

NHS South of Tyne and Wear

serving Gateshead Primary Care Trust, South Tyneside Primary Care Trust and
Sunderland Teaching Primary Care Trust

SHARED CARE GUIDELINE

For

Hydroxychloroquine for the Management of mild, chronic inflammation, polyarthritis, SLE assosciated musculoskeletal and skin manifestations, palindromic arthritis and as an adjunct to other DMARDs.

Implementation Date: 5th October 2010

Review Date: 12th June 2012

This guidance has been prepared and approved for use within Gateshead, South Tyneside and Sunderland in consultation with Primary and Secondary Care Trusts and Local Medical Committees.

The guideline sets out the details of the transfer of prescribing and respective responsibilities of GPs and specialist services within shared care prescribing arrangements. It is intended to provide sufficient information to allow GPs to prescribe these treatments within a shared care setting

Further copies are available from

Lynn Cunningham	SOTW Medicines	Clarendon
	Management Team	Windmill Way
		Hebburn
		Tyne & Wear NE311AT
		Tel 0191 283 1348

Approved by:

Committee	Date
Gateshead Medicines Management	
Committee	
South Tyneside Prescribing Committee	
Sunderland Primary Care Prescribing	
Group	
South of Tyne and Wear Medicines	5.10.2010
Management Committee	

Name of drug:	Hydroxych	loroquine	Form and strength:	Tablets 200mg
Brand name:	Plaquenil		BNF Code:	10.1.3
Conditions(s) to	o be treated		inflammation, poly musculoskeletal a	ne is given for mild, chronic arthritis, SLE assosciated and skin manifestations, is and as an adjunct to
Excluded patie	nts	Patients v	where GP refused	shared care
Eligibility criteri shared care	a for	all patient		
Initiation		Treatment	will be initiated by	the hospital
Duration of trea	atment	has to sto	p	lvise GP when treatment
Usual Maintena and Dose Range	ance Dose	maintenar Normally of	nce dose. dose will be reduced	onths then 200mg daily as d to half after 3-6 months dvise of dose changes
Maximum Dose	<u> </u>	6.5mg/kg/	dav	
Available Stren (Colours)		White 200		
Preparations		Tablets		
Cost 28 days (Drug Tariff)	£5 / month	<u> </u>	
Adverse effects	6	visual field uncommo exceeded discontinu such as ha be transie Blurring of accommod	d defects can occur, n if the recommend. In its early form it a ation of hydroxychloaloes, blurring of vis	ed daily dose is not appears reversible on proquine. Disturbances sion or photophobia may be on stopping treatment.
		pigmentar membrane been repo stopping to dermatitis pustulosis Hydroxych porphyria Other adv disturband abdomina symptoms	rted. These usually reatment. Isolated of and acute generalis (AGEP) have been aloroquine can also and may precipitate erse effects includes such as nauseal cramps and rarely, are often transient ely on reducing the offerse.	rand hair loss have also resolve readily on eases of exfoliative sed exanthematous reported. precipitate or exacerbate attacks of psoriasis. le gastrointestinal diarrhoea, anorexia, vomiting. These but usually resolve

Contra-indications	Known hypersensitivity to the product Pre-existing maculopathy Caution: history of epilepsy, porphyria and the elderly. Breast feeding May interact with quinine, digoxin, amiodarone,
Drug-interactions	aminoglycosides, mefloquine, cimetidine, and ciclosporin. Avoid antacids within 4 hours of dose Live vaccines must be avoided.
Renal impairment and liver disease	Caution and monitoring of hydroxychloroquine in renal and hepatic impairment
Pregnancy and breast feeding	Normally hydroxychloroquine is discontinued during pregnancy but may be continued in some patients with lupus. Hydroxychloroquine is contra-indicated in lactation
Monitoring	Patient should have near visual acuity in each eye recorded (with reading glasses if worn) at the hospital using Royal College of Ophthalmology Chart. Annual review to be performed either using this chart or by an optometrist. The patient will be referred to an Ophthalmologist if visual impairment or eye disease is detected at the baseline assessment or if there is a change in acuity or blurred vision (hydroxychloroquine should be stopped). With 3 years of continuous use there is little or no incidence of retinopathy. Patients requiring long term therapy (5 years) should be discussed with an ophthalmologist to determine whether further monitoring is required. If patient reports change in vision, discontinue hydroxychloroquine and discuss with rheumatology. Recommend flu vaccination and consider pneumococcal vaccine

Responsibilities

DISEASE MONITORING

Clinical response to therapy will be assessed by the hospital physician in all cases and communicated to the GP

RESPONSIBILITY FOR PRESCRIBING

On initiation of therapy the patient will be given a one month supply of hydroxychloroquine by secondary care. Responsibility thereafter for prescribing may be transferred to the patients GP depending on the locality in which the GP is based and the secondary care centre the patient attends. This is detailed below.

