Quick Guide to Laboratory Values

Use this convenient cheat-sheet to help you monitor laboratory values related to fluid and electrolyte status, acid-base balance, and hematology. Remember, normal value ranges may vary according to techniques used in different laboratories.

SERUM ELECTROLYTES	
Electrolyte (Range)	Nursing Considerations
Electrolyte (Range) Calcium (Ca ²⁺) 8.5-10.5 mg/dL	Nursing Considerations Hypocalcemia • Signs and symptoms • Tetany (neuromuscular irritability) is the hallmark symptom (may include paresthesia, bronchospasm, laryngospasm, carpopedal spasm [Trousseau's sign], Chvostek's sign [facial muscle contractions elicited by tapping facial nerve on ipsilateral side], tingling sensations of the fingers, mouth, and feet, increased deep tendon reflexes [DTRs], generalized seizures). • Psychiatric manifestations such as emotional instability, depression, anxiety, hallucinations and psychosis. • ECG changes may include prolonged QT interval and arrhythmias. • Papilledema with or without intracranial hypertension. • Anxiety, depression, fatigue. • Can also be asymptomatic. • Implement seizure precautions and close monitoring for cardiac arrhythmias and respiratory depression. • Treatment: initiate intravenous calcium repletion • Signs and symptoms • Lethargy, confusion, nausea, vomiting, anorexia, constipation, muscle weakness, depressed DTRs, polyuria, polydipsia, dehydration • Severe hypercalcemia can provoke supraventricular or ventricular arrhythmias. Monitor cardiac rate and rhythm.
	Watch for digitalis toxicity.



Chloride (Cl ⁻)	Hypochloremia
97-107 mEq/L	Signs and symptoms
	 Muscle spasms, alkalosis, and depressed respirations
	 May be precipitated or exacerbated by GI losses (vomiting,
	diarrhea).
	Hyperchloremia
	Monitor for metabolic acidosis.
	 Associated with large volume 0.9% normal saline
	resuscitation.
Magnesium (Mg ²⁺)	Hypomagnesemia
1.8-3 mg/dL	Signs and symptoms
	 Cardiac/ventricular arrhythmias (ventricular
	arrythmias (torsades de pointes) and atrial
	fibrillation) tetany tremor weakness anathy
	delirium seizures and coma
	 Bick factors: chronic diarrhea, PPL use alcoholism diuretic
	 Monitor cardiac rate and rhythm
	 Monitor for digitalis toxicity
	Often associated with hypokalemia. Treat both
	Often associated with hypokalemia. Treat both simultaneously.
	simultaneously.
	Hypermaanesemia
	• Signs and sumptoms
	• Signs and symptoms
	 Cardiaa manifestational humatansian humatansian
	o Cardiac mannestations: hypotension, bradycardia,
	Complete neart block, cardiac arrest
	 Neurologic manifestations: letnargy/somnolence, degraded DTBs, weight associations associations
	decreased DTRs, muscle paralysis, coma, respiratory
	muscle weakness (shallow respirations, aphea)
	Avoid magnesium-containing medications in patients with
	compromised renal function.
	 Monitor cardiac rate and rhythm.
	 Monitor neurologic status, including DTRs.
Phosphate (PO ₄ ⁻)	Hypophosphatemia
2.5-4.5 mg/dL	 Signs and symptoms (rare unless PO₄⁻ less than 1mg/dL)
	 Muscle weakness, rhabdomyolysis
	 Treatment indicated when PO₄⁻ less than 2mg/dL.
	Oral replacement preferred.
	 IV indicated if PO₄⁻ less than 1mg/dL; administer slowly.
	When administering IV phosphate products, measure serum

	phosphate levels every 6-8 hours.
	 Monitor for hypocalcemia, renal failure, arrhythmias, and
	diarrhea (with oral replacement).
	Acute or chronic hypophosphatemia can cause
	rhabdomyolysis.
	Hyperphosphatemia
	Signs and symptoms
	 Typically, asymptomatic
	 Clinical features are due to accompanying
	hypocalcemia (see above)
	 Severe hyperphosphatemia can be life threatening.
	 Risk factors include advanced renal insufficiency,
	rhabdomyolysis, tumor-lysis syndrome, and over ingestion of
	phosphate containing laxatives)
	Soft tissue calcification can be a long-term complication of
	chronically elevated serum phosphate levels.
Potassium (K ⁺)	Hypokalemia
3.5-5 mEq/L	Signs and symptoms
	 Muscle cramps/weakness, rhabdomyolysis,
	respiratory muscle weakness, decreased bowel
	motility, cardiac arrhythmias, hypotension, mental
	status changes, speech changes.
	Characteristic ECG findings include ST segment depression,
	flattened T wave and U wave.
	Monitor cardiac rate and rhythm.
	 Common causes include GI losses (diarrhea/vomiting) and
	diuretic therapy.
	 Educate patient on using laxatives and diuretics only
	as prescribed.
	Monitor potassium levels in patients on digoxin; hypokalemia
	will potentiate its effects.
	 Prolonged hypokalemia can lead to structural and functional
	changes in the kidneys.
	Hyperkalemia
	Signs and symptoms
	 Irritability/anxiety, paresthesia, ascending muscle
	weakness, cardiac arrhythmias, cardiac conduction
	abnormalities, lethargy, GI symptoms (nausea and
	intestinal colic)
	Characteristic ECG findings include tall, peaked T waves with
	shortened QT interval, prolonged PR interval, wide QRS



