

# INTERPRETING NEONATAL LABS

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Anna Holcomb, DNP, APRN-NP, NNP-BC

# OBJECTIVES

- Identify value ranges for common neonatal lab tests
- Discuss blood gas values and interpretations with disease states
- Discuss common infection laboratory values
- Discuss basics of hematology with CBC interpretation



# COMMON NEONATAL LABS

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Blood gases (ABG/VBG/CBG/cord gases)

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Complete metabolic panel (CMP), Basic metabolic panel (BMP)

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Complete blood count (CBC)

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Urinalysis

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Cultures

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Respiratory Viral Panel (RVP)

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Drug levels

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Thyroid studies

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State newborn screens

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CSF studies

**Table 3.1**

## Common Laboratory Tests in the NICU by System

Fluids	Respiratory	Cardiac	GI	Renal	Endocrine/Metabolic	Neurologic	Hematologic	Other
<b>Electrolytes</b>	Blood gas	Iso-enzymes	Alkaline phosphatase	BUN (serum)	Glucose	Phenobarbital level	CBC with differential	Chromosome studies
<b>Nutrition</b>	pH	Digoxin level	Triglyceride levels	Creatinine (serum)	Thyroxine level Thyroid-stimulating hormone level	Dilantin level	Hct/Hgb	Bilirubin Total/indirect/direct
Electrolytes (serum)			AST (SGOT)	Specific gravity	Cortisol level	Ammonia (serum)	Platelet count Platelet antibody	CRP
Calcium (serum)	Paco <sub>2</sub>		ALT (SGPT)	Urine sodium	Insulin level	Amino acid (serum)	Reticulocyte count	Drug levels
Magnesium (serum)	Pao <sub>2</sub>		GGT (GGTP)	Urine potassium	Growth hormone level	Amino acid (urine)	Direct Coombs' test/ DAT	Urine/meconium/ cordtoxicology screen
Phosphorus (serum)	Bicarbonate (HCO <sub>3</sub> <sup>-</sup> )		Bilirubin (total and direct)	Urine osmolality	Testosterone level	Organic acids (urine)	Blood type and Rh	Culture and sensitivity (all sources)
Alkaline phosphatase	Base excess		Ammonia (serum)		Aldosterone (serum and urine) level	Glycine (serum and CSF)	G6PD	RPR/VDRL
Glucose (serum)	Theophylline level		α <sub>1</sub> -antitrypsin	Uric acid (serum)	17-hydroxy corticosteroids	Pyridoxine level	Sedimentation rate	Occult blood (fecal)
Vitamin levels	Caffeine level		Trypsin (stool)	Urinalysis	Parathyroid hormone level	Lactic acid/lactate level	Folic acid	PCR (serum)
Serum osmolality	Cultures (tracheal/sputum)		Pyridoxine level	Urine culture/sensitivity	State blood spot screening		Fibrinogen PT PTT Immunoglobulins Methemoglobin	
Albumin (serum)								
Total protein								

ALT (SGPT), Alanine aminotransferase (serum glutamic-pyruvic transaminase); AST (SGOT), aspartate aminotransferase (serum glutamic-oxaloacetic transaminase); BUN, blood urea nitrogen; CBC, complete blood count; CRP, C-reactive protein; CSF, cerebrospinal fluid; DAT, direct antiglobulin test; GI, gastrointestinal; GGT (GGTP), γ-glutamyltransferase (γ-glutamyl transpeptidase); G6PD, glucose-6-phosphate dehydrogenase; Hct, hematocrit; Hgb, hemoglobin; NICU, neonatal intensive care unit; PCR, polymerase chain reaction; PT, prothrombin time; PTT, partial thromboplastin time; RPR, rapid plasma reagin; VDRL, venereal disease research laboratory.

**Box 13.1****Laboratory Interpretation: Is It Reliable? Is It Believable?**

<b>Yes</b>	<b>No</b>
Implement an appropriate clinical intervention, if needed. Consider repeating the laboratory test if the clinical status continues to warrant a laboratory test.	Consider repeating the laboratory test when the appropriate correction is made.
<b>Result Too Low</b>	<b>Result Too High</b>
Is the sample diluted?  Was the sample handled correctly? Was the timing of the test an issue? Was medical therapy not implemented or inadequate? Was the specimen site condition a factor? Was the point-of-care testing (POCT) device calibrated? Was there interference from a past medical therapy? Was the sample handled correctly? Was medical therapy not implemented or inadequate?	Does the blood gas need to be temperature corrected (as with neuroprotective hypothermia therapy)? Was the laboratory specimen drawn too early?

The goal of laboratory testing is to diagnose and guide management of disease

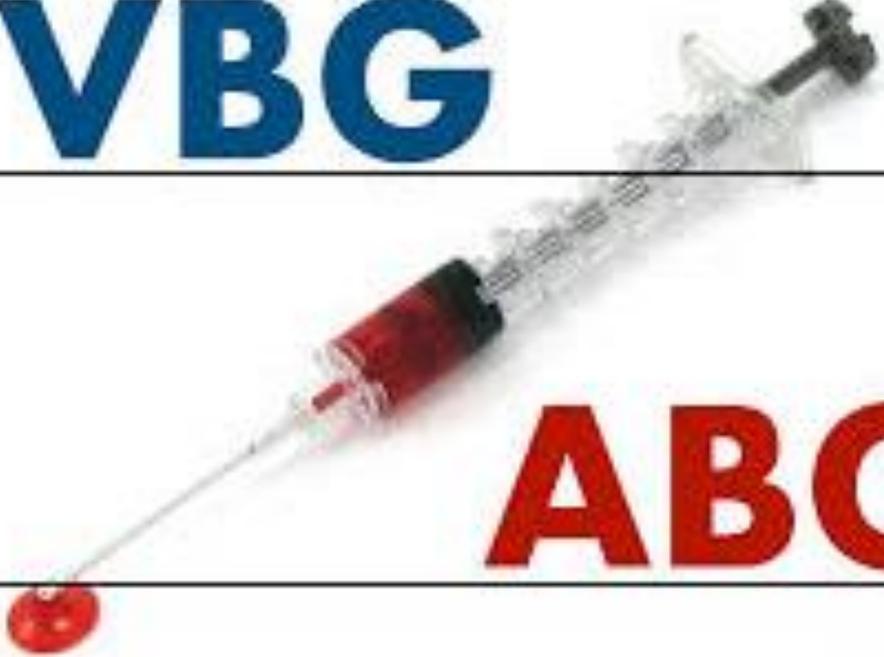
Careful laboratory data interpretation is essential in providing accurate therapeutic interventions

Does the lab result match the clinical picture?

