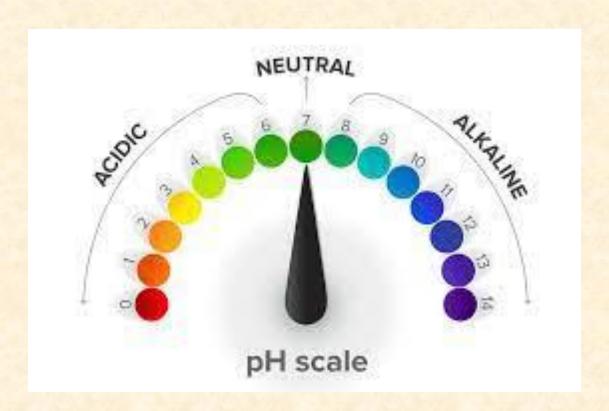
ACID BASE BALANCE BY DR/ HEBA KAREEM



Acid-Base balance

- ☐ A base releases hydroxyl ions (OH-) in aqueous solution.

This results in <u>increase</u> in pH of the solution

□ NaOH — Na⁺ + OH⁻

Amphoteric substances

Some substances, such as amino acids &proteins,

act acids as well as bases

Maintenance of blood pH

- □ The normal pH of the blood is maintained in the narrow range of <u>7.35-7.45</u> (slightly alkaline).
- The body has developed three lines of defense to regulate the body's acid-base balance.
- □ 1- Blood <u>buffers</u>
- ☐ 2-Respiratory mechanism
- □ 3-Renal mechanism
- Blood buffers:
- □ A buffer may be defined as a solution of a weak acid & its salt with a strongbase

Blood contains three buffer systems

- □ Bicarbonate buffer
- □ Phosphate buffer
- □ Protein buffer
- **■** Bicarbonate buffer system:
- Sodium bicarbonate & carbonic acid (NaHCO₃- H₂CO₃) is the most <u>predominant</u> buffer system of ECF.
- □ Carbonic acid dissociates into hydrogen and bicarbonate ions.

$$H_2CO_3 \longleftrightarrow H^+ + HCO_3$$

- ☐ The blood pH 7.4, the ratio of bicarbonate to carbonic acid is 20: 1
- □ The bicarbonate concentration is much higher (20times) than carbonic acid in the blood.
- ☐ This is referred to as <u>alkali reserve</u>.

Respiratory mechanism for pH regulation

- □ A <u>rapid</u> mechanism.
- □ This is achieved by regulating the concentration of carbonic acid (H₂CO₃) in the blood.

The large volumes of CO_2 produced by the cellular metabolic activity. All of this CO_2 is eliminated from the body in the expired air via the lungs

The respiratory centre is highly sensitive to changes in the pH of blood.

Decrease in blood pH causes hyperventilation to blow off co₂& reducing the H₂CO₃ concentration Respiratory control of blood pH is rapid but only a short term regulatory process, since hyperventilation cannot proceed for long.

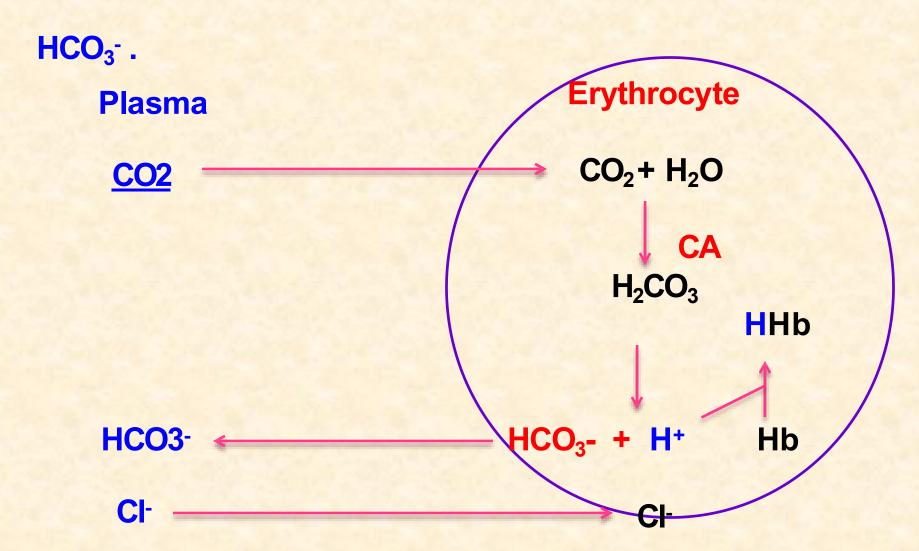
Hemoglobin as a buffer

- □ Hemoglobin binds to H⁺ ions & helps to transport CO₂ as HCO₃-with a minimum change in pH.
- In the lungs, hemoglobin combines with O₂, H⁺ ions are removed which combine with HCO₃⁻to form H₂CO₃
 & is dissociates to release CO₂to be exhaled.