	complex and in severe cases, ventricular standstill.
	Monitor cardiac rate and mythin. Avoid notacsium supplements
	• Avoid potassium-spaning didretics, potassium supplements,
	of sait substitutes in patients with renarms unclency.
	• Use ACE inhibitors cautiously, as they cause K retention.
Sodium (Na⁺)	Hyponatremia
135-145 mEq/L	 Signs and symptoms
	 Neurologic: lethargy, weakness, irritability,
	confusion, tremors, myoclonus, seizures
	 Other: hypotension, GI symptoms (anorexia, nausea,
	vomiting, abdominal cramping)
	 Cerebral edema can occur in rapid reduction of
	serum sodium concentration
	 Correction should be slow (4 to 6 mEq/L in first 24 hours) to
	avoid osmotic demyelination syndrome; monitor serum Na ⁺
	levels and neurologic status frequently.
	 Avoid large water supplements to patients receiving isotonic
	tube feedings.
	 Implement seizure precautions in severe cases.
	 Monitor fluid losses and gains.
	 Hypernatremia Signs and symptoms Excessive thirst, dehydration, dry mucous membranes, oliguria, mental status changes including lethargy, disorientation, restlessness, elevated body temperature. Can cause rapid decrease in brain volume potentially leading to rupture of the cerebral veins, subarachnoid hemorrhages, and possibly irreversible neurologic damage. Monitor fluid losses and gains; urine and plasma osmolality may assist in establishing etiology. Give sufficient free water with tube feedings or salt-free IV fluids to keep serum Na⁺ and BUN within normal limits.
	ACID-BASE STATUS
Arterial Blood Gas (ABG)	
Component	Nursing Considerations
(Range)	
рН	 Identification of the specific acid–base disturbance is
7.35-7.45	important in identifying the underlying cause of the disorder



	and determining appropriate treatment.
	• A pH less than 7.35 indicates acidosis and a pH greater than
	7.45 indicates alkalosis.
PaCO ₂	• The PaCO ₂ is influenced almost entirely by respiratory
- 35-45 mmHg	activity.
5	• When the $PaCO_2$ is low, carbonic acid leaves the body in
	excessive amounts: when the $PaCO_2$ is high there are
	excessive amounts, when the race $\frac{1}{2}$ is high, there are
	The bicarbonate level of the ABG reflects the bicarbonate
22-26 mFg/l	level of the body
22 20 1129/2	 The kidneys are involved in either reabsorbing bicarbonate
	• The kidneys are involved in either reabsorbing bicarbonate
	maintain acid base balance
Laboratory Value	
(Bange)	Nursing Considerations
Blood urea nitrogen (BLIN)	 Increased BLIN may be seen in patients with impaired renal
10-20 mg/dl	function
10 20 mg/uL	 Increased PLIN may be caused by hypotension/check, heart
	failure, salt and water deplotion, debudration, diabetic
	ketoosidosis gostrointostional homorrhago, and hurns
	Retoacidosis, gasti onitestional hemorriage, and burns.
Creatinine	 Increased creatining lovels may be seen in patients with
0.7-1.4 mg/dl	 Increased creatining levels may be seen in patients with impaired repair function due to decreased blood flow to the
0.7-1.4 mg/dL	kidney (heart failure, sheek liver disease, debydration), urinary
	tract obstruction, intrinsic kidnov disease, denyulation, unitary
	demonuterentrice), or contain modications
	giomerationeprintis), or certain medications.
	 Acute kidney injury (Aki) is diagnosed when baseline creatinine increases abrumtly by 0.2 mg/dl, or more, even if creatining
	increases abruptly by 0.3 mg/dL or more, even if creatinine
	remains in the normal range.
	HEMATOLOGIC STUDIES
Laboratory Value	
	Nursing Considerations
	 Increased homoglabin lovals may be asyred by hyperia, bish
	 Increased nemoglobin levels may be caused by hypoxia, high altitude living, or homospheretration of the blood from.
IVIAIES: 13-18 g/dL	altitude living, or nemoconcentration of the blood from
Females: 12-16 g/dL	denyaration.
	Decreased levels of hemoglobin (anemia) may be due to
	hemorrhage/blood loss, hemodilution, nutritional
	deficiencies, chronic disease, underlying malignancy,

