

# ACID-BASE BALANCE

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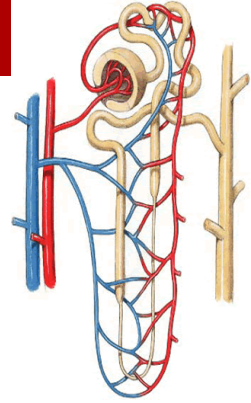
# Objectives

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- Define: acid and base.
- Explain what is meant by strong and weak acids and bases.
- List and identify the names/formulas for the common strong acids and strong bases.
- To explain the role of Henderson-Hasselbalch equation in acid-base regulation.
- To define buffer system and discuss the role of blood buffers and to explain their relevant roles in the body.
- To describe the role of kidneys in the regulation of acid-base balance.
- To describe the role of lungs in the regulation of acid-base balance.

# Contents

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- What are acids and bases?
- What is meant by a weak and a strong acid or base?
- 
- What is the normal pH of body fluids?
- Why is it important to keep body fluid pH within certain limits?
- What are the body's defense mechanisms against changes in blood pH: body buffers, the lungs and the kidney.
- Understand the role of the kidney in regulating pH of body fluids.

# Acid-Base Balance

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- Acid-base balance is concerned with the precise regulation of free (unbound) hydrogen ion ( $H^+$ ) concentration in body fluids.
- Normally,  $[H^+] = 0.00004 \text{ mEq/L}$  ( $40 \text{ nEq/L}$ ).
- ***Why is it important to control  $[H^+]$ ?***

# Why Should $[H^+]$ be Tightly Controlled?

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- Slight deviations in  $[H^+]$  have profound effects on enzyme and protein activity and thus the body's metabolic activity in general.
- Changes in  $[H^+]$  affects  $K^+$  levels in the body.

# Why is the Body's $[H^+]$ Constantly Changing?

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*A number of processes can alter  $[H^+]$  concentration in the body, such as;*

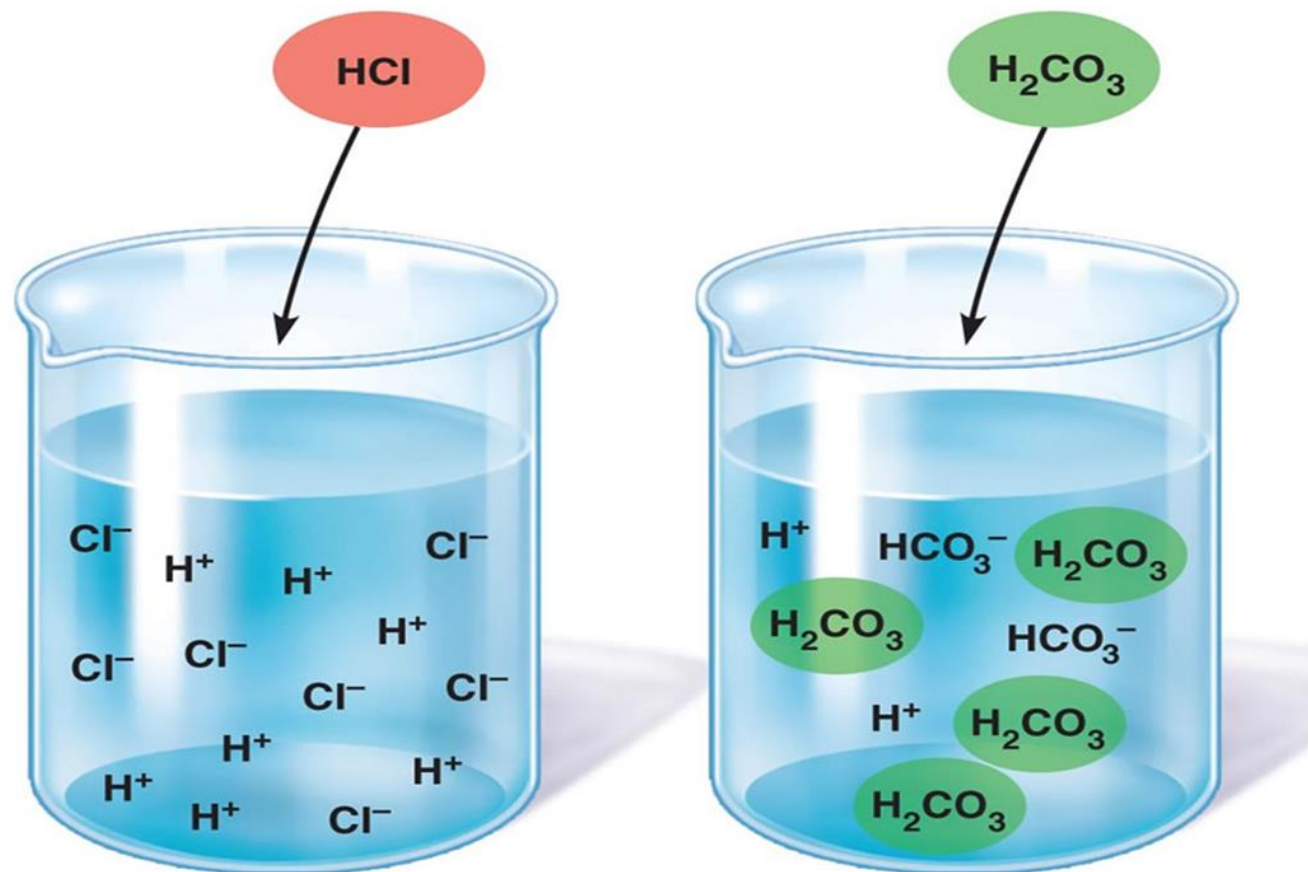
1. Metabolism of ingested food.
2. GI secretions.
3. Generation of acids & bases from amino acid/protein metabolism.
4. Changes in  $CO_2$  production.

# Acid-Base Fundamentals

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- **An Acid** = a molecule that can release  $H^+$  in a solution.
  - $H_2CO_3$  (carbonic acid)
  - $HCl$  (hydrochloric acid)
- **A base** = a molecule that accepts  $H^+$  in a solution.
  - Bicarbonate ions ( $HCO_3^-$ ).
  - Hydrogen phosphate ( $HPO_4^{-2}$ )
- ***What is the difference between carbonic & hydrochloric acid?***

# Strong vs Weak Acids & Bases



**Strong acids** dissociate rapidly and release large amounts of  $\text{H}^+$  in solution

**Weak acids** dissociate incompletely and less strongly releasing small amounts of  $\text{H}^+$  in solution



# Weak Acids

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*Acid*  $\leftrightarrow$  *Conjugate base* +  $H^{+}$

The extent to which a given acid dissociates in solution is constant. And is known as the **dissociation constant (K)**.

