

Consultant in Emergency Medicine North Manchester General Hospital ED

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Introduction

- ► Common cause of ED presentations; 597 exacerbations of COPD at NMGH from Oct 07-Oct 08
- ▶ 900 000 in UK diagnosed with COPD, 450 000 not yet diagnosed, increasing in ♀, static in ♂
- ▶ 30 000 deaths in 1999. Increasing ♀, reducing ♂ last 30yrs
- ▶ Guidelines on COPD from BTS 1997, then NICE 2004
- ► Airflow obstruction with reduced FEV_I (80%) and reduced FEV_I/FVC ratio <0.7

Pathophysiology I

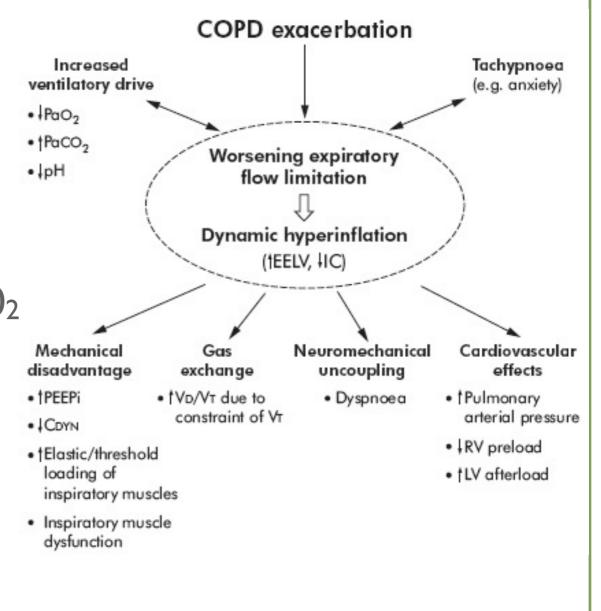




- Smoking accounts for 80-90%; risk in α₁-antitrypsin deficiency/occupation
- Early bronchial epithelium metaplasia, incr gland number & size
- Later acute & chronic inflammation/acinar expansion, destruction and coalescence. Mucous
- Reduced elasticity and surface area-hyperinflation
- ► Impedance to (expiratory. ++) airflow, incr. resistance/small caliber

Pathophysiology II

- Chronic hypoxia leads to polycythaemia, pulmonary hypertension and cor pulmonale
- May retain CO₂, central receptors lose their respiratory drive and rely on hypoxia. Giving high flow O₂ may then lead to worsening CO₂, (chronic bronchitic blue bloaters)
- Some have normal CO2, low/ normal 02, (emphysematous 'pink puffers')



