## LITHIUM – Monitoring Summary

Appendix Three – Monitoring					
Parameters.					
Parameter	Frequency	Result	Action for Primary Care		
As well as responding to absolute values in laboratory tests, a rapid change or a consistent trend in any value should prompt caution and extra vigilance.					
Plasma lithium level taken 10-	At least every 12 weeks for the	12-hour plasma lithium level.	Assess adherence, including discussion with patient and check of GP		
14 hours post-dose. NB:	first year, then every 6 months.	Below target range	clinical systems. Offer advice on adherence if appropriate (e.g. daily		
samples should be taken as	More frequent long-term	NB: range for each patient to be	routines, reminders). Ensure level was taken 12 hours after lithium		
close to 12-hours post-dose as	monitoring may be advised by	determined by the specialist team.	dose.		
possible.	the specialist team in some	Note that local reference ranges may	Contact specialist team for advice if suspected that the dose is too		
Record results in the	circumstances (e.g. elderly,	vary	low.		
patient's record as well as	renal impairment, altered	Above target range	Ensure level was taken 12 hours after lithium dose and that the		
patient-held purple lithium	laboratory parameters, poor	NB: range for each patient to be	correct dose has been prescribed and taken. Check for interactions,		
pack, or other suitable	symptom control or adherence,	determined by the specialist team.	hydration, patient's physical and mental status, and features of		
recording mechanism.	concurrent interacting	Note that local reference ranges may	toxicity. Repeat level if necessary.		
It is advisable to document the	medicines) or if most recent 12-	vary	Withhold lithium if there are features of toxicity. Contact specialist		
actual time interval between	hour plasma lithium level is at		team for advice in all cases.		
the last dose and the blood	the threshold of target range.		If ≥2.0mmol/L – consider sending patient to A&E, based on clinical		
sample	Consider additional monitoring		presentation (e.g. features of toxicity) and inform specialist team.		
	whenever there is a change in	Within target range but toxicity	Contact specialist team for advice.		
	the patient's circumstances,	suspected	Referral to secondary care may be required depending on the		
	e.g. intercurrent illness.	NB: range for each patient to be	severity of symptoms and the certainty of toxicity. Use clinical		
		determined by the specialist team.	judgement to determine the urgency of referral.		
		Note that local reference ranges may			
		vary			
		Within target range but marked	Establish whether level was taken 12 hours after lithium dose.		
		change since last level (and there has	Repeat level with an urgency determined by clinical judgement.		
		been no dose change)	Assess adherence, including discussion with patient and check of GP		
		NB: range for each patient to be	clinical systems. Offer advice on adherence if appropriate (e.g. daily		
		determined by the specialist team.	routines, reminders).		
		Note that local reference ranges may	More frequent monitoring may be required.		
		vary	more requent monitoring may be required.		
• TFTs	Every 6 months.	Thyroid function	Contact specialist team for advice.		
	More frequent monitoring	Altered TFTs without symptoms	During lithium treatment, TFTs are commonly abnormal; the TSH		
	(particularly renal function)	Altered if is without symptoms	can rise early in treatment but settle with time.		

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	may be advised by the specialist team in some circumstances (e.g. elderly, renal impairment, altered TFTs, concurrent interacting medicines).	Subclinical <u>hypo</u> thyroidism • Raised TSH • Normal T4 Clinical features not overtly manifest Overt <u>hypo</u> thyroidism • High TSH • Low T4 Symptomatic <u>Hyper</u> thyroidism	Note that the symptoms of hypothyroidism can be difficult to discriminate from depression and common side effects of lithium.         Contact specialist team for advice, which may include input from endocrinology services.         Thyroid hormone replacement is usually indicated and often continued throughout the course of lithium treatment.         Contact specialist team for advice, which may include input from endocrinology services.
Parameter	Frequency	Result	Action for Primary Care
• eGFR	<b>Every 6 months.</b> More frequent monitoring (particularly renal function) may be advised by the specialist team in some circumstances (e.g. elderly, renal impairment, altered TFTs, concurrent interacting medicines).	<ul> <li>eGFR &lt;45ml/min</li> <li>rapidly falling eGFR</li> <li>gradual decline in eGFR</li> </ul>	The response to impaired or deteriorating renal function should be individualised. Contact specialist team for advice, which may include input from nephrology services. A cardiovascular risk profile may guide specialist advice and should be provided if available. Use clinical judgement to determine the urgency of consultation. Anticipate the need for increased monitoring as trends in renal function are more useful than absolute values. In the elderly or those at the extremes of muscle mass, creatinine clearance provides a better estimate of renal function than eGFR. Adjustments to dose may be advised. If renal function is significantly compromised, lithium may no longer be an appropriate treatment and specialists will advise accordingly.
		Renal function Polyuria and polydipsia	Polyuria is common with lithium and often well tolerated. Advise the patient to maintain adequate fluid intake and advocate excellent oral hygiene.