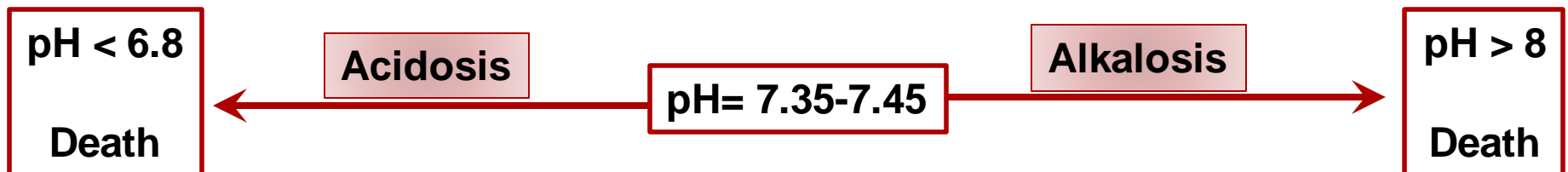
$$K = \frac{[H^{+}][A^{-}]}{[AH]}$$



# [H<sup>+</sup>] & the pH

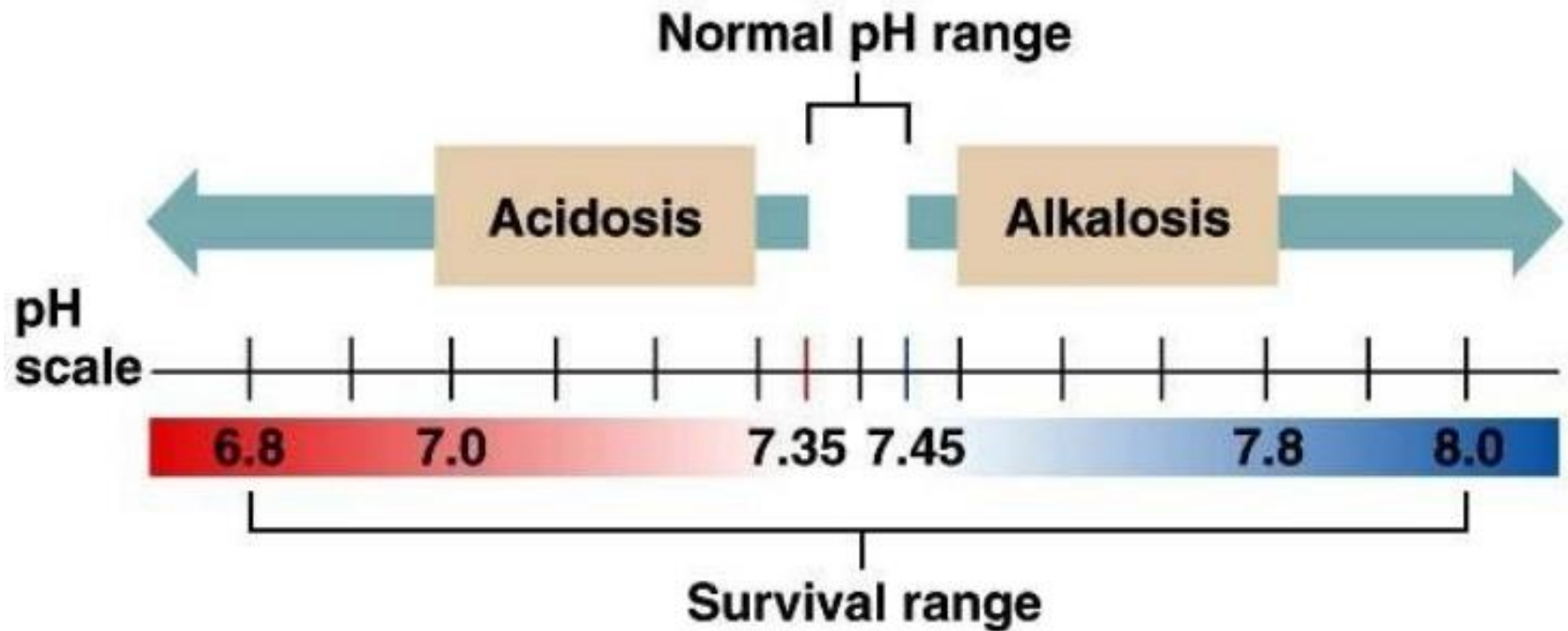
- H<sup>+</sup> ion concentrations are expressed as pH.
- $\text{pH} = -\text{Log} [\text{H}^+]$ 
  - If the [H<sup>+</sup>] increase → pH will decrease (more acidic)
  - If the [H<sup>+</sup>] decrease → pH will increase (more alkaline)
- ***What is the normal pH of the ECF?***

Normally pH= 7.35-7.45



# The Body and pH

## pH of arterial blood



# pH and H<sup>+</sup> Concentration of Body Fluids

	[H <sup>+</sup> ] (mEq/L)	pH
<b>Extracellular fluid</b>		
<b>Arterial blood</b>	$4.0 \times 10^{-5}$	<b>7.4</b>
<b>Venous blood</b>	$4.5 \times 10^{-5}$	<b>7.35</b>
<b>IF</b>	$4.5 \times 10^{-5}$	<b>7.35</b>
<b>Intracellular fluid</b>	$1 \times 10^{-3}$ to $4 \times 10^{-5}$	<b>6-7.4</b>
<b>Urine</b>	$3 \times 10^{-2}$ to $1 \times 10^{-5}$	<b>4.5-8</b>
<b>Gastric HCl</b>	<b>160</b>	<b>0.8</b>

# Acid Production by the Body

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- The body produces large amounts of acids on daily basis as by products of metabolism.
  - Metabolism of dietary proteins.
  - Anaerobic metabolism of carbs and fat.
- Acids in the body are of two kinds:
  1. Volatile ( $\text{CO}_2$ )
  2. Non-volatile “fixed” (sulfuric acid, lactic acid) (daily acid load  $\approx$  50-100 mEq/day) (0.8 mEq/kg/d).

# The Body's Defense Against Changes in $[H^+]$

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## Three main systems:

### 1. *Body fluid buffers.*

Works within seconds.

### 2. *Lungs*

Works within minutes.

### 3. *Kidneys*

Works within hours-days.

The most powerful of the three.

