Hypokalemia

Nephkids electrolyteworkshop

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Definition

• Severe hypokalemia – Potassium level less than 2.5 mEq/L

Moderate hypokalemia – Potassium level between 2.5 and 3 mEq/L

Mild hypokalemia – Potassium level between 3 and 3.5 mEq/L

Etiology

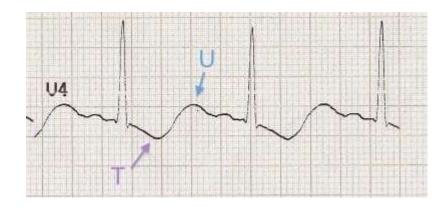
Decreased intake	Increased intracellular uptake	Increased loss-extra renal	Increased loss-renal	Endocrine
Severe acute malnutrition	Metabolic alkalosisInsulinBeta adrenergic agents	DiarrheaEmesisCystic fibrosis	DiureticsDKATubulo interstitial disease	 Aldosterone- secreting adenomas Glucocorticoid remediable aldosteronism
• Anorexia	 Heavy metals(barium) Anti psychotic drugs Hypokalemic periodic paralysis 		 Bartter syndrome Gitelman syndrome Renal tubular acidosis Amphotericin Liddle syndrome 	 Apparent mineralocorticoid excess (AME) 11-beta-hydroxylase deficiency 17-alpha-hydroxylase deficiency

Clinical features

- Asymptomatic
- Muscle weakness- head lag, hypotonia, ascending paralysis, respiratory failure, death
- Ileus, constipation, abdominal tenderness
- Cramps and fasciculation
- Arrhythmias (more risk with hypomagnesemia)

ECG changes

- PR prolongation
- Flattening of T waves
- ST depression.
- U waves can emerge after the T waves(best seen in the precordial leads V4-V6)

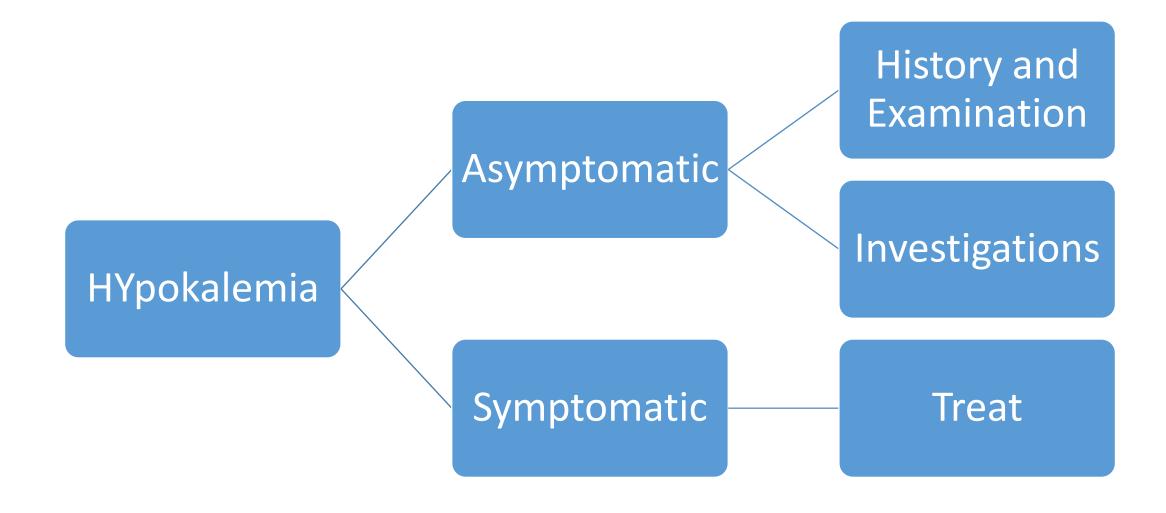


Step -1 Rule out Spurious hypokalemia

• Due to sampling errors - Recent line flush ,IV fluids near sampling site

• It can also occur if the WBC count is very high as in leukemia, if plasma for analysis is left at room temperature where the K is taken up by the cells.

