

Hyperkalemia

Inpatient

Center for



Pre-Pathway Validation

Is this Hyperkalemia?

Hyperkalemia is an elevated plasma or serum potassium level. Diagnosis is based on K+ >5.5 mEq/L on a non-hemolyzed specimen. Please note reference range is for patients \geq 90 days of age; a level up to 5.9 mEq/L can be normal in infants.

Disease Severity

Mild hyperkalemia: 5.5-5.9 mEq/L Moderate: 6-6.9 mEq/L Severe: ≥7 mEq/L

Common diagnoses associated with hyperkalemia

- latrogenic potassium supplementation (fluids or medications)
- Acute kidney injury
- Chronic kidney disease
- Decreased effective arterial volume (dehydration)
- Tubular dysfunction
- Hemolysis
- Malignancy
- Infection
- Sickle cell disease
- Metabolic acidosis

Pathway Inclusion Criteria

 Patients ≥90 days with potassium >5.5 mEq/L on nonhemolyzed sample.

Pathway Exclusion Criteria

- Neonates <90 days
- Pt in NICU, PICU, or CTICU
- Renal disease requiring dialysis
- Tumor lysis syndrome
- Diabetic Ketoacidosis

Always stop potassium containing fluids in children with moderate to severe hyperkalemia. Also hold medications that could increase potassium, unless risk outweigh benefit. **Differential Diagnosis**

- Pseudohyperkalemia i.e. a high serum potassium level that does not reflect the true in vivo level.
- Causes:

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- In vitro hemolysis
- Fist clenching during phlebotomy
- Undue delay in processing blood samples
- Inappropriate storage temperature of blood samples
- Potassium contamination
 of blood samples
- Always ensure specimen is not hemolyzed prior to proceeding to treatment

Expected EKG Changes by Potassium Level

Changes may not progress in a stepwise or predictable fashion

5.5-6.4 mEq/L

- Peaked T waves
- Normal or decreased QT
- Prolonged PR interval.

6.5-8.0 mEq/L

- Widening of QRS complex
- Prolonged PR interval
- Broad, low amplitude P waves
- QT prolongation
- ST elevation or depression

<u>>8 mEq/L</u>

- P waves disappear
- Marked widening of QRS
- "Sine wave" pattern
- High risk of ventricular fibrillation or asystole.



Return to Algorithm

Adapted from Sood et al. 2007

Symptoms of Hyperkalemia

Symptoms are rare, even in severe hyperkalemia

- Palpitations
- Nausea
- Muscle pain
- Parethesias

Medications and fluids that can increase potassium

- IV fluids that contain K (including TPN, LR, and IV potassium replacement)
- Oral potassium supplements (KCL, KPhos, K iodide, KAce, KBicarb etc.)
- Penicillin K
- Potassium sparing diuretics (spironolactone, amiloride)
- Calcium channel blockers (amlodipine, nifedipine)
- Angiotensin-converting enzyme (ACE) inhibitors (lisinopril) and Angiotensin II Receptor Blockers (ARBs)
- NSAIDs
- Heparin
- Pentamidine
- Trimethoprim
- Propranolol
- Immunosuppressants decision to hold immunosuppression should be carefully evaluated with guidance from appropriate/managing provider
 - Cyclosporine
 - Tacrolimus

Medications Risk vs. Benefit

		Benefit	Risk
First Line Medications	Calcium Gluconate	 Rapid onset of action to stabilize cardiac membrane and prevent cardiac arrhythmias Can be administered via peripheral IV 	(1) Does not decrease potassium serum concentration(2) Duration lasts 30-60 min, repeat doses may be necessary
	Albuterol	 Lowers serum potassium by shifting K intracellularly Available for rapid administration via inhalation Onset of action within 30 min Increases the efficacy of insulin in lowering potassium 	(1) Effect lasts only 2 hours, repeat dosing is often necessary(2) Rebound hyperkalemia is common
	Insulin + Dextrose	 Lowers serum potassium by shifting K intracellularly Produces a reliable shift of potassium into cells within 15 minutes (peak 30-60 min) that lasts 2-6 hours Works synergistically with albuterol 	(1) Hypoglycemia may occur despite administration of dextrose; blood glucose should be monitored every hour while on insulin
	K-Cocktail	 Standard mixture of calcium gluconate, insulin, dextrose, and sodium acetate that reliably shifts K intracellularly Continuous infusion can maintain serum K up to 2 mEq/L lower over 12-24 H 	 (1) Central line is preferred administration route due to high glucose concentration (though it may be administered peripherally) (2) Rapid drop in potassium may result in hypokalemia
Second Line Medications	Calcium chloride	 Rapid onset of action to stabilize cardiac membrane and prevent cardiac arrhythmias 	 Does not decrease potassium serum concentration Duration lasts 30-60 min, repeat doses may be necessary Central line required for administration
	Furosemide (Lasix)	(1) Total body excretion of potassium	 Typically, several hours until notable effect makes it not ideal for acute treatment Amount of potassium excreted is unpredictable Patient must be producing urine for effect
	Sodium Polystyrene sulfonate (Kayexalate [™])	(1) Binds potassium to remove via the gut	 Delayed and highly variable onset of action Intestinal side effects are common Can cause gut necrosis; should be avoided in infants and those with possible altered gut mobility