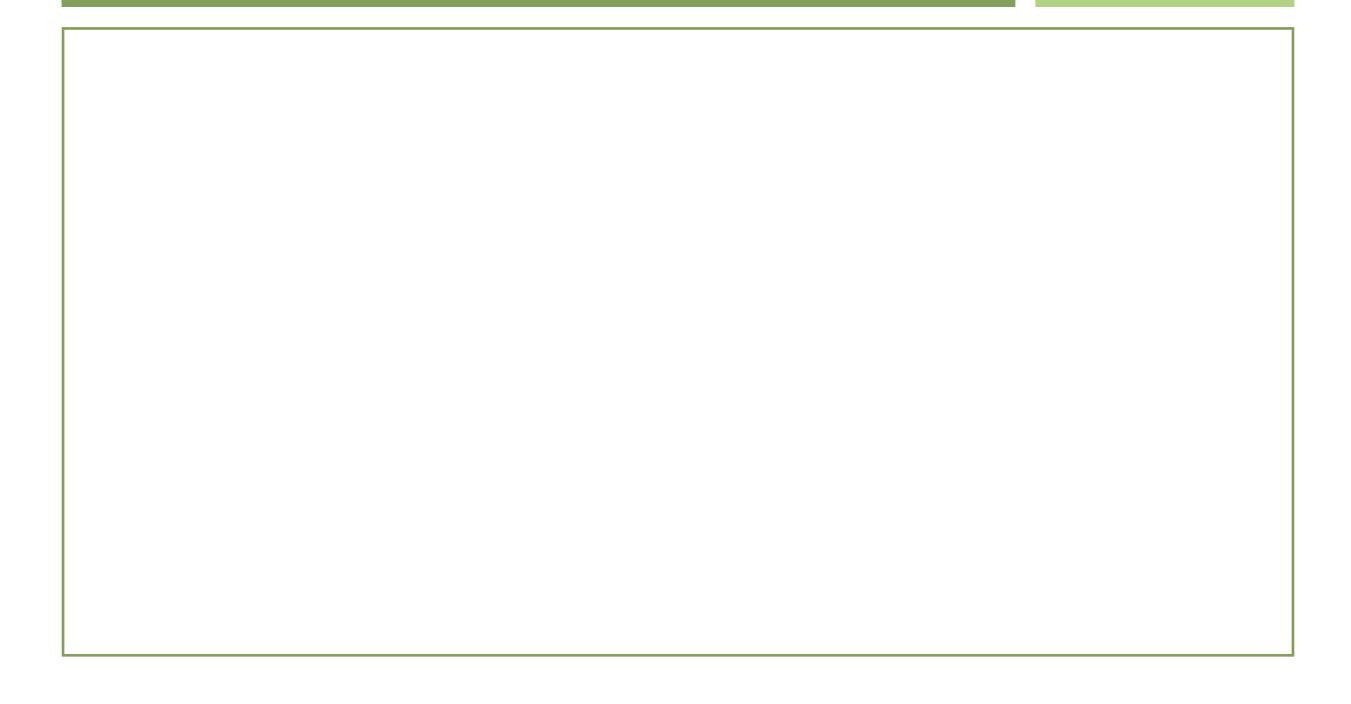
Clinical Presentation

- Smoking history
- Gradual onset of increasing exertional dyspnoea
- ► Cough, >3 consecutive months, >2 consecutive years
- ► Tachypnoea, hyperinflation, wheeze, creps
- ► Flattened diaphragms on CXR. May have infection, chronic changes, pneumothorax, bullae
- ► Exacerbation: respiratory distress, altered GCS: ? infection, ?resp failure, ?Cor pulmonale, ? ↓ CO2
- ► ABGs....

pH=7.34	pH=7.10	pH=7.5
p0 ₂ =8.5 RA	p0 ₂ =7 RA	p0 ₂ =10 RA
pC0 ₂ =9.5	pC0 ₂ =15	pC0 ₂ =3.5
HC0 ₃ =35	HC0 ₃ =33	HC0 ₃ =23

	ABGs	2
pH=7.34	2 pH=7.10	3 pH≡7.5
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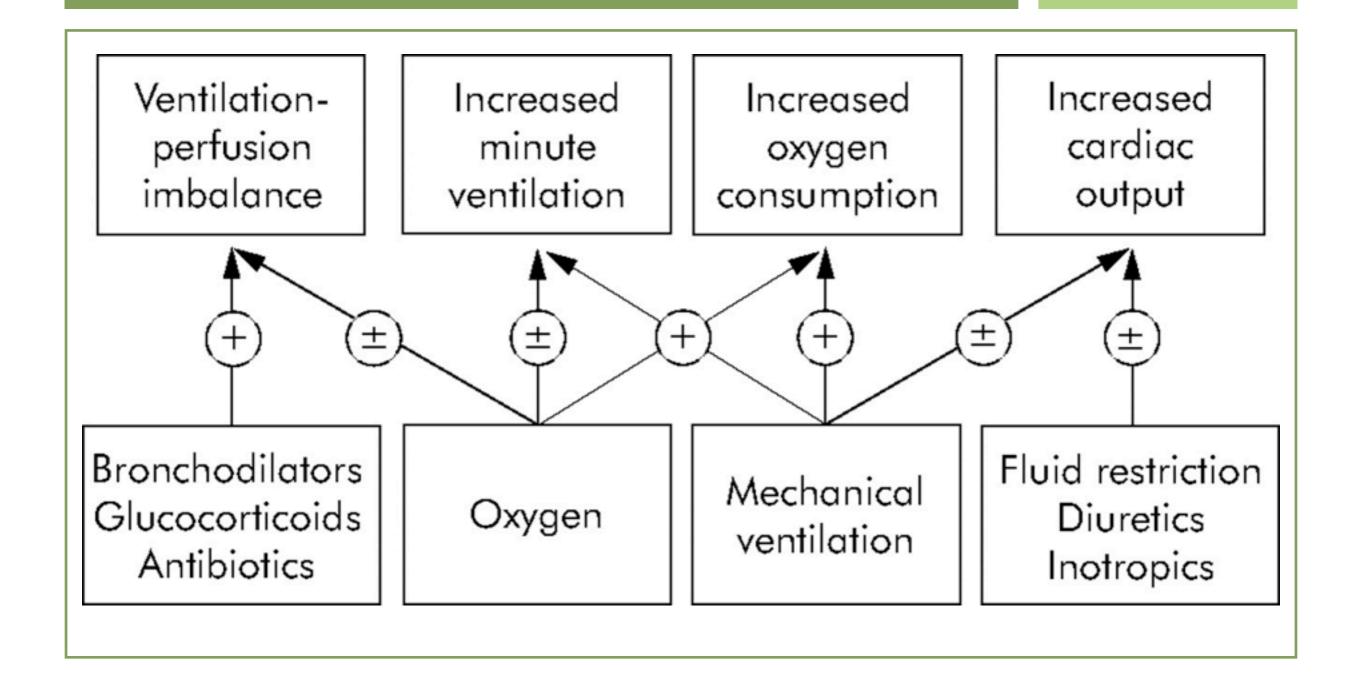
On-going treatment GOLD guidelines