Approach to hypokalemia



Management in symptomatic patients

Care of ABC

• Fluid boluses may be needed if volume depleted, IV potassium

• 12 lead ECG, continuous cardiac monitoring

Therapeutic goals

Prevent or treat life-threatening complications

Correct the potassium deficit

- Address the underlying cause.
- Therapeutic urgency depends on the severity of hypokalemia, the existence of comorbid conditions, and the rate of decline of serum potassium levels

Management in symptomatic patients

• IV in the presence of cardiac dysrhythmias, digitalis toxicity

KCl IV 0.5 to 1 mEq/kg of body weight per hour under cardiac monitoring.

• The goal is to raise the potassium level by 0.3 to 0.5 mEq/L.

 An infusion with a potassium concentration of no more than 40 -60 mEq/L is given through peripheral line.

May repeat as needed based on lab values

Caution

Associated with pain and phlebitis when administered through a peripheral vein

• Do NOT administer undiluted or by IV push. Rapid potassium infusion can cause cardiac arrest.

Always add K in a new bag

• IV fluids should not contain dextrose as it can stimulate insulin secretion

Monitoring

Continuous ECG monitoring

 Serum concentrations should be evaluated 1 to 2 hours after completion of infusion and then every 2 to 4 hours untill symptoms resolve

• Watch for rebound hyperkalemia when rates exceed 20 mmol/h or when cellular redistribution is the cause of hypokalemia.[

Once K normal evaluate

History

- Polyuria, polydipsia (impaired concentrating capacity)
- Loose stools, emesis
- Drug intake
- Renal disease
- Dark urine(rhabdomyolysis, myoglobinuria)
- Symptoms of thyrotoxicosis
- Family history

Physical examination

- HR (bradycardia) and rhythm. Inappropriate low HR in shock
- Blood pressure
- Hypotonia, phantom hernia, head lag, diminished bowel sounds
- Deep tendon reflexes
- Volume status
- Rickets

Hypokalemia etiology based on BP

With hypertension

- Liddle syndrome
- Conn syndrome
- Cushing syndrome
- 11 beta hydroxylase deficiency
- Licorice excess

Without hypertension

- Diuretics
- Diarrhea
- Vomiting
- Bartter syndrome
- Gitelman syndrome
- RTA -I & II

Investigations

• First line: sodium, potassium, chloride, bicarbonate, calcium, magnesium, urea, creatinine, venous blood gas

 Second line: Urine K(Random urinary potassium levels, Spot potassium-tocreatinine ratios, 24 hours), TTKG, urinary chloride, USG kidney

