

Treating Acid Base Disturbances and Correcting Electrolyte Abnormalities

Paul Wisniewski, DO

Cutting Edge Surgical Medical Group

June 30, 2026

Objectives

Management and Identification
on Acid Base Disorders

Identification of Electrolyte
Abnormalities

Correction of Electrolyte
abnormalities

Disclosures

None



Metabolic Acidosis

Sodium Bicarbonate:

- **Indication:** pH < 7.1 or severe acidosis with renal failure.
- Normal pH 7.35-7.45
- **Dose:** 1–2 mEq/kg IV bolus, or continuous infusion (150 mEq in 1L D5W at 100–200 mL/hr) titrated to effect (Kraut & Madias, 2010).
- Calculate the base deficit = $.3 \text{ (BE) } * (\text{ Ideal body wt}) / 50 \text{ meq per amp}$

Renal Replacement Therapy (RRT):

- **Indication:** Refractory acidosis or renal failure.
- **Mode:** CRRT or intermittent hemodialysis.



Respiratory Acidosis

Noninvasive Ventilation (BiPAP) or Intubation:

- Indicated based on mental status, respiratory effort, and gas exchange.
- $HP < 7.2$

Naloxone (if opioid-related):

- **Dose:** 0.04–0.4 mg IV, titrate to respiratory response.



Metabolic Alkalosis

Normal Saline (0.9% NaCl):

- **Indication:** Chloride-responsive alkalosis (e.g., vomiting, diuretics).
- **Dose:** 1–2 L IV bolus followed by maintenance rate as needed.

Potassium Chloride (KCl):

- **Dose:** Oral 40–100 mEq/day or IV 10–20 mEq/hr, max 40 mEq/hr via central line with cardiac monitoring.

Acetazolamide:

- **Indication:** Volume overload with alkalosis.
- **Dose:** 250–500 mg IV or PO once or BID.



Respiratory Alkalosis

Benzodiazepines (e.g., lorazepam):

- **Dose:** 1–2 mg IV/PO as needed for anxiety-induced hyperventilation.
- pH >7.45

Treat underlying cause
(e.g., antibiotics for sepsis).



pH Range and Associated Mortality Risk

pH Range	Clinical Interpretation	Associated Mortality Risk	Key Clinical Concerns
< 7.00	Severe Acidemia	> 80%	Cardiovascular collapse, arrhythmias, impaired contractility
7.00 – 7.19	Moderate to Severe Acidemia	50–60%	High lactate, shock, poor tissue perfusion
7.20 – 7.34	Mild Acidemia	20–30%	Often in sepsis, renal failure
7.35 – 7.45	Normal	< 5%	Optimal physiologic range
7.46 – 7.54	Mild Alkalemia	10–15%	May be asymptomatic or early compensatory response
7.55 – 7.59	Moderate Alkalemia	20–35%	Arrhythmias, hypokalemia, hypoventilation
≥ 7.60	Severe Alkalemia	> 45–50%	Cerebral vasoconstriction, seizures, decreased coronary perfusion



Sodium

Hyponatremia

3% Hypertonic Saline:

- **Indication:** Severe or symptomatic hyponatremia ($\text{Na}^+ < 120 \text{ mEq/L}$ with seizures or confusion).
- **Dose:** 100 mL IV over 10 min $\times$ 3 doses PRN, then infusion 0.5–2 mL/kg/hr.
- **Correction goal:** $\leq 10 \text{ mEq/L}$ per 24 hours (Sterns et al., 2022).

Tolvaptan (SIADH, euvolemic hyponatremia):

- **Dose:** 15 mg PO daily, titrate to 30–60 mg/day.

Sodium

Hypernatremia

D5W or 0.45% NaCl:

- **Dose:** Free water deficit = $0.6 \times \text{weight (kg)} \times [(\text{serum Na}^+ / 140) - 1]$; replace over 48–72 hours.



Potassium

Hypokalemia

Oral Potassium Chloride:

- **Dose:** 20–40 mEq PO BID-TID; max 100 mEq/day.

IV Potassium Chloride:

- **Dose:** 10–20 mEq/hr via peripheral line; up to 40 mEq/hr via central line with telemetry.

Magnesium Sulfate (if concurrent hypomagnesemia):

- **Dose:** 1–2 g IV over 1 hour, repeat as needed.



Potassium

Hyperkalemia

Calcium Gluconate 10%:

- **Dose:** 10 mL IV over 5–10 min; may repeat every 30–60 minutes.

Insulin + Dextrose:

- **Dose:** Regular insulin 10 units IV + 25–50 mL of D50W over 5 minutes.

Albuterol:

- **Dose:** 10–20 mg nebulized over 10 minutes.