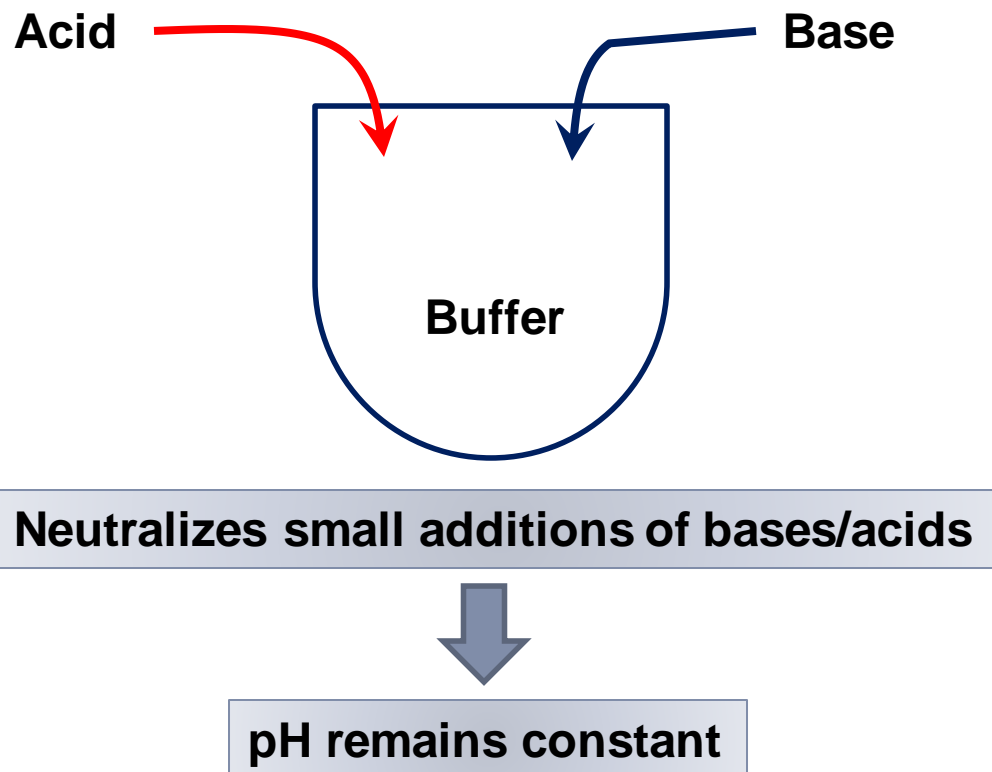
# **BODY FLUID BUFFERS**

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# What is a Buffer?

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**A buffer** = a solution that resists changes in pH upon addition of small amount of acids or bases.





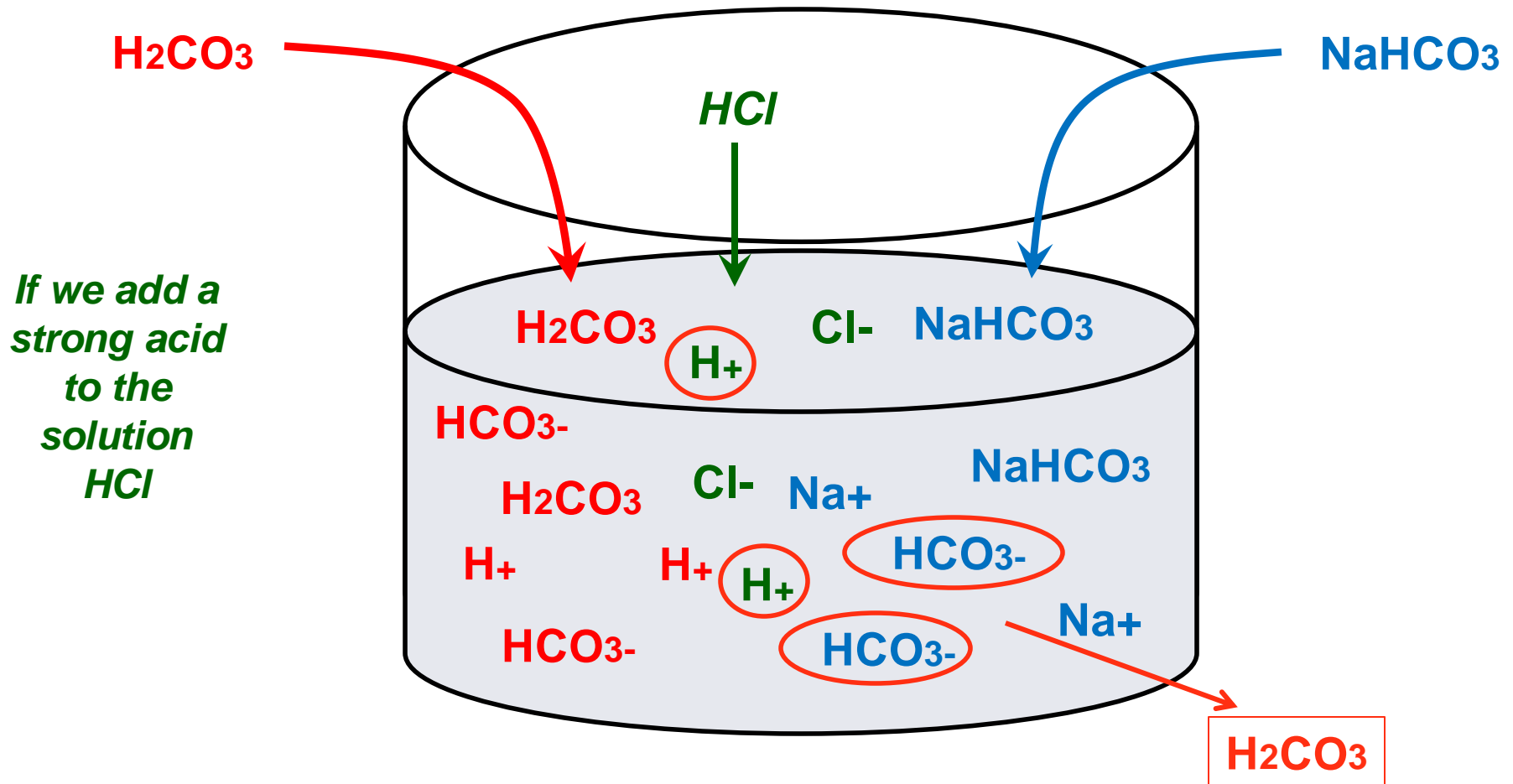
# How do Buffers work?