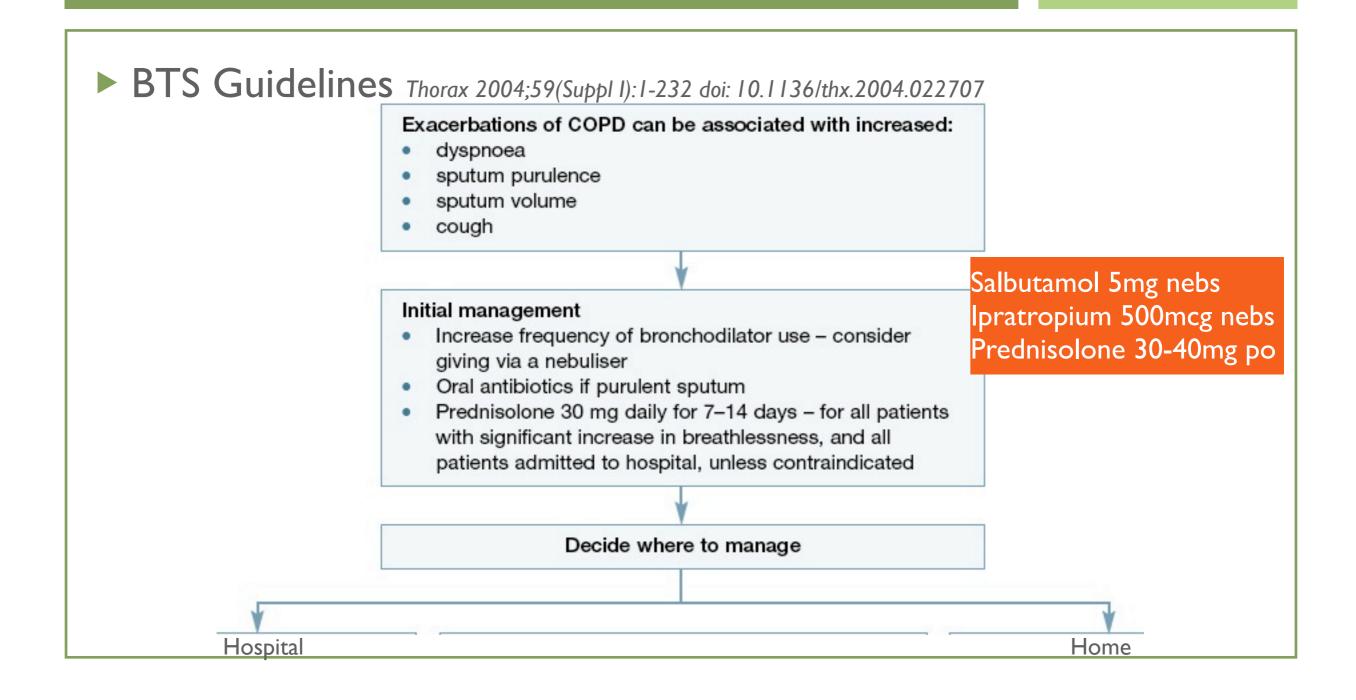
Severity	Test Values	Clinical Management
0. At Risk	Normal spirometry Chronic symptoms (cough, sputum production)	
I. Mild COPD	FEV ₁ /FVC <70% predicted	■ Short acting bronchodilator when needed
	FEV ₁ >80% predicted With or without chronic symptoms (cough, sputum production)	
II. Moderate COPD	FEV ₁ /FVC <70% predicted 30%< FEV ₁ <80% predicted (IIA - 50%< FEV ₁ <80% predicted IIB - 30%< FEV ₁ <80% predicted) With or without chronic symptoms (cough, sputum production, dyspnoea)	 Regular treatment with one or more bronchodilators Inhaled steroids if significant symptoms and lung function response indicates reversible element to disease (or in IIB if repeated exacerbations) Pulmonary rehabilitation
III. Severe COPD	FEV ₁ / FVC <70% predicted FEV ₁ <30% predicted or FEV ₁ <50%predicted plus respiratory failure or clinical signs of right heart failure	 Regular treatment with one or more bronchodilators Inhaled steroids if reversible element, or if repeated exacerbations Pulmonary rehabilitation Treatment of complications, such as heart failure Long term oxygen therapy if respiratory failure Consider surgical treatments

