

NHS Grampian Staff Guideline For The Management Of Acute Hypokalaemia In Adults

Co-ordinators:	Consultation Group:	Approver:
Medicines Information Pharmacist	See relevant page in guidance	Medicine Guidelines and Policies Group
Signature:		Signature:
2	· · · · · · · · · · · · · · · · · · ·	
Identifier:	Review Date:	Date Approved:
NHSG/Hypokal/MGPG1201	November 2024	November 2021

Uncontrolled when printed

Version 4

Executive Sign-Off

This document has been endorsed by the Director of Pharmacy and Medicines Management

Signature:

Title: NHS Grampian Staff Guideline For The Management Of

Acute Hypokalaemia In Adults

Unique Identifier: NHSG/Hypokal/MGPG Version 4

Replaces: NHSG/Hypokal/MGPG905, Version 3.2

Across NHS Boards	Organisation Wide	Directorate	Clinical Service	Sub Department
				Area

This controlled document shall not be copied in part or whole without the express permission of the author or the author's representative.

Lead Author/Co-ordinator: Medicines Information Pharmacist

Subject (as per document registration categories):

Clinical Guidelines

Key word(s): Guideline hypokalaemia potassium management adults

serum K sodium chloride KCL glucose

Process Document: Policy,

Protocol, Procedure or

Guideline

Guideline

Document application: NHS Grampian

Purpose/description: To guide the management of hypokalaemia in adults.

Responsibilities for implementation:

Organisational: Chief Executive and Management Teams

Corporate: Senior Managers

Departmental: Heads of Service/Clinical Leads

Area: Line Managers

Hospital/Interface services: Assistant General Managers and Group Clinical Directors

Operational Management Unit Operational Managers

Unit:

Policy statement: It is the responsibility of all staff to ensure that they are

working to the most up to date and relevant policies,

protocols procedures.

Review: This policy will be reviewed in three years or sooner if

current treatment recommendations change.

Responsibilities for review of this document: Medicines Information Pharmacist

Responsibilities for ensuring registration of this document on the NHS Grampian Information/ Document Silo:

Pharmacy and Medicines Directorate

Physical location of the original of this document:

Medicines Information Office, ARI

Job/group title of those who have control over this document:

Medicines Information Pharmacist

Responsibilities for disseminating document as per distribution list:

Medicines Information Pharmacist

Revision History:

Revision Date	Previous Revision Date	Summary of Changes (Descriptive summary of the changes made)	Changes Marked* (Identify page numbers and section heading)
June 2021	May 2018	Clarified wording around continuous ECG	P3 - Table
August 2021	May 2018	Added detail on choice of dose	P2 - Dosage
August 2021	May 2018	Reduced maximum rate for peripheral line	P3 – Table

^{*} Changes marked should detail the section(s) of the document that have been amended, i.e. page number and section heading.



NHS Grampian Staff Guideline for the Management of Acute Hypokalaemia in Adults

This guideline is for use within Primary or Secondary Care in NHS Grampian. Intravenous potassium replacement should only be used in an acute setting, as outlined below.

The NHS Grampian reference range for serum potassium in patients over 16 years of age is 3.5 - 5.3mmol/L.

POTASSIUM SUPPLEMENTS SHOULD NOT BE GIVEN IN SEVERE RENAL IMPAIRMENT, OR IF SERUM POTASSIUM IS GREATER THAN 5.0mmol/L¹

Table 1 – Serum Potassium Classification of Hypokalaemia

Normal	Mild	Moderate	Severe
3.5 - 5.3mmol/L	3.0 - 3.4mmol/L	2.5 - 2.9mmol/L	< 2.5mmol/L

Causes of Hypokalaemia^{2,3}

- High urine flow rate and distal sodium delivery, e.g. loop diuretics and thiazides, uncontrolled diabetes.
- External losses from the GI tract, e.g. vomiting, aspiration, fistulae, chronic diarrhoea, laxative abuse, gastric suction.
- Hypomagnesaemia potassium can be difficult to correct until magnesium levels are normalised.
- Mediated by mineralocorticoid receptor (often associated with hypertension)
 e.g. primary aldosteronism, secondary aldosteronism, Cushing's syndrome,
 steroid therapy, carbenoxolone and glycyrrhizinic acid (liquorice.
- Other medications: theophylline intoxication, insulin, B2 agonists (e.g. salbutamol and terbutaline).
- Reduced intake, e.g. inadequate dietary intake, potassium-free intravenous fluids.
- Sequestration of fluid in bowel, e.g. ileus, intestinal obstruction.
- Defective proximal reabsorption of potassium, e.g. recovery phase of acute tubular necrosis, after relief of urinary tract obstruction, proximal renal tubular acidosis, tubular damage by drugs, e.g. amphotericin.
- Shift of potassium into cells, e.g. metabolic alkalosis.
- Refeeding Syndrome.
- Artefactual: Prolonged contact of serum with cells (i.e. delayed sample transit) may cause measured levels to fall as well as rise.