# LABORATORY INTERPRETATION

# VBG

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# ABG

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## BLOOD GAS

- Most common laboratory measurement to assess the ventilation/acid-base balance/oxygenation of a neonate.
  - Sample types: venous, arterial, capillary
  - Sample methods:
    - Venous: venous puncture/umbilical vein draw
      - Capillary: capillary sampling (heel stick)
      - Arterial: umbilical artery draw/arterial puncture/peripheral arterial line
      - Arterial preferred sample as it has a more accurate measurement of the PaO<sub>2</sub>
- Cord gases: arterial and venous (arterial more accurate picture of infant condition)

# BLOOD GAS-PH

Normal arterial blood gas ph: 7.35-7.45

Monitor acid-base balance: normal cell function is dependent on regulation of the hydrogen ion ( $H^+$ ) concentration in the blood.

Changes in ph indicate a change in the acid-base balance of the infant = change in cellular metabolism/function

## **BUFFER SYSTEMS: Ways to regulate ( $H^+$ )**

1. Exhalation of carbon dioxide ( $CO_2$ ) via the lungs
2. Excretion of  $H^+$  by the kidney/reabsorption of  $HCO_3^-$

Acidosis is ph  $<7.35$

Alkalosis is ph  $>7.45$

The ph value is going to give you acid or basic state but the other values can lead you to why!



# BLOOD GAS- PCO2

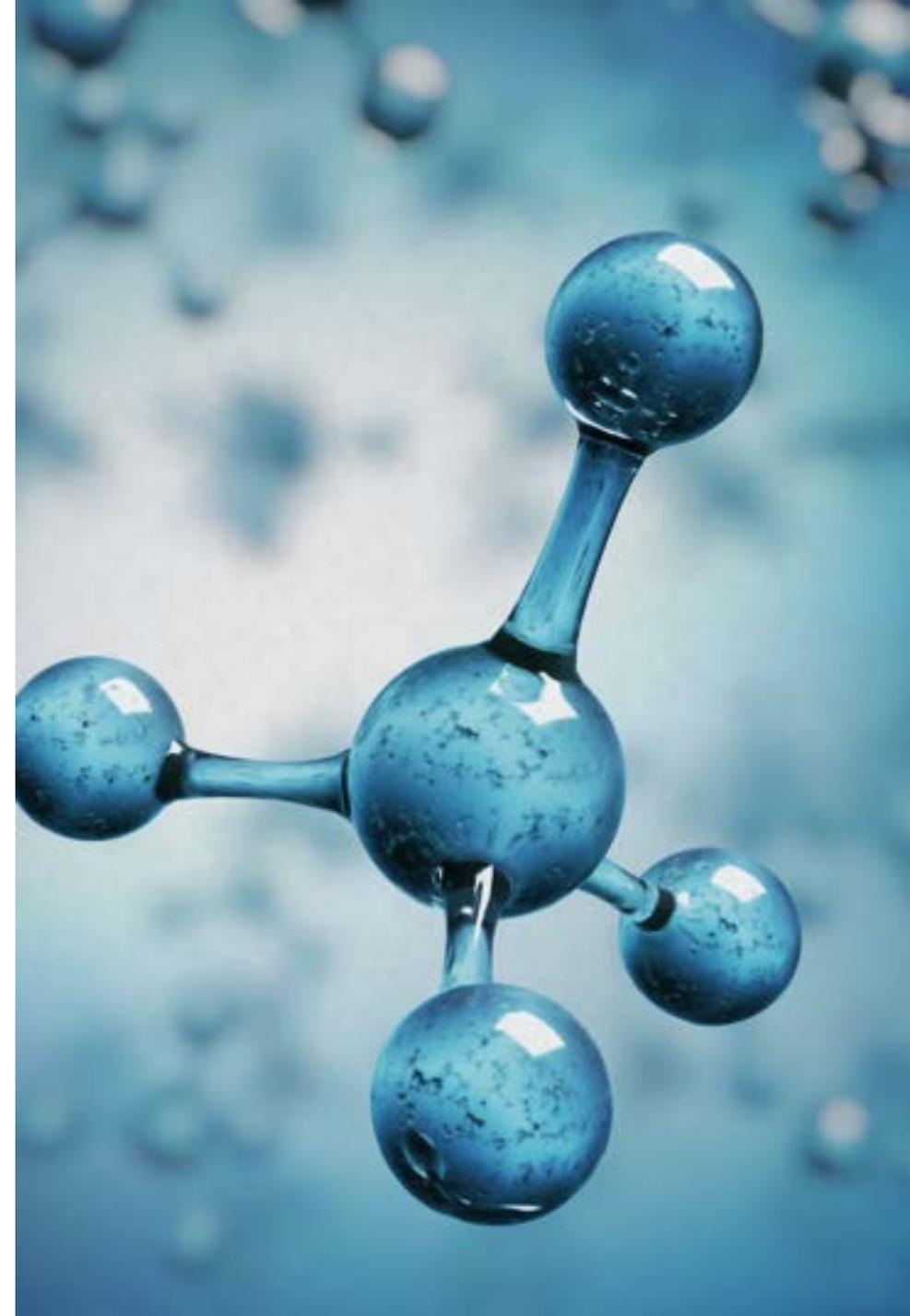
Carbon dioxide (CO<sub>2</sub>)-waste product of cellular metabolism

Three ways carbon dioxide is transported in the blood stream from the peripheral tissues and back to the lungs

1. **dissolved gas (pCO<sub>2</sub>)** 2. bicarbonate 3. carbaminohemoglobin bound to hemoglobin (and other proteins)

Moves via diffusion from peripheral tissues to blood stream

Most CO<sub>2</sub> combines with H<sub>2</sub>O in the RBC to form carbonic acid which immediately dissociates into bicarbonate ions, thus most of the bodies CO<sub>2</sub> is transported via bicarbonate



# BLOOD GAS-PAO<sub>2</sub>

Normal arterial level: 50-80 mm Hg



Assessing **Hypoxemia** (oxygen deficiency in arterial blood mostly due to respiratory conditions)

Oxygen enters lungs during inspiration & diffuses across alveolar-capillary membrane

Oxygen (O<sub>2</sub>) transported to the tissues either as

dissolved in the blood plasma (3%) and reported as partial pressure of oxygen (PaO<sub>2</sub>)

OR

Bound to hemoglobin (97%) reported as the O<sub>2</sub> saturation (SpO<sub>2</sub>)

As the amount of O<sub>2</sub> is increased in the blood (higher PaO<sub>2</sub>) hemoglobin (that is available) becomes more saturated

Affecting factors: fetal hemoglobin (affinity for O<sub>2</sub>), high O<sub>2</sub> consumption states (sepsis, increased temperature, decreased pH)

# BLOOD GAS-HCO<sub>3</sub>

Normal arterial level of HCO<sub>3</sub><sup>-</sup>: 22-26 mEq/L

HCO<sub>3</sub><sup>-</sup> = Bicarbonate: compound capable of accepting H<sup>+</sup> (buffer)

Amount of HCO<sub>3</sub><sup>-</sup> in the body is controlled by the kidneys = kidneys reabsorb

Rate of neonate ability to reabsorb HCO<sub>3</sub><sup>-</sup> is 1/3 of an adults

Primary indicator of metabolic acid-base imbalances

In normal circumstances: kidneys retain or reabsorb HCO<sub>3</sub><sup>-</sup> based on pH and CO<sub>2</sub> in plasma.

