

Emergency (ED) Acute Exacerbation of Chronic Obstructive Pulmonary Disease (AECOPD) Adult Order Set

Evidence-based, severity-guided support for managing Acute Exacerbations of COPD, with recommendations on aerosol delivery, pharmacologic therapy, oxygen use, non-invasive support, and environmental impact.

PATIENT INFORMATION									
Last Name (Legal)		Fi	First Name (Legal)						
Preferred Name Last		First			DOB (do	d-mm-y	d-mm-yyyy)		
PHN UL	.I Sa	ame as PHN				MRN			
Administrative Gender Male	Fe	emale N	Non-b	inary	Prefe	r not to	disclose	Unknown	
SEVERITY-BASED DECISION SUPP	ORT	– use <u>ONE</u> o	f the f	ollowing	validate	d COPI	O scoring tool	S	
GOLD									
MILD		МС	ODER	ATE			SEVE	RE	
Signs and symptoms: □ Dyspnea VAS <5 □ RR < 24 Breaths/min □ HR < 95 bpm □ Resting SaO2 ≥ 92% on ambient air or home O2 prescription □ CRP <10mg/L if obtained Severity:		Signs and symptoms: □ Dyspnea VAS ≥ 5 □ RR ≥ 24 Breaths/min □ HR ≥ 95 bpm □ Resting SaO2 < 92% on ambient air or home O2 prescription □ CRP 10mg ≥ /L if obtained □ If obtained ABG may show hypoxemia (PaO2 ≤ 60mmHg) and/or Hypercapnia (PaCO2 >45mm HG) but no acidosis			emia apnia	Signs and symptoms: Dyspnea RR, HR, SaO2m and CRP is same as moderate ABG shows new onset/worsening hypercapnia AND acidosis (PaCO2 > 45mmHg and pH <7.35)			
DECAF							_		
DECAF Score[3]		0		•	1		2	3	
Dyspnoea (MRC)		1–3		!	5		4		
Eosinopenia (<0.05 × 10^9/L)		No `		Y	es				
Consolidation on CXR		No N		Y	es				
Acidaemia (pH < 7.30)		No Y		Y	es				
Atrial fibrillation		No Yes		es					
Total Score:	Dispos	ition: □ Discharge H	Home: 0-	1 □ Conside	r Admission	to Floor: 2	□ Consider escala	ted care/monitoring: 3-6	
Assessed by (print)	Des	signation	Sig	nature			Date/Time (do	d/mm/yyyy hhmm)	



MC	MONITORING / LABS						
	Continuous SpO ₂		CBC		SARS-CoV-2 PCR		
	Continuous Heart Rate Monitoring		Basic Metabolic Panel		Influenza PCR		
	Blood Pressure Monitoring		Comprehensive Metabolic Panel		Chest X-Ray		
	ABG		Sputum Culture		D-dimer		
	VBG		Respiratory Panel		Troponin		

ACUTE MANAGEMENT (first hour)