Signs and Symptoms of Hypokalaemia^{2,3}

(Mild hypokalaemia rarely causes symptoms)

- Muscular weakness (possibly paralysis and respiratory failure) and cramping
- Reduced intestinal motility or paralytic ileus
- Polyuria
- ECG changes, ventricular arrhythmias or asystole.

Dosage, Administration and Monitoring^{2,7}

- When hypokalaemia is severe with marked clinical features, or unresponsive to oral therapy, potassium must be replaced intravenously. The speed of replacement is based on clinical symptoms.
- The presence of life-threatening emergency such as serious cardiac dysrhythmia or paralysis requires rapid correction. Otherwise, slow intravenous replacement is preferable to avoid induction of hyperkalaemia.
- The dose required is dependent upon the clinical presentation of the patient. Correction of the underlying cause may suffice when hypokalaemia is mild. For those with severe/symptomatic hypokalaemia, 40mmol/L can be given intravenously; a second or third dose may be required. A recommended maximum dose is 2-3mmol/kg of potassium in 24 hours. The recommended maximum should not be considered a target dose; such high doses may be inappropriate in larger patients given the likely infusion volume required. See Table 2 for more information on dosing.
- Consider changing loop diuretics to potassium-sparing agents.
- Caution should be used in patients with renal impairment or when ACE inhibitors or potassium-sparing diuretics are being administered concomitantly

Side Effects of Potassium Administration 1,4

Oral

- Nausea and vomiting, abdominal pain, diarrhoea, flatulence if these occur, give dose with or after food.
- Hyperkalaemia.

Intravenous⁶

- Thrombophlebitis and pain.
- Tissue damage in the case of extravasation.
- Cardiac arrhythmias (ECG monitoring if rapid administration (i.e.
 >20mmol/hour) or high concentration used (i.e. >40mmol/L)), heart block, cardiac arrest.
- Hyperkalaemia.

Suggested monitoring intervals³

Serum potassium range 3.0-3.5 mmol/L Monitor serum potassium twice weekly until stable. Once stable or potassium >4.5mmol/L, reassess need for supplementation.

Serum potassium ≤2.9 mmol/L When serum potassium is < 3mmol/L, intravenous supplementation is usually required. This must only be administered in a hospital setting. Monitor serum potassium following initial therapy, and then at least daily until serum potassium >2.9mmol/L then manage as above. More frequent monitoring may be required depending on the patient's clinical condition. See <u>Table 2</u>.

Table 2: Administration and Dosage 1-8

Please note: In the table below K = potassium and CI = chloride

Route of	Presentation	Adult Dose	Notes
administration Oral			
Oral	Potassium chloride and bicarbonate effervescent tablets (Sando K)	Serum K range 3.0 to 3.5mmol/L TWO tablets THREE times per day (72mmol K per day) Serum K range 2.5 to 2.9mmol/L THREE tablets THREE times per day (108mmol K per day)	Each tablet contains 12mmol K and 8mmol Cl. Tablets dissolve in water which minimises local high concentrations and therefore possibly less risk of adverse effects. May be unpalatable. Give with food to minimise GI irritation.
D. Giorfeei en	Potassium chloride 7.5% sugar free syrup (Kay-Cee-L)	Serum K range 3.0 to 3.5mmol/L 20mL THREE times per day (60mmol K per day) Serum K range 2.5 to 2.9mmol/L 30mL THREE times per day (90mmol K per day)	Liquid contains 1mmol/mL of K and Cl. Take with food.
IV infusion			
Intravenous supplements are indicated if the patient cannot eat, is vomiting, is unlikely to absorb potassium orally or has profound	For available preparations see <u>Table 3</u>	Serum K range 2.5 to 3.5mmol/L Standard concentration via PERIPHERAL line Rate of administration is based on the clinical picture. Maximum rate of administration: 10mmol/hour Maximum concentration: 20mmol/500mL	Hypokalaemia should be interpreted in terms of fluid balance. Any disturbances in acid-base balance or hypomagnesaemia should be corrected where appropriate. Continuous ECG monitoring is mandatory for infusion
hypokalaemia		Serum K range <2.5mmol/L High concentration via PERIPHERAL line Use in fluid restriction or emergencies only. Give via a large vein under the direction of a consultant. Prescribe concentration required, up to 40mmol/500mL. Maximum concentration: 40mmol/500mL. Usual maximum rate 20mmol/hour Maximum of 2 hours duration then review. Further therapy is guided by serum potassium levels. High concentration via CENTRAL line For use in ICU or HDU only. For safety, prescribe as dilute as the patient's fluid status will reasonably allow. Maximum concentration 1mmol/mL. Usual maximum rate 20mmol/hour Maximum of 2 hours duration then review. Further therapy is guided by serum potassium levels.	

