# Derbyshire Medicines Management, Prescribing and Guidelines DERBYSHIRE PRIMARY CARE FORMULARY

**Chapter 3: RESPIRATORY SYSTEM** 

Updated: March 2025

# The following prescribing guidelines are relevant to the respiratory chapter and can be found here

- Children's referral guideline for SLIT- Grazax and Acarizax (Derby Children's Hospital)
- Anaphylaxis treatment for adults and children
- Asthma management in adults/ Asthma management in children
- Chronic Obstructive Pulmonary Disease guideline
- Greener Inhaler Prescribing guidance
- Oxygen guideline

#### Relevant Resources

- · COPD detailing aid
- Greener inhaler choice: Flowchart/ Be Greener and Breathe Better patient information leaflet
- Stepping-down combination asthma inhaler therapy: Adults over 17 years of age
- Optimisation of inhaled corticosteroid (ICS) in COPD
- Adult Asthma Diagnosis Algorithm North Derbyshire, South Derbyshire
- Respiratory action plans Asthma & Lung UK
- RightBreathe inhaler information/training videos
- Inhaled corticosteroid equivalent doses

# Reducing the carbon impact of inhalers

Metered dose inhalers (MDIs), also known as pressurised metered dose inhalers (pMDIs), including breath-actuated MDIs, contain propellants known as hydrofluorocarbons (HFCs) which are powerful greenhouse gases and can contribute to global warming. Dry powder inhalers (DPIs) and soft mist inhalers (SMIs) such as Respimat do not contain propellant, so they have a lower carbon footprint than other inhalers.

NHS Derby and Derbyshire ICB/System partners support the prescribing of inhalers with a reduced carbon footprint such as DPIs and SMIs, wherever clinically appropriate and acceptable to the patient, in accordance with <a href="NHS targets">NHS targets</a>. Medication reviews, Structured Medication Reviews or planned Asthma Reviews taking place in primary care should consider moving or facilitating patients to lower carbon options where it is clinically appropriate to do so. See <a href="Greener inhaler prescribing guidance">Greener inhaler prescribing guidance</a>.

All formulary dry powder inhalers contain lactose and are contraindicated in patients with IgE mediated hypersensitivity to lactose or milk proteins. Refer to The SPC for full prescribing information

MHRA July 2018 Pressurised metered dose inhalers (pMDIs): risk of airway obstruction from aspiration of loose objects. Remind patients to check and remove the mouthpiece cover properly before inhaling a dose and to shake the inhaler to remove loose objects that may have become trapped in the inhaler during storage.

#### 3.1 Bronchodilators

#### 3.1.1 Adrenoreceptor agonists

0:1:1 Adictionaceptor agonists	'		
Short acting beta agonist (SABA)	Traffic Light Classification	Licenced	Additional information
Salbutamol preparation			The metered dose inhaler (MDI) is usually
100 micrograms CFC Free inhaler	GREEN	Adults and	the most cost-effective delivery device for
(MDI) ( <b>Salamol</b> )		children ≥4 years	salbutamol and should be considered for
(iii2i) (Galamei)		of age	those patients requiring a SABA and as a
100 micrograms salbutamol	GREEN		matter of routine for all paediatric patients.
Easyhaler (DPI)			Salamol MDI is the preferred choice
	GREEN		salbutamol inhaler due to lower carbon
200micrograms Ventolin			footprint compared to other salbutamol
Accuhaler (DPI)			MDIs. Patients who are unable to use a
/ tesariater (21 1)	GREEN		standard salbutamol MDI may find a dry
100 microgramBreath-actuated	GILLIN		powder device or breath-actuated inhaler
CFC free inhaler (MDI) (Salamol)			more acceptable. The brand must be
Ci Ci i de ililialei (MDI) (Salaliloi)			