Generation of HCO3 by RBC

- □ Due to lack of aerobic metabolic pathways, RBC produce very little CO₂.
- □ The plasma CO₂ diffuses into RBCalong the concentration gradient, it combines with water to form H₂CO₃ by Carbonic anhydrase.
- □ In RBC, H₂CO₃ dissociates to produce H⁺ & HCO₃⁻
- ☐ The H⁺ ions are buffered by Hemoglobin.
- □ As the concentration of HCO₃-increases in the RBC, it diffuses into plasma along with concentration gradient, in exchange for Cl-ions, to maintain electrical neutrality.

This is referred to as chloride shift, helps to generate



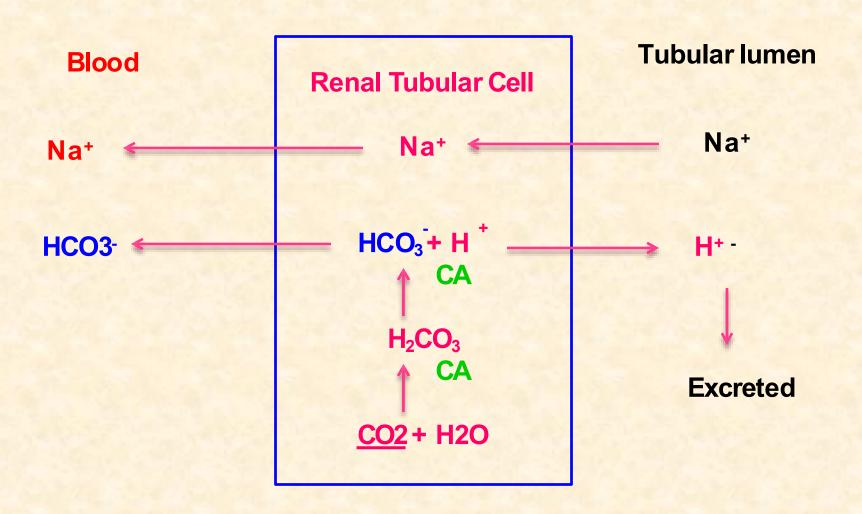
Renal mechanism for pH regulation

- □ The kidneys plays an important role in the regulation of pH through:
- □ 1-Excretion of H⁺ ions
- □ 2-Reabsorption of Bicarbonate
- □ 3-Excretion of titratable acid
- □ 4-Excretion of ammonium ions

Excretion of H+ ions

- □ Kidney is the <u>only route</u> through which the H⁺ can be eliminated from the body.
- □ H+ excretion occurs in the proximal convoluted tubules & is coupled with generation of HCO₃-.
- □ Carbonic anhydrase catalyses the production of carbonic acid (H₂CO₃) from CO₂&H₂O in renal tubular cells.
- □ H₂CO₃then dissociates to H⁺ & HCO₃-
- □ H⁺ ions are secreted into tubular lumen in exchange for Na⁺
- □ Na⁺ in association with HCO₃-is reabsorbed into blood

