LOWER RESPIRATORY TRACT INFECTIONS Acute Exacerbation of COPD

(CKS Chronic obstructive pulmonary disease)

Clinical features

An acute exacerbation of chronic obstructive pulmonary disease (COPD) is a sustained worsening of a person's symptoms from their usual stable state (beyond normal day-to-day variations) which is acute in onset. Commonly reported symptoms include:

- Increased breathlessness.
- Increased cough.
- Increased sputum production and change in sputum colour.

Core pathogens

Respiratory viruses (30%), bacterial (30-50%) – *Streptococcus pneumoniae, Haemophilus influenzae* (amoxicillin sensitive and resistant strains), *Moraxella catarrhalis*, and atypical pathogens such as *Mycoplasma pneumoniae* and *Chlamydophilia pneumoniae*.

Management

Nottinghamshire COPD Self-management plan here

Viral infections may cause acute exacerbations, but if purulent sputum is being produced bacterial infection is possible.

If there are no contraindications, consider oral corticosteroids for people with a significant increase in breathlessness that interferes with daily activities.

- Offer 30 mg oral prednisolone once daily for 5 days discuss adverse effects of prolonged therapy.
- Consider the need for bone protection for people requiring frequent courses of oral corticosteroids (≥3 courses per year).

Antibiotics are most valuable in patients with purulent sputum *and* increased shortness of breath **and/or** increased sputum volume.

Consider the need for an antibiotic taking into account:

- Severity of symptoms (particularly sputum colour changes, increase in volume or thickness beyond normal).
- Risk of complications.
- Previous sputum culture and susceptibility results (send sputum sample if possible).
- Risk of antimicrobial resistance and current antibiotic prophylaxis (treatment should be with an antibiotic from a different class).

NICE recommend as part of self-management that patients are given a course of antibiotics and oral corticosteroids to keep at home and commence if their sputum becomes purulent (see <u>Nottinghamshire guidance for prescribers on</u> <u>COPD Exacerbation Rescue Medication Pack</u>).

Risk factors for antibiotic resistant organisms include:

- Severe COPD
- Co-morbid disease
- Frequent exacerbations and/or hospital admissions
- Multiple courses of antibiotics, or antibiotics within last 3 months.
- Previous resistant organisms in sputum culture.

Treatment options for adults 18 years and older:

Antibiotic ¹	Dosage	Duration
Empirical treatment g	uided by most recent sputum culture and s	uscentibilities
Einpinear treatment guided by most recent sputum culture and susceptionities		
Amovicillin	Adult: 500mg three times a day	5 days
Amoxiciiin	Addit. Sooning three times a day	Juays
Alternative first line choices (if penicillin contraindicated or not tolerated)		
Doxycycline ²	Adult: 200mg day one, then 100mg	5 days
(Not suitable in pregnancy)	once daily.	
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Clarithromycin ³ (if penicillin and	Adult: 500mg twice a day	5 days
doxycycline not suitable). Not in		
patients taking azithromycin		
prophylaxis.		
Second line choice – If there is no improvement in symptoms on first choice taken for at least 2 to 3 days send a		
sputum sample for culture and susceptibility testing. Use alternative first line (from a different class) if suitable.		
If higher risk of treatment failure, treat options according to sputum culture:		
guided by microbiology sensitivities		
Dive	Adult: 625mg three times a day	5 days
Amovicillin	Adults F00mg three times a day	
Amoxiciiiii	Adult: 500mg tillee tilles a day	
(UNLY In reported sensitivity to Co-		
Co-trimoxazole	Adult: 960mg twice a day	5 days
	с ,	
Levofloxacin	Adult: 500mg once a day	5 days
	(ONLY Increase frequency to twice a day	
Note: Fluoroquinolones should only be	if reported sensitivity to Levofloxacin is	
used when other antibiotics are	"DS" or "I" **)	
inappropriate. If a penicillin allergy is		
recorded, the exact nature of the		
whether other beta lactams (e.g.		
cephalosporins) have been previously		
tolerated.		
Fluoroquinolones can cause long-lasting		
(up to months or years), disabling, and		
potentially irreversible side effects,		
sometimes affecting multiple systems,		
organ classes, and senses. Please refer		
nere for further information on MIRRA		
Increased risk of tendon damage if co-		
prescribed corticosteroid and		
fluoroquinolone.		
** "DS" or "I" = dose dependent susceptible. This means there is a high likelihood of therapeutic success if antibiotic		
exposure is optimised by using higher doses or increasing dosing frequency.		
Microbiology interpreting Sensitivity Results		
¹ See <u>BNF</u> and <u>BNFC</u> for appropriate use and dosing in specific populations, e.g., hepatic, renal impairment, pregnancy,		
breastfeeding.		
² Doxycycline is not suitable for pregnant women.		

³Withhold statins whilst on clarithromycin course. Avoid if the patient is at risk of QTc prolongation.