Nebuliser solution 2.5mg/2.5ml			specified to avoid confusion with these devices.  Nebulised bronchodilators should not be prescribed unless a formal nebuliser assessment has been carried out. Do not use 5mg salbutamol nebules for COPD, as evidence does not support doses above 2.5mg. Nebulisers should be used with extreme caution in children and only under the care of a respiratory paediatrician (MHRA Aug 2022).
Long-acting beta agonist (LABA)	Traffic Light Classification	Licenced	Additional information
Formoterol preparations  12 micrograms Easyhaler (DPI)	GREEN	Adults and children ≥6 years of age	First-line LABA
12 micrograms CFC free inhaler (MDI) (Atimos Modulite)	GREEN	Adults and children ≥12 years of age	Alternative first-line LABA for patients requiring an MDI
Salmeterol preparations  • 25 micrograms CFC Free inhaler (MDI) (Soltel)	GREEN 2 <sup>nd</sup> line LABA	Soltel - Adults and children >12 years of age Serevent - Adults and children ≥4 years of age	Soltel is currently the formulary choice N.B. All brands apart from Serevent contain soya lecithin – contra-indicated in peanut or soya allergy). If the patient has a soya or peanut allergy, then prescribe as the brand Serevent

#### 3.1.2 Antimuscarinic bronchodilators

Short acting antimuscarinic bronchodilators (SAMA)	Traffic Light Classification	Licenced	Additional information
<ul><li>Ipratropium</li><li>20 microgram inhaler CFC free (MDI)</li></ul>	GREEN	COPD and asthma Adults and children (including those <6 years of age)	Asthma – MDI is not recommended by SIGN/BTS or NICE for routine asthma management. Nebulised solution may be used as an add-on treatment for a severe asthma attack.

- 1. Ipratropium nebuliser solution for COPD should only be used in severe COPD patients [FEV1<30%] after initiation by a specialist.
- 2. Long acting muscarinic antagonist (LAMA) monotherapy is no longer recommended first line in the management of COPD- see <u>local guidance</u>. When initiating a LAMA, take care to ensure the SAMA is stopped. All the single component LAMAs are available in combined LABA/LAMA inhalers but some patients may need a single component LAMA therapy or may not wish to switch.
- 3. Tiotropium (GREY) is the preferred LAMA for existing stable patients on single component LAMA treatment.

Long-acting muscarinic antagonists (LAMA)	Licensed	Traffic Light Classification	Additional Information
Tiotropium  (Preferred LAMA for existing stable patients on single component LAMA treatment.)	COPD	GREY	18 microgram inhalation powder (DPI): Advise patients to NEVER insert the capsule directly into the mouthpiece, follow instructions provided with the inhaler.  Trokide is the most cost effective brand. Trokide inhaler device should be replaced every 3 months.  Tiogiva inhaler device should be replaced every 6 months.

	Asthma	GREY after consultant/specialist initiation	2.5microgram inhalation solution cartridge CFC free (soft mist inhaler) (Spiriva Respimat):
			Respimat inhaler is a re-usable device with cartridge. Each re-usable inhaler may be used with up to six cartridges.
Glycopyrronium	COPD	GREY 2 <sup>nd</sup> line LAMA	Seebri Breezhaler: In patients with severe renal impairment (eGFR below 30mL/min/1.73m²), including those with end-stage renal disease requiring dialysis. These patients should be monitored closely for potential adverse reactions.
Umeclidinium	COPD	GREY 2 <sup>nd</sup> line	ONCE daily
Aclidinium	COPD	GREY 3 <sup>rd</sup> line	TWICE daily  Each delivered dose contains 375mcg of aclidinium bromide, equivalent to 322mcg.

MHRA Feb 2015 prescribers should take the risk of cardiovascular side effects into account when prescribing inhaled tiotropium to patients with certain cardiac conditions, who were excluded from clinical trials of tiotropium. This also applies to all LAMAs (See individual SPCs for further details detail).

Combination inhalers N.B. Prescribe by brand

LABA/LAMA combinations	Traffic Light Classification	Licenced	Dosing	Additional information
Tiotropium and olodaterol (Spiolto Respimat) (SMI)  Glycopyrronium and Indacaterol (Ultibro Breezhaler) (DPI)  Umeclidinium and vilanterol (Anoro Ellipta) (DPI)  Aclidinium and formoterol (Duaklir Genuair) (DPI)  Glycopyrronium + formoterol (Bevespi Aerosphere) (MDI)	GREEN 1st line LABA /LAMA  GREEN 1st line LABA /LAMA	COPD COPD COPD COPD	OD OD BD BD	Respimat inhalers are re-usable device with cartridge. Each re-usable inhaler may be used with up to six cartridges.  Choice should be based on patient tolerance, ease of use, and environmental impact of the inhaler device.  LABA/LAMA combination inhaler is recommended for COPD patients who remain breathless or have exacerbations despite SABA or SAMA treatment and present with no asthmatic features or features suggestive of steroid responsiveness.

