

This patient group direction (PGD) must only be used by registered health professionals who have been named and authorised by their organisation to practice under it. The most recent and in date final signed version of the PGD should be used.

Patient Group Direction (PGD)

For the administration or supply of

Salbutamol, Nebulised 2.5mg/2.5ml and Inhaler 100mcg metered dose inhalation

By registered health care professionals for

Reversibility testing

Throughout the Manx Care and those contracted by the Manx Care where appropriate within practice

PGD NUMBER 105

1. Change history

Version number	Change details	Date
1	Original PGD ratified	June 2021

Reference number: 105 Valid from: 06/2021 Review date: 06/2023

2. Medicines practice guideline 2: Patient group directions

Refer to the relevant sections of NICE medicines practice guideline 2: *Patient group directions* as stated in the blank template notes. For further information about PGD signatories, see the NHS and Manx Care <u>PGD</u> website FAQs

3. PGD development

Refer to the <u>NICE PGD competency framework for people developing PGDs</u>

Job Title & organisation	Name	Signature	Date
Author of the PGD			
Member of the PGD working group			

4. PGD authorisation

Refer to the <u>NICE PGD competency framework for people authorising PGDs</u>

Job Title	Name	Signature	Date
Medical Director			
Chief Pharmacist/ Pharmaceutical Adviser			
Senior Paramedic			
Director of Nursing			
GP Adviser			
Senior Microbiologist (if PGD contains antimicrobials)			

Reference number: 105 Valid from: 06/2021 Review date: 06/2023

5. PGD adoption by the provider

Refer to the <u>NICE PGD competency framework for people authorising PGDs</u>

Job title and organisation	Signature	Date	Applicable or not applicable to area

6. Training and competency of registered healthcare professionals, employed or contracted by the Manx Care, GP practice or Hospice

Refer to the <u>NICE PGD competency framework for health professionals using PGDs</u>

	Requirements of registered Healthcare professionals working	
	under the PGD	
Qualifications and professional registration	 Registered healthcare professionals, working within or contracted by the Manx Care, GP practice or Hospice who are permitted staff groups outlined within the current PGD policy Pharmacists must be practising in Manx Care authorised premises i.e. contracted pharmacy premises 	
Initial training	 Knowledge of current guidelines and the administration of the drug specified in this PGD/BNF and of the inclusion and exclusion criteria Training which enables the practitioner to make a clinical assessment to establish the need for the medication covered by this PGD Local training in the use of PGD's 	
Competency	Staff will be assessed on their knowledge of drugs and clinical	
assessment	assessment as part the competency framework for registered health professionals using PGD's	
Ongoing training and competency	The registered health care professionals should make sure they are aware of any changes to the recommendations for this medication; it is the responsibility of the registered health care professionals to keep up to date with continuing professional development. PGD updates will be held every two years	

Reference number: 105 Valid from: 06/2021 Review date: 06/2023

7. Clinical Conditions

Clinical condition or	Reversibility testing		
situation to which this			
PGD applies			
Inclusion criteria	Nebulisation or metered dose inhaler with spacer administration to		
	determine reversibility in asthma or improvement in COPD		
	Patients 1 month and over		
Exclusion criteria	Neonates (Under 1 month)		
	Follow British Thoracic Society Guidelines on Asthma management.		
	Lactation		
	Pregnancy		
	Patients already taking non selective beta blocking drugs (e.g.		
	propranolol)		
	Severe cardiac disease		
Cautions (including any	Arrhythmias		
relevant action to be	Cardiovascular disease		
taken)	Diabetes (risk of hyperglycaemia and ketoacidosis, especially		
	with intravenous use)		
	Hypertension		
	Hyperthyroidism		
	Hypokalaemia		
	Susceptibility to QT-interval prolongation		
	Tachycardia		
Arrangements for referral	Patient should be referred to a more experienced clinical		
for medical advice	practitioner for further assessment		
Action to be taken if	Patient should be referred to a more experienced clinical		
patient excluded	practitioner for further assessment		
Action to be taken if	A verbal explanation should be given to the patient on: the need		
patient declines	for the medication and any possible effects or potential risks		
treatment	which may occur as a result of refusing treatment		
	This information must be documented in the patients' health		
	records		
	Any patient who declines care must have demonstrated capacity		
	to do so		
	Where appropriate care should be escalated		

8. Details of the medicine

Name, form and strength	Salbutamol, Nebulised 2.5mg/2.5ml and Inhaler 100mcg metered
of medicine	dose inhalation
Legal category	Prescription Only Medicine (POM)
Indicate any off-label use	Metered dose inhalation is off-label
(if relevant)	Nebuliser dose is off-label in under 4 years
Route/method of	Inhalation of nebuliser solution or inhalation from MDI inhaler via
administration	spacer

Reference number: 105 Valid from: 06/2021 Review date: 06/2023

Dose and frequency	1 x Salbutamol 2.5mg nebuliser	
	4 x Salbutamol 100mcg (MDI using a spacer)	
Quantity to be	Administered:	
administered or supplied	400mcg salbutamol by MDI & spacer	
duminister ed or supplied	OR	
	 2.5mg salbutamol via nebuliser 	
	 Wait 20 minutes and then redo the spirometry tests and see 	
	whether there has been any reversibility (improvement)	
	following the salbutamol	
	Tollowing the Salbutanion	
	Supplied:	
	 30 days twice a day 2.5mg/2.5ml nebule to be used if 	
	, , , , , , , , , , , , , , , , , , , ,	
Maximum or minimum	bronchospasm occurs post hypertonic saline administration	
	One episode of care	
treatment period	1 x salbutamol 2.5mg nebuliser	
	4 x salbutamol 100mcg total of 400mcg MDI	
Storage	Room temperature	
Adverse effects	Arrhythmias	
	Headache	
	Hypokalaemia (with high doses)	
	Muscle cramps/spasms	
	Nasopharyngitis	
	Nausea	
	Palpitations	
	Rash	
	Tremor	
Records to be kept	The administration of any medication given under a PGD must be	
	recorded within the patient's medical records	

9. Patient information

Verbal/Written information to be given to patient or carer	 Where medication is being supplied under a PGD, written patient information leaflet must also be supplied A patient information leaflet is available on request 	
Follow-up advice to be	If symptoms do not improve or worsen or you become unwell, seek	
given to patient or carer	medical advice immediately	

Reference number: 105 Valid from: 06/2021 Review date: 06/2023

10. Appendix A

References

- 1. British National Formulary (BNF) available online: https://bnf.nice.org.uk
- 2. Nursing and Midwifery (2018) "The code" available online: https://www.nmc.org.uk
- 3. Current Health Care Professions Council standards of practice
- 4. General Pharmaceutical Council standards
- 5. The General Optical Council
- 6. Electronic medicines compendium available online: https://www.medicines.org.uk
- 7. British Thoracic Society and Association for Respiratory Technology and Physiology. Guidelines for the measurement of respiratory function. Respiratory Medicine 1994 88; 165-194.
- 8. National Institute for Health and Clinical Excellence. Management of chronic obstructive pulmonary disease in adults in primary and secondary care (partial update). 2010. www.nice.org.uk/CG101

11. Appendix B

Health professionals agreed to practice

- Each registered healthcare professional will hold their own Competency framework which will be signed and agreed by their mentor
- A mentor is defined within the Manx Care policy as any ward/area managers, sisters, senior nurses, GPs, pharmacists or senior paramedics who has completed the PGD training themselves

Reference number: 105 Valid from: 06/2021 Review date: 06/2023