- Plasma more acidotic more HCO<sub>3</sub><sup>-</sup> is recovered and reabsorbed by tubular cells
- Plasma alkalotic less HCO<sub>3</sub><sup>-</sup> is retained or reabsorbed.

Decrease in HCO<sub>3</sub><sup>-</sup> = acidosis/Increase = alkalosis

# WHAT'S A BUFFER?

Examples:  $\text{HCO}_3^-$ -(bicarbonate, Plasma proteins, Hemoglobin

- First line of defense against excess  $\text{H}^+$  concentration in the blood.
- Acid is an  $\text{H}^+$  donor, Base is an  $\text{H}^+$  receptor
- Bicarbonate is major buffer, teams with  $\text{H}^+$  to form carbonic acid, which dissociates into water and  $\text{CO}_2$  to be eliminated

# BLOOD GAS-BASE DEFICIT/EXCESS

Normal arterial base excess: -2 - +2

Base excess= calculated value that considers the amount of  $\text{HCO}_3^-$ -generation/retention by the kidneys (metabolic clue).

Answers question: How much a change in  $\text{PaCO}_2$  will cause a change in  $\text{HCO}_3^-$ -

Base excess measures moles of acid needed to return one liter of blood to pH of 7.40

Base deficit are the moles of acid needing to be excreted to return.....

+ number indicates alkalosis

- number indicates acidosis

# NORMAL ARTERIAL BLOOD GAS VALUES

<b>pH</b>	<b>7.35-7.45</b>
<b>PaCO<sub>2</sub></b>	<b>35-45 mm Hg</b>
<b>PaO<sub>2</sub></b>	<b>80-100 mmHg</b>
<b>HCO<sub>3</sub><sup>-</sup></b>	<b>22-26 mEq/L</b>
<b>Base excess</b>	<b>-2 - +2</b>

# BLOOD GASES-ANALYSIS

Respiratory acidosis ( ↓ pH)	PCO <sub>2</sub> >45
Respiratory alkalosis ( ↑ pH)	PCO <sub>2</sub> <35
Metabolic acidosis ( ↓ pH)	HCO <sub>3</sub> <sup>-</sup> <22/base deficit
Metabolic alkalosis ( ↑ pH)	HCO <sub>3</sub> <sup>-</sup> >26/base excess

# AT BIRTH WHAT AFFECTS BLOOD GAS VALUES?

- Respiratory distress syndrome (surfactant insufficiency)
- Meconium aspiration syndrome
- Chorioamnionitis
- Maternal magnesium/general anesthesia
- Congenital heart defects
- Multiple congenital anomalies
- HIE (hypoxic-ischemic encephalopathy)
- Pneumothorax

**Table 26.2****Causes of Acidosis and Alkalosis**

<b>Cause</b>	<b>Mechanism</b>
<b>Respiratory Acidosis (<math>\uparrow</math> <math>P_{aCO_2}</math>, <math>\downarrow</math> pH)</b>	
CNS depression	Maternal narcotics during labor, asphyxia, intracranial hemorrhage, neuromuscular disorder, CNS dysmaturity (apnea of prematurity)
Decreased ventilation-perfusion ratio	Obstructed airway, meconium aspiration, choanal atresia
Decreased lung compliance	Respiratory distress syndrome, pulmonary insufficiency, diaphragmatic hernia
Injury to the thorax	Phrenic nerve paralysis, pneumothorax
<b>Metabolic Acidosis (<math>\downarrow</math> <math>HCO_3^-</math>, pH, and Base Deficit [Negative Value])</b>	
Decreased tissue perfusion	Increased lactic acid production
Sepsis, congestive heart failure	Increased lactic acid production
Renal failure	Increased organic acids
Renal tubular acidosis	Renal loss of base
Diarrhea	Gastrointestinal loss of base
<b>Respiratory Alkalosis (<math>\downarrow</math> <math>P_{aCO_2}</math>, <math>\uparrow</math> pH)</b>	
Iatrogenic	Excessive mechanical ventilation
Hypoxemia	Increase in alveolar ventilation
CNS irritation (pain)	Increase in alveolar ventilation
<b>Metabolic Alkalosis (<math>\uparrow</math> <math>HCO_3^-</math>, pH, and Base Excess [Positive Value])</b>	
Gastric suction	Loss of acid
Vomiting	Loss of acid
Diuretic therapy	Renal losses of $H^+$ ion
Iatrogenic	Administration of $HCO_3^-$ (base added)

CNS, Central nervous system.

