



# **Advanced Blood Gas Analysis From Physiology to Practice**

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# OBJECTIVES

Indications for BGA

Composition of a BG report

Sample types

Pitfalls in BGA

Steps in BG interpretation

Case scenarios

Clinical Mx of acid base abnormalities

# Indications for BG analysis

- To obtain information about ventilation (PCO<sub>2</sub>) oxygenation (PO<sub>2</sub>)
- To obtain information about acid base balance or electrolyte disturbance
- To evaluate response to a clinical intervention



Analyzer #: 8  
 Analyzed on 11-11-2023 at 09:11 AM  
 Sample #: 24897  
 Operator ID:  
 Patient ID: 173861  
 FIO<sub>2</sub> %: 20.9  
 Patient Temperature C: 37.0  
 Sample Type: Arterial

Time Drawn: \_\_\_\_\_

Results - Measured at 37°C

pH	7.373		
pCO <sub>2</sub>	40.4	mmHg	
pO <sub>2</sub>	41.9	mmHg	
SO <sub>2</sub> %			UC
Hct	41	%	
Na <sup>+</sup>	144.0	mmol/L	
K <sup>+</sup>	4.00	mmol/L	
Ca <sup>++</sup>	0.81	mmol/L	
Glu		mg/dL	UC
Lac		mmol/L	UC

Results - Calculated

HCO <sub>3</sub> <sup>-</sup>	23.7	mmol/L	
TCO <sub>2</sub>	24.9	mmol/L	
BE <sub>ecf</sub>	-1.7	mmol/L	
BE <sub>b</sub>	-0.8	mmol/L	
SBC	23.2	mmol/L	
A	100.9	mmHg	
A-aDO <sub>2</sub>	59.1	mmHg	
a/A	0.4		
RI	1.4		
PO <sub>2</sub> /FIO <sub>2</sub>	200.4	mmHg	
SO <sub>2</sub> %	76.0		
Hb	13.7	g/dL	
nCa <sup>++</sup>	0.80	mmol/L	

System Status

33 SO<sub>2</sub>% Overload

# Composition of BG report

## Directly measured variables

- pH
- pCO<sub>2</sub>
- pO<sub>2</sub>
- SpO<sub>2</sub>
- Electrolytes –Na/K/Ca/Cl
- Hct
- Glucose
- Lactate

## Indirectly/Calculated variables

- HCO<sub>3</sub>/Std HCO<sub>3</sub>
- Base Excess-Blood/ECF (Std)
- Hb
- a/A
- P/F

# Actual Bicarbonate



- ▶ Derived from pH and  $p\text{CO}_2$  using the Henderson-Hasselbach equation.
- ▶ Normal range 21 - 26 mmol.
- ▶ **H-H equation:  $\text{pH} = 6.1 + \log [\text{HCO}_3^-] / 0.03 p\text{CO}_2$**
- ▶ 6.1 is the pKa (negative logarithm of the acid dissociation constant) for carbonic acid ( $\text{H}_2\text{CO}_3$ )

# Standard Bicarbonate



- ▶ The calculated concentration of  $\text{HCO}_3^-$  if the sample it were at 'standard conditions'.
- ▶ At a temperature of  $37^\circ\text{C}$  and a  $\text{pCO}_2$  of 5.3 kPa (40mmHg; the middle of the  $\text{pCO}_2$  reference range).
- ▶ By assuming a  $\text{pCO}_2$  of 5.3kPa, the standard bicarbonate compensates for any abnormality in  $\text{HCO}_3^-$  caused by a respiratory cause.

## Base Excess (Buffer Base)

- ▶ Base excess and base deficit refer to an excess or deficit, respectively, in the amount of base present in the blood.
- ▶ Usually reported as a concentration in units of mEq/L (milliequivalents of solute per litre of solvent).
- ▶ Positive numbers indicating an excess of base and negative a deficit.

## Base Excess (Buffer Base)

- ▶ Base excess beyond the reference range indicates:
- ▶ metabolic alkalosis if too high (more than +2 mEq/L)
- ▶ metabolic acidosis if too low (less than -2 mEq/L)

**Base Excess** : Amount of Acid or Alkali required to return plasma **in vitro** to normal pH under standard conditions.

**Standard BE:** BE calculated for Anaemic Blood (Hb = 5Gm%).

- ▣ Since Hb effectively buffers plasma & ECF to a large extent.
- ▣ Quantity of Acid or Alkali required to return plasma **in-vivo** to a normal pH under standard conditions



## Types of samples

- Arterial
- Venous –peripheral/central
- Capillary

What should be your choice?