Doses of Medications to treat Hyperkalemia

First Line Medications						
Calcium gluconate (preferred)	60-100 mg/kg/dose IV (MAX 3000 mg) over 10 minutes	Repeat doses may be necessary				
Calcium chloride (if central line)	20 mg/kg/dose (MAX 1000 mg) over 10 minutes	Repeat doses may be necessary				
Insulin regular + dextrose	Insulin: 0.1 unit/kg/dose IV (MAX 10 units) over 1-2 minutes WITH Dextrose: 0.5-1 g/kg IV (5-10 mL/kg if using D10%) over 30 minutes	Consider max dose of 5 units of insulin in children with chronic kidney disease				
	< 25 kg: 2.5-5 mg nebulized over 10 minutes	Repeat Q2H PRN				
Albuterol	25-50 kg: 5-10 mg nebulized over 10 minutes	Repeat Q2H PRN				
	> 50 kg: 10-20 mg nebulized over 10 minutes	Repeat Q2H PRN				
K-Cocktail (Calcium gluconate 3000 mg, Dextrose 27%, Insulin Regular 30 units, Sodium acetate 100 mEq)	2 mL/kg/hr for the first hour followed by 0.5-1 mL/kg/hr	Due to its hyperosmotic content, the K-Cocktail is best administered by central line, but it may be given peripherally temporarily for urgent treatment				
Second Line Medications use with consultation of ICU or Nephrology Only						
Sodium Polystyrene Sulfonate (Kayexalate [™])	1 g/kg/dose enterally (MAX 30 g)	Q6H PRN				
Furosemide (Lasix [™])	0.5-1 mg/kg/dose IV (MAX 40 mg)	Q6H PRN				

Signs of Deterioration & Escalation of Care

Signs of deterioration:

- Tachycardia, arrhythmia
- Altered mental status
- Development of symptomatic hyperkalemia (nausea, palpitations, myalgias, paresthesias)
- A rise in potassium level despite treatment
- Sustained K > 7 mEq/L despite treatment
- Decreasing urine output or urine output < 1 ml/kg/hr (excluding known anuric or oliguric patients)
- Progression of EKG changes (see "Expected EKG changes by potassium level") regardless of potassium level

Escalation plan:

- Consult ACT team
- Consult Nephrology
- Ensure IV access
- Ensure full disclosure telemetry or serial EKG if not available
- Consider additional treatments: K-cocktail, IV fluid bolus, furosemide, Kayexalate[™] under guidance of experts

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Quality Measures

Goals:

- 1. Prevent or counteract life-threatening cardiac conduction disturbances due to hyperkalemia.
- 2. Efficiently and safely lower potassium to normal level with the least amount of blood draws

Metrics:

Process measures:

- 1. IP Order Set use
- 2. Rate of EKGs obtained for patients with a non-hemolyzed potassium of 6 mEq/L or greater.
- 3. Rate of calcium gluconate or calcium chloride administration for patients with a non-hemolyzed potassium greater than 7 mEq/L.

Outcome measures:

- 1. Time until potassium normalization (potassium less than or equal to 5.5 mEq/L).
- 2. Rate of patients with a potassium greater than 8 mEq/L.
- 3. Mortality rate

Balancing measure:

- 1. Rate of hypoglycemia (blood glucose less than 60) for hyperkalemic patients treated with insulin.
- 2. Number of potassium lab draws per encounter.

Team & Process

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Clinical Pathway Development

This clinical pathway was developed using the process described in the NCH Clinical Pathway Development Manual Version 6, 2022. Clinical Pathways at Nationwide Children's Hospital (NCH) are standards which provide general guidance to clinicians. Patient choice, clinician judgment, and other relevant factors in diagnosing and treating patients remain central to the selection of diagnostic tests and therapy. The ordering provider assumes all risks associates with care decisions. NCH assumes no responsibility for any adverse consequences, errors, or omissions that may arise from the use or reliance on these guidelines. NCH's clinical pathways are reviewed periodically for consistency with new evidence; however, new developments may not be represented, and NCH makes no guarantees, representations, or warranties with respect to the information provided in this clinical pathway.

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