Roflumilast is GREY- specialist initiation. Roflumilast is a phosphodiesterase type-4 inhibitor with anti-inflammatory properties. It is used as an add-on to bronchodilator therapy in adults with severe COPD with chronic bronchitis as per NICE TA461. Ongoing GP prescribing and care of patients on roflumilast should only be considered if patient is stable and free from adverse reactions, after a minimum of 3 months roflumilast treatment under the respiratory specialist. For more details see local COPD guideline.

#### 3.1.3 Theophylline

### Theophylline SR tablets (Uniphyllin Continus) 200mg, 300mg, 400mg

- 1. Common interactions for theophylline which increase clearance and it may therefore be necessary to increase dosage to ensure therapeutic effect include barbiturates, carbamazepine, lithium, phenytoin, rifampicin, primidone, ritonavir. (See SPC for full details)
  - The following reduce clearance, and a reduced dosage may therefore be necessary to avoid side-effects: allopurinol, cimetidine, macrolide antibiotics (e.g., erythromycin), furosemide, and oral contraceptives. (See SPC for full details)
- 2. Smoking can increase theophylline clearance and increased doses of theophylline are therefore required; dose adjustments are likely to be necessary if smoking started or stopped during treatment.
- 3. Plasma-theophylline concentration is increased in heart failure, hepatic impairment, and in viral infections. Plasma-theophylline concentration is decreased in smokers and by alcohol consumption. Differences in the half-life of theophylline are important because the toxic dose is close to the therapeutic dose.
- 4. See SPS for monitoring information.

#### 3.1.5 Peak flow meters, inhaler devices and nebulisers

Appropriate peak flow meter

Appropriate spacer device

EasyChamber Spacer Volumatic

## See Right Breathe website for information on spacer compatibility.

- 1. Standard-range peak flow meters are suitable for both adults and children; low-range peak flow meters are appropriate for severely restricted airflow in adults and children.
  - a. Standard = 60-800L/minute
  - b. Low = 30-400L/minute
- 2. EasyChamber fits most MDIs and is compatible with formulary choices although may not be licensed, refer to Right Breathe's website for more information.
- 3. Follow usage and cleaning instructions supplied with the spacer. Spacer devices should be allowed to air dry to prevent the build -up of static which can alter airflow. Most spacers should be replaced every 12 months and may be recyclable, however this varies from brand to brand. See Right Breathe for further information.
  - <u>Drug Safety Update</u>, July 2008 **Spacers should not be regarded as interchangeable** patients whose asthma is well-controlled and who are using a spacer should always use the same type of spacer and not switch between spacers. Different spacers may deliver different amounts of inhaled corticosteroid, which may have implications for both safety and efficacy.
- 4. All patients with an acute asthma attack at home, and symptoms not controlled by the maximum daily licensed dose of anti-inflammatory reliever (AIR) therapy or maintenance and reliever therapy (MART) or up to 10 puffs of salbutamol via a pMDI and spacer, should seek urgent medical attention. (BTS/SIGN/NICE Jan 2025).
- 5. MHRA Aug 2022: home use of nebulisers in paediatric asthma should be initiated and managed only by specialists (under a treatment plan). Use of a nebuliser purchased independently of medical advice for use in the home to deliver nebulised asthma rescue medications to children can mask a deterioration in the underlying disease and may increase the risk of potentially fatal delays in seeking medical attention if asthma deteriorates.