# Disadvantages of Arterial Puncture

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Failure and/or pain

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Haematoma or haemorrhage

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Arteriospasm

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Arterial occlusion

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Vasovagal response

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Pseudoaneurysm

# Self Fill Syringe





# Venous Blood Gas



Done when you need to determine mainly  
 $\text{pH}$ ,  $\text{pCO}_2$ ,  $\text{HCO}_3$



**Peripheral**

Drawn without a tourniquet



**Central**

ICU/HDU setting to  
determine  $\text{ScvO}_2$

	Arterial	Venous	Arterio-venous (A-V) difference
pH	7.35-7.45	7.31-7.41	~ 0.04
pCO <sub>2</sub> (kPa)	4.7 - 6.0	5.5 - 6.8	~ 0.6
pCO <sub>2</sub> (mmHg)	35 -45	41 - 51	~ 6
Bicarbonate (mmol/L)	22-28	23-29	~ 1
PO <sub>2</sub> (kPa)	10.6 - 13.3	4.0 -5.3	~ 8.0
pO <sub>2</sub> (mmHg)	80-100	30 -40	~ 55
sO <sub>2</sub> (%)	> 95	75	> 20

# ABG vs VBG ?

## pH

- Good correlation

## pCO<sub>2</sub>

- Good correlation in normocapnia
- Correlation dissociates when PaCO<sub>2</sub> >45mmHg

## HCO<sub>3</sub>

- Good correlation

## Lactate

- Good correlation, dissociates >2

## PO<sub>2</sub>

- PO<sub>2</sub> values compare poorly

# When do we need ABG ?

- To accurately determine PaO<sub>2</sub> / PaCO<sub>2</sub> in a ventilated patients
- To accurately determine parameters in a haemodynamically unstable patients