The GP should not prescribe unless the monitoring has been carried out and the GP is satisfied that it is safe to continue treatment.

Practices in Sunderland PCT

PATIENTS REFERRED TO SUNDERLAND (helpline 0191 5656256 ext 47533–Mon- Fri 9am –5pm).

All the monitoring, and dosing will be carried out entirely by the rheumatology clinic with the responsibility of secondary care to inform the GP of any abnormalities. The GP will be responsible for prescribing in agreement with the Secondary care service.

PATIENTS REFERRED TO GATESHEAD (helpline 0191 445 5240 Mon- Fri 9am- 5pm or consultant secretary)

All monitoring, prescribing and dosing to be carried out by GP with support from secondary care specialist as required

PATIENTS REFERRED ELSEWHERE

All monitoring, prescribing and dosing to be carried out by GP with support from secondary care specialist as required

Practices in Gateshead PCT

PATIENTS REFERRED TO GATESHEAD (helpline 0191 445 5240 Mon- Fri 9am- 5pm or consultant secretary

Secondary care staff will carry out base line monitoring prior to initiating therapy. GPs will carry out ongoing prescribing. Secondary care staff will carry out monitoring and advise GPs of changes to dose or monitoring intervals.

PATIENTS REFERRED ELSEWHERE

All monitoring, prescribing and dosing to be carried out by GP with support from secondary care specialist as required

	PATIENTS REFI	uth Tyneside PCT ERRED TO GATESHEAD (helpline Mon- Fri 9am- 5pm or consultant
	All monitoring, pr	rescribing and dosing to be carried out ort from secondary care specialist as
	Most monitoring by the rheumatol secondary care to the GP will be rewith the Secondar responsibility may between special will then be responsed.	erred to Sunderland (helpline at 47533–Mon- Fri 9am –5pm). and dosing will be carried out entirely logy clinic with the responsibility of o inform the GP of any abnormalities. esponsible for prescribing in agreement ary care service. Once stable, by be transferred to GP as agreed st and GP at time of transfer. The GP consible for monitoring, dosing and
Communications	Consultant	Please refer to the standard letter from the patient's consultant. For Gateshead patients a copy of the Gateshead GP information sheet should be enclosed with the letter
	G.P.	If the GP is unwilling to accept prescribing responsibility for an individual patient the consultant should be informed within 1 month of receipt of the shared care request. In such cases the GP must inform the consultant of all relevant medical information regarding the patient and any changes to the patient's medication irrespective of indication.
	Patient	The patient will have received an information leaflet from the hospital.
Re- referral criteria		
Contact details	Consultant:	
	Additional information for Gateshead patients is available at www.gatesheadhealth.nhs.uk/rheumatology	
Agreed Date		.gatesileauneaiin.iiii5.uk/meumat0l0gy
Agreed Date	Expiry date	

Reference to full prescribing information e.g. SPC

Appendix 2 Shared Care Request Form

- Consultant to complete FIRST SECTION of form
- GP to complete SECOND section and RETURN to ACUTE TRUST CLINICIAN TEAM if NOT accepting shared care

Section 1	
Consultant	
Hospital address	
Contact Phone Number	
	,
Patient's name	
Address	
This patient is stabilised on	
Dose	
Prescription for 28 days supply given on	
Compliance aid	YES/NO
Monitored by	
Designated community pharmacy	
Their treatment has been explained to them and	a review has been arranged for
Appointments to continue everyr	months

Section 2			
Patient's name			
Address			
I do NOT ACCEPT the	e proposed Shared-Care Agreement for this patient		
My reasons for not accept Please complete this sect	ing: ion		
Signeddate			
Please return to the Acute Trust Clinician team at :			