#### 3.2 Inhaled Corticosteroids

**N.B. Prescribe by brand**; Use spacer device with MDI for all doses of inhaled corticosteroid. **See adults and children's asthma guidance** for inhaled corticosteroid doses

Corticosteroids (ICS)	Licenced	Traffic Light Classification	Additional information
Soprobec MDI 50, 100, 200, 250 micrograms	Asthma- adults and children (only 50 & 100 microgram inhalers are licensed for children &	GREEN	Soprabec is therapeutically equivalent to Clenil.
Kelhale MDI 50,100     micrograms (extra fine particles)	adolescents) Asthma- adults >18 years	GREEN	Kelhale is therapeutically equivalent to Qvar.
QVAR MDI 50, 100     micrograms(extra fine particles)	Asthma- ≥5 years of age	GREEN	Kelhale/Qvar twice as potent as Soprobec (e.g. Kelhale/Qvar 50 micrograms is equivalent to Soprobec100micrograms)
Budesonide • Easyhaler (DPI) 100, 200,400	Asthma- ≥6 years of age	GREEN	dose equivalent to standard particle beclomethasone (Soprobec/ Clenil)
Fluticasone propionate  Flixotide Accuhaler (DPI)100 microgram  Flixotide Evohaler (MDI) 50 microgram	Asthma- >4 years of age	GREEN for children GREY for adults	As per children's asthma guidance  Fluticasone is twice as potent as Soprobec/Clenil (e.g. fluticasone 50 microgram is equivalent to Soprobec100micrograms)  All other formulations of fluticasone propionate are classified as GREY on the formulary.

**Combination inhalers N.B.** Prescribe by brand. See asthma and COPD <u>guideline</u> and <u>Inhaled corticosteroid equivalent doses</u>.

LABA/ICS combination	Licenced	Traffic Light Classification	Additional information
Budesonide and formoterol	Asthma (≥6	GREEN 1st line	
Fobumix easyhaler DPI	years of age)	LABA/ICS combination inhaler	Fobumix 80/4.5, 160/4.5, 320/9 is equivalent to
80/4.5,  • Fobumix easyhaler DPI	Asthma (≥12 years of age) & COPD	GREEN 1st line LABA/ICS combination inhaler	Fobumix 80/4.5 licensed for asthma only.
160/4.5, 320/9	33. 2	GREEN	
	Asthma		WockAIR is the most cost effective DPI where
WockAIR DPI 160/4.5, 320/9	(≥12years of age) & COPD	GREEN	appropriate but is only available in two strengths
	· ,		WockAIR and DuoResp 160/4.5, 320/9 is
<ul> <li>DuoResp Spiromax DPI 160/4.5, 320/9</li> </ul>	Asthma (≥12 years of age) &	GREEN	equivalent to Symbicort 200/6, 400/12 respectively.
	COPD	GREEN	
Symbicort Turbohaler DPI	Asthma (≥6	GREEN	
100/6	years of age)		
Symbicort Turbohaler DPI	Asthma (≥12	GREEN	
200/6, 400/12	years of age) & COPD		Symbicort pMDI contains propellant HFA227ea which has a significantly higher
	COPD		carbon footprint than other propellants and
Symbicort MDI 200/6	COPD	GREY	so should be avoided where possible.
Beclometasone (extra fine) and			
formoterol			
Fostair Nexthaler DPI 100/6	Asthma (≥18	GREEN	
Factoria Navithalas DDI 200/C	years of age) & COPD		
<ul> <li>Fostair Nexthaler DPI 200/6</li> </ul>	COPD		

