

Bon Secours Richmond
Pharmacy and Therapeutics Committees
Potassium Chloride Infusions Policy Update for Adults
6/2007

Recommendations:

Small Volume Infusion		
	Peripheral Line	Central Line
Recommended Infusion Rate	10 meq/hour	10 meq/hour
Maximum Infusion Rate	20 meq/hour	20 meq/hour
Cardiac Monitoring Required		40 meq/hour*
Recommended Concentration	10 meq/100 ml	10 meq/50 ml
Maximum concentration	0.2 meq/ml 10 meq /50 ml 20 meq /100 ml	0.4 meq/ml 20 meq/50 ml

*If potassium < 3 meq/liter and the patient is symptomatic 40 meq/hour may be administered to intensive care patients. Hourly serum potassium determinations should be drawn to avoid severe hyperkalemia and/or cardiac arrest. Symptoms of hypokalemia include: fatigue, malaise, generalized muscle weakness, respiratory failure, paralysis; EKG changes include T wave flattening or inversion, U waves, or ST segment depression, and arrhythmia's.

Small volume infusions of Potassium chloride are available from the pharmacy in premixed bags of:

- 10 meq/100 ml (Preferred peripheral concentration)
- 10 meq/50 ml (Preferred central concentration)
- 20 meq/100 ml (Cardiac monitoring required, peripheral or central use)
- 20 meq/50 ml (Cardiac monitoring required and central line)

Large Volume Infusion		
	Peripheral Line	Central Line
Maximum concentration large volume solution	0.04 meq/ml 40 meq/L	0.08 meq/ml 80 meq/L
TPN Maximum Concentration	40 meq/l	80 meq/l

Recommended maximum dose should not usually exceed 10 meq/hour or 200 meq for a 24 hour period if the serum potassium level is greater than 2.5 meq/liter per product package insert

Background:

- All potassium infusions must be administered via an infusion pump.
- Potassium is never administered IV push or undiluted from a vial.
- The oral route of administration should be used when feasible except in cases of life-threatening hypokalemia.
- As a general rule, in patients with normal renal function, 10 meq of potassium will increase serum potassium by 0.1 meq/l although studies have demonstrated changes from 0.1-0.3 meq/l. Magnesium levels should be checked and corrected if low concomitantly with the potassium.
- Peak serum levels occur at the end of the infusion and stabilize 15 minutes post infusion.
- Total body content of potassium is approximately 3500 meq (35-45 meq/kg) of which 90% is intracellular.

Serum Potassium (meq/l)	Total Body Deficit (Assuming Normal pH) meq
3	200
2.5	300
2	400
1	400+

- For every 0.1 increase in serum pH, serum potassium will fall by approximately 0.6 and vice versa.
- Patients must be on continuous EKG monitoring if receiving > 10 meq/hour of potassium. Potassium replacement therapy should be guided primarily by serial electrocardiograms. Plasma potassium levels are not necessarily indicative of tissue potassium levels.
- Concentrations of 0.3 meq/ml and higher should be exclusively administered via central route per product insert.
- The premixed small volume infusions should be given over 1 hour. Premixed bags are available from the manufacturer in 30 meq/100ml and 40 meq/100 ml. Pharmacy does not carry these.
- Current product information recommends whenever possible use central line infusion rather than peripheral line, Literature suggests the use of a central line with highly concentrated infusions (30 meq/100 ml, 40 meq/100 ml, 20 meq/50 ml) or with an infusion rate greater than 10 meq/hour. Total dose over 24 hours should not exceed 200 meq. If potassium < 2 meq/liter, can use as much as 400 meq administered over 24 hours.¹
- **PREMIXED POTASSIUM CHLORIDE INFUSIONS WILL BE STOCKED IN PYXIS STATIONS.**
 - In 1998 the Joint Commission suggests that health care organizations not make concentrated potassium chloride available outside of the pharmacy.
 - There are numerous reports in the literature that illustrate the severe consequences of improper use of potassium injection. All the cases have one common factor or root cause: Potassium chloride injection concentrate was kept in medication stock in patient care areas.
- Symptoms of hypokalemia include: fatigue, malaise, generalized muscle weakness, respiratory failure, paralysis; EKG changes include T wave flattening or inversion, U waves, or ST segment depression, and arrhythmia's.

References:

1. May RJ, Phillips SM, Williams DB. Improving the safety of potassium intravenous infusions. P&T 1995;February:104-16
2. Davis NM. Potassium Perils. AJN. 1995;March:14.
3. Dickerson, Roland. Guidelines for the Intravenous Management of Hypophosphatemia, Hypomagnesemia, Hypokalemia, and Hypocalcemia. Hospital Pharmacy. 2001;36:1201-1208.
4. Electrolyte Infusion Guidelines. www.hosp.uky.edu/pharmacy/formulary/criteria/electrolyte.htm. August 2006. Retrieved 6/2007.
5. Kruse JA, Carlson RW. Rapid correction of hypokalemia using concentrated intravenous potassium chloride infusions. Arch Intern Med. 1990;150:613-17
6. Hamill RJ, Robinson LM, Wexler HR, Moote C. Efficacy and safety of potassium infusion therapy in hypokalemic critically ill patients. Crit Care Med. 1991;19:694-99
7. Potassium Chloride Injection. Package Insert. Abbott Laboratories.
8. Kruse JA. Concentrated Potassium Chloride Infusions in Critically Ill Patients with Hypokalemia. J Clin Pharmacol 1994;34:1077-1082
9. Cohn JN. New Guidelines for Potassium Replacement in Clinical Practice. Arch Inter Med. 2000;160:2429-36
10. Swanson DW. Implementing an IV potassium policy. Hospital Pharmacist. 2003;10:348
11. University of Texas Medical Branch, Use of Intravenous Potassium Solutions In Adult Patients
12. University of Kentucky Pharmacy Services, Electrolyte Infusion Guidelines at UK Hospital
13. Vanderbilt University Medical Center, Intravenous Potassium