# What are the preferred sites for ABG sampling

Radial

Brachial

Femoral

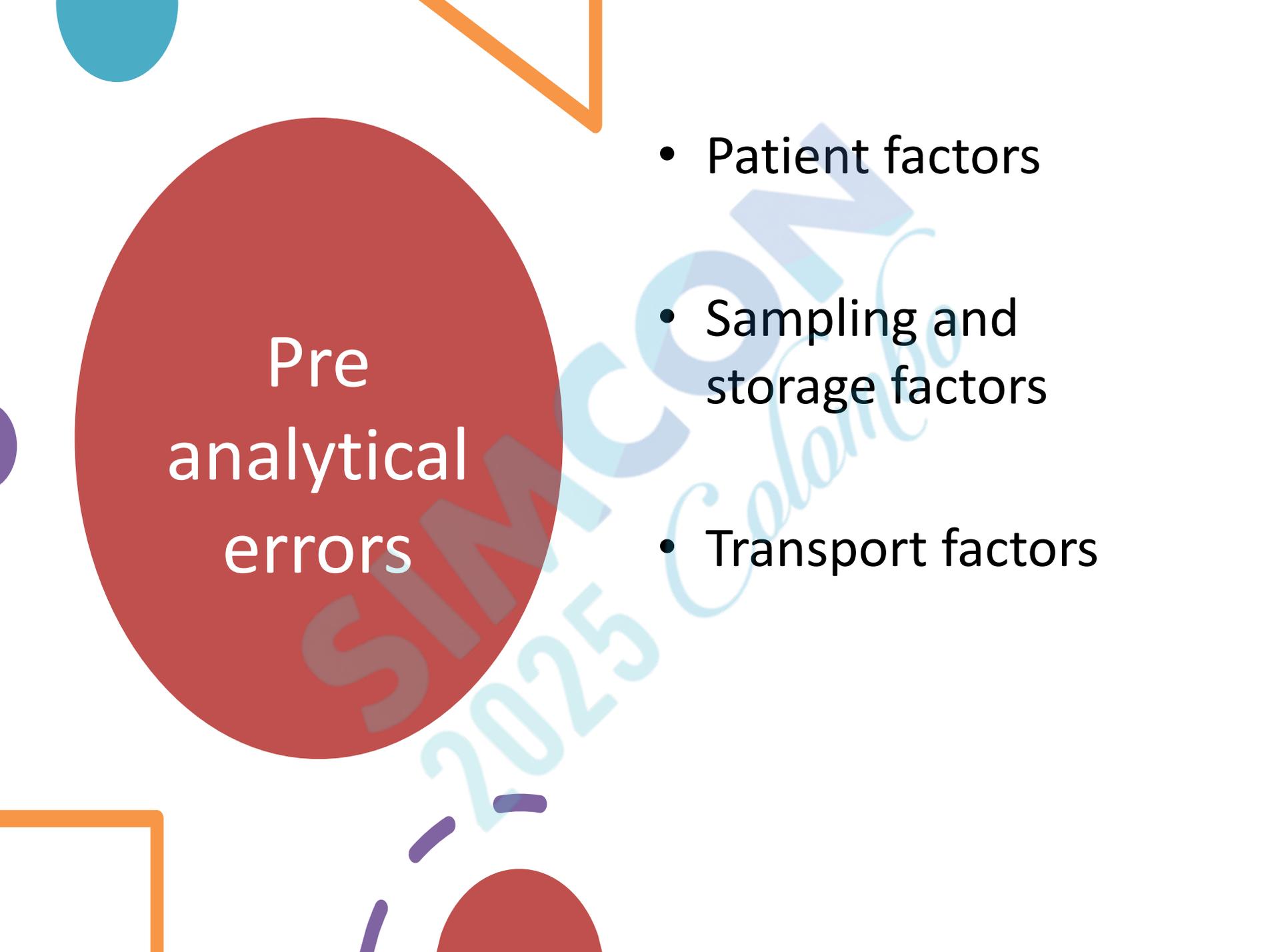
Preferably arterial line

# Pitfalls in BG Analysis

Pre analytical

Analytical

Post analytical



# Pre analytical errors

- Patient factors
- Sampling and storage factors
- Transport factors

Table III: A summary of the sources of error and their consequences during blood gas analysis

Error	Consequences
<b>Pre-analytical phase</b>	
Test over-utilisation	Iatrogenic anaemia
Entrainment of air bubbles	PO <sub>2</sub> /PCO <sub>2</sub> equilibration with environment False hypocalcaemia
Storage on ice	PO <sub>2</sub> equilibration with environment
Excessive time to analysis	Significant deviations in PO <sub>2</sub> /PCO <sub>2</sub>
Excessive/incorrect heparin	Analyte dilution Deviations in cation measurements
Aspiration of sample from IV fluid containing catheter	Dilution of analytes Falsely elevated or reduced lactate level depending on fluid type
Haemolysis	False hyperkalaemia, hyponatraemia, hypocalcaemia
Transport via pneumatic tube system	PO <sub>2</sub> /PCO <sub>2</sub> equilibration with air bubbles Haemolysis
<b>Analytical phase</b>	
	Amplification of errors from previous phase
Improper QC/maintenance	Unreliable results
Electrolytes with ionisation similar to standard electrolytes	Falsely elevated electrolyte with similar ionisation
Antifreeze intoxication	False hyperlactataemia
Aspirin intoxication	False hyperchloraemia
<b>Post analytical phase</b>	
	Amplification of errors from previous phases
Use of different analysers for serial analysis	Improper trends in pH, PO <sub>2</sub> , PCO <sub>2</sub>
Application of traditional approach (Henderson-Hasselbalch)	Amplification of errors from previous phases Too simple to identify complex acid-base disturbances
Application of physicochemical and semiquantitative approaches (Stewart and FencI)	Amplification of errors from previous phases Cumbersome May only be applicable to critical patients
Use of incorrect reference range for species	Incorrect interpretation of results

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**Analytical phase**

Improper QC/maintenance

Electrolytes with ionisation similar to standard electrolytes

Antifreeze intoxication

Aspirin intoxication

Amplification of errors from previous phase

Unreliable results

Falsely elevated electrolyte with similar ionisation

False hyperlactataemia

False hyperchloraemia

---

**Post analytical phase**

Use of different analysers for serial analysis

Amplification of errors from previous phases

Application of traditional approach (Henderson-Hasselbalch)

Improper trends in pH, PO<sub>2</sub>, PCO<sub>2</sub>

Application of physicochemical and semiquantitative approaches (Stewart and Fencel)

Amplification of errors from previous phases  
Too simple to identify complex acid-base disturbances

Use of incorrect reference range for species

Amplification of errors from previous phases  
Cumbersome  
May only be applicable to critical patients

Incorrect interpretation of results

# Interpretation of Blood Gases

1. Look at the **pH**
2. What is the **primary disorder**
3. In **Respiratory disorder** compensation **acute or chronic**
4. In **Metabolic disorder** ,is the compensation **pure**
5. Is there a **mixed** metab/resp.disorder
6. In MA -Is there a **HAGMA**
7. If HAGMA calculate -**Delta ratio**
8. Final **Interpretation**/Clinical diagnosis

# Case 1 :Known COPD on inhalers , presents with 3 days history of worsening SOB

pH	6.95
pCO <sub>2</sub>	80
HCO <sub>3</sub>	40.2
Na	134
Cl	102
PaO <sub>2</sub>	78

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# Step 1

- **Look at the pH**

<7.35- acidosis

>7.45-alkalosis

**Normal 7.35-7.45** –does not rule out a complex acid base disorder

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## Step 2

**What is the primary disorder**

Look at pH and  $p\text{CO}_2$  –

**moved same direction-Metabolic**

**moved opposite direction-Respiratory**

This is not true always !!

