Potassium - ORAL

Newborn use only

Alert	High risk medication in A PINCH Medicines list under New South Wales Clinical Excellence Commission.	
	Perrigo brand contains 1 mg of methyl hydroxybenzoate/1 mL. Uricosal brand also contains	
	hydroxybenzoate. Avoid exposure of >99 mg/kg/day of sodium benzoate in neonates. (1)	
	Oral potassium chloride and potassium citrate solutions are high in osmolality with a reported osmolality	
	of 2200 mOsm/kg (Cytra-K, Cypress Pharmaceuticals, NJ). Therefore, it is recommended to be given with	
	feeds. (4)	
Indication	Potassium chloride:	
	Treatment and prevention of hypokalaemia	
	Potassium citrate and citric acid:	
	Treatment of hypokalaemia in the presence of simultaneous metabolic acidosis	
Action	Intracellular cation. Essential in the maintenance of body fluid composition and electrolyte balance.	
Drug type	Electrolyte	
Trade name	1. Potassium chloride oral mixture 10% w/v by Perrigo	
	2. Potassium citrate and citric acid oral mixture (Uricosal)	
	3. Potassium citrate mixture APF	
Presentation	1. Potassium Chloride oral mixture 10% w/v by Perrigo – 500 mL bottle. Potassium content:	
	20mmol/15mL = 1.33 mmol/1 mL.	
	2. Potassium citrate and citric acid oral mixture (Uricosal): Potassium content: 1.9 mmol/1 mL and	
	citrate monohydrate component: 40mg/1 mL.	
	3. Potassium citrate mixture 200 mg/mL Australian Pharmaceutical Formulary (APF) - compounded in-	
	house by pharmacy – Refer to local hospital policies.	
Dose	0.5-1.5 mmol/kg/dose 6-12 hourly (1-6 mmol/kg/day) *	
	*Always prescribe as millimol (mmol) of elemental potassium.	
Dose adjustment	Adjust dose based on serum potassium concentrations.	
•	Renal impairment- increased risk of hyperkalaemia. Avoid in severe renal impairment.	
Maximum dose		
Total cumulative		
dose		
Route	Oral	
Preparation	No preparation required.	
Administration	Give oral doses with feeds to minimise gastric irritation.	
Monitoring	Close monitoring of serum potassium concentrations is needed to avoid hyperkalaemia.	
· · · · · · · · · · · · · · · · · · ·	Clinical status including urine output, creatinine, electrolytes.	
Contraindications	Potassium chloride: Hypersensitivity to any component of the formulation, hyperkalaemia, renal failure,	
	cardiac disease, conditions in which potassium retention is present.	
	Potassium citrate: Hypersensitivity to any ingredient of the formulation, severe renal insufficiency with	
	oliguria or azotaemia, potassium restricted diet, untreated Addison's disease, acute dehydration, anuria,	
	severe myocardial damage, hyperkalaemia.	
Precautions	Use with caution in patients with renal impairment, cardiac disease, acid/base disorders, or potassium-	
	altering medicines/conditions/disorders.	
Drug interactions	Use with caution in patients receiving potassium-sparing diuretics (e.g. spironolactone), medications	
	known to increase risk of hyperkalaemia (e.g. ACE inhibitors) and medications that contain potassium.	
Adverse	Vomiting, abdominal pain, flatulence, GI bleeding, GI obstruction, skin rash, hyperkalaemia.	
reactions	, and a substitution of the substitution of th	
Compatibility	Not applicable.	
Incompatibility	Not applicable.	
Stability	Refer to the product label.	
Storage	Store below 25°C.	
Jiorage	Protect from light.	
Evcinients	Potassium chloride oral mixture 10% w/v by Perrigo – contains glycerol (126 g/100 mL), methyl	
Excipients		
	hydroxybenzoate (100 mg/100 mL), citric acid (0.25 g per 100 mL)(5)	
	Uricosal and APF mixture contains 0.5 mg/1 mL of hydroxybenzoate.(6) Uricosal brand also contains	
	sucrose.	

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Special	
comments	
Evidence	Efficacy Treatment of hypokalaemia There are no reported trials on the efficacy and safety of potassium therapy in hypokalaemia in
	neonates.
	Limited evidence in infants and children suggests enteral potassium replacement may be an equally efficacious alternative first-line therapy in treating hypokalaemia.(2) (LOE II GOR C) Merchant et al (2) performed an open-label randomised trial to study the serum potassium changes with enteral versus IV potassium in hypokalaemic infants and children (aged 1 month to 15 years). In the oral potassium chloride group, the concentration used was 2.66 mmol/1 mL. The parenteral/enteral dose used was 0.1-0.3 mmol/kg dose for serum potassium of 3.5-4.4 mmol/L; 0.5 mmol/kg/dose for serum potassium of 3.0-3.4 mmol/L and 0.7-1.0 mmol/kg/dose for serum potassium of <3.0 mmol/L. There was no statistically significant difference in change in potassium levels after either enteral or parenteral route.
	In Merchant's trial of enteral and intravenous potassium, no mortality was reported in either arm. A few episodes of vomiting were reported in enteral route (2) Pharmacokinetics
	Almost all of potassium ingested through diet is absorbed. The kidneys excrete more than 90% of daily intake and are the organs primarily responsible for the elimination of potassium.(3)
Practice points	The preferred administration of K ⁺ is via the oral/enteral route. However, in the presence of severe symptomatic hypokalaemia and gastrointestinal problems such as ileus, the intravenous route may be used.(3) The normal daily required intake of K ⁺ is 1–2 mEq/kg/day.
	The choice of the type of K ⁺ salt depends on the clinical situation. Potassium chloride is usually appropriate if hypovolemia is present. In the presence of simultaneous metabolic acidosis, other K ⁺ salts producing K ⁺ bicarbonate, K ⁺ citrate, and K ⁺ acetate may be given. The correction of total body K ⁺ deficit may take days and even weeks. In cases of treatment resistant hypokalaemia, hypomagnesemia should be considered. In these cases, K ⁺ levels normalise following magnesium treatment.(3)
References	1. Meyers RS, Thackray J, Matson KL, McPherson C, Lubsch L, Hellinga RC, Hoff DS. Key Potentially Inappropriate Drugs in Pediatrics: The KIDs List. The Journal of Pediatric Pharmacology and Therapeutics. 2020;25(3):175-91.
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