PHARMACOLOGIC THERAPY - Bronchodilators

	MILD-MODERATE	SEVERE Requiring HFNO/NIV
	□ Salbutamol pMDI 100mcg/puff with spacerPuffs q20 min x3 PRN Shortness of breath	□ Salbutamol pMDI 100mcg/puff with SpacerPuffs q20min x3 PRN Shortness of breath
	☐ Ipratropium pMDI 20mcg/puff with spacer Puffs x3	☐ Ipratropium pMDI 20mcg/puff with spacer chamber Puffs x3
OPTIONS	 For patients unable to coordinate breaths or generate adequate inspiratory flow, VMN should be considered [23,26,31] pMDI should be delivered with a spacer to increase deposition 	 For patients unable to coordinate breaths or generate adequate inspiratory flow, VMN should be considered [23,26,31] pMDI should be delivered with a spacer to increase deposition
Ē		
	☐ Salbutamol via VMN + Aerosol Reservoir mg q20min x3 PRN Shortness of breath	□ Salbutamol via VMN in-line via HFNO or NIVmg q20min x3 PRN Shortness of breath
DELIVERY	☐ Ipratropium Bromide via VMN + Aerosol Reservoir 0.5 mg x1 PRN Shortness of breath	☐ Ipratropium Bromide via VMN in-line via HFNO or NIV 0.5 mg x3 PRN Shortness of breath
ATION	☐ Salbutamol via JN mg q20min x3 PRN Shortness of breath	☐ Salbutamol via JN mg q20min x3 PRN Shortness of breath
MEDICATION	☐ Ipratropium Bromide via JN 0.5 mg x3 Shortness of breath	☐ Ipratropium Bromide via JN 0.5 mg x 3 Shortness of breath
~	JN may be inferior to VMN (clinical outcomes & deposition) and not superior to pMDI I1 2:311	For patients on HFNO or NIV: [25] - not recommended to disrupt oxygen delivery to deliver aerosol treatment - concurrent aerosol treatment with mask/mouthpiece not recommended (in-line delivery recommended) - adding flow to the circuit via JN is not recommended due to changes to FIO2 and nuisance alarms

Infection Prevention

In patients with respiratory infections, it is preferred to use pMDI due to risk of secondary exposure. [33]

If nebulizer is needed due to patient inability to coordinate breaths, or lack of inspiratory flow, VMN with mouthpiece & filter or in-line with viral filter is preferred over JN to reduce the risk of secondary transmission. [33]

In patients receiving HFNO it is recommended to place a surgical mask over cannula to reduce the risk of transmission. [33]

Environmental Sustainability

VMN + Ultra: Enables continuous delivery in-line with HFNO/BiPAP; reusable; Less plastic waste than disposable jet nebulizers [6-10] [11,12]

pMDI + Spacer: Lower plastic waste and energy use vs disposable jet nebulizers; reusable spacers last months.x



Antibiotics

Initiate antibiotics if any of the following are present:

- Increased dyspnea, sputum volume, and sputum purulence (all three)
- sputum purulence and either dyspnea or sputum volume
- Requirement for mechanical ventilation (invasive or noninvasive)

Route	Medication/dose	Select	Continuation
Oral	Amoxicillin/clavulanate*	□ 875mg/125mg X1	□ 875mg/125mg q12h X5 days
Oral	Doxycycline	□ 200 mg X1	□ 100 mg BID X5 days
Oral	Azithromycin	□ 500 mg X1	☐ 500 mg Qday X4 days
Oral	Levofloxacin*	□ 750 mg X1	☐ 750 mg Qday X4 days
IV	Ceftriaxone	□ 2000 mg X1	□ 1gm qday X4 days
IV	Cefepime*	□ 2000 mg X1	□ 2000mg q8h
IV	Levofloxacin*	□ 750 mg X1	□ 750 mg Qday X4 days
IV	Piperacillin/Tazobactam*	□ 4.5g X1	□ 4.5gm q8h X4 days

^{*}Should be dose adjusted based on renal function

- Antibiotics should be chosen based on local resistance patterns
- Antibiotics should be continued for no more than 5 days for simple COPD
- Critically ill patients should be dosed with the most aggressive regimen possible

Ordering Prescriber (print)	Designation	Signature	Date/Time (dd/mm/yyyy hhmm)

^{*}GOLD guidelines recommend 40mg of prednisone or equivalent. Methylprednisolone 4mg=prednisone 5mg



REASSESSMENT / MAINTENANCE (post 1-hour)

Reassessed	COPD	Severity	/ :

PHARMACOLOGIC THERAPY (Continuation) - Bronchodilators

(continued device selection should be based on clinical considerations from the acute table)