Excretion of H+ ions

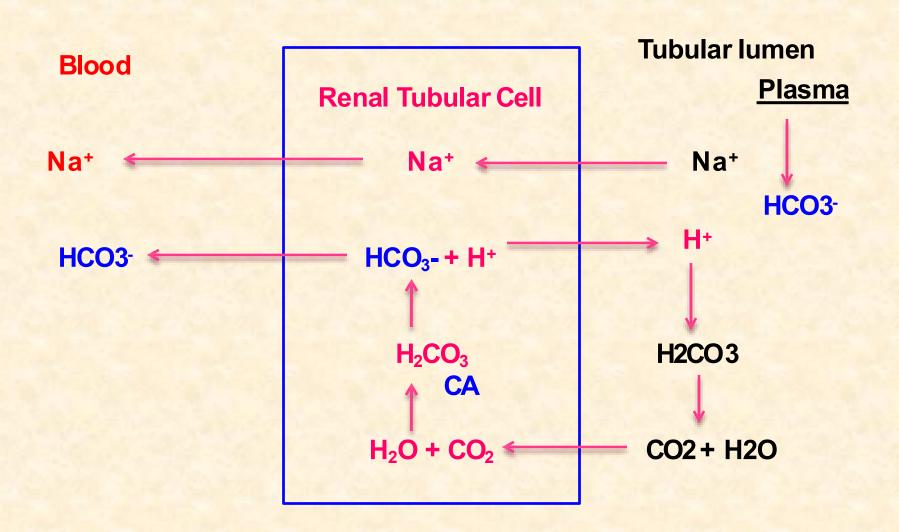


Reabsorption of Bicarbonate

- ☐ This mechanism is responsible to conserve blood HCO₃-, with simultaneous excretion of H⁺ ions.
- ☐ Bicarbonate freely diffuses from plasma into tubular lumen.
- □ HCO_3 -combines with H⁺, secreted by tubular cells, to form H_2CO_3 .
- \Box H₂CO₃ is then cleaved to form CO₂ and H₂O.
- □ As the CO₂ concentration builds up in the lumen, it diffuses into the tubular cells along the concentration gradient.

- □ In the tubular cell, CO₂again combines with H₂O to form H₂CO₃ which then dissociates into H⁺ &HCO₃-
- ☐ The H⁺ is secreted into the lumen in exchange for Na⁺.
- □ The HCO₃-is reabsorbed into plasma in association with Na⁺.
- □ Reabsorption of HCO₃- is a cyclic process without net excretion of H⁺ or generation of new HCO₃-

Reabsorption of bicarbonate



Excretion of titratable acid

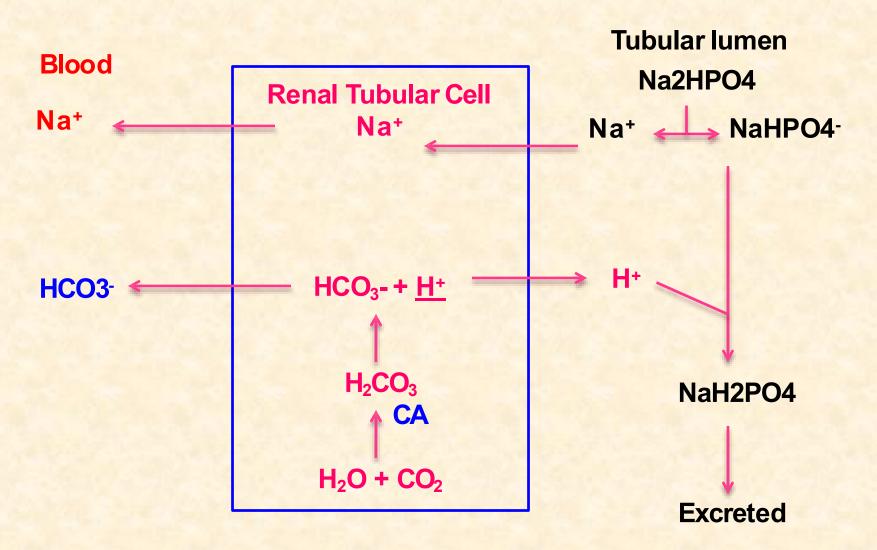
- □ Titratable acidity is a measure of acid excreted into urine by the kidney.
- □ Titratable acidity refers to the number of milliliters of N/10 NaOH required to titrate 1liter of urine to pH 7.4.
- ☐ Titratable acidity reflects the H⁺ ions excreted into urine.

- ☐ H⁺ ions are secreted into the tubular lumen in exchange for Na⁺ ion.
- □ This Na⁺ is obtained from the base, disodium hydrogen phosphate (Na₂HPO₄).
- □ This combines with H⁺ to produce the acid, sodium dihydrogen phosphate (NaH2PO4), in which form the major quantity of titratable acid in urine is present.
- □ Tubular fluid moves down the renal tubules,

more and more H+ ions are added, resulting in the acidification of

urine. Causes a fall in the pH of urine as low as 45.