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	hereditary disorders, or a hemolytic reaction.
Hematocrit *Typically, three times the hemoglobin level Males: 42-52% Females: 35-47%	 Increased hematocrit values are seen in severe fluid volume deficit and shock (when hemoconcentration rises considerably). Decreased hematocrit values are seen with blood loss, hemolytic reactions after transfusion of incompatible blood, fluid overload, and in similar conditions in which decreased levels of hemoglobin are seen.
Platelet count 150,000-450,000/mm ³	 Increased platelet levels (thrombocythemia or thrombocytosis) may be caused by a bone marrow disorder or malignancy, infection or inflammation, anemia, previous splenectomy, or certain medications. Decreased platelet levels (thrombocytopenia) may be a result of bone marrow suppression, sepsis, sequestration from an enlarged spleen, increased platelet destruction (seen with autoimmune syndromes or drug-induced reactions), or decreased platelet production (related to infections or malignancy). Liver disease, renal disorders, and pregnancy can also cause thrombocytopenia.
	COAGULATION STUDIES
Laboratory Value (Range)	COAGULATION STUDIES Nursing Considerations
Laboratory Value (Range) Prothrombin time (PT) 9.5-12 seconds	 COAGULATION STUDIES Nursing Considerations The PT measures the activity of the extrinsic pathway of the clotting cascade. Prothrombin is a protein made by the liver. Elevated PT may indicate liver dysfunction, Vitamin K deficiency, or coagulation factor deficiency (e.g., factor VII)
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Laboratory Value (Range)Prothrombin time (PT)9.5-12 secondsPartial thromboplastin time (activated) (PTT) 20-39 secondsInternational normalized ratio (INR) 1.0; 2-3.5 for patients taking warfarin sodium (varies based on diagnosis)	COAGULATION STUDIES Nursing Considerations • The PT measures the activity of the extrinsic pathway of the clotting cascade. • Prothrombin is a protein made by the liver. • Elevated PT may indicate liver dysfunction, Vitamin K deficiency, or coagulation factor deficiency (e.g., factor VII) • The PTT is a measure of the activity of the intrinsic pathway of the clotting cascade. • The PTT is used to monitor the effects of unfractionated heparin. • The INR is used to monitor the effectiveness of warfarin therapy. • As INR increases, time for blood to clot increases.
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Total protein 6-8 g/100 mL Albumin 3.5-5 g/100 mL	 Proteins influence the colloid osmotic pressure. Includes albumin and globulin. Makes up 60% of total protein. Keeps fluid from leaking out of blood vessels. Changes in serum albumin affect total serum calcium. Degreesed albumin can be due to malautritian or liver.
	 Decreased abumin can be due to mainturnion or liver disease and can lead to edema, ascites, and pulmonary edema.
	SERUM OSMOLALITY
Laboratory Value (Range)	Nursing Considerations
Osmolality 280-300 mOsm/L water	 Increased osmolality may be caused by severe dehydration, free water loss, diabetes insipidus, hypernatremia, hyperglycemia, stroke or head injury, renal tubular necrosis, or ingestion of methanol or ethylene glycol (antifreeze). Decreased osmolality may be caused by volume excess, SIADH, renal failure, diuretic use, adrenal insufficiency, hyponatremia, overhydration, or paraneoplastic syndrome associated with lung cancer.
	URINE TESTS
Laboratory Value (Range)	Nursing Considerations
pH (urine) 4.6-8.2	 Decreased urine pH may be caused by metabolic acidosis, diabetic ketoacidosis, or diarrhea. Increased urine pH may be caused by respiratory alkalosis, potassium depletion, or chronic renal failure.
Specific gravity (urine) 1.010-1.025	 The urine specific gravity range depends on the patient's state of hydration and varies with urine volume and the load of solutes to be excreted. Increased urine specific gravity may be seen with dehydration, vomiting, diarrhea, infection, and heart failure. Decreased urine specific gravity can occur with renal damage.

References:

Farinde, A. (2019, May 14). Lab values, normal adult. *Medscape*. <u>https://emedicine.medscape.com/article/2172316-overview</u>

Hinkle, J., & Cheever, K. (2018). Brunner & Suddarth's Textbook of Medical-Surgical Nursing, Fourteenth Edition. Philadelphia: Lippincott Williams & Wilkins.

Mount, D. B. (2024, February 12). Clinical manifestations and treatment of hypokalemia in adults. *UpToDate*. <u>https://www.uptodate.com/contents/clinical-manifestations-and-treatment-of-hypokalemia-in-adults</u>

Mount, D. B. (2022, December 7). Clinical manifestations of hyperkalemia in adults. *UpToDate*. <u>https://www.uptodate.com/contents/clinical-manifestations-of-hyperkalemia-in-adults</u>

Sterns, R.H. (2023, January 23). Manifestations of hyponatremia and hypernatremia in adults. *UpToDate*. <u>Manifestations of hyponatremia and hypernatremia in adults</u> - <u>UpToDate</u>

Yu, A. S., & Aditi, G. (2022, May 19). Hypermagnesemia: Causes, symptoms, and treatment. *UpToDate*. <u>https://www.uptodate.com/contents/hypermagnesemia-causes-symptoms-and-treatment</u>

Yu, A. S., & Stubbs, J. R. (2023, April). Hypophosphatemia: Evaluation and treatment. *UpToDate*. <u>https://www.uptodate.com/contents/hypophosphatemia-evaluation-and-treatment</u>