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- A buffer is a mixture of a weak acid and a weak base that are in equilibrium.
- To be more accurate, its either made of:
  - A weak acid and its conjugated base ( $\text{H}_2\text{CO}_3$  &  $\text{NaHCO}_3$ ).
  - A weak base and its conjugated acid ( $\text{NH}_3$  &  $\text{NH}_4^+$ ).
- ***How does a buffer do its job?***

# How does a buffer do its job?

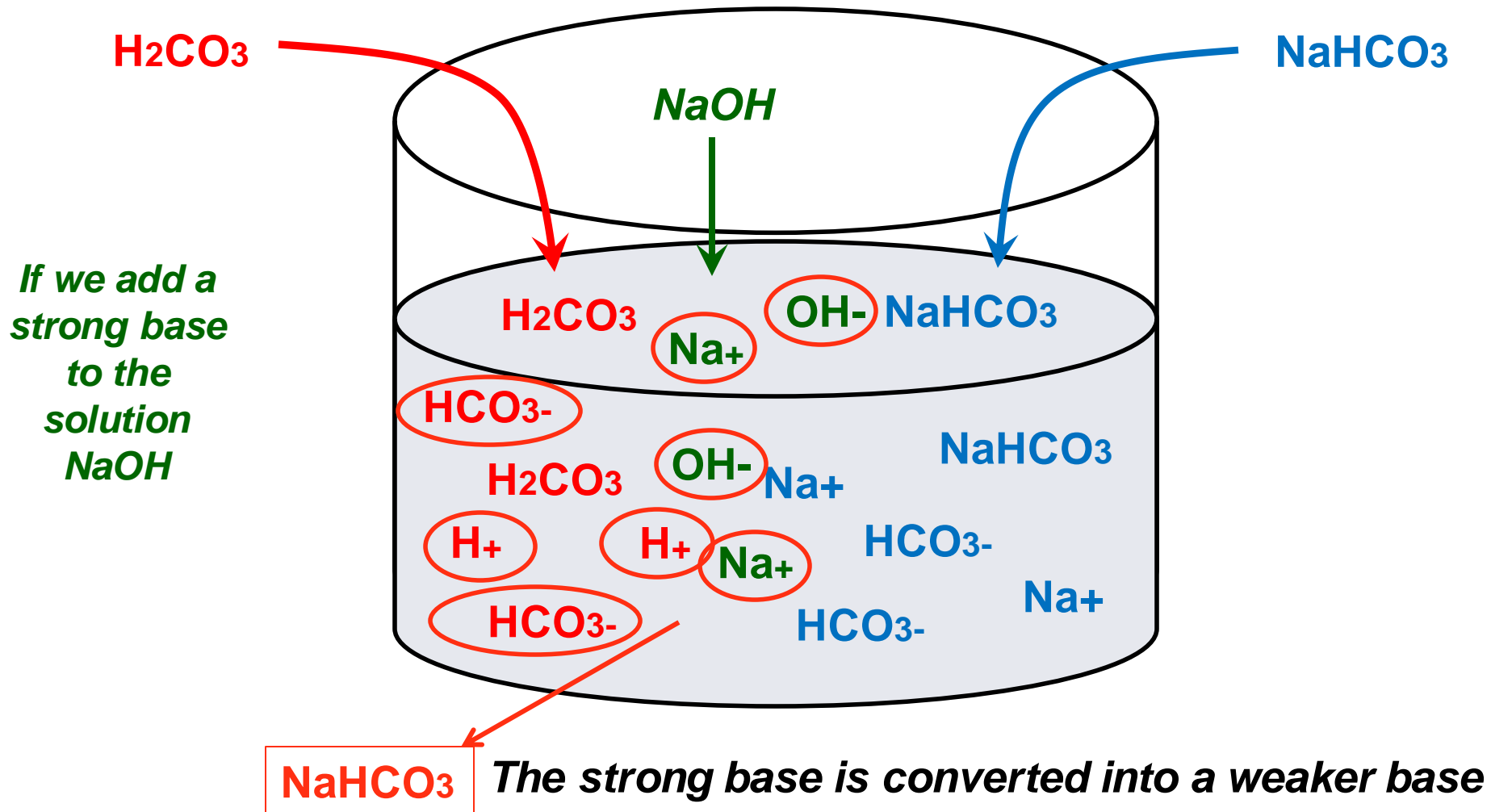
## Bicarbonate buffer solution



**The strong acid is converted into a weaker acid**

# How does a buffer do its job?

## Bicarbonate buffer solution



# Chemical Buffer Systems in the Body

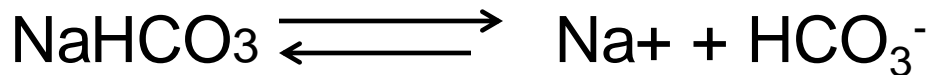
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- ***There are 3 chemical buffers in the body;***
  1. The Bicarbonate buffer system.
  2. The phosphate buffer system.
  3. Proteins.
- They are the 1<sup>st</sup> line of defence against changes in pH i.e.  $[H^+]$ , act within seconds.
- Some are more powerful extracellularly and others are more powerful intracellularly.

# The Bicarbonate Buffer System

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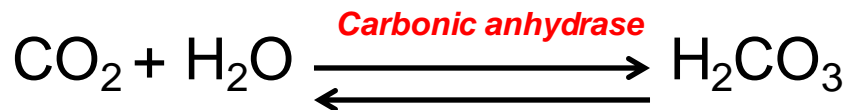
- Composed of:
  - A weak acid ( $\text{H}_2\text{CO}_3$ ).
  - Its conjugated base ( $\text{NaHCO}_3$ ).



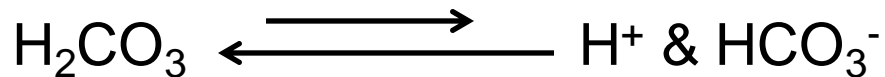
# The Bicarbonate Buffer System

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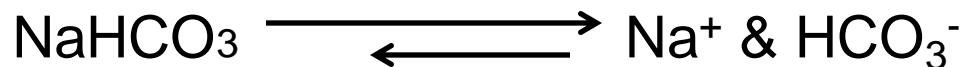
1.  $H_2CO_3$  forms in the body by the reaction of  $CO_2$  &  $H_2O$