Mixed Met/Res acidosis  $\downarrow$  pH-( $\downarrow$   $\text{HCO}_3^-$ )+ ( $\uparrow$   $\text{CO}_2$ )

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## Step 3 – Is the compensation acute/chronic

- **Acute**- starts in 5-10 min ,modest
- **Chronic**-Starts in 3-5 days

- **Respiratory acidosis**

Acute-  $\uparrow 1$  HCO<sub>3</sub> for every 10 rise in pCO<sub>2</sub>

Chronic -  $\uparrow 4$  HCO<sub>3</sub> for every 10 rise in pCO<sub>2</sub>

- **Respiratory alkalosis**

Acute -  $\downarrow 2$  HCO<sub>3</sub> for every 10 decrease in pCO<sub>2</sub>

Chronic-  $\downarrow 5$  HCO<sub>3</sub> for every 10 decrease in pCO<sub>2</sub>

( Respiratory 1,4-2,5 Rule)

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7. If HAGMA calculate **-Delta ratio**
8. **Final Interpretation/Clinical diagnosis**

# Step 8–Interpretation/Clinical diagnosis

- Determining the final interpretation of BGA
  - a) Partially compensated high anion gap pure metab. acidosis
  - b) Partially compensated chronic respiratory acidosis
  - c) Mixed metab./respiratory acidosis
- Clinical diagnosis
  - a) DKA
  - b) COPD exacerbation
  - c) Severe pneumonia with sepsis

# A1

- Partially compensated  $\text{CO}_2$  retaining chronic respiratory acidosis
- COPD exacerbation

# Management

1. V 28 Oxygen
2. Start EWS chart
3. Reassess WOB
4. Start BiPAP IPAP -15 EPAP -5
5. Repeat BGA

## Case 2: Advanced level female student was brought by schoolteachers with acute SOB

pH	7.56
pCO <sub>2</sub>	20
HCO <sub>3</sub>	19.8
Na	131
Cl	101

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
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## A2

- Partially compensated acute respiratory alkalosis
- Voluntary hyper ventilation

# What's your Management

- Breath into a paper bag ?
- Reassure

**Case 3:** Known type 1 diabetic off insulin for 3 days presents with vomiting and altered behavior.

pH	7.03
pCO <sub>2</sub>	18.9
HCO <sub>3</sub>	7.1
Na <sup>+</sup>	134
Cl <sup>-</sup>	96

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder compensation is pure or mixed
5. Is there a mixed metab/resp.disorder
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## Step 4 – In MA is the compensation pure

- **Metabolic acidosis** (Winters formula)

Expected  $p\text{CO}_2$ -  $(1.5 \times \text{HCO}_3)+8$  (+/- 2)

- **Metabolic alkalosis**

Expected  $p\text{CO}_2$ -  $(0.7 \times \text{HCO}_3)+20$  (+/- 5)

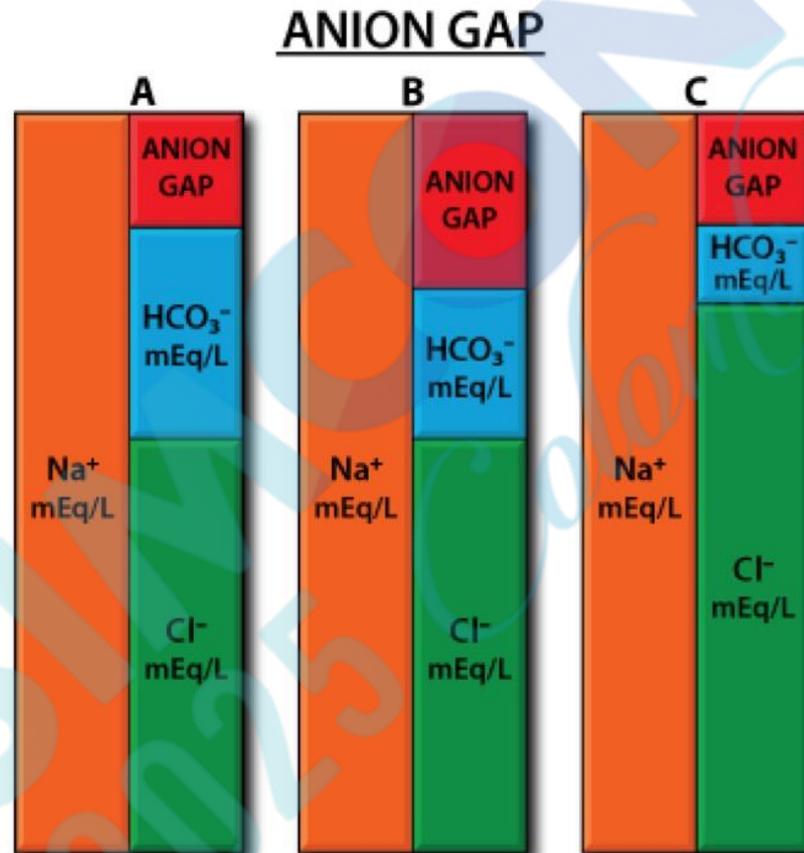
Lowest  $p\text{CO}_2$  - 8-12

Maximum  $p\text{CO}_2$  - 55

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# Anion Gap



A. NORMAL ION DISTRIBUTION

B. METABOLIC ACIDOSIS due to acid accumulation;  
decreased HCO<sub>3</sub><sup>-</sup>, increased anion gap

C. METABOLIC ACIDOSIS due to HCO<sub>3</sub><sup>-</sup> loss;  
decreased HCO<sub>3</sub><sup>-</sup>, normal anion gap, increased Cl<sup>-</sup>

