PAN MERSEY APC

PAN MERSEY AREA PRESCRIBING COMMITTEE
SHARED CARE FRAMEWORK
FIRST APC BOARD DATE: 27 SEP 2017
LAST APC BOARD DATE: 28 NOV 2018

NHS Pan Mersey Area Prescribing Committee

SODIUM AUROTHIOMALATE for patients within adult services

1. Background	The precise mode of action of sodium aurothiomalate is not yet known. Treatment with gold has been shown to be accompanied by a fall in ESR and C-reactive protein, an increase in serum histidine and sulphydryl levels and a reduction in serum immunoglobulins, rheumatoid factor titres and Clq-binding activity. Indications, dose adjustments and monitoring requirements for disease modifying drugs (DMDs) (licensed and unlicensed indications) included in this Framework are in line with national guidance published by the British Society for Rheumatology 2017.
2. Licensed Indications	• Rheumatoid arthritis • Juvenile idiopathic arthritis
3. Locally agreed off-label use	N/A
4. Initiation and ongoing dose regime	**Transfer of monitoring and prescribing to Primary care is normally after 3 months** **The duration of treatment will be determined by the specialist based on clinical response and tolerability.** **Adults** An initial test dose of 10 mg should be given in the first week followed by weekly doses of 50 mg until signs of remission occur (normally after a total dose of 300mg to 500mg). At this point 50 mg doses should be given at two week intervals until full remission occurs. With full remission, the interval between injections should be increased progressively to three, four and then, after 18 months to 2 years, to six weeks. If after reaching a total dose of 1 g (excluding the test dose), no major improvement has occurred and the patient has not shown any signs of gold toxicity, six 100 mg injections may be administered at weekly intervals. If no sign of remission occurs after this time other forms of treatment are to be considered.
	All dose or formulation adjustments will be the responsibility of the initiating specialist unless directions have been discussed and agreed with the primary care clinician

<table>
<tr><td></td><td colspan="2">Dose increases should be monitored by FBC creatinine/eGFR, ALT and/or AST and albumin every 2 weeks for 6 weeks after the dose increase, then revert back to previous schedule.</td></tr>
<tr><td></td><td colspan="2">Termination of treatment will be the responsibility of the specialist. Treatment continues up to a maximum of 5 years after remission</td></tr>
<tr><td>5. Baseline investigations, initial monitoring and dose titration to be undertaken by specialist</td><td colspan="2">Baseline
Height, weight, BP, FBC, creatinine/eGFR, ALT and /or AST, albumin, urinalysis.
Vaccinations against pneumococcus and influenza are recommended.
Shingles vaccine (Zostavax) is recommended as per the JCVI for eligible patients.
Specialist to highlight in the first clinic letter notifying the GP of the decision to initiate DMDs that the GP will need to give the shingles vaccine if the patient is older than 69 years and the pneumococcal vaccine if this has not already been given. The GP should also be advised to add the patient to the influenza vaccine list.
Patients should be assessed for comorbidities that may influence DMD choice, including evaluation of respiratory disease and screening for occult viral infection.
Initiation
FBC, creatinine/eGFR, ALT and /or AST and albumin every 2 weeks until on stable dose for 6 weeks;
Once on stable dose, monthly FBC, creatinine/eGFR, ALT and /or AST and albumin for 3 months.
Thereafter, FBC, creatinine/eGFR, ALT and/or AST and albumin at least every 12 weeks.

N.B. Patients receiving gold therapy should have urinalysis for blood and protein prior to each dose</td></tr>
<tr><td rowspan="3">6. Ongoing monitoring requirements to be undertaken by primary care.</td><td>Monitoring</td><td>Frequency</td></tr>
<tr><td>Urinalysis for blood and protein</td><td>Before each injection</td></tr>
<tr><td>FBC, Creatinine/Egfr, ALT and/or AST, Albumin