Table 3: Parenteral potassium preparations in sodium chloride 0.9% available in NHS Grampian

Preparation (infusion fluid and potassium chloride (KCL))	Millimoles (grams) of potassium chloride per bag	
	500mL bag	1000mL bag
Sodium chloride 0.9% + 0.15% KCL	10mmol (0.75g)	Not stocked
Sodium chloride 0.9% + 0.3% KCL	20mmol (1.5g)	Not stocked
Sodium chloride 0.9% + 0.6% KCL	40mmol (3g)	Not stocked

Please note: Solutions containing glucose are less suitable for the initial correction of hypokalaemia. Glucose containing solutions may reduce serum potassium concentrations.

Table 4: Parenteral potassium preparations containing glucose available in NHS Grampian

Preparation (infusion fluid and potassium chloride (KCL))	Millimoles (grams) of potassium chloride per bag	
	500mL bag	1000mL bag
Sodium chloride 0.18% + Glucose 4% + 0.15% KCL	Not stocked	20mmol (1.5g)
Sodium chloride 0.18% + Glucose 4% + 0.3% KCL	Not stocked	40mmol (3g)
Sodium chloride 0.45% + Glucose 5% + 0.15% KCL	10mmol (0.75g)	Not stocked
Sodium chloride 0.45% + Glucose 5% + 0.3% KCL	20mmol (1.5g)	Not stocked
Glucose 5% + 0.15% KCL	Not stocked	20mmol (1.5g)
Glucose 5% + 0.3% KCL	20mmol (1.5g)	Not stocked
Glucose 5% + 0.6% KCL	40mmol (3g)	Not stocked
Glucose 10% + 0.15% KCL	10mmol (0.75g)	Not stocked
Glucose 10% + 0.3% KCL	20mmol (1.5g)	Not stocked

Consultation Group

Sarah O'Beirne Lead Pharmacist Medicines Information Lynne Davidson Clinical Pharmacist Cardiology Dr Adelle Dawson Cardiology Consultant Dr Iain Macleod ITU Consultant

References

- 1. British National Formulary. Online. Accessed via medicinescomplete.com on 06/11/2020
- 2. The Merck Manual, R. Porter, K. Kaplan, 19th Edition, 2011
- 3. Handbook of Clinical Medicine 8th Edition
- 4. Summary of Product Characteristics Sando K, access online at http://www.medicines.org.uk/emc/medicine/812 on 05/01/2021

- 5. Summary of Product Characteristics Kay-Cee-L Syrup, accessed online at https://mhraproducts4853.blob.core.windows.net/docs/b52817499cad6c2f30b95 abe0fdfc6741fc5b4f0 on 05/01/2021
- 6. Medusa Injectable Medicines Guide, potassium chloride monograph. Accessed online at http://medusa.wales.nhs.uk/ on 18/02/2021
- 7. UKMi Q&A How should intravenous (IV) potassium chloride be administered in adults? G. Woodland. Published 20th September 2020. Accessed online at https://www.sps.nhs.uk/articles/how-should-intravenous-iv-potassium-chloride-be-administered-in-adults/
- 8. Minimum Infusion Volumes for fluid restricted critically ill patients. UKCPA Critical Care Group. 4th Edition. December 2012. Available online at https://www.scottishintensivecare.org.uk/uploads/2014-07-24-19-56-30-Minimuminfusionvolumesinl-40262.pdf