• Endocrine: Renin, aldosterone, 17 alpha hydroxylase, 11 beta hydroxylase

Genetic testing

Hypokalemia based on acid base status

Metabolic acidosis

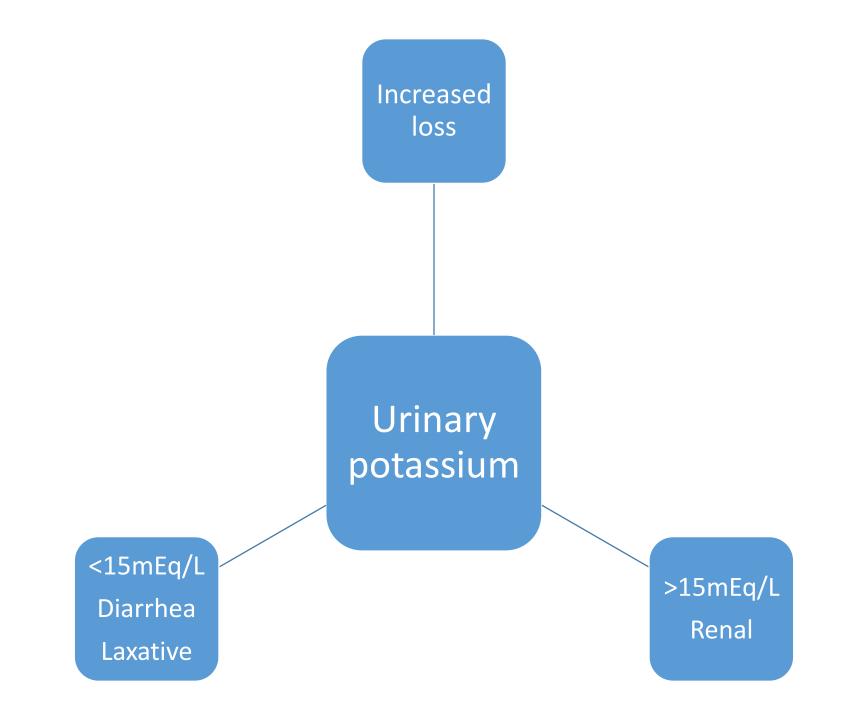
- RTA
- Diarrhea
- DKA

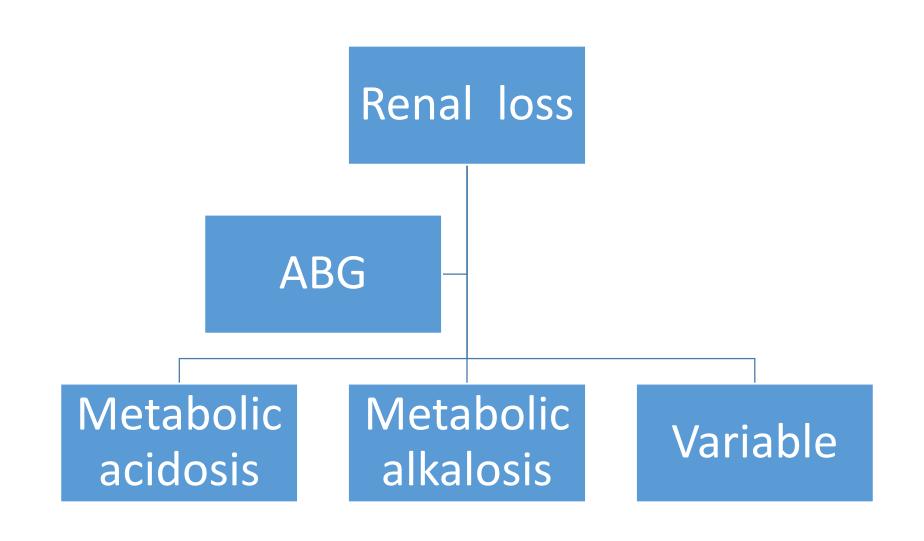
Metabolic alkalosis

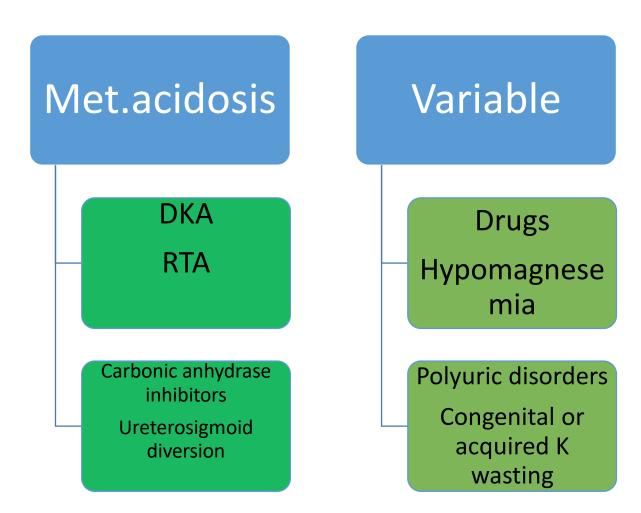
- Conn syndrome
- Vomiting
- Diuretic use
- Heriditary renal channelopathies (Liddle, Gitelman, Bartter syn)