COPD Exacerbation

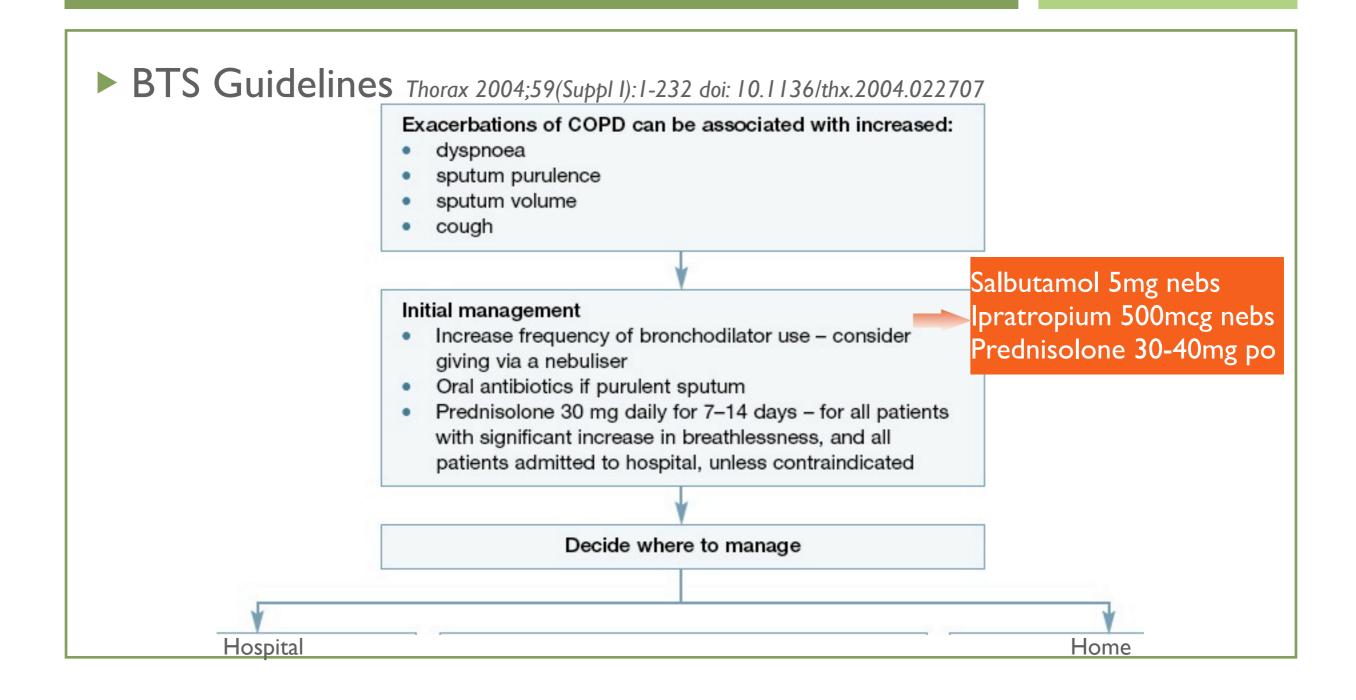
Thorax 2006;61:535-544;



COPD Exacerbation



COPD Exacerbation



Hospital

Investigations

- Chest X-ray
- Arterial blood gases (record inspired oxygen concentration)
- ECG
- Full blood count and urea and electrolytes
- Theophylline level if patient on theophylline at admission
- Sputum microscopy and culture if purulent

Further management

- Give oxygen to keep Sao₂ above 90%
- Assess need for noninvasive ventilation:
 - consider respiratory stimulant if NIV not available
 - assess need for intubation
- Consider intravenous theophyllines if poor response to nebulised bronchodilators

Consider hospital-at-home or assisted-discharge scheme

Prior to discharge

- Establish on optimal therapy
- Arrange multidisciplinary assessment if necessary