			Contact specialist team for advice, which may include input from nephrology services. In some instances, dose adjustment or specific treatments may be advocated.
<ul> <li>U&amp;Es</li> <li>Calcium</li> </ul>	Every 6 months. More frequent monitoring (particularly renal function) may be advised by the specialist team in some circumstances (e.g. elderly, renal impairment, altered TFTs, concurrent interacting medicines).	U&Es or calcium out of range	Check that the most recent 12-hour plasma lithium level is in the desired range and act accordingly if not. Determine whether there are symptoms and signs related to the electrolyte disturbance or lithium toxicity. Consider arranging an ECG in those at risk for QT prolongation. Contact specialist team for advice. Changes in calcium levels may reflect parathyroid dysfunction and input from endocrinology services may be indicated.
• Height, weight, and BMI.	<b>Every 6 months.</b> More frequent monitoring (particularly renal function) may be advised by the specialist team in some circumstances (e.g. elderly, renal impairment, altered TFTs, concurrent interacting medicines).	Weight and BMI Outside healthy range	<ul> <li>Provide appropriate support on multicomponent interventions to increase physical activity levels, improve eating behaviour and quality of diet. Remind patient of the importance of maintaining adequate fluid intake and avoiding dehydration while exercising.</li> <li>Consider measuring waist circumference for individualised monitoring.</li> <li>Patients should be instructed to avoid sudden changes in diet, especially avoiding low sodium diets. Lithium levels are influenced by body weight and so for patients being supported to lose weight, lithium levels may need to be checked more frequently (akin to other situations of caution). Use clinical judgement, lithium levels and the rate of weight loss when determining the frequency of blood tests.</li> <li>If rapid weight gain – discuss with specialist</li> </ul>
Additional monitoring – bipolar o	disorder		
Diet, nutritional status and level of physical activity, Cardiovascular status including pulse and BP, LFTs, Metabolic status including fasting blood glucose, HbA <sub>1c</sub> and blood lipid profile.		Annually as part of physical health check recommended in NICE <u>CG185 Bipolar disorder: assessment and</u> <u>management</u> .	
		Any physical health problems should be treated by the appropriate primary care health professional and communicated to the specialist team within 14 days.	

Title	LITHIUM - Monitoring Summary		
Description of policy	To inform healthcare professionals		
Scope	Norfolk and Waveney Integrated Care System		
Prepared by	Norfolk and Waveney ICB Medicines Optimisation Team		
Impact Assessment (Equalities and Environmental)	Please indicate impact assessment outcome: Positive impact Adverse impact - low - action plan completed as per guidance		
	Adverse impact - medium - action plan completed as per guidance Adverse impact - high - action plan completed as per guidance No impact <b>No policy will be approved without a completed equality impact assessment</b>		
Other relevant approved documents	Lithium shared care agreement		
Evidence base / Legislation	Level of Evidence: <i>A. based on national research-based evidence and is considered best evidence</i> <i>B. mix of national and local consensus</i> <i>C. based on local good practice and consensus in the absence of national research based information.</i>		
Dissemination	Is there any reason why any part of this document should not be available on the public web site?		
Approved by	Norfolk & Waveney Therapeutics Advisory Group (TAG) July 2025		
Authorised by	Medicines Optimisation Programme Board on behalf of the ICS (July 2025)		
Review date and by whom	Medicines Optimisation Team – July 2027		
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Version Number	Author	Purpose / Change	Date
1.0	MO Team	To support prescribing	July 2025