# Step 5 –In MA calculate anion gap

- Metabolic acidosis

$$\text{Anion Gap} = (\text{Na}^+) - (\text{Cl}^-) - (\text{HCO}_3^-)$$

Normal 10-14 (12)

# Anion Gap

- $AG = (Na^+) - (HCO_3^- + Cl^-)$
- Three clinical applications
  - 1) Presence or absence of an AG assists in determining the cause of the metabolic acidosis.
  - 2) The AG is useful in determining the presence of a mixed acid/base disturbance by calculating the Delta Ratio
  - 3) The AG can be useful in detecting selected disorders that occur with a low, not high AG. (Lithium toxicity, Multiple Myeloma with production of cationic paraproteins)

# CAUSES OF INCREASED ANION GAP METABOLIC ACIDOSIS

- **M: M**ethanol
- **U: U**remia
- **D: D**KA
- **P: P**araldehyde
- **I: I**ron / Isoniazid
- **L: L**actic acidosis
- **E: E**thylene glycol
- **S: S**alicylates

**“MUDPILES”**



# Interpretation of Blood Gases

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# 7. In HAGMA Calculate Delta ratio

- Used in HAGMA to see whether change in  $\text{HCO}_3^-$  is appropriate (i.e. whether there is a coexistent NAGMA or metabolic alkalosis)
- Delta ratio = Increase in Anion Gap/ Decrease in  $\text{HCO}_3^-$
- $(\text{change in anion gap}) / (\text{change in bicarbonate})$

$$(\text{AG}-12) / (24 - \text{Bicarbonate})$$

## Calculate Delta ratio

$$\frac{\text{(change in anion gap)}}{\text{(change in bicarbonate)}} \\ \text{(AG-12)} / \text{(24-Bicarbonate)}$$

- Value 1-2 -Uncomplicated HAGMA
- Value <1 –Combined HAGMA plus NAGMA
- Value >2- Combined HAGMA/Metab Alkalosis

# Interpretation of Blood Gases

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# A3

- Partially compensated high anion gap pure metabolic acidosis
- DKA

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# Management

IV Fluids

IV Insulin

Potassium replacement

Treat the precipitant

Role of IV  $\text{HCO}_3^-$  ?



Always treat the aetiology



Role of HCO<sub>3</sub>



Indications ?



Disadvantages?



Bolus/Infusion ?



Concentration -8.4% vs 1.26%



Role of RRT

# Treatment of Metabolic acidosis

- Always treat the aetiology
- Role of  $\text{HCO}_3^-$
- Indications ?
- Disadvantages?
- Bolus/Infusion ?
- Concentration -8.4% vs 1.26%
- Role of RRT

## Case 4: Known patient with CKD presents with SOB and diarrhoea.

pH	7.12
pCO <sub>2</sub>	31
HCO <sub>3</sub>	15
Na	131
Cl	108

1. Look at the pH
2. What is the primary disorder
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## Step 4 – In MA is the compensation pure

- **Metabolic acidosis** (Winters formula)

Expected  $p\text{CO}_2 = (1.5 \times \text{HCO}_3) + 8 \text{ (+/- 2)}$

- **Metabolic alkalosis**

Expected  $p\text{CO}_2 = (0.7 \times \text{HCO}_3) + 20 \text{ (+/- 5)}$

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# Step 5 –In MA calculate anion gap

- Metabolic acidosis

$$\text{Anion Gap} = (\text{Na}^+) - (\text{Cl}^-) - (\text{HCO}_3^-)$$

Normal 10-14 (12)

# Causes of Non-Anion Gap Metabolic Acido



## USED PART

**U:** Urinary diversion/ Ureteroentero

**S:** Small Bowel Fistula

**E:** Extra Chloride

**D:** Diarrhea

**P:** Pancreatic fistula

**A:** Addison, Acetazolamide

**R:** RTA

**T:** Tenofovir, Topiramate

# Interpretation of Blood Gases

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# A4

- Partially compensated non anion gap metabolic acidosis
- Secondary to diarrhoea