Factors to consider when deciding where to manage patient

Factor	Favours treatment at home	Favours treatment in hospital
Able to cope at home	Yes	No
Breathlessness	Mild	Severe
General condition	Good	Poor/ deteriorating
Level of activity	Good	Poor/confined to bed
Cyanosis	No	Yes
Worsening peripheral oedema	No	Yes
Level of consciousness	Normal	Impaired
Already receiving LTOT	No	Yes
Social circumstances	Good	Living alone/ not coping
Acute confusion	No	Yes
Rapid rate of onset	No	Yes
Significant comorbidity (particularly cardiac and insulin dependent	35.50	
diabetes)	No	Yes
Sao ₂ < 90%	No	Yes
Changes on the chest radiograph	No	Present
Arterial pH level	≥7.35	<7.35
Arterial Pao ₂	≥7 kPa	<7 kPa

Home

Investigations

- Sputum culture not normally recommended
- Pulse oximetry if severe exacerbation

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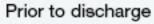


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NIV



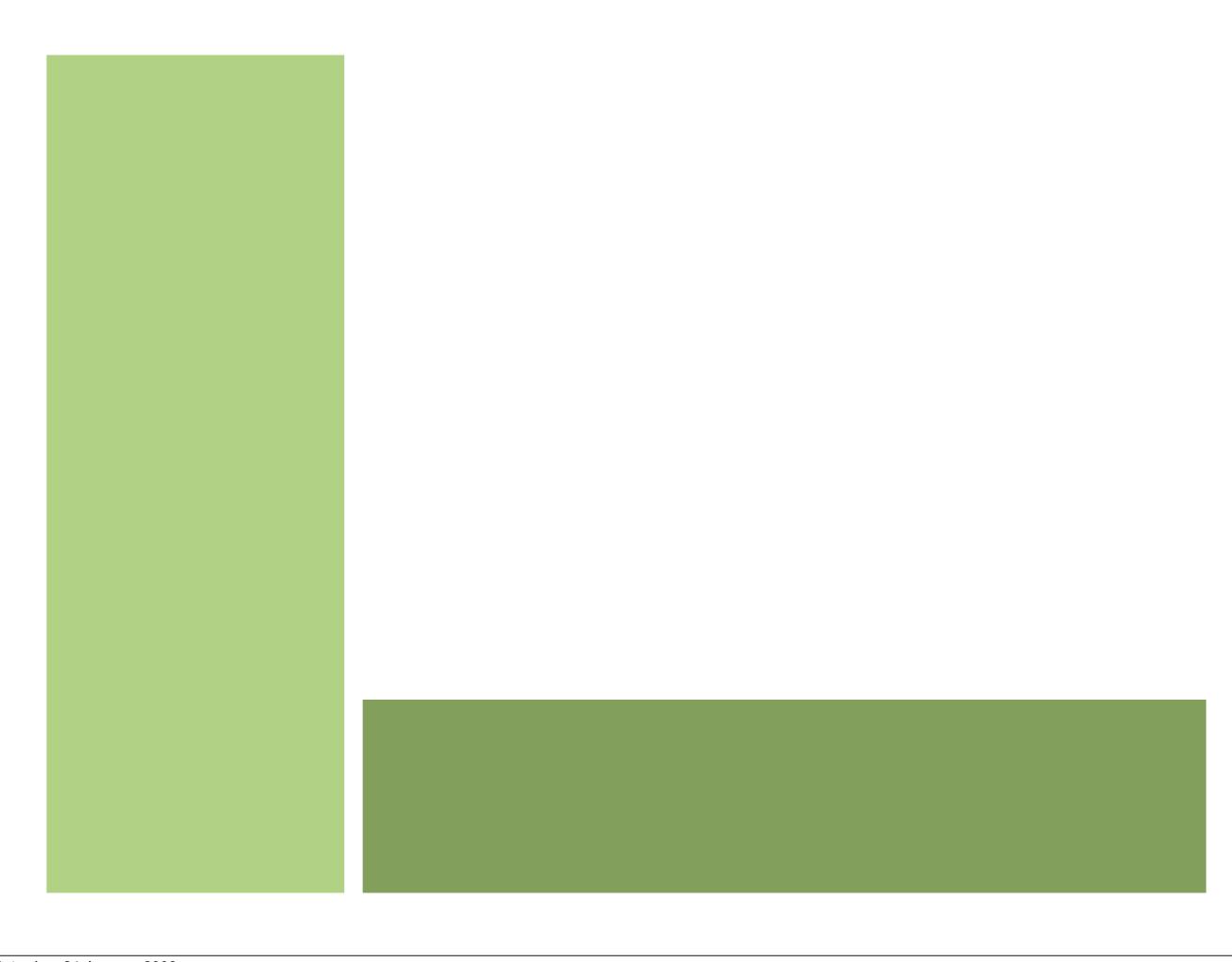
- In patients who are tiring, poor ABGs (pH <7.35, pCO₂ >6, pO₂ <8), poor response to therapy</p>
- ► Have to be making spontaneous effort
- ► Provide recruitment of collapsed (mucous plugged, atelectatic) alveoli
- ▶ Reduce work of breathing by helping with compliance, reduced muscle work, help overcome auto PEEP by providing PEEP, reduce pulmonary oedema (cor pulmonale), and provide high 0₂ and help reduce C0₂
- ► Waters circuit can provide temporary relief until NIV machine

