad day or a COPD flare	Actions
 More breathless than usual I have less energy for my daily activities Increased or thicker phlegrn/mucus Using quick relief inhaler/nebulizer more often More swelling in ankles More coughing than usual I feel like I have a "chest cold" Poor sleep and my symptoms woke me up My appetite is not good My medicine is not helping 	Continue daily medication Use quick relief inhaler every hours Start an oral corticosteroid (specily name, dose, and duration) Start an antibiotic (specily name, dose, and duration) Use oxygen as prescribed Get plenty of rest Use pursed lip breathing Avoid secondhand smoke, e-cigarette aerosol, and other inhaled irritants Call provider immediately if symptoms do not improve
Red Zone: I need urgent medical care	Actions
Severe shortness of breath even at rest Not able to do any activity because of breathing Not able to sleep because of breathing Fever or shaking chills Feeling confused or very drowsy Chest pains Coughing up blood	Call 911 or seek medical care immediately While getting help, immediately do the following:

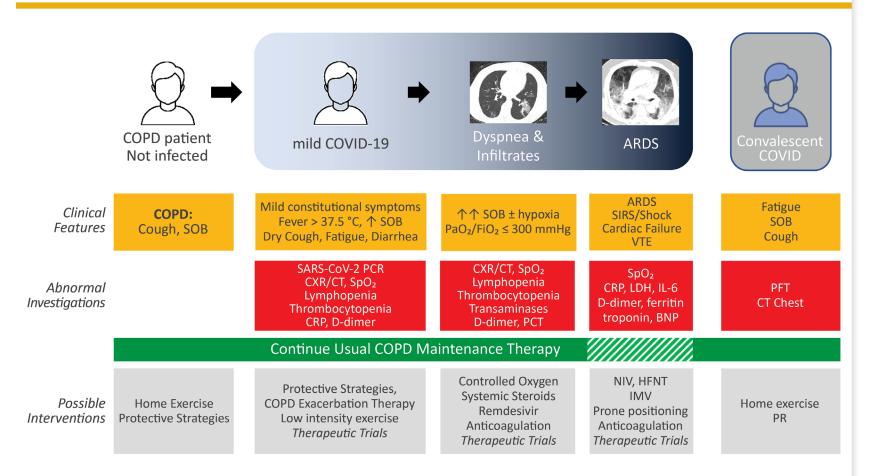
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For Reference COPD and COVID



(ARDS, Adult respiratory distress syndrome; BNP, brain natriuretic peptide; CRP, C reactive protein; CT, computed tomography; CXR, chest radiograph; HFNT, high flow nasal therapy; IL-6, interleukin 6; IMV, invasive mechanical ventilation; LDH, lactate dehydrogenase; NIV, non-invasive ventilation; PCT, procalcitonin; PFT, pulmonary function tests; PR, pulmonary rehabilitation; SOB, Shortness of breath; SpO₂, peripheral oxygen saturation; VTE, venous thromboembolism)

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Halpin et al. 2020. Global Initiative for the Diagnosis, Management, and Prevention of Chronic Obstructive Lung Disease: The 2020 GOLD Science Committee Report on COVID-19 & COPD. Published Ahead of Print: https://www.atsjournals.org/doi/abs/10.1164/rccm.202009-3533SO The American Journal of Respiratory and Critical Care Medicine is an official journal of the American Thoracic Society

For Reference

https://www.lung.org/get media/c7657648-a30f-4465-af92fc762411922e/copdaction-plan.pdf.pdf

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