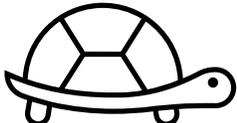
# COMPENSATION

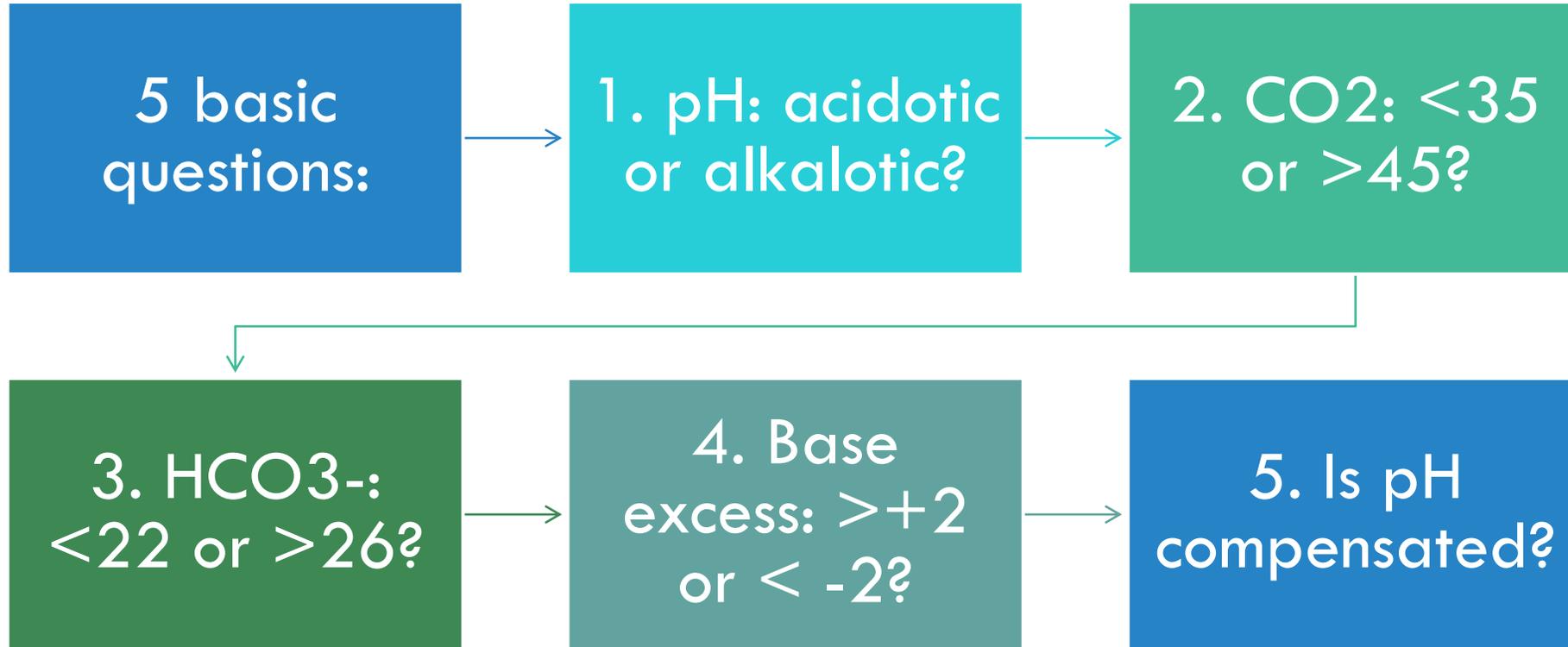
Cause of acid/base imbalance is either respiratory or metabolic

Compensation = Body attempt at maintaining a normal pH level/maintain homeostasis

Respiratory system and kidneys working together to maintain pH in normal range.

Respiratory system = FAST

Kidneys = 



# BLOOD GAS ANALYSIS

## ABG Quick Interpretation

Parameter	Acidosis	Normal	Alkalosis	Reflects
pH	< 7.35	7.35-7.45	> 7.45	Acid/Base Status of Body
pCO <sub>2</sub>	> 45	35-45	< 35	Respiratory Component
HCO <sub>3</sub>	< 22	22-26	> 26	Metabolic Component

**Facts:** Body will not overcompensate when it comes to acid/base balance so:

- pH midpoint is 7.4
- If pH on report is < 7.4 = original problem was acidosis in nature
- If pH on report is > 7.4 = original problem was alkalosis in nature

**Examples:**

1. pCO<sub>2</sub> is High and pH is 7.37 = compensated Respiratory Acidosis because in spite of high pCO<sub>2</sub> which would indicate Acidosis the pH is within normal range indicating that the metabolic component has kicked in and caused pH to shift more towards the midpoint of 7.4 and therefore compensated for the respiratory acidosis.
2. pH is 7.44 and the HCO<sub>3</sub> is 33 it is a Metabolic Alkalosis with Respiratory Compensation bringing the pH back to WNL but not past midpoint.

pH	pCO <sub>2</sub>	HCO <sub>3</sub>	Interpretation*
↑	↓	WNL	Respiratory Alkalosis
↑	WNL	↑	Metabolic Alkalosis
↓	↑	WNL	Respiratory Acidosis
↓	WNL	↓	Metabolic Acidosis
< 7.4	↑	↑	Compensated Resp. Acidosis
> 7.4	↓	↓	Compensated Resp. Alkalosis
< 7.4	↓	↓	Compensated Meta. Acidosis
> 7.4	↑	↑	Compensated Meta. Alkalosis

\*Where compensation is not involved you can look at the arrows:

- Arrows are in same direction = metabolic conditions
- Arrows in opposite direction = respiratory conditions

# BLOOD GAS ANALYSIS PRACTICE

## Uncompensated Respiratory Acidosis

ph 7.22

PCO<sub>2</sub> 65

PaO<sub>2</sub> 47

HCO<sub>3</sub> 22

Base deficit -6.3

## Uncompensated Metabolic Acidosis

ph 7.18

PCO<sub>2</sub> 50

PaO<sub>2</sub> 68

HCO<sub>3</sub> 15

Base deficit -8.4

# BLOOD GAS ANALYSIS PRACTICE

Compensated respiratory  
acidosis

pH 7.36

PCO<sub>2</sub> 66

PaO<sub>2</sub> 54

HCO<sub>3</sub> 32

Base excess +4.5

Mixed respiratory/metabolic  
acidosis

pH 6.81

PCO<sub>2</sub> 97

PaO<sub>2</sub> 23

HCO<sub>3</sub> 15

Base deficit -27

# BLOOD GAS ANALYSIS PRACTICE

Uncompensated Mixed acidosis

ph 7.22

pCO<sub>2</sub> 78

PaO<sub>2</sub> 45

HCO<sub>3</sub> 17

Base deficit -6.7

Uncompensated respiratory acidosis

ph 7.17

PCO<sub>2</sub> 84

PaO<sub>2</sub> 48

HCO<sub>3</sub> 22

Base deficit -7

# COMPLETE METABOLIC PANEL

Blood sample test that measures 14 different substances in the blood. It provides important information about the body's chemical balance and metabolism

- Glucose
- Sodium
- Potassium
- Chloride
- Calcium

*Phosphorus & Magnesium*

**CO2/Anion Gap**

**BUN**

**Creatinine**

**Total Protein/Albumin/Globulin**

**Alkaline Phosphatase**

**ALT (SGPT)**

**AST (SGOT)**

**Magnesium/Phosphorus**

# COMPLETE METABOLIC PANEL-TCO2

Most of the body's CO<sub>2</sub> (cellular waste) is in the form of HCO<sub>3</sub> (bicarbonate)

Bicarbonate is a compound with the ability to accept H<sup>+</sup>, thus a buffer or base.

**CO<sub>2</sub> on the CMP is a measure of Bicarbonate as an indicator of the level of CO<sub>2</sub> in your body**

The test measures all types of carbon dioxide in your blood: bicarbonate, carbonic acid, and dissolved CO<sub>2</sub>.