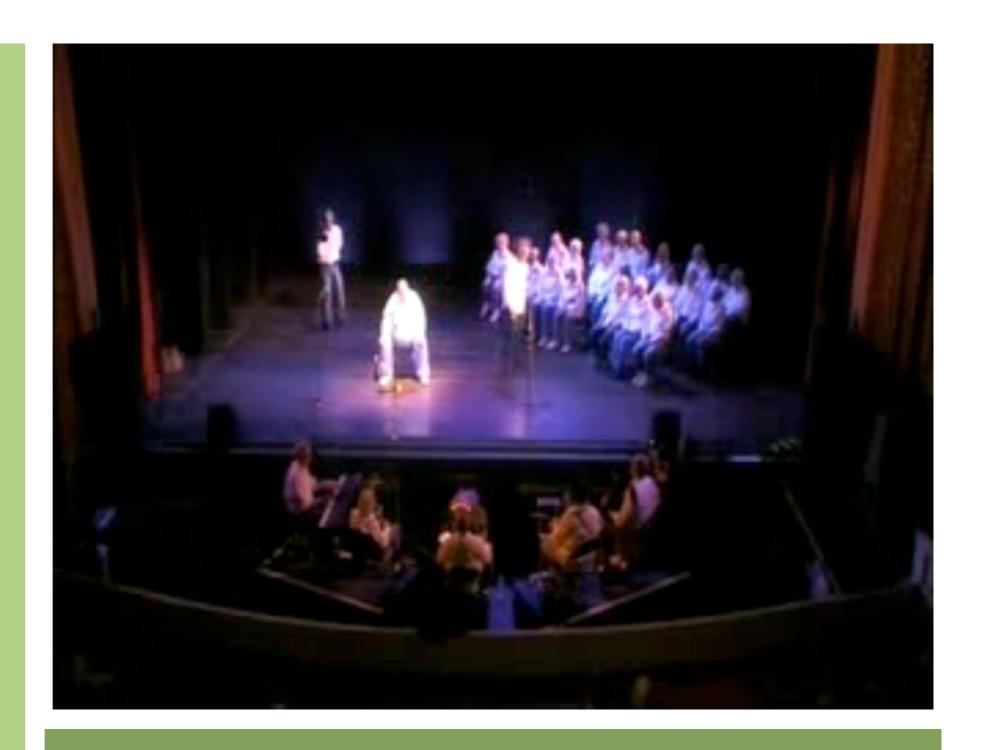
ARASTeam

- Only with previous diagnosed COPD on spirometry (already known to chest team/ARAS)
- Are able to do home follow up and support to avoid admission (if possible)
- Mostly referral from MEU by medics

Inclusion Criteria	Yes	No
Has clinically diagnosed COPD, and is currently exacerbating		
Min Mental Score >6		
pH > 7.35	16	
Systolic BP > 100mmHg		
Oxygen saturation >90% (with supplementary oxygen if needed at 2-4 litres or 28% via mask)		
Has adequate social support and telephone		

Exclusion Criteria	Yes	No
Pneumothorax/cardiac signs on CXR		
CURB score >2		
Pulse Rate <60/min or >110/min		
Concomitant medical conditions ie. LVF, angina requiring inpatient stay		
Acute ECG changes		
Requires full time nursing		
Has refused to consider home care		





Questions?

Summary

- ▶ Common
- Worsening with increase sputum & symptoms
- Nebulise salbutamol/ipratropium and steroids
- Least 02 flow (28%, 35% etc) to keep sats at about 90%
- ► NIV for in extremis patients