2.  $H_2CO_3$  ionizes weakly to form small amounts of  $H^+$  &  $HCO_3^-$

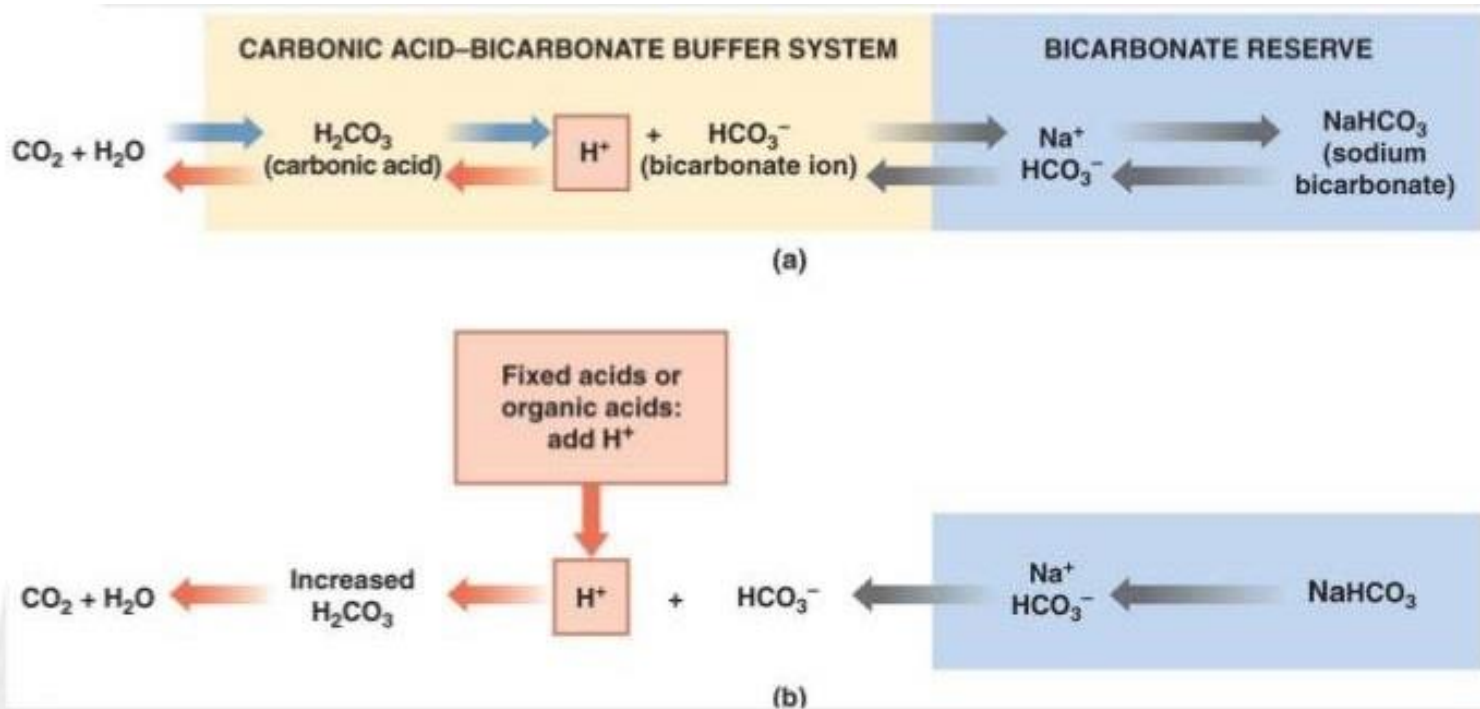


3. The second component is  $NaHCO_3$  which dissociates to form  $Na^+$  &  $HCO_3^-$



# The Bicarbonate Buffer System

*Putting it all together;*



The Carbonic Acid-Bicarbonate Buffer System

This is the main ECF buffer system

***What happens if you add a base or an acid to the system?***

# The Henderson-Hasselbalch Equation

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## *What is the HHE?*

- It is an equation that enables the calculation of pH of a solution.

## *What is it?*

$$pH = pK + \log \frac{HCO_3^-}{0.03 \times P_{CO_2}}$$

K = dissociation constant, pK = 6.1

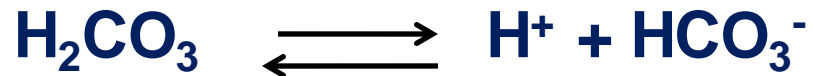
0.03 = solubility of CO<sub>2</sub>



# The Henderson-Hasselbalch Equation

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*How was it derived?*



1.  $\text{H}_2\text{CO}_3$  and its dissociated ions are always in equilibrium → the products of the reaction on one side of the equation are proportional to the product on the other side.

$$[\text{H}_2\text{CO}_3] \propto [\text{H}^+] \times [\text{HCO}_3^-]$$

2. Since  $\text{H}_2\text{CO}_3$  is a weak acid, it will not dissociate completely and the concentration of its products will depend on its dissociation constant (K)

$$K \times [\text{H}_2\text{CO}_3] = [\text{H}^+] \times [\text{HCO}_3^-]$$

# The Henderson-Hasselbalch Equation

3. Based on the previous equation,  $[H^+]$  can be expressed as follows;

$$[H^+] = K \times \frac{[H_2CO_3]}{[HCO_3^-]}$$

4. Because  $H_2CO_3$  can rapidly dissociate into  $CO_2$  and  $H_2O$ . And since  $CO_2$  is much easier to measure it can replace  $H_2CO_3$  in the equation;

$$[H^+] = K \times \frac{[CO_2]}{[HCO_3^-]}$$

*This is Henderson's equation  
(1908)*

*It means that;*

$\uparrow [CO_2] \rightarrow \uparrow [H^+]$

$\uparrow [HCO_3^-] \rightarrow \downarrow [H^+]$

# The Henderson-Hasselbalch Equation

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5. In 1909, Sorensen created the pH scale to express  $[H^+]$

$$pH = -\log[H^+]$$

6. In 1916, Hasselbalch decided to merge Henderson's equation with Sorensen's pH scale creating what we now know as the *"Henderson-Hasselbalch equation"*.

$$[H^+] = K \times \frac{[CO_2]}{[HCO_3^-]} \implies -\log[H^+] = -\log\left(K \times \frac{[CO_2]}{[HCO_3^-]}\right)$$

$$-\log[H^+] = -\log K - \log\left(\frac{[CO_2]}{[HCO_3^-]}\right)$$

# The Henderson-Hasselbalch Equation

$$-\log[H^+] = -\log K - \log\left(\frac{[CO_2]}{[HCO_3^-]}\right)$$



$$pH = pK + \log\frac{[HCO_3]}{[CO_2]}$$



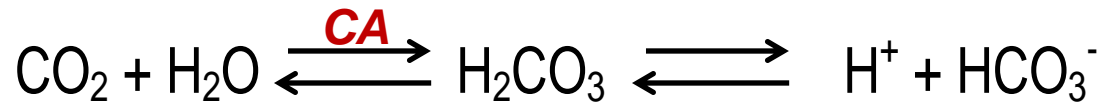
*This is Henderson-Hasselbach equation  
(1908)*