## BLOOD GAS- PCO<sub>2</sub>

Carbon dioxide (CO<sub>2</sub>)-waste product of cellular metabolism

Three ways carbon dioxide is transported in the blood stream from the peripheral tissues and back to the lungs

1. dissolved gas (pCO<sub>2</sub>) 2. bicarbonate 3. carbaminohemoglobin bound to hemoglobin (and other proteins)

Moves via diffusion from peripheral tissues to blood stream

Most CO<sub>2</sub> combines with H<sub>2</sub>O in the RBC to form carbonic acid which immediately dissociates into bicarbonate, thus most of the bodies CO<sub>2</sub> is transported via bicarbonate

Result	Specimen	Action List
<b>CO<sub>2</sub> 22 mmol/L</b>		
Normal Low	<b>16</b>	Normal High <b>25</b>
Critical Low	<b>&lt;13</b>	Critical High

<input type="checkbox"/> Glucose		* 99 117
<input type="checkbox"/> Sodium		141
<input type="checkbox"/> Potassium		(U) 4.9
<input type="checkbox"/> Chloride		(H) 100
<input type="checkbox"/> CO <sub>2</sub>		19
<input type="checkbox"/> Anion Gap		14
<input type="checkbox"/> BUN		(H) 30
<input type="checkbox"/> Creatinine		0.86
<input type="checkbox"/> eGFR		* N/A
<input type="checkbox"/> Calcium		9.6
<input type="checkbox"/> Phosphorus		6.1
<input type="checkbox"/> Total Protein		(L) 4.1
<input type="checkbox"/> Albumin		2.9
<input type="checkbox"/> Globulin		1.2
<input type="checkbox"/> Total Bilirubin		* 3.0
<input type="checkbox"/> Alk Phos		244
<input type="checkbox"/> ALT(SGPT)		<5
<input type="checkbox"/> AST(SGOT)		67
<input type="checkbox"/> Magnesium		2.6

**Anion Gap** 12 mmol/L

Normal Low 7 Normal High 16

**Anion Gap** 12 mmol/L

Normal Low 7 Normal High 16

**Albumin** 3.0 g/dL

Normal Low 2.9 Normal High 5.5

Result	Specimen	Action List
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<b>Total Protein</b> 4.7 g/dL	(LOW)	
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Normal Low 5.4 Normal High 7.4

**Anion gap:** The anion gap is defined as serum sodium (Na) concentration minus the sum of serum chloride ( $\text{Cl}^-$ ) and serum bicarbonate ( $\text{HCO}_3^-$ ) concentrations. (math equation points to negative to positive gap and certain physiologic conditions)

**Total protein:** This is a measurement of the total amount of albumin and globulins, which are proteins in your blood.

**Albumin:** This is a protein that your liver makes. It transports important substances through your bloodstream and keeps fluid from leaking out of your blood vessels.

**Globulin:** a group of proteins called globulins in blood serum. Serum is the liquid part of blood.

# COMPLETE METABOLIC PANEL

# COMPLETE METABOLIC PANEL



**Alkaline phosphatase:** is an enzyme that's found throughout your body. ALP blood tests measure the level of ALP in your blood that comes from liver and bones.

**Blood urea nitrogen (BUN):** This is a measurement of urea nitrogen, which is a waste product that your kidneys help remove from the bloodstream.

**Creatinine:** This is a byproduct of muscle activity. It's a waste product that your kidneys filter and remove from your blood.

BUN 21 mg/dL

Normal Low 6 Normal High 22

Alk Phos 221 U/L

Normal Low 70 Normal High 350

Creatinine 0.50 mg/dL

Normal Low 0.50 Normal High 1.20

**ALT 9 U/L**

Normal Low 0 Normal High 40

**AST 32 U/L**

Normal Low 15 Normal High 120

**Phosphorus 4.1 mg/dL (LOW)**

Normal Low 4.8 Normal High 8.2

**Magnesium 2.1 mg/dL**

Normal Low 1.7 Normal High 2.6

**Alanine transaminase (ALT)** is an enzyme that mainly exists in your liver

**Aspartate transferase (AST)** is an enzyme that's found in your liver, heart, pancreas, muscles and other tissues in your body

**Magnesium/Phosphorus:** bone health, checked with premature infants for bone/metabolic health

# COMPLETE METABOLIC PANEL

# AMMONIA

Produced from the deamination of amino acids during protein metabolism and is the byproduct of bacteria protein breakdown.

Metabolized by the liver

Elevated in liver failure/acute or chronic liver disease

Elevated in inborn errors of metabolism

Normal values 90-150 mcg/dL

General Hematology Results	
<input type="checkbox"/>	WBC
<input type="checkbox"/>	RBC
<input type="checkbox"/>	HGB
<input type="checkbox"/>	HCT
<input type="checkbox"/>	MCV
<input type="checkbox"/>	MCH
<input type="checkbox"/>	MCHC
<input type="checkbox"/>	RDW
<input type="checkbox"/>	RDW-SD
<input type="checkbox"/>	MPV
<input type="checkbox"/>	Platelet
<input type="checkbox"/>	PRELIM ANC
<input type="checkbox"/>	Neutrophils
<input type="checkbox"/>	Band
<input type="checkbox"/>	Lymph
<input type="checkbox"/>	Reactive Lymphs
<input type="checkbox"/>	Mono
<input type="checkbox"/>	Eos
<input type="checkbox"/>	Baso
<input type="checkbox"/>	Immature Neutrophils
<input type="checkbox"/>	Meta
<input type="checkbox"/>	Myelo
<input type="checkbox"/>	nRBC
<input type="checkbox"/>	ABS Neut
<input type="checkbox"/>	ABS Lymph
<input type="checkbox"/>	ABS Imm Neut
<input type="checkbox"/>	ABS Mono
<input type="checkbox"/>	ABS Eos
<input type="checkbox"/>	ABS Basos
<input type="checkbox"/>	ABS nRBC



# COMPLETE BLOOD COUNT

A CBC does many tests to measure and study red blood cells, white blood cells and platelets.

- **Red blood cells** carry oxygen throughout the body.
- **White blood cells** are part of your immune system. They help your body fight infection.
- **Platelets** help your body clot

**CBC without differential** counts the total number of white blood cells.

**CBC with differential.** There are five kinds of white blood cells. The differential looks at how many of each kind of white blood cell you have.