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# Causes of Low Anion Gap

- Increased 'unmeasured' cations
  - • Hypermagnesemia
  - • Lithium toxicity
  - • Paraproteinemia
    - Myeloma
    - Waldenstrom's macroglobulinaemia

Case 5 :Known CCF on large doses of frusemide was brought in with SOB, generalized weakness.

pH	7.51
pCO <sub>2</sub>	47
HCO <sub>3</sub>	40
Na	128
Cl	102
K	2.8

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder is compensation pure
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# A5

- Partially compensated pure metabolic alkalosis
- Diuretic induced hypokalaemia

## Case 6 :Known COPD and DM presents with fever, cough and SOB with reduced LOC.

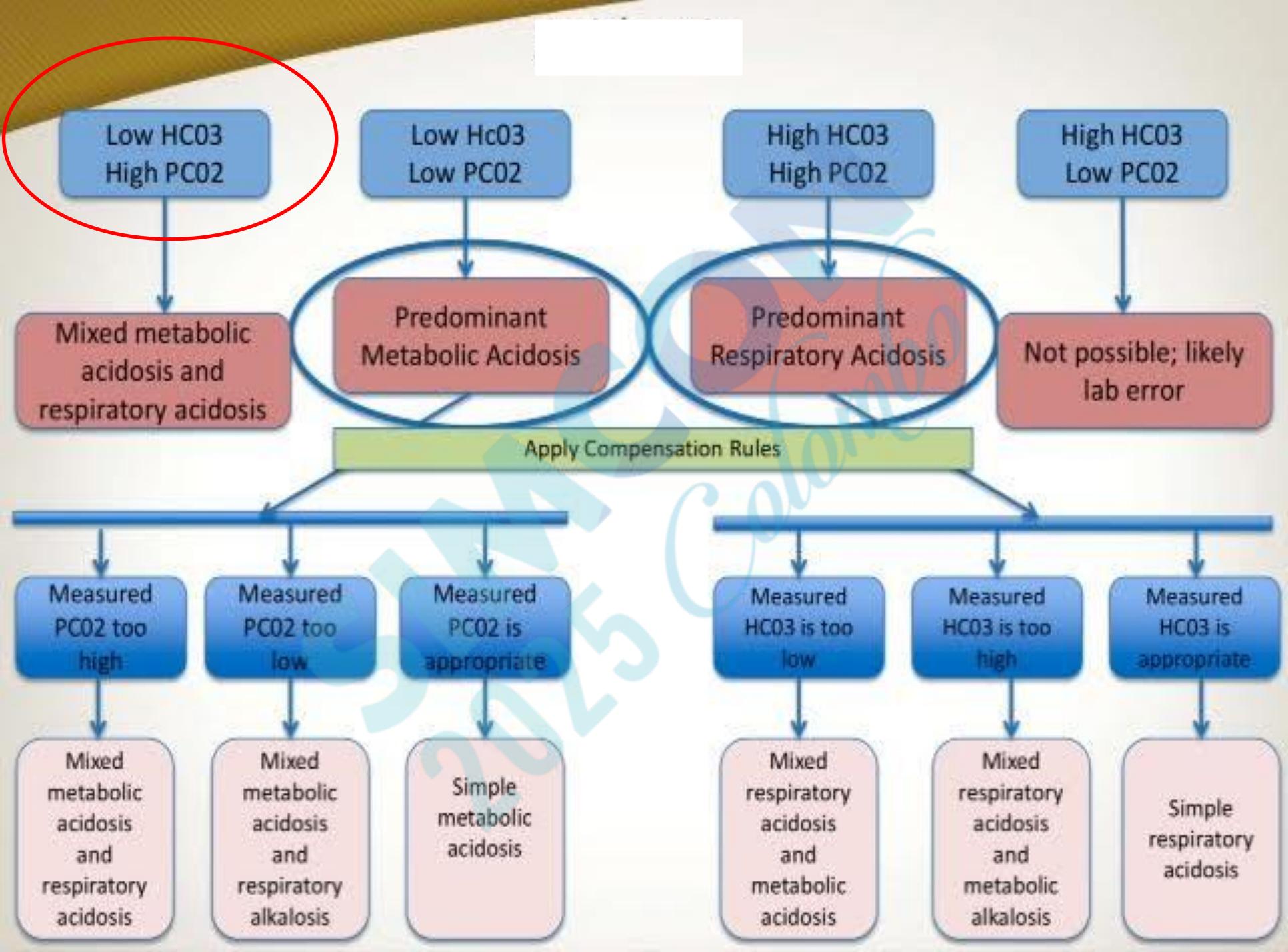
pH	6.84
pCO <sub>2</sub>	71
HCO <sub>3</sub>	18
Na	134
Cl	92
CBS	141
Lactate	08

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder compensation is pure or mixed
5. Is there a mixed metab/resp.disorder
6. In MA -Is there a HAGMA
7. If HAGMA calculate (Delta ratio)
8. Final Interpretation/Clinical diagnosis