7. Since it is much easier to measure  $PCO_2$  rather than dissolved  $[CO_2]$  and because dissolved  $CO_2$  is proportional to  $PCO_2$  multiplied by the solubility of  $CO_2$  (0.03 mmol/mmHg)  $\rightarrow [CO_2]$  was replaced by  $PCO_2 \times 0.03$

$$pH = pK + \log\frac{[HCO_3]}{0.03 \times PCO_2}$$



# The Bicarbonate Buffer System



$$\text{pH} = \text{pK} + \log \frac{[\text{HCO}_3^-]}{0.03 \times \text{PCO}_2}$$

## • *What do we understand from this equation?*

1.  $\text{pH} \propto \frac{\text{HCO}_3^-}{\text{PCO}_2}$

**Regulated by kidneys** (referring to  $\text{HCO}_3^-$ )

**Regulated by lungs** (referring to  $\text{PCO}_2$ )

*Each element of the buffer system is regulated*

- $\uparrow\uparrow \text{HCO}_3^-$  will  $\uparrow\uparrow \text{pH}$
- $\uparrow\uparrow \text{PCO}_2$  will  $\downarrow\downarrow \text{pH}$

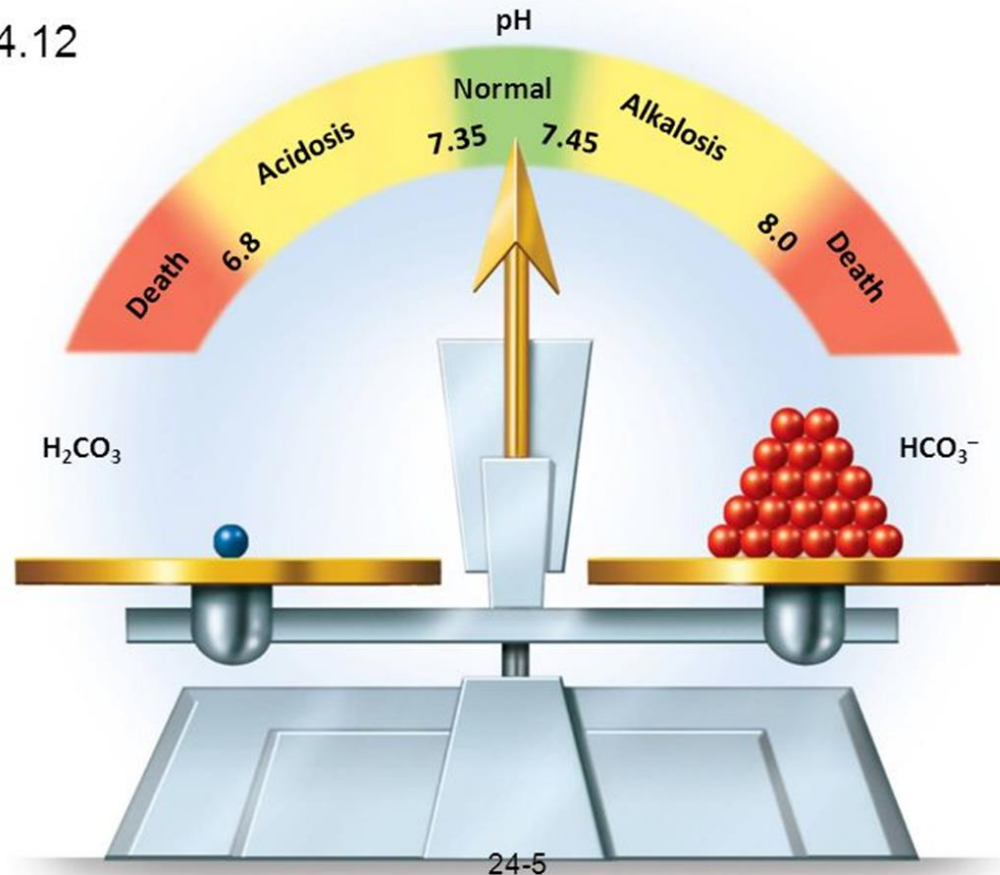


Ratio of  $\frac{HCO_3^-}{PCO_2}$  is  $\approx 20:1$

## Acid-Base Balance

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Figure 24.12



# Other Buffering Systems

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## *The phosphate buffer:*

- Plays a major role in buffering intracellular & renal tubular fluid.
- Composed of;
  - $\text{H}_2\text{PO}_4^-$  (dihydrogen phosphate)
  - $\text{HPO}_4^{2-}$  (Hydrogen phosphate)

## *Proteins:*

- Contributes to buffering inside cells.
- E.g. Hb.



# Summary of Body's Buffering Systems

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- Buffer systems do not work independently in body fluids but actually work together.
- A change in the balance in one buffer system, changes the balance of the other systems.
- Buffers do not reverse the pH change, they only limit it.
- Buffers do not correct changes in  $[H^+]$  or  $[HCO_3^-]$ , they only limit the effect of change on body pH until their concentration is properly adjusted by either the lungs or the kidney.