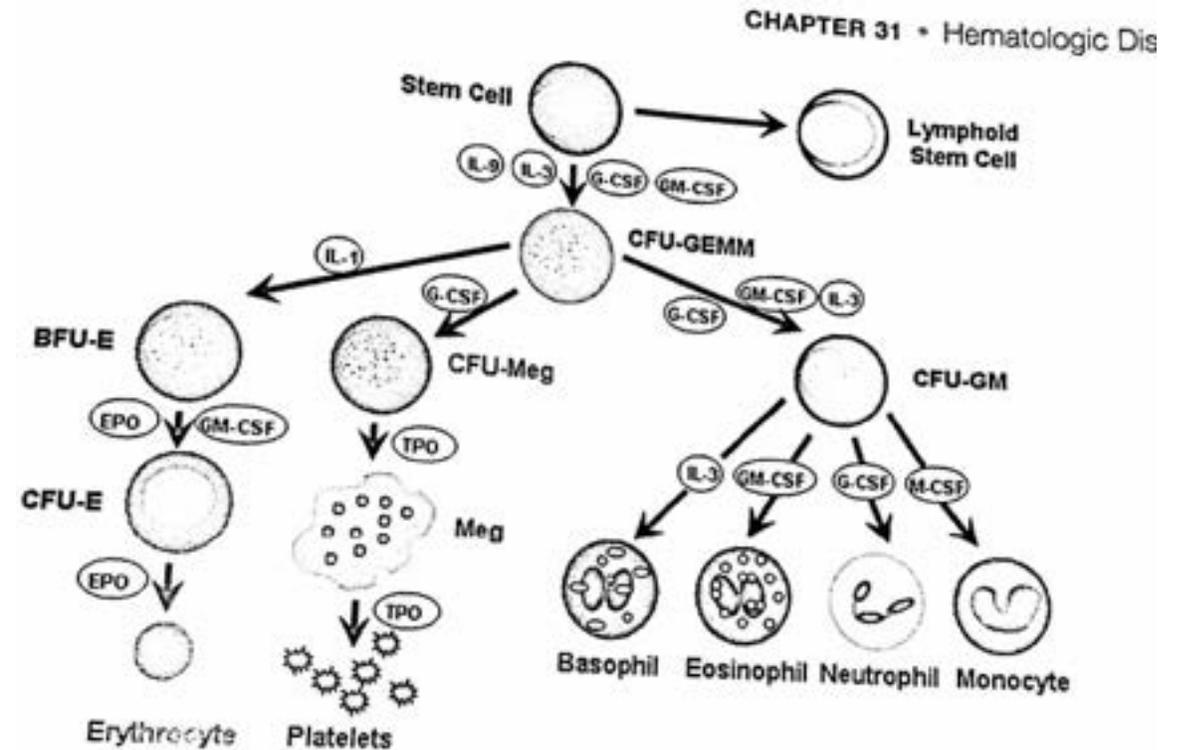
**Hemoglobin** tests measure hemoglobin, the protein in red blood cells that carries oxygen.

**Hematocrit** describes the concentration of red blood cells in your blood

Aniso
Poly
Macro
Ovalocytes
Teardrops
Echinocytes
Plat Estimate
Giant Platelet
Pathologist Review Submitted
Path Review

# HEMATOPOIESIS

Formation,  
production,  
maintenance of  
blood cells  
Occurs in the bone  
marrow



Hematopoiesis and selected growth factors. *BFU-E*, Burst-forming unit, erythroid; *CFU-E*, colony-forming unit-erythroid; *CFU-GEMM*, colony-forming unit-granulocyte, erythrocyte, monocyte, megakaryocyte; *CFU-GM*, colony-forming unit-granulocyte-macrophage; *CFU-Meg*, colony-forming unit-megakaryocyte; *EPO*, erythropoietin; *G-CSF*, granulocyte colony-stimulating factor; *GM-CSF*, granulocyte-macrophage colony-stimulating factor; *IL-1*, interleukin-1; *IL-3*, interleukin-3; *IL-9*, interleukin-9; *M-CSF*, macrophage colony-stimulating factor; *Meg*, megakaryocyte; *TPO*, thrombopoietin. (From Israels, L.G., & Israels, S. J. [2002]. *Mechanisms in hematology* [3rd ed., p. 402]. Toronto: McGraw-Hill Health Sciences.)

# COMPLETE BLOOD COUNT

**Erythropoiesis:** production of erythrocytes (RBCs)

- Regulated by the hormone Erythropoietin-regulated by hypoxia sensing mechanisms in the kidneys

**RBC:** oxygen transport to tissues, CO<sub>2</sub> transport

**RBC count:** number of circulating mature RBC's per cubic millimeter

RBC lifespan for term infant = 60-70 days, preterm infant = 35-50 days

**nRBC:** nucleated RBC are circulating immature RBCs

increased number can indicate oxidative (cellular) stress

**RBC indices:** measure of RBC size and hemoglobin content used for designation of anemias

# CBC-RED CELL INDICES

**Mean corpuscular volume (MCV):** average size and volume of a single RBC.

**Mean corpuscular hemoglobin (MCH):** average amount (by weight) of hemoglobin in each RBC

**Mean corpuscular hemoglobin concentration (MCHC):** average concentration of hemoglobin per single RBC

**Red blood cell distribution width (RDW):** measures how varied RBC are in size and volume.

**Mean platelet volume (MPV):** measures the average size of your platelets.

All help define different anemias

General Hematology Results	
WBC	8.6
RBC	5.07
HGB	(H) 20.7
HCT	57
MCV	112
MCH	(H) 41
MCHC	(H) 36.5
RDW	(H) 16.6
RDW-SD	(H) 68
MPV	10.2
Platelet	(L) 132
Neutrophils	48.5
Lymph	33.8
Mono	12.5
Eos	1.5
Baso	1.4
Immature Neutrophils	(H) 2.3
nRBC	(H) 12.7
ABS Neut	4.2
ABS Lymph	2.9
ABS Imm Neut	(H) 0.20
ABS Mono	1.1
ABS Eos	0.1
ABS Basos	0.1
ABS nRBC	1.1
Poly	Slight
Macro	Moderate
Plat Estimate	* Normal
Platelet Clumps	* Present

