

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 of

Ceftriaxone 1g Powder for Solution for Injection Vials

By registered health care professionals for

Bacterial Meningitis/Meningococcal Septicaemia

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

PGD NUMBER 81

1. Change history

Version number	Change details	Date
1	Original PGD ratified	June 2021
2	Minor wording amendments	March 2022

Reference number: 81 Valid from: 03/2022 Review date: 03/2024

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 website FAQs</u>

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)			

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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 PGDs
Competency	Staff will be assessed on their knowledge of drugs and clinical
assessment	assessment as part the competency framework for registered health professionals using PGDs
Ongoing training and	The registered health care professionals should make sure they
competency	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

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7. Clinical Conditions

Clinical condition or	Suspected bacterial meningitis or meningococcal septicaemia
situation to which this	
PGD applies	
Inclusion criteria	Adults and children presenting with signs of meningitis or
	meningococcal septicaemia
Exclusion criteria	Allergy to Cephalosporins
	History of Immediate Hypersensitivity to Penicillin and other
	Beta-Lactams
	Neonates
	Concomitant treatment with intravenous calcium (including
	total parenteral nutrition containing calcium) in premature
	and full-term neonates—risk of precipitation in urine and lungs
	(fatal reactions) (in neonates)
	Full-term neonates with jaundice, hypoalbuminaemia,
	acidosis, unconjugated hyperbilirubinaemia, or impaired
	bilirubin binding — risk of developing bilirubin encephalopathy
	(in neonates); premature neonates less than 41 weeks
	corrected gestational age (in neonates)
	Concomitant treatment with intravenous calcium (including)
	total parenteral nutrition containing calcium) (in adults)
Cautions (including any	Breast feeding - low concentration in breast milk
relevant action to be	History of hypercalciuria and kidney stones
taken)	May cause confusion
	Pregnancy
	Severe liver impairment
	Severe renal impairment
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
patient declines	need for the medication and any possible effects or potential
treatment	risks 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

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8. Details of the medicine

Name, form and strength	Ceftriaxone Sodium 1g powder for solution for injection vial		
of medicine	Description Only Madician (DOM)		
Legal category	Prescription Only Medicine (POM)		
Indicate any <u>off-label use</u>	None		
(if relevant)			
Route/method of	Intravenous injection or infusion		
administration	Deep intramuscular injection – split dose across two separate		
	sites if administering over 1g		
	Intraosseous (IO)		
Dose and frequency	Child 1 month-11 years (body weight up to 50kg)		
	80mg/kg max 2g once only given via intravenous infusion over		
	30mins, or deep intramuscular injection		
	Child 9-11 years (body weight 50kg and above)		
	2g once only given intravenous injection, or infusion over 30mins, or		
	deep intramuscular injection		
	Child 12-17 years		
	2g once only given intravenous injection, or infusion over 30mins, or		
	deep intramuscular injection		
	Adult		
	2g once only given intravenous injection, or infusion over 30mins, or		
	deep intramuscular injection		
Quantity to be	Single dose		
administered			
Maximum or minimum	Once only		
treatment period			
Storage	Room temperature		
Adverse effects	Abdominal pain Leucopenia		
	Agranulocytosis Pseudomembranous enterocolitis		
	Angioedema Nephritis tubulointerstitial		
	Anaphylactic (reversible)		
	reaction Diarrhoea • Severe cutaneous adverse reactions		
	• Dizziness (SCARs)		
	Eosinophilia Skin reactions		
	Haemolytic anaemia		
	Headache Vomiting		
	Nausea Vulvovaginal candidiasis		
	Neutropenia		
Records to be kept	The administration of any medication given under a PGD must be		
	recorded within the patient's medical records		
	1 coorded within the patient 5 incured records		

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9. Patient information

Verbal/Written	Verbal information must be given to patients and or carers for
information to be given	all medication being administered under a PGD
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,
given to patient or carer	seek medical advice immediately

10. Appendix A

References

- 1. British National Formulary (BNF) available online: https://bnf.nice.org.uk
- 2. Nursing and Midwifery "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

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

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