# **RESPIRATORY REGULATION OF ACID-BASE BALANCE**

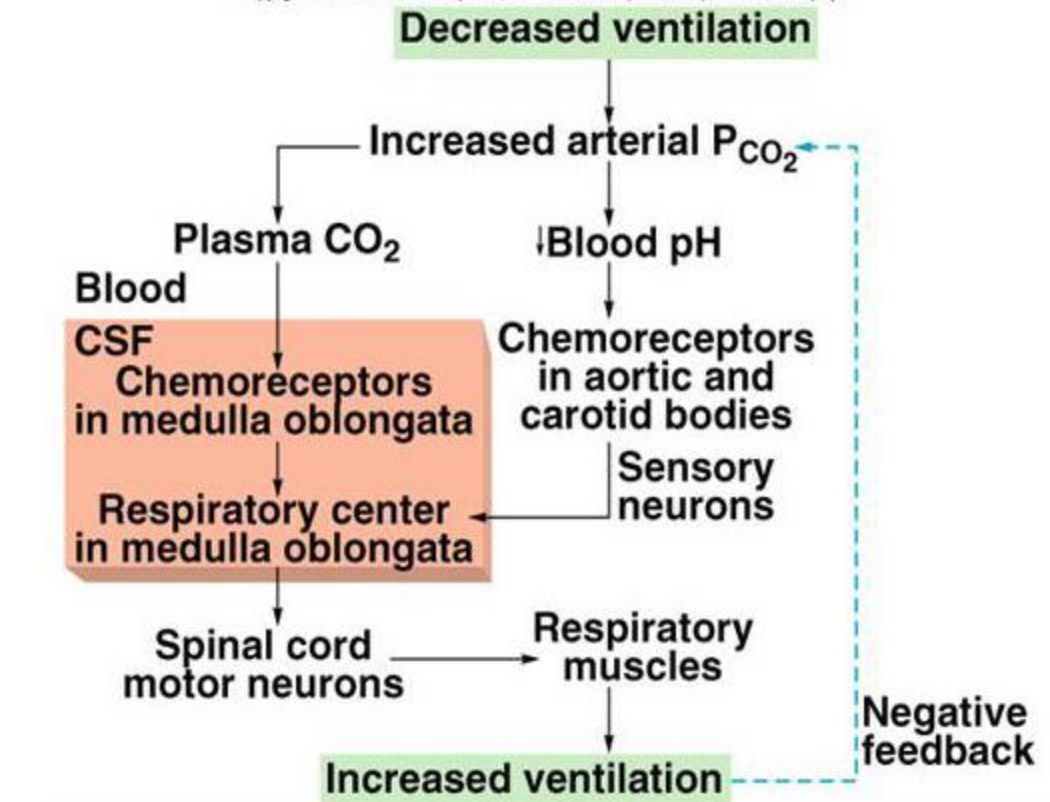
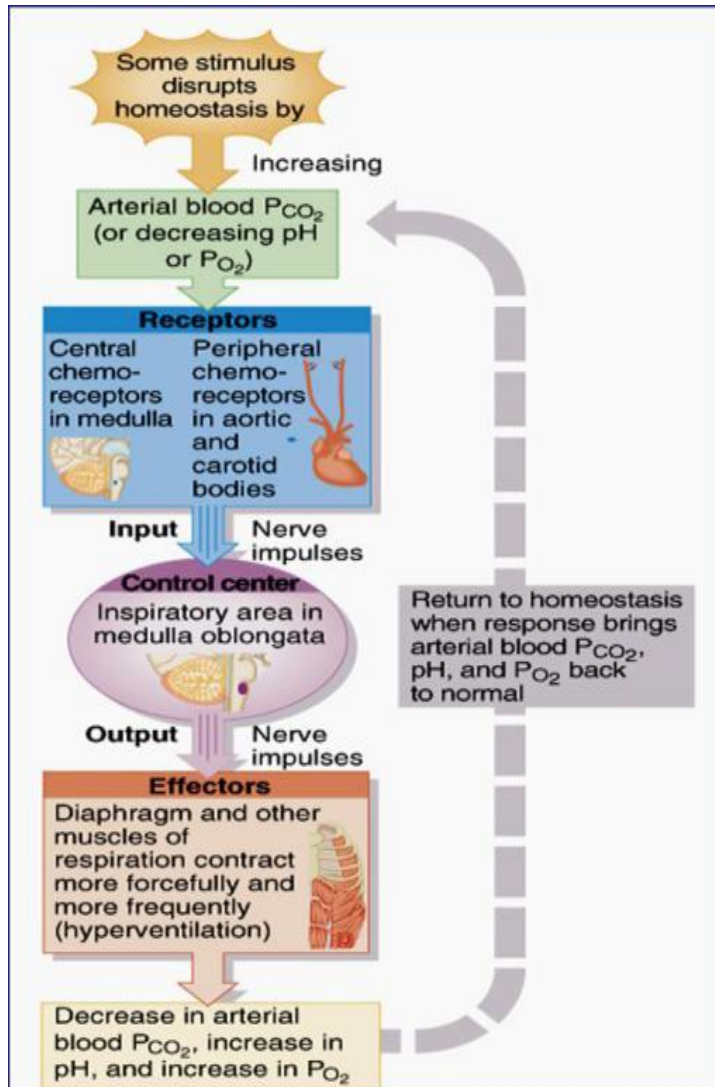
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# Respiratory Regulation of A/B

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- 2<sup>nd</sup> line of defence against acid-base disturbances in the body.
- **HOW?**
  - By modulating CO<sub>2</sub> excretion.
- ↑↑ [H<sup>+</sup>] → ↑↑ ventilation (RR) → ↓↓ PCO<sub>2</sub>
- ↓↓ [H<sup>+</sup>] → ↓↓ ventilation (RR) → accumulation of CO<sub>2</sub> → ↑↑ PCO<sub>2</sub>.
- Normally, PCO<sub>2</sub> = 40 mmHg (35-45 mmHg)