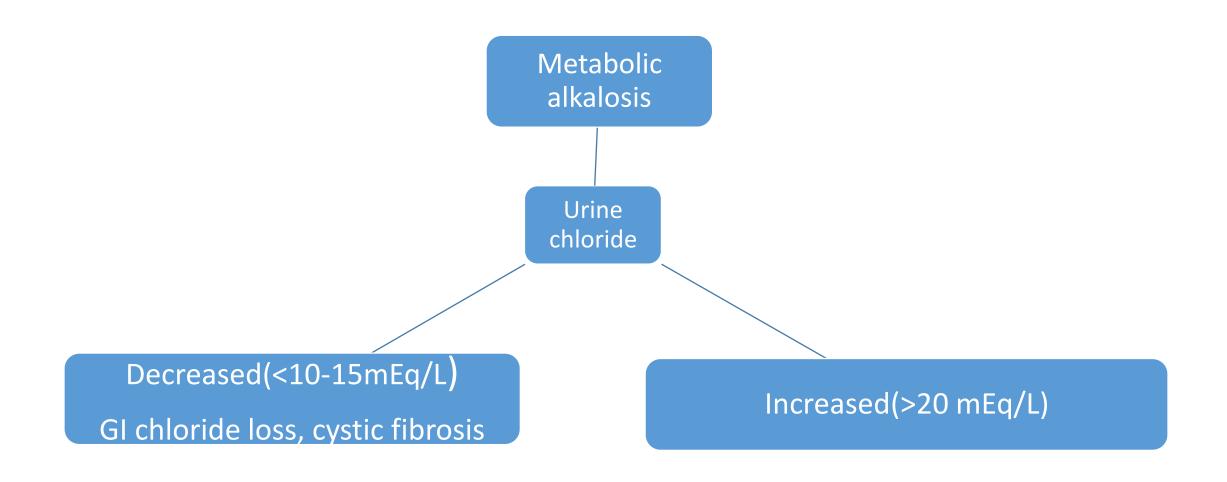
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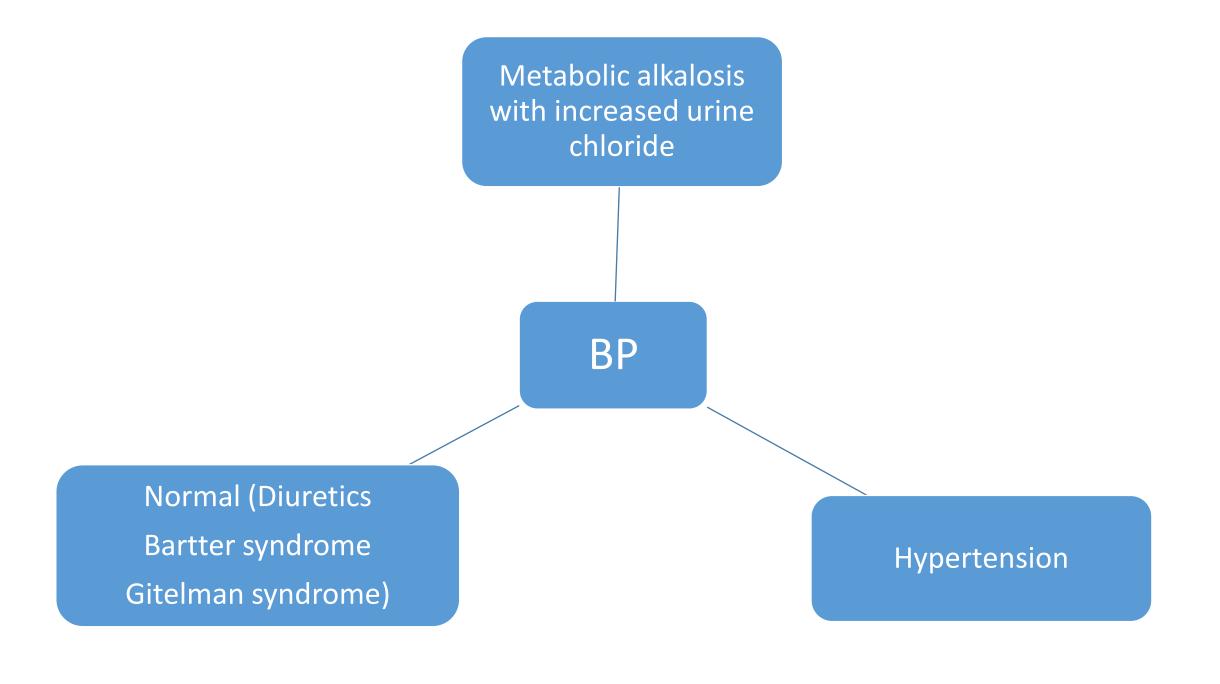
Decreased intake Increased loss Transcellular shift