CRP and ESR (rheumatology patients only)</td><td>After the initial monitoring period (see section 5), every 12 weeks, or more frequently in patients at higher risk of toxicity as advised by the specialist team. NB: Some of the initial monitoring (likely to be 1-2 months of monthly monitoring) may take place in primary care. The exact frequency of the monitoring to be communicated by the specialist in all cases.</td></tr>
<tr><td rowspan="4">7. Pharmaceutical aspects</td><td>Route of administration</td><td>Intramuscular injection</td></tr>
<tr><td>Formulation</td><td>Sodium aurothiomalate 20mg/ml 0.5ml amps (10mg) and 100mg/ml 0.5ml amps (50mg)</td></tr>
<tr><td>Administration details</td><td>It should be given by deep intramuscular injection and the area gently massaged.</td></tr>
<tr><td>Other important information</td><td>Sodium aurothiomalate should be discontinued in the presence of blood</td></tr>
</table>

			disorders, gastro-intestinal bleeding (associated with ulcerative enterocolitis), or unexplained proteinuria (associated with immune complex nephritis) which is repeatedly above 300 mg/litre.
8. Contraindications **Please note this does not replace the Summary of Product Characteristics (SPC) and should be read in conjunction with it.**	Pregnancy Patients with gross renal or hepatic disease, a history of blood dyscrasia, exfoliative dermatitis or systemic lupus erythematosus.		
9. Significant drug interactions	For a comprehensive list consult the BNF or Summary of Product Characteristics. SPC Seek advice from the initiating Specialist if there are any concerns about interactions.		
10. Adverse Effects and managements	Result		Action
	Abnormal bruising or severe sore throat		Stop drug until FBC results available, contact Specialist Nurse (SN)
	Fall in WCC <3.5 x 10^9/l Fall in neutrophils <1.6 x 10^9/l Fall in platelets <140 x 10^9/l		Stop drug. SN for advice and management.
	Increased MCV >105f/l		Check folate, B12 & TSH. Treat if abnormal but contact SN for advice if normal.
	Unexplained reduction in albumin <30g/l		Stop drug. Contact SN
	Abnormal LFTs – AST or ALT > 100 U/l		Stop drug. Contact SN
	Rash		Stop drug and contact SN.
	Mouth ulcers		Stop drug and contact SN.
	Nausea, vomiting, diarrhoea		Discuss with SN
	Increase in serum creatinine >30% over period of 12 months or less OR decline in eGFR > 25%		Contact SN if there is new or unexplained renal impairment
	Proteinuria:	Trace	Continue drug
		+	Send MSU to exclude infection
		++/+++	Send MSU. If negative - stop drug and contact SN
	Haematuria:	Trace	Continue drug
		+	Send MSU to exclude infection
		++/+++	Stop drug. Send MSU and contact SN
11. Advice to patients and carers	The specialist will counsel the patient with regard to the benefits and risks of treatment and will provide the patient with any relevant information and advice, including patient information leaflets on		

	individual drugs.
12. Pregnancy and breast feeding	Female patients receiving sodium aurothiomalate should be instructed to avoid pregnancy. Pregnant patients should not be treated with sodium aurothiomalate. Significant amounts of gold are excreted in breast milk so lactating mothers under treatment should not breast feed their infants.
13. Specialist contact information	See appendix 2
14. Additional information	**Where patient care is transferred from one specialist service or GP practice to another, a new shared care agreement must be completed.**
15. References	BSR monitoring guidelines
16. To be read in conjunction with the following documents.	1. Policy for Shared Care 2. Shared care agreement. When two or more DMDs are initiated, one shared care agreement form should be completed for all relevant drugs.

Appendix 1

Policy for Shared Care

Shared care is only appropriate if it provides an optimum solution for the patient and it meets the criteria outlined in the Shared Care section of the Pan Mersey **Definitions and Criteria for Categorisation of Medicines in the Pan Mersey Formulary** document.

Before prescribing responsibilities are transferred to primary care:

- Prescribing responsibility will only be transferred when the consultant and the patient's GP agree that the patient's condition is stable.
- All information required by the shared care framework for the individual medicine has been provided to the patient's GP.
- Patients will only be referred to the GP once the GP has agreed to the Shared Care Agreement and returned signed copies.

Inherent in any shared care agreement is the understanding that participation is at the discretion of the GP, subject to the availability of sufficient information to support clinical confidence.