## Case 6 :Known COPD and DM presents with fever, cough and SOB with reduced LOC.

pH	6.84
pCO <sub>2</sub>	71
HCO <sub>3</sub>	18
Na	134
Cl	92
CBS	141
Lactate	08

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder compensation is pure or mixed
5. Is there a mixed metab/resp.disorder
6. In MA -Is there a HAGMA
7. If HAGMA calculate (Delta ratio)
8. Final Interpretation/Clinical diagnosis



# Step 4-Is there a mixed Metab./Resp. disorder

- Mixed acidosis (metab.+ resp.)
- Metab. acidosis with resp. alkalosis
- Metabolic alkalosis with respiratory acidosis
- Mixed alkalosis
- Metabolic alkalosis with metabolic acidosis

**Table 6: Selected mixed and complex acid-base disturbances**

Disorder	Characteristics	Selected situations
Respiratory acidosis with metabolic acidosis	$\downarrow$ in pH $\downarrow$ in $\text{HCO}_3^-$ $\uparrow$ in $\text{PaCO}_2$	<ul style="list-style-type: none"><li>• Cardiac arrest</li><li>• Intoxications</li><li>• Multi-organ failure</li></ul>
Respiratory alkalosis with metabolic alkalosis	$\uparrow$ in pH $\uparrow$ in $\text{HCO}_3^-$ $\downarrow$ in $\text{PaCO}_2$	<ul style="list-style-type: none"><li>• Cirrhosis with diuretics</li><li>• Pregnancy with vomiting</li><li>• Over ventilation of COPD</li></ul>
Respiratory acidosis with metabolic alkalosis	pH in normal range $\uparrow$ in $\text{PaCO}_2$ , $\uparrow$ in $\text{HCO}_3^-$	<ul style="list-style-type: none"><li>• COPD with diuretics, vomiting, NG suction</li><li>• Severe hypokalemia</li></ul>
Respiratory alkalosis with metabolic acidosis	pH in normal range $\downarrow$ in $\text{PaCO}_2$ $\downarrow$ in $\text{HCO}_3^-$	<ul style="list-style-type: none"><li>• Sepsis</li><li>• Salicylate toxicity</li><li>• Renal failure with CHF or pneumonia</li><li>• Advanced liver disease</li></ul>
Metabolic acidosis with metabolic alkalosis	pH in normal range $\text{HCO}_3^-$ normal	<ul style="list-style-type: none"><li>• Uremia or ketoacidosis with vomiting, NG suction, diuretics, etc.</li></ul>

# A6

- Mixed metabolic and respiratory acidosis
- High anion gap metabolic acidosis (lactic acidosis)
- COPD exacerbation

## Management

- Fluid Resuscitate
- IV Antibiotics
- V Oxygen
- BiPAP
- Repeat ABG

Case 7. Known diabetic off treatment for 2 weeks presents with fever with chills and rigors and a red swollen warm right lower limb. He was tachypnic / hypotensive and receiving a fluid bolus

pH	7.37
pCO <sub>2</sub>	27
HCO <sub>3</sub>	18
Na	134
Cl	92
CBS	341
Lactate	06

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder compensation is pure or mixed
5. Is there a mixed metab/resp.disorder
6. In MA -Is there a HAGMA
7. If HAGMA calculate (Delta ratio)
8. Final Interpretation/Clinical diagnosis