# Respiratory Regulation of CO<sub>2</sub>



# **RENAL REGULATION OF ACID-BASE BALANCE**

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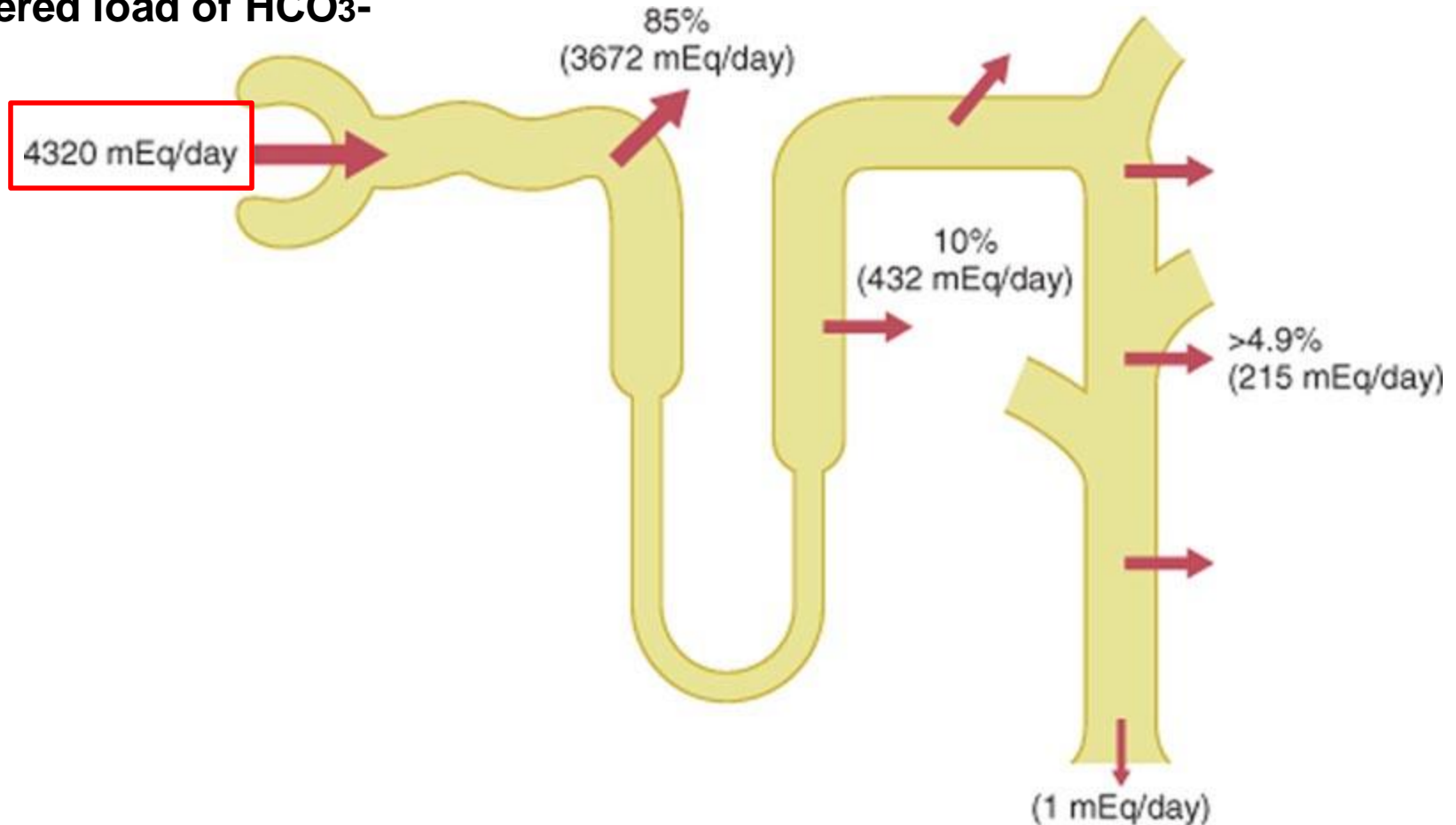
# Renal Regulation of A/B Balance

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- 3<sup>rd</sup> line of defence against acid-base disturbances and the most powerful.
- It regulates by excreting either an acidic or basic urine.
- **HOW?**
  1. Secreting  $H^+$
  2. Reabsorbing  $HCO_3^-$
  3. Generating “new” bicarbonate ions.

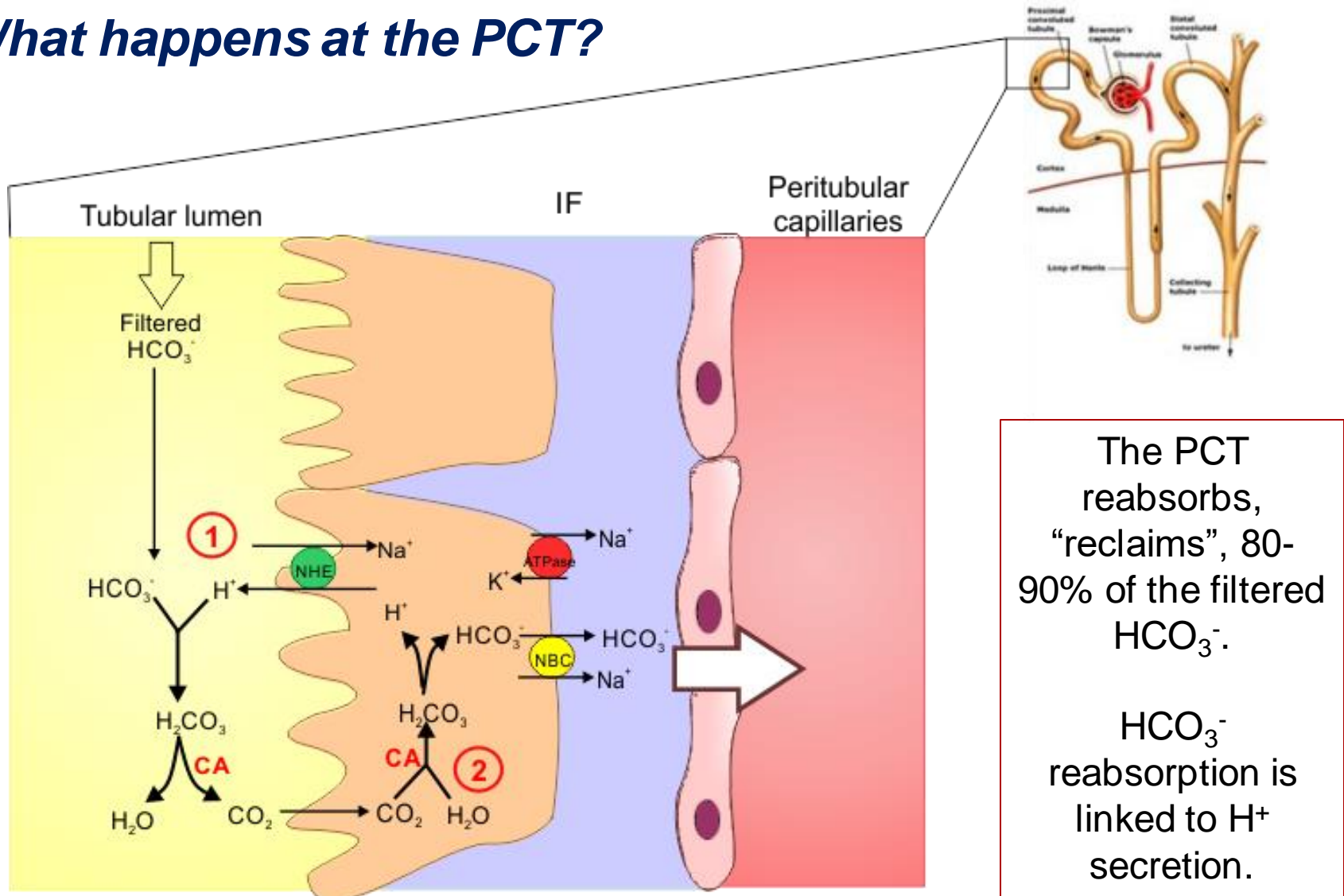
# Overview $\text{HCO}_3^-$ Reabsorption by the Renal Tubules

Filtered load of  $\text{HCO}_3^-$



# How is $\text{HCO}_3^-$ Reabsorbed by the tubules?

*What happens at the PCT?*





# How Does the Proximal Tubule Reabsorb $\text{HCO}_3^-$ ?