Sodium Bicarbonate (if metabolic acidosis):

- **Dose:** 50–100 mEq IV over 5–10 minutes.



Potassium

Hyperkalemia (Cont.)

Furosemide:

- **Dose:** 20–40 mg IV; titrate for diuresis.

Sodium Polystyrene Sulfonate (Kayexalate):

- **Dose:** 15–30 g PO/PR; avoid in ileus or post-op.

Dialysis:

- Indicated in refractory or renal failure-associated hyperkalemia.

Calcium

Hypocalcemia

Calcium Gluconate:

- **Dose:** 1–2 g IV over 10–20 minutes; repeat PRN.

Calcium Chloride (for severe/symptomatic cases):

- **Dose:** 1 g IV over 10 minutes via central line only.

Vitamin D (Calcitriol):

- **Dose:** 0.25–1 mcg PO daily for chronic hypocalcemia or renal disease.

Calcium

Hypercalcemia

Normal Saline IV:

- **Dose:** 200–300 mL/hr for volume repletion.

Furosemide (after rehydration):

- **Dose:** 20–40 mg IV every 12 hours.

Calcitonin:

- **Dose:** 4 IU/kg SC or IM q12h; effect in 4–6 hours.

Zoledronic Acid (bisphosphonate):

- **Dose:** 4 mg IV over 15 minutes, single dose.



Magnesium

Hypomagnesemia

Magnesium Sulfate:

- **Mild (Mg^{2+} 1.2–1.7 mg/dL):**
1–2 g IV over 1–2 hours.
- **Severe (<1.2 mg/dL or seizures/arrhythmia):** 4–6 g IV over 4–12 hours.



Magnesium

Hypermagnesemia

Calcium Gluconate 10%:

- **Dose:** 1–2 g IV over 10 minutes to reverse cardiac/muscular effects.

Loop diuretics + IV fluids to enhance renal excretion.

Dialysis if anuric or severe toxicity.



Phosphate

Hypophosphatemia

Oral phosphate (Neutra-Phos):

- **Dose:** 250 mg PO BID-TID.

IV Sodium or Potassium Phosphate:

- **Mild (2.0–2.5 mg/dL):** 0.08–0.16 mmol/kg IV.
- **Severe (<1.0 mg/dL):** 0.32–0.64 mmol/kg IV over 4–6 hours.
- Kphos 10-30 mmol = (30-90 meqK+)



Phosphate

Hyperphosphatemia

Sevelamer (Renvela):

- **Dose:** 800–1600 mg PO TID with meals.

Calcium Acetate:

- **Dose:** 667 mg PO TID with meals.

Dialysis if refractory or ESRD.



Best Practice Recommendations

Use a **systematic approach**: Assess volume status, acid-base status, and electrolyte trends.

Correct coexisting abnormalities simultaneously (e.g., Mg and K).

Avoid overcorrection: Follow published correction limits to prevent iatrogenic harm.

Consider **consultation with nephrology or critical care** in complex cases.

Regular monitoring: Electrolytes, ABG, renal function every 4–6 hours in unstable patients.





- Questions



References

1. Kraut, J. A., & Madias, N. E. (2010). Metabolic acidosis: pathophysiology, diagnosis and management. *Nature Reviews Nephrology*, 6(5), 274–285. <https://doi.org/10.1038/nrneph.2010.33>
2. Cooper, D. J., et al. (1990). Bicarbonate does not improve hemodynamics or reduce vasopressor requirement in lactic acidosis. *Annals of Internal Medicine*, 112(7), 492–498.
3. Adroque, H. J., & Madias, N. E. (1998). Management of life-threatening acid-base disorders. *New England Journal of Medicine*, 338(1), 26–34.
4. Palmer, B. F. (2004). Managing hyperkalemia caused by inhibitors of the renin-angiotensin-aldosterone system. *New England Journal of Medicine*, 351(6), 585–592.
5. Sterns, R. H., et al. (2022). Hyponatremia: evaluation and management. *Kidney International*, 101(5), 928–939.
6. Verbalis, J. G., et al. (2013). Diagnosis, evaluation, and treatment of hyponatremia: expert panel recommendations. *American Journal of Medicine*, 126(10), S1–S42.
7. Weiner, I. D., & Wingo, C. S. (1997). Hypokalemia—consequences, causes, and correction. *Journal of the American Society of Nephrology*, 8(7), 1179–1188.
8. Baird, G. S. (2012). Ionized calcium. *Clinica Chimica Acta*, 413(5-6), 696–701.
9. Swaminathan, R. (2003). Magnesium metabolism and its disorders. *Clinical Biochemist Reviews*, 24(2), 47–66.
10. Block, G. A., et al. (2004). Mineral metabolism, mortality, and morbidity in maintenance hemodialysis.