# A7

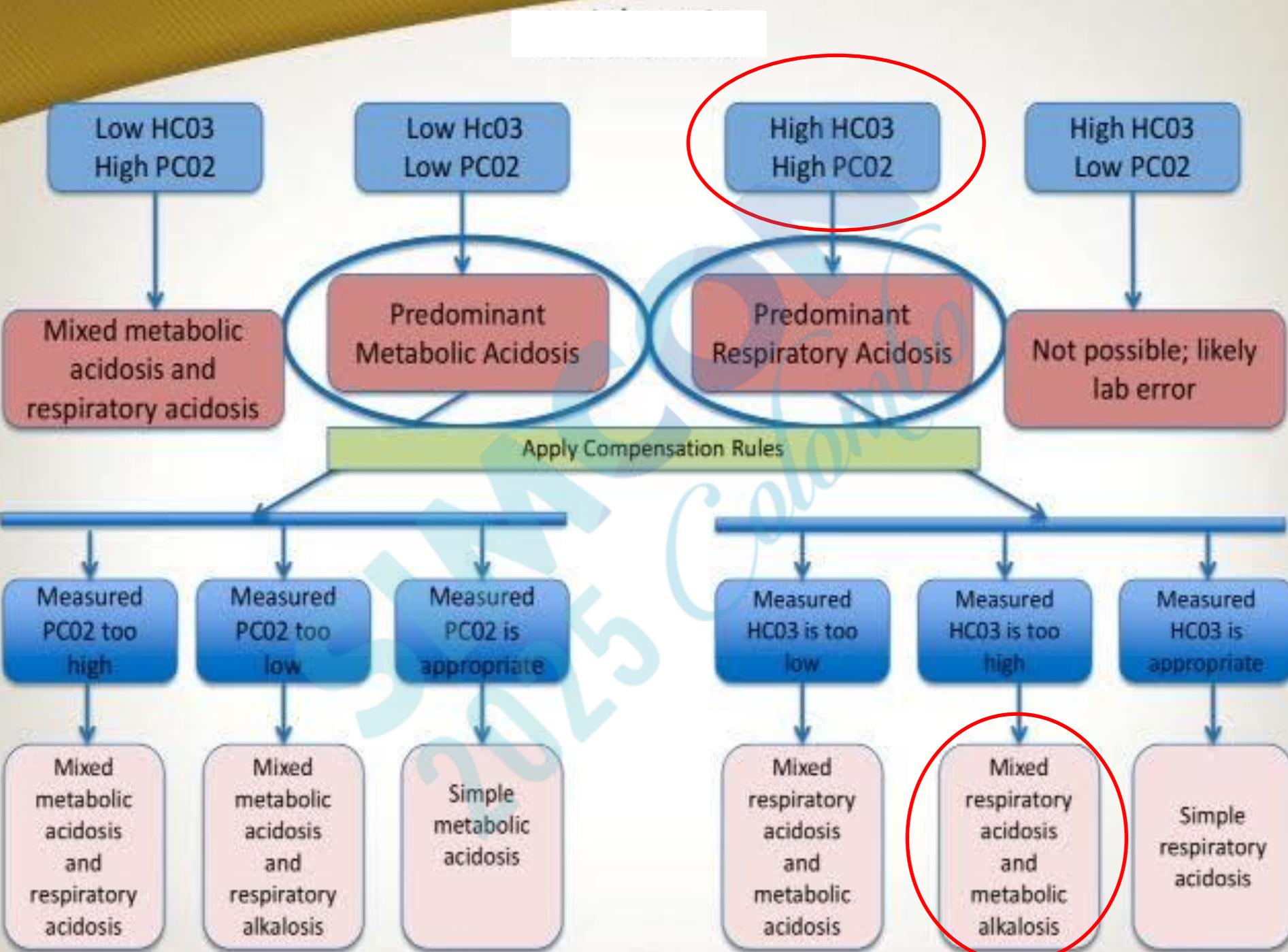
- Metabolic acidosis with respiratory alkalosis

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Case 8. Known COPD patient admitted with worsening SOB and abdominal fullness .He had a NG tube inserted which drained out 800 ml within 6hrs

pH	7.38
pCO <sub>2</sub>	61
HCO <sub>3</sub>	36
Na	134
Cl	92
K	2.8

1. Look at the pH
2. What is the primary disorder
3. In Respiratory disorder compensation acute or chronic
4. In Metabolic disorder compensation is pure or mixed
5. Is there a mixed metab/resp.disorder
6. In MA -Is there a HAGMA
7. If HAGMA calculate (Delta ratio)
8. Final Interpretation/Clinical diagnosis



Low HCO<sub>3</sub>  
High PCO<sub>2</sub>

Low HCO<sub>3</sub>  
Low PCO<sub>2</sub>

High HCO<sub>3</sub>  
High PCO<sub>2</sub>

High HCO<sub>3</sub>  
Low PCO<sub>2</sub>

Mixed metabolic acidosis and respiratory acidosis

Predominant Metabolic Acidosis

Predominant Respiratory Acidosis

Not possible; likely lab error

Apply Compensation Rules

Measured PCO<sub>2</sub> too high

Measured PCO<sub>2</sub> too low

Measured PCO<sub>2</sub> is appropriate

Measured HCO<sub>3</sub> is too low

Measured HCO<sub>3</sub> is too high

Measured HCO<sub>3</sub> is appropriate

Mixed metabolic acidosis and respiratory acidosis

Mixed metabolic acidosis and respiratory alkalosis

Simple metabolic acidosis

Mixed respiratory acidosis and metabolic acidosis

Mixed respiratory acidosis and metabolic alkalosis

Simple respiratory acidosis

# A8

- Respiratory acidosis and metabolic alkalosis

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## Management

- V O2
- BiPAP
- IV K replacement
- Repeat BG



# Summary

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Know the basics of blood gas analysis

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Minimize errors in the blood gas sampling process

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Develop a systematic approach of your own to analyse blood gases

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Apply compensation formulas that are familiar to you and Delta ratio to find out complex mixed gas disorders

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Mixed gas disorders are the commonest that you see in clinical practice

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Plan evidence-based management and repeat BGA