	MILD - MODERATE			SEVERE				
	SCHEDULED DOSES							
	☐ Salbutamol pMDI 100mcg/puff	with spacerPuffs	q6h	☐ Salbutamol pMDI 10	00mcg/puff with SpacerPuffs q6h			
	☐ Ipratropium pMDI 20mcg/puff w	vith spacerPuffs o	q6h	☐ Ipratropium pMDI 20	Omcg/puff with spacerPuffs q6h			
SNOI	☐ Salbutamol via VMN + Aerosol Reservior mg q6h			☐ Salbutamol via VMN in-line via HFNO or NIVmg q6h				
Y OP1	☐ Ipratropium Bromide via VMN + q6h	· Aerosoi Reservoir	0.5 mg	☐ Ipratropium Bromide via VMN in-line via HFNO or NIV 0.5 mg q6h				
LIVER	☐ Salbutamol via JN mg q6h							
N DE	☐ Ipratropium Bromide via JN 0.5	mg q6h						
ATIC	AS-NEEDED DOSES							
MEDICATION DELIVERY OPTIONS	☐ Salbutamol pMDI 100mcg/puff q1h PRN Shortness of breath	with spacerPเ	☐ Salbutamol pMDI 100mcg/puff with Spacer Puffs q1h PRN Shortness of breath					
	☐ Salbutamol via VMN + Aeroso Shortness of breath	I Reservoirmg q1	hr PRN	☐ Salbutamol via VMN + Aerosol Reservoir mg q1hr l Shortness of breath				
	□ Salbutamol via JN mg q20i breath ×3	min PRN Shortness o	f					
Orde	Ordering Prescriber (print) Designation Signate			ure	Date/Time (dd/mm/yyyy hhmm)			



RESPIRATORY SUPPORT / SUPPLEMENTAL OXYGEN

Target $SpO_2 \ge 94\%$ (peds) / $\ge 92\%$ (adult)
Room Air
Nasal CannulaL/min
HFNC:L/min (Peds: 1.5–2 L/kg/min; Adults: 30–60 L/min)
- Inline Aerogen Ultra VMN for bronchodilator delivery
NIV/BiPAP: IPAP/ EPAP
- Inline Aerogen Ultra VMN via T-piece or mask adapter

Considerations

- HFNO with Cannula (Moderate)
 - o In-line with Fisher&Paykel Airvo2 or 3 in combination with the Airvo Neb humidifier adaptor
 - o In-line with the Vapotherm HVT 2.0 Aerosol Adapter
 - If High-flow Nasal Oxygen is being delivered via standalone humidification Aerogen should be on the Dry side of the humidifier at the inlet
 - Higher delivery occurs when the patients inspiratory flow is matched to or greater than flow from the HFNO device (consider reducing the flow of the highflow device)
- Optimal Placement for NIV (Severe)
 - Single Limb Circuit: Between a non-vented mask and the patient side of the leak port (non-vented masks not recommended).
 - Dual Limb Circuit: Optimal position would 15cm back from the Wye at the inspiratory limb or between the Wye and the patient, and pre-humidifier
- Reassessment
 - Response to NIV should be monitored at least hourly
 - Follow institutional guidelines for need of escalation

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	nsider discharge in patients fitting into the mild category: Peak flow > 80% predicted O2>95%, RR <20, n	Ю
desati	ration during walk test and able to use pMDIs at home.	
□ Ac	mit to Ward	
□ Ad	mit to ICU	

DEFINITION	DEFINITIONS				
pMDI	Pressurized Metered Dose Inhaler				
VMN	Aerogen Solo Vibrating Mesh Nebulizer				
Ultra	Aerogen Ultra Aerosol Reservoir with aerosol mask or valved mouthpiece				
HFNO	High-Flow Nasal Oxygen				
NIV	Non-Invasive Ventilation				
DECAF	Dyspnea, Eosinopenia, Consolidation on chest x-ray, Acidemia (pH<7.3), Atrial fibrillation				
BiPAP	Bilevel Positive Airway Pressure				
EPAP	Expiratory Positive Airway Pressure				
FiO2	Fraction of inspired Oxygen				



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