# COMPLETE BLOOD COUNT



**WBC count:** number of circulating WBC's per cubic millimeter

**Platelets:** small nonnucleated disk-shaped cells aid in hemostasis, coagulation, and thrombus formation

**Hemoglobin:** major iron-containing component of the RBCs

- Carries oxygen from lungs to tissue cells

**Hematocrit:** percentage of RBCs in a unit volume of blood

**Absolute Neutrophil count:** how many neutrophils (infection fighting WBCs) are in a sample of your blood.

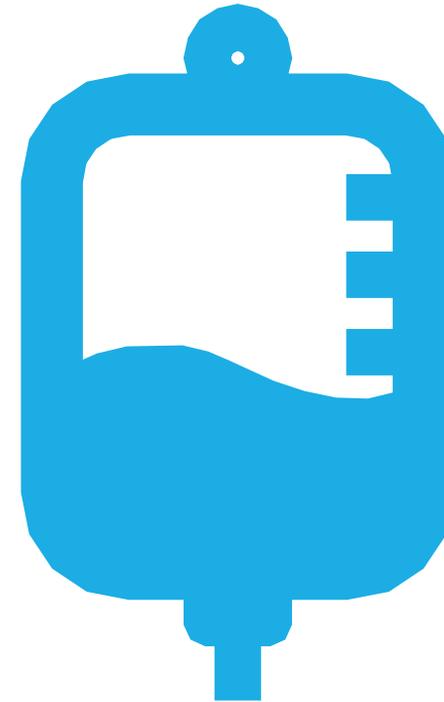
**Reticulocytes:** immature RBC

- In absence of stress mature 1-2 days in the bone marrow and 1 more day in circulation before fully mature

Low hemoglobin concentration and/or decreased number of RBC (HCT)=diminished oxygen carrying capacity of the blood and level of oxygen available to the tissues.

3 main cause at birth:

1. Blood loss
2. RBC shorted life span
3. Underproduction of erythrocytes



HCT/HGB (ANEMIA)

# HCT/HGB (ANEMIA)

Hemorrhage (blood loss)	Shortened RBC survival	Under production of RBC
<p>Fetal-maternal (acute or chronic)</p> <p>Twin to twin transfusion (mono/mono)</p> <p>Placenta/cord issues</p> <p>Internal hemorrhage: IVH or ICH, organ rupture, pulmonary bleed</p> <p>External: phlebotomy, iatrogenic</p>	<p>Hemolysis: ABO incompatibility, Rh sensitization, enzymatic defects (ex. G6PD), hemoglobin disorders (ex. A-thalassemia)</p>	<p>PREMATURITY-considered physiologic/naturally occurring</p> <p>Decreased erythropoietin production</p>

# POLYCYTHEMIA

(100,000 TO 400,000/ MM<sup>3</sup> FIRST 10 DAYS OF LIFE)

- Excess of circulating RBCs
  - Hct > 65% (venous sample)
  - Impairs peripheral circulation r/t blood viscosity
    - Can lead to reduction of blood flow to organs
- Active Polycythemia
  - Excess production of RBCs
- Passive Polycythemia
  - Transfusions
  - Twin to twin
    - Monochorionic monoamniotic
    - Blood is exchanged unequally between the fetuses with the donor twin being anemic and the recipient twin polycythemia
  - Delayed cord clamping
  - Fetal stress

# THROMBOCYTOPENIA (< 150,000)

- Caused by decreased production, increased destruction, sequestration or loss
- **Most common bleeding disorder in newborn**
- Bacterial/viral infections/NEC
- Hypoxia/birth asphyxia
- Disseminated Intravascular Coagulation (DIC)
- Persistent Pulmonary Hypertension of Newborn (PPHN)
- Polycythemia
- Congenital infections
- Congenital syndromes (Down syndrome)
- Preeclampsia-falsely low
- Giant hemangiomas
- Late indicator of sepsis
- Watch for wide-spread petechia

# TRANSFUSION GUIDELINES

## Transfusion Criteria

Post-natal Age	Respiratory Support* Hct (Hgb)	No Respiratory Support Hct (Hgb)
0 through 6 Days of life	32 (11.0)	29 (10.0)
7 through 13 days of life	29 (10.0)	25 (8.5)
14 days of life or greater	25 (8.5)	21 (7.0)

\*Respiratory support was defined as mechanical ventilation, continuous positive airway pressure (CPAP),  $FiO_2 > 0.35$ , or high flow nasal cannula of 1 liter/minute or greater (includes room air nasal cannula of 1 liter/min or more).

## Transfusion Criteria

	Population	Threshold
Thrombocytopenia	Stable preterm or term infant	25,000/ $\mu$ L
	ELBW (BW<1000g) in the first week of life	25,000/ $\mu$ L
	Clinically unstable* infant	25,000/ $\mu$ L
	Immediately pre-/post-op	50,000/ $\mu$ L
	ECMO or active bleeding	100,000/ $\mu$ L
Normal platelet count	Massive transfusion protocol initiated	
	Active bleeding with known qualitative platelet defect	

\*Clinically unstable defined as evidence of sepsis, necrotizing enterocolitis, intracranial hemorrhage, hypotension requiring vasopressors, or other hemodynamic instability

# COMPLETE BLOOD COUNT-WBC TYPES

- Mature in the bone marrow and lymphatic tissue
- WBC's leave circulation to enter the tissues where they function as part of the immunologic system in reaction to foreign proteins.
- WBCs=granulocytes, lymphocytes, and monocytes

Granulocytes	Lymphocytes-immune response	Monocytes-macrophages
Basophils Eosinophils Neutrophils	Thymus derived T cells Bone marrow derived B cells	Circulating immature macrophages

# COMPLETE BLOOD COUNT

## CBC-White Blood Cells with differential

- Protect against infective organisms and foreign substances
- Leukocytosis and leukopenia can be problematic
- 5 main types of WBCs

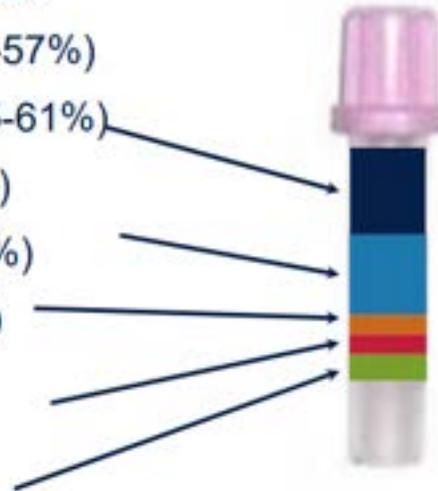
- **Neutrophils** (31-57%)

- Lymphocytes (35-61%)