Specialist Responsibilities in Shared Care

- To initiate the medicine, prescribe, monitor for toxicity and efficacy as described by the shared care framework until the patient is stabilised.
- To ensure the patient or their carer:
    - Is counselled with regard to the risks and benefits of the medicine.
    - Provide any necessary written information to the patient with regard to the individual medicine including patient information leaflets on individual drugs.
    - Obtain and document informed consent from the patient when any medicines is prescribed for an off-label indication for any condition
- To be familiar with the shared care framework.
- To provide all information to the patient's GP as required by the shared care framework when prescribing responsibility is initially transferred and at any subsequent times as necessary for safe and effective treatment of the patient.
- To assess the patient regularly as necessary for the duration of therapy.
- To review the patient promptly if required by the GP concerned.

- To meet any additional requirements as required by the individual medicine shared care framework.
- To communicate failure of a patient to attend a routine hospital review and advise the GP of appropriate action to be taken.
- **Addition of a second DMD**: Following the addition of a new drug to an existing regime covered by a Shared Care Agreement, the Specialist must initiate, prescribe and monitor the new drug in accordance with the relevant shared care agreement including subsequent review and inform the GP of this. A new Shared Care Agreement must then be initiated for the new drug.

Primary Care Responsibilities in Shared Care

- To reply to a written request for Shared Care within 21 days ensuring both copies of the Shared Care Agreement are signed if appropriate.

If agreeing to shared care, the GP is asked to:

- To provide prescribe or manage and monitor the medicine as advised by the Specialist and in line with the individual Shared Care Framework.
- To review the patient as required by the Shared Care Framework
- To make appropriate and contemporaneous records of prescribing and/or monitoring and to note the existence of the Shared Care Agreement on the patient`s clinical record. A READ code of “6652 Shared Care- Specialist/GP” can be used.
- To be familiar with the individual Shared Care Framework.
- To report any adverse effects of treatment to the specialist team.
- To inform the Specialist of any relevant change in the patient’s circumstances.
- To seek Specialist advice as appropriate.
- To meet any additional requirements as required by the individual Shared Care Framework.
- To respond to Specialist communication relating to any change or addition to the patients treatment covered by the Shared Care Agreement.

Appendix 2: Shared Care Agreement

Disease modifying drugs (DMDs)

Request by Specialist Clinician for the patient's GP to enter into a shared care agreement

Part 1
To be signed by Consultant / Prescribing member of Specialist Team

Date _____

Name of patient_____

Address _____

_____

Patient NHS No _____

Patient hospital unit No _____

Diagnosed condition _____

If using addressograph label please attach one to each copy

Dear Dr_____

I request that you prescribe

(1) _____

(2) _____

(3) _____

(4) _____

for the above patient in accordance with the enclosed shared care framework.

Last Prescription Issued: ….. / ….. / ….. Next Supply Due: ….. / ….. / …..

Date of last blood test: ….. / ….. / ….. Date of next blood test: ….. / ….. / …..

Frequency of blood test: ……

I confirm that the patient has been stabilised and reviewed on the above regime in accordance with the Shared Care Framework and Policy. ☐

I confirm that if this is a Shared Care Agreement for a drug indication which is unlicensed or off label, informed consent has been received. ☐ N/A

Details of Specialist Clinicians

Name _____Date_____

Consultant / Prescribing member of the Specialist Team ***circle or underline as appropriate***

Signature _____

In all cases, please also provide the name and contact details of the Consultant.

When the request for shared care is made by a prescriber who is not the consultant, it is the supervising consultant who takes medico-legal responsibility for the agreement.

Consultant: _____

Contact details*:*

Telephone number: _____Ext:_____

Address for return of documentation _____

_____

_____

Part 2
To be completed by Primary Care Clinician

I agree to prescribe _____for the above patient in accordance with the enclosed shared care framework.

GP signature _____Date _____

GP name _____Please print

GP: *Please sign and return a copy* ***within 21 calendar days*** *to the address above*

OR

GP- If you do not agree to prescribe, please delete the section above and provide any supporting information as appropriate below: