6-4-18 -> J. YOUD

CALDERDALE & HUDDERSFIELD NHS FOUNDATION TRUST A. KEASKIN AUTHORISATION AND RECORD OF AGREEMENT OF NAMED HEALTH PROFESSIONALS TO SUPPLY MEDICINES UNDER: M. DAVIES

PATIENT GROUP DIRECTION FOR THE SUPPLY OF FLUCLOXACILLIN BY REGISTERED HEALTH PROFESSIONALS IN EMERGENCY DEPARTMENTS



1. PGD AUTHORISATION

Position	Name	Signature	Date
Acting Clinical Director of Pharmacy	Fiona Smith	Auth	28/3/18
Executive Director of Nursing	Brendan Brown	Jundan M	28/23/18
Medical Director	David Birkenhead	0. 800	22/3/18
Chairman of Medicines Management Committee	Anu Rajgopal	Q-2	29/3/18

Date of Patient Group Direction: March 2018

If revision please tick box √

Valid Until: March 2020

Review Date: September 2019

Approved by the Trust Medicine Management Committee on: 24T-1 MAY 7018

PATIENT GROUP DIRECTION FOR THE SUPPLY OF FLUCLOXACILLIN

BY

REGISTERED HEALTH PROFESSIONALS IN

EMERGENCY DEPARTMENTS

2. CLINICAL CONDITION

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Indication	Potential or established soft tissue infection Refer to Trust antibiotic policy.	
Relevant National and Local Guidelines/Information sources	Licensed useRecognised first line antibiotic	
Description of Patients included in treatment	Adults Children 1yr and over	
Description of Patients excluded from treatment under the terms of this PGD	 Allergy to Penicillin Severe renal failure Hepatic impairment 	
Action if excluded	Refer to doctor, Advanced Clinical Practitioner or consider other antibiotics	
Action if patient self excludes/declines	Refer to doctor, Advanced Clinical Practitioner or consider other antibiotics	

3. TREATMENT

Name, form and strength of	Flucloxacillin 500mg capsule	
medicine	Suspension 125mg/5ml, 250mg/5ml	
Legal Status <i>GSL, P, POM</i>	POM	
Dose	Adult - 500mg	
	Child 1-2 years – 125mg	
	Child 2-10 years – 125-250mg	
	Child 10-18 years - 250-500mg	
Frequency of administration	QDS	
Method and route of administration	Oral	
Supporting facilities required		
Quantity to supply/administer	Course length 7 days	
Duration of treatment	500mg caps qds – 28	
	125mg/5ml qds – 200ml	
	250mg/5ml qds – 200ml	
Potential side effects	Nausea/vomiting, diarrhoea, gastric upset, hypersensitivity reaction, hepatitis and cholestatic jaundice	

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Advice to patient/carer Managing & Reporting	 Complete prescribed course Take before food If rash/pruritis develops stop taking If jaundice develops during or after course, stop taking & consult doctor Suspension – shake the bottle All suspected adverse drug reactions occurring after
Adverse Events	 treatment following this PGD must be reported to a senior medical practitioner responsible for the area in which the direction is in use. The healthcare professional administering/supplying from the PGD must also report the ADR using Trust incident reporting procedure All serious adverse drug reactions should be reported to the MHRA / CSM using the Yellow Card System. Yellow cards and guidance on its use are available at the back of the BNF or at www.yellowcard.gov.uk
Follow up	As required by clinical condition
When to refer to doctor	Any reaction to medication or exacerbation of symptoms
Treatment record Specify method of recording supply/administration sufficient for audit trail	 Electronic Patient Record in ED Prescription Name, dose, and frequency of drug, Volume/ quantity supplied Advice given, verbal or written Signed and dated

4. STAFF

Professional Qualifications	Registered Nurse or Registered Paramedic Current NMC or HCPC Registration	
Any Exceptions to above	Bank and Agency Staff	
Specialist competencies, qualifications and experience	Emergency Nurse Practitioner Programme (ED) or Advanced Clinical Practitioner Programme (inc. Trainees) Trust PGD training	
Continuing training & education	Update in line with clinical guidance	

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5. MANAGEMENT AND MONITORING

Records to be kept for Audit Purposes	 STORAGE AND RETRIEVAL Pharmacy will retain the original signed version of the PGDs Adult – 8 years Children (under 18 years) As the requirement is until child is 25 years old or for eight years after child's death and PGDs are not child specific – this would be indefinitely (at least a minimum of 43 years) Division/Author is responsible for keeping the record/retrieval method of those authorised to work under a PGD/signature sheet to comply with the above
Date of writing	March 2018
Name of manager holding record of names of those authorised to work under this PGD	Louise Croxall – Matron, ED

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Names of all authors of PGD (to include a Dr or Dentist)	Print Name: Janet Youd
	Title: Emergency Nurse Consultant
	Signature: Jonet Youd Date: 2013)18
	Print Name: Dr Mark Davies .
	Title: Emergency Medicine Consultant
	Signature: Date: 21 3 18
Lead Pharmacist involved in preparation of PGD	Print Name: Lisa Hodgson
propulation of 1 OB	
	Signature:
	Date: .23/.3/.18
Approval of Clinical Director	Print Name: Mark Davies
	Signature:
	Date:2(S Y

PATIENT GROUP DIRECTION FOR THE SUPPLY OF FLUCLOXACILLIN BY REGISTERED HEALTH PROFESSIONALS IN EMERGENCY DEPARTMENTS

This Patient Group Direction is to be read and agreed and this Authorisation and Record of Agreement signed by all Health Professionals who will administer and/or supply treatment using it. It is the responsibility of each professional to practice only within the bounds of their own competence

A copy of the Patient Group Direction and the original, signed Record of Agreement must be held together by the Ward/Departmental Manager/Community Team Leader.

'I confirm that I have read and understood the content of this Patient Group Direction and that I am willing to work under it within my Professional Code of Practice/Conduct.'

Name of Health Professional	Designation e.g. RGN	Signature of Health Professional	Signature of Ward/Departmental/Area Manager	Date