<ul> <li>Proxor, Bibecfo, Luforbec MDI 100/6</li> <li>Proxor, Bibecfo, Luforbec MDI 200/6</li> </ul>	Asthma (≥18 years of age)  Asthma (≥18 years of age) & COPD	GREEN 1st line for patients requiring an MDI	100 micrograms of beclometasone dipropionate via Proxor/Luforbec/Bibecfo/Fostair products are equivalent to 250 micrograms of beclometasone dipropionate in standard particle CFC-free inhalers.
Fluticasone propionate and salmeterol  Combisal MDI 50/25  Seretide MDI 50  Seretide Accuhaler DPI 100  Fixkoh Airmaster DPI 50/100	Asthma ≥4 years of age Asthma ≥12 years of age	GREEN for children GREY for adults	See children's asthma guidance.  Combisal MDI is the cost-effective alternative for Seretide evohaler.  Children receiving Seretide/Combisal 50 MDI or Seretide/ Fixkoh 100 Accuhaler should be reviewed at age 18 year and considered for the formulary choice ICS/LABA combination
<ul> <li>Combisal MDI 125/25, 250/25</li> <li>Fixkoh DPI 500/50</li> <li>Fusacomb Easyhaler DPI 500/50</li> </ul>	Asthma ≥12 years of age  Asthma >12 years of age & COPD  Asthma(≥12 years of age) & COPD	place of therapy for adults	inhalers.  Fluticasone/salmeterol combination has limited place of therapy in adults- following choices in local COPD guidance only.  Fixkoh DPI/Fusacomb Easyhaler are cost-effective alternative for Seretide Accuhaler 500
Triple combination (ICS+ LABA+ LAMA)	Licenced	Traffic Light Classification	Additional information
Fluticasone, vilanterol and umeclidinium (Trelegy) (DPI)	COPD	GREY 1st line	Indicated for the maintenance treatment of
Beclometasone, formoterol	COPD	triple combination GREY 1 <sup>st</sup> line	moderate to severe COPD.  Triple therapy is reserved for exceptional use in severe disease in the presence of persistent
Beclometasone, formoterol and glycopyrronium (Trimbow NEXThaler DPI)		combination  GREY 1 <sup>st</sup> line triple combination	moderate to severe COPD.  Triple therapy is reserved for exceptional use
Beclometasone, formoterol and glycopyrronium (Trimbow NEXThaler DPI)  Beclometasone, formoterol and glycopyrronium (Trimbow) (MDI)	COPD Asthma in adults (172/5/9 microg licensed for asthma only)	combination  GREY 1st line triple combination  GREY after consultant/ specialist initiation for asthma.	moderate to severe COPD.  Triple therapy is reserved for exceptional use in severe disease in the presence of persistent exacerbations or patient is still limited by symptoms despite other treatments.  Triple therapy in a single inhaler may be preferable for people who have difficulty using more than one device or who find their medication regimen difficult or confusing and have trouble complying with treatment. However triple therapy lack flexibility and makes it difficult to amend the individual
Beclometasone, formoterol and glycopyrronium (Trimbow NEXThaler DPI)  Beclometasone, formoterol and glycopyrronium (Trimbow)	COPD Asthma in adults (172/5/9 microg licensed for	combination  GREY 1st line triple combination  GREY after consultant/ specialist initiation for	moderate to severe COPD.  Triple therapy is reserved for exceptional use in severe disease in the presence of persistent exacerbations or patient is still limited by symptoms despite other treatments.  Triple therapy in a single inhaler may be preferable for people who have difficulty using more than one device or who find their medication regimen difficult or confusing and have trouble complying with treatment.  However triple therapy lack flexibility and

- 1. All doses of inhaled steroid delivered via an MDI should be via a spacer device. This form of administration improves delivery of drug to the airways reducing local effects e.g., oral candida and also reduces deposition in the oropharynx (thereby reducing local adverse effects and the amount of systemic absorption).
- 2. **SIDE EFFECTS**: Inhaled steroids do not usually cause adrenocortical suppression at normal doses (up to about 1500 micrograms/day of beclometasone (standard particle size)). Oral candidiasis may be related to dose and dose frequency. Patients should be advised to rinse the mouth after use of high

- dose steroids to help minimize this. Dysphonia can occur. The use of a large volume spacer device (Volumatic) may reduce these local adverse effects.
- 3. Inhaled corticosteroids and adrenal suppression in children 'Adrenal suppression may be under-recognised'. Prescribers are reminded that:
  - It is important to monitor therapy regularly and titrate down to the lowest dose at which effective control of asthma is maintained.
  - Growth (height and weight centile) should be monitored at least annually in children with asthma.
  - In paediatric practice adrenal suppression has been seen at doses as low as 800 micrograms/day (standard particle size). All children on these doses should have careful growth monitoring and a written management plan advising about the risk of adrenal suppression.
- 4. If a doctor considers that a child's asthma is not controlled despite an increase to a high dose of inhaled corticosteroid plus a LABA, the child should be referred to a specialist in the management of paediatric asthma.
- 6. MHRA, 2017. Advise patients to report any blurred vision or other visual disturbances due to rare risk of central serous chorioretinopathy with corticosteroids.
- 7. The role of Maintenance and reliever therapy (MART) has a limited place for patient therapy and selection. MART is suitable for patients who despite good adherence to regular maintenance doses of a combination ICS/LABA inhaler, have;
  - 1. Inadequate asthma control and are in frequent need of reliever medication.
  - 2. Asthma exacerbations in the past requiring medical intervention.
- Inhaled Budesonide for treatment of COVID-19- Inhaled budesonide should no longer be considered as a treatment for individuals with COVID-19 infection other than within the context of a clinical trial. People already using budesonide for conditions other than COVID-19 should continue treatment if they test positive for COVID-19. See NICE NG191 Covid-19 rapid guideline: managing COVID-19 (updated May 2024)