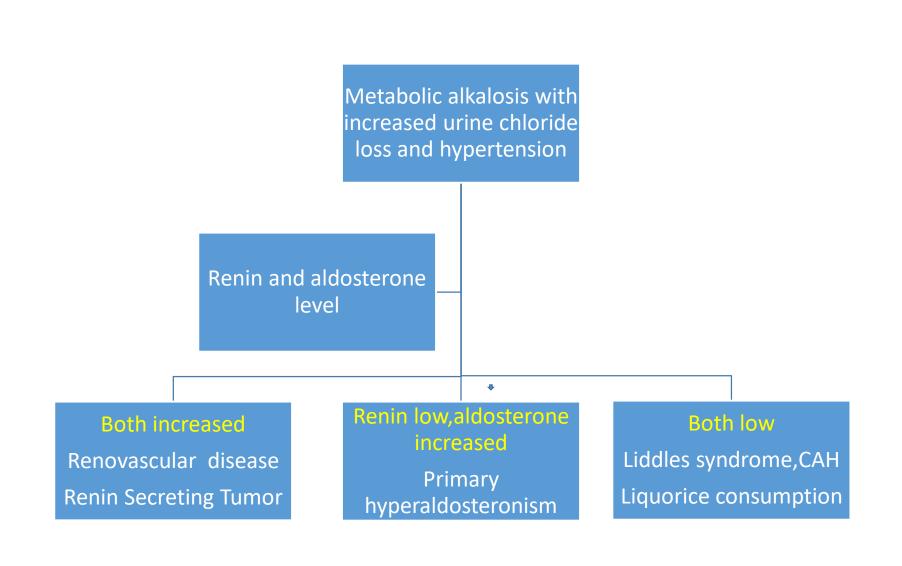












Management of asymptomatic patients

3 and 3.5 mEq/L-

Correction of the underlying cause and dietary potassium

 Maintenance amount of potassium to IV fluids 20mEq/L

Less than 3 mEq/L

Oral replacement.

• IV only for those who are unable to take oral medications.

In asymptomatic patients with chronic hypokalemia(RTA) potassium supplementation may be needed

Oral potassium

- Potassium chloride, phosphate, acetate, citrate, bicarbonate.
- Initial: 1 to 2 mEq/kg/day in divided doses
- Titrate to desired clinical response
- Usual range: 1 to 5 mEq/kg/day
- Not to exceed 1 to 2 mEq/kg as a single dose up to 40 mEq/dose

Magnesium sulphate

• In hypomagnesemia

25-50mg/kg IV over 30 minutes

• Watch for bradycardia, hypotension

Other treatment

• Potassium-sparing diuretic such as amiloride in Bartter, Gitelman

• Hyperaldosteronism - spironolactone or eplerenone