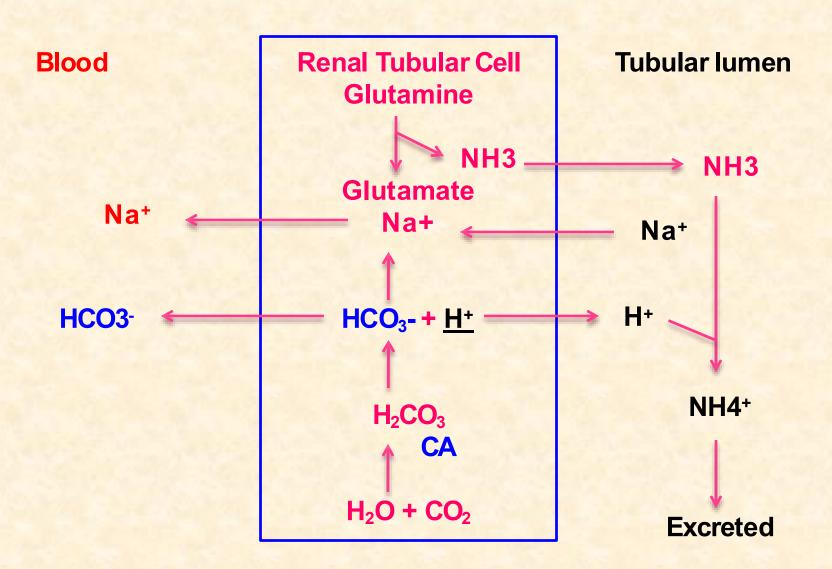
Excretion of titratable acid

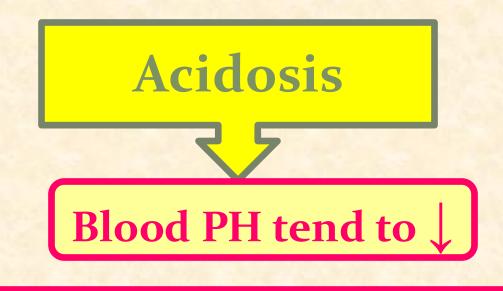


Excretion of ammonium ions

- □ The H+ ion combines with NH3 to form ammonium ion (NH4+).
- ☐ The renal tubular cells deaminate glutamine to glutamate and NH3 by the action of enzyme glutaminase.
- ☐ The liberated NH3 diffuses into the tubular lumen where it combines with H+ to form NH4+.
- □ Ammonium ions cannot diffuse back into tubular cells and excreted into urine.

Excretion of ammonium ions





Results from formation of **excessive acids**More than the capacity of the body to eliminate them



Respiratory acidosis

Metabolic acidosis

Respiratory acidosis

↑CO2 (CO2 RETENTION) due to

- Bronchial asthma
- Chronic bronchitis
- Emphysema
- Pneumonia
- Respiratory centre inhibition
- Asphexia
- ↑CO2 → ↑ blood H2CO3

Respiratory acidosis

```
↑CO2 ↑ blood H2CO3 HCO3 not changed
  \rightarrow \downarrow HCO3^{-}/H2CO3 (N=20:1)
   \rightarrow \downarrowblood PH
  (Uncompensated respiratory acidosis [acidemia])
How to compensate?
Kidney reabsorbs more HCO3<sup>-</sup>
Till normal HCO3<sup>-</sup>/H2CO3 (20:1)
→ PH reach 7.4
```

Metabolic acidosis

```
\uparrow acids or \downarrow bases (HCO3<sup>-</sup>) in blood
               → blood HCO3<sup>-</sup>
blood H2CO3 not changed
          \rightarrow \downarrow \text{HCO3}^{-}/\text{H2CO3} (N=20:1)
          \rightarrow \downarrowblood PH
  (Uncompensated metabolic acidosis [acidemia])
How to compensate?
\downarrow PH\rightarrow ++ chemoreceptors in respiratory
centre→ hyperventilation→ loss of
CO2 \rightarrow \downarrow H2CO3
Till normal HCO3<sup>-</sup>/H2CO3 (20:1)
→ PH reach 7.4 (Compensated metabolic acidosis)
```