## 3.3.2 Leukotriene receptor antagonists

See adult and children's asthma quidance

Montelukast tabs 10mg, chewable tabs 4mg, 5mg, granules 4mg

- 1. Granules are more expensive and should be reserved for children between 6 months and 2 years old or if chewable tablets not suitable. 4mg chewable tablets are licensed in children 2-5 years old; 5mg chewable tablets are licensed in children 6-14 years old; 10mg tablets licensed for adults and children 15 years and over
- 2. MHRA April 2024 Montelukast: risk of neuropsychiatric reactions
  - be alert for neuropsychiatric reactions in patients taking montelukast; events have been reported in adults, adolescents, and children
  - discontinue montelukast if patients experience new or worsening symptoms of neuropsychiatric reactions
  - advise patients and their caregivers to carefully read the list of neuropsychiatric reactions in the Patient Information Leaflet and to seek medical advice immediately should they occur.
  - Patients, parents and carers should be warned of possible adverse reactions affecting sleep, behaviour and mood.

#### 3.4.1 Antihistamines

For treatments of minor self-limiting conditions such as mild to moderate hay fever, <u>self-care</u> is encouraged. Treatments are available to purchase over-the-counter.

Cetirizine 10mg tabs
Loratadine 10mg tabs, oral solution 5mg/5ml
Chlorphenamine tabs 4mg, oral solution 2mg/5ml

Non-sedating Non-sedating Sedating 1. JAPC has classified alimemazine as **Do Not Prescribe (DNP)** due to lack of cost-effectiveness. Suitable alternative that may be considered is promethazine (GREY).

## 3.4.2 Allergen immunotherapy

- 1. Omalizumab (NICE TA278) for allergic asthma is RED
- 2. Mepolizumab (NICE TA671) for eosinophilic asthma is RED

## 3.4.3 Allergic emergencies

**Jext** auto-injector 150 micrograms, 300 micrograms PIL **EpiPen** auto-injector 0.3mg, **EpiPen Jr** auto-injector 0.15mg PIL /EpiPen Jnr PIL

- MHRA June 2023- Adrenaline auto-injectors (AAIs): new <u>guidance</u> and resources for safe use-including an easy step-by-step guide on what to do in an emergency and updated advice on body positioning. A toolkit of resources is available for health and social care professionals to support the safe and effective use of AAIs. Use the materials to inform patients and caregivers what do if they suspect anaphylaxis and how to use adrenaline auto-injectors (AAIs)
- 2. <u>MHRA 2017</u> recommends two adrenaline auto-injectors should be prescribed which patients should carry at all times.
- 3. Adrenaline has a narrow therapeutic index. Primary care clinicians should prescribe in line with product licensing as summarised in the table below:-

	Weight range	Dose
Jext	15-30kg	150 micrograms
	>30kg	300 micrograms
Epipen	Between 7.5 to 25kg	150 micrograms (Epipen Junior)
	>25kg	300 micrograms

4. For educational material produced by manufacturer see links:-, <u>EpiPen</u>, <u>Jext</u> Other resources see links:, <u>Epipen</u>, <u>Jext</u> (key differences between devices)

## 3.7 Mucolytics

Carbocisteine375mg caps, 250mg/5ml oral solutionGREENAcetylcysteine600mg Sugar-free effervescent tabletsGREEN

- 1. NICE NG115 COPD in over 16s: diagnosis and management:
  - Consider mucolytic drug therapy for people with a chronic cough productive of sputum.
  - Only continue mucolytic therapy if there is symptomatic improvement (for example, reduction in frequency of cough and sputum production).
  - Do not routinely use mucolytic drugs to prevent exacerbations in people with stable COPD.