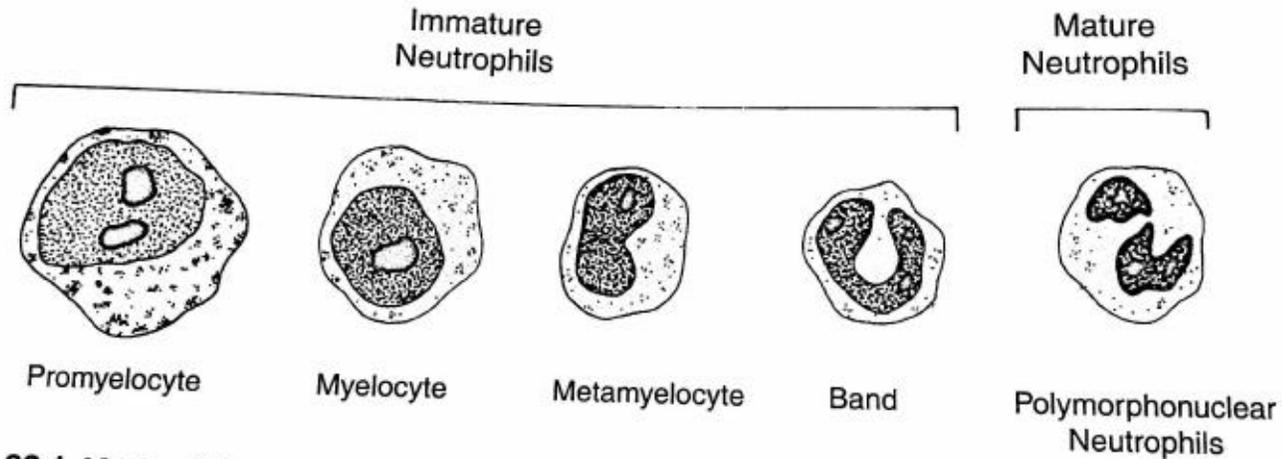
- Monocytes(4-7%)

- Eosinophils (2-4%)

- Basophils (0-1%)



## Neutrophil: Stages of Maturation



**Fig. 32.1** Neutrophils represent a percentage of the total white blood cell count and are reported as the differential on a complete blood cell count.

# NEUTROPHIL: STAGES OF MATURATION

White blood cell count (WBC): 5000-30,000

Not always good indicator of infection

Neutrophilia: inconsistent response to infection, often normal with serious infection

Neutropenia: risk for infection based on not enough infection fighting cells.  $<1800 \text{ cells/mm}^3$ ?

# COMPLETE BLOOD COUNT

## How do you know its infection?

- Early onset <72 hours
  - From maternal origin through vertical transmission through amniotic fluid or from bacteria from a vaginal delivery
- Most common group B streptococci/e coli
- We follow maternal risk factors (GBS status, gestational age, chorioamnionitis, prolonged rupture of membranes.
- Vital sign trends and clinical status with WBC value
- I/T ratios (immature-total): increase in the I/T ratio is known as a left shift, reflects an increase in immature neutrophils circulating.
  - Best served as a negative predictive value-if its normal it's probably not infection

# DISSEMINATED INTRAVASCULAR COAGULATION (DIC) PANEL

Acquired hemorrhagic disorder associated with an underlying disease manifested as uncontrolled activation of coagulation and fibrinolysis.

- Consumption of clotting factors through cascade that starts with damaged or diseased tissue
- Microvascular thrombosis that this lyses and increases bleeding

## Values include:

- Partial thrombin (PT), Partial thrombin time (PTT), Thrombin Time, Fibrinogen, D-dimer, Platelet Count and Schistocyte DIC-not on neonatal panel
- Clotting times are lengthened, Fibrinogen is low (<200), D-dimer elevated=clots being lysed, platelet count is low
- Aggressively treat underlying disease/condition
- Replaced clotting factors: PRBC, Fresh Frozen Plasma (all clotting factors), Cryoprecipitate (fibrinogen, factor VIII), platelets

# C-REACTIVE PROTEIN (CRP)

- Acute phase reactant synthesized in the liver
- Appears in blood during an acute inflammatory process/associated with tissue injury
- Most recent research does not support the use of CRP for identifying infections
- Not specific for infection-rules out the false negatives
- Not sensitive for infection-just because it's elevated doesn't mean it's infection
- Change in practice is hard!

# LACTATE

- Evaluates lactic acid
- Measures perfusion and adequate oxygen function to the cells
- Increased when hypoxia present = tissue injury/necrosis
- Increased with sepsis r/t anaerobic metabolism
- CHD and sepsis

# INFECTION RELATED LAB

## Cerebral Spinal Fluid

Obtained via lumbar puncture or ventricular tap/reservoir tap

Bacterial/Viral meningitis

Cell count, Glucose, Protein, Gram stain/culture and/or viral studies

## Urine

Urine culture and/or urine analysis

Culture obtained via straight catheter specimen or indwelling foley

- CFU >50,000

Higher incidence in longer LOS

Urine analysis obtained with bag

CMV testing (urine/CSF)

# INFECTION RELATED LAB

## Respiratory Viral Panel (RVP)

Nasal swab

Includes- COVID-19, Influenza A, B, RSV, Adenovirus, Human Metapneumovirus, Parainfluenza Viruses 1-4, and Rhinovirus

Droplet isolation while waiting for results

If positive isolation continues based on pathogen identified

## Blood culture

Obtained sterilely via indwelling line or peripheral puncture

Ideally 1-2ml for best bacterial count

99% positive by 36-48H

Repeat until negative culture is obtained

Consider peripheral site if indwelling line draw is positive

# OTHER

Therapeutic drug monitoring

- antibiotic peak/troughs/random levels
- Phenobarbital level (therapeutic around 40mcg/ml)

Anti Xa level: used to guide anticoagulation treatment ie) Lovenox

# THYROID STUDIES

- Hypothyroidism: Permanent (congenital) or acquired (transient)
- TSH
  - Stimulates thyroid to produce T4 and T3
- T4
  - Needed for pre and postnatal brain maturation

Thyroid hormones responsible for almost every tissue in the body

# CONGENITAL HYPOTHYROIDISM

- Most preventable and treatable cause of mental retardation – symptoms don't often present until 6 weeks of age
  - Deficiency in thyroid hormone
  - Tested for on state newborn screens
- TSH – thyroid stimulating hormone used to predict congenital hypothyroidism
- T4 – tests newborns for hypothalamic-pituitary-hypothyroidism
- Treatment – thyroxine

# NEWBORN STATE SCREENS

60 conditions screened: Rare genetic/metabolic/endocrine disorders

Required by law for every baby born in Nebraska to be screened

Healthy term infants: collected between 24-48 hours old

NICU: #1: drawn on admission #2: drawn 48-72 hours, #3: 28 days of life (if <2kg)

Why important?

- Conditions usually don't have signs or symptoms recognizable until internal damage is done
- Conditions are treatable, early treatment can prevent serious problems

SCREENING: false positives are common, if positive result specific testing for condition recommended or repeat dried blood spot card

[Nebraska | Newborn Screening \(hrsa.gov\)](https://www.hrsa.gov/newborn-screening/nebraska)

<http://dhhs.ne.gov/Newborn%20Screening/2018%20Practitioners%20Manual%205th%20Edition.pdf>

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