Causes of Metabolic acidosis

1- **†blood** acids

↑production

↓excretion

☐ failure of
excretion by the
kidney in chronic
renal failure

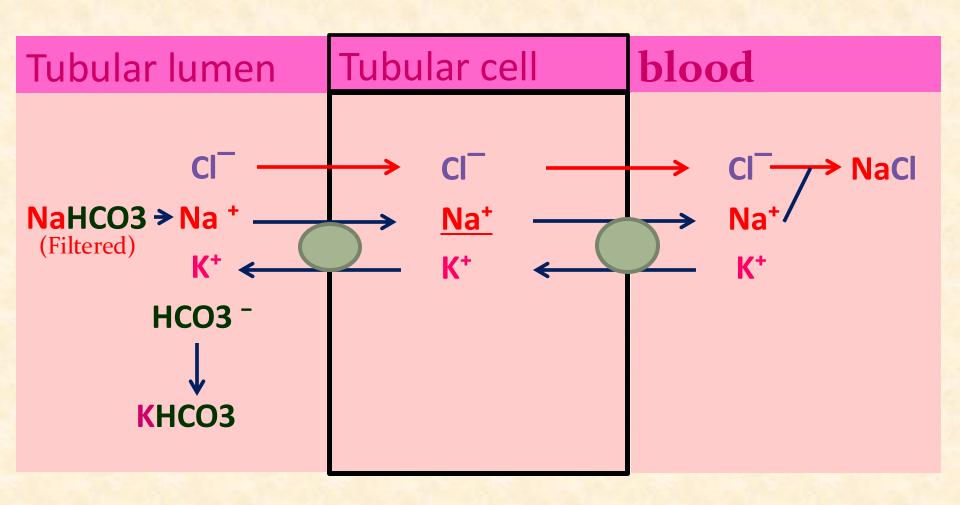
Causes of Metabolic acidosis

1- \(\gamma\) base loss

- Diarrhea: Intestinal juices are alkaline being rich in Na⁺ & K⁺ bicarbonate
- Vomiting: due to low intestinal obstruction
- Hyperkalemia:
 - * Trenal tubular reabsorption of Na⁺ in exchange with K⁺
 - → stop of Na⁺/ H⁺ exchange
- * Na⁺ reabsorption will be in the form of NaCl not NaHCO3 > HCO3 will be excreted in the form of KHCO3 in urine.
 - **HCO3** loss in urine → metabolic acidosis (Alkaline urine)
 - ↑ Cl in blood→ hyperchloremic acidosis (Acidic blood)
- The alkaline urine& acidic blood is called paradoxical acidosis

Causes of Metabolic acidosis

1- **†blood** acids





Blood PH tend to 1



More than the capacity of the body to neutralize & eliminate them



Respiratory alkalosis

Metabolic alkalosis

Respiratory alkalosis

个 CO2 loss due to fever encephalitis high altitude late stages of salicylate poisoning hystrical hyperventilation \downarrow CO2 \rightarrow \downarrow blood H2CO3

Respiratory alkalosis

```
↓CO2 → blood H2CO3

↓HCO3 not changed
  \rightarrow \uparrow HCO3<sup>-</sup>/H2CO3 (N=20:1)
  \rightarrow \uparrow blood PH
  (Uncompensated respiratory alkalosis [alkalemia])
How to compensate?
-- of renal tubular reabsorption of HCO3
Kidney excretes more HCO3<sup>-</sup>
Till normal HCO3<sup>-</sup>/H2CO3 (20:1)
\rightarrow PH reach 7.4
(Compensated respiratory alkalosis)
Urine will be alkaline because of ↑ secretion of K<sup>+</sup> & HCO3
in urine
```

Metabolic alkalosis

- ↑ bases or ↓ acids in blood
 - → ↑ blood HCO3⁻
 blood H2CO3 not changed
 - \rightarrow \uparrow HCO3⁻/H2CO3 (N=20:1)
 - → **↑blood PH**

(Uncompensated metabolic alkalosis [acidemia])

How to compensate?

- ↑ PH \rightarrow - chemoreceptors in respiratory centre \rightarrow hypoventilation \rightarrow CO2 retention \rightarrow ↑H2CO3
- Till normal HCO3⁻/H2CO3 (20:1)
- →PH reach 7.4 (Compensated metabolic alkalosis)

Causes of Metabolic alkalosis

1- ↑absorption of bases

- Intake of high vegetable and fruit diet: They contain Bicarbonate salts and citrate salts. Citrate salts will be transformed into bicarbonate salts by krebs cycle
- □ Intake of drugs containing bicarbonate & citrate salts (drugs used for treatment of hyperacidity & peptic ulcer)

Causes of Metabolic alkalosis

2- \(\psi\) loss of acids

- Prolonged <u>suction</u> of gastric juice
- Vomiting due to high intestinal obstruction
- Hypokalemia:
 - *

 √ renal tubular reabsorption of Na⁺ in exchange with K⁺
 - → instead there is Na⁺/ H⁺ exchange
- * Na⁺ reabsorption will be in the form of NaHCO3 not NaCl \rightarrow # Cl⁻ loss in urine in the form of NH4Cl \rightarrow hypochloremia and acidic urine
- ↑ NaHCO3 in blood→ alkalosis(alkaline blood)
 The acidic urine& alkaline blood is called paradoxical alkalosis
- □ <u>Cushing syndrome</u>: →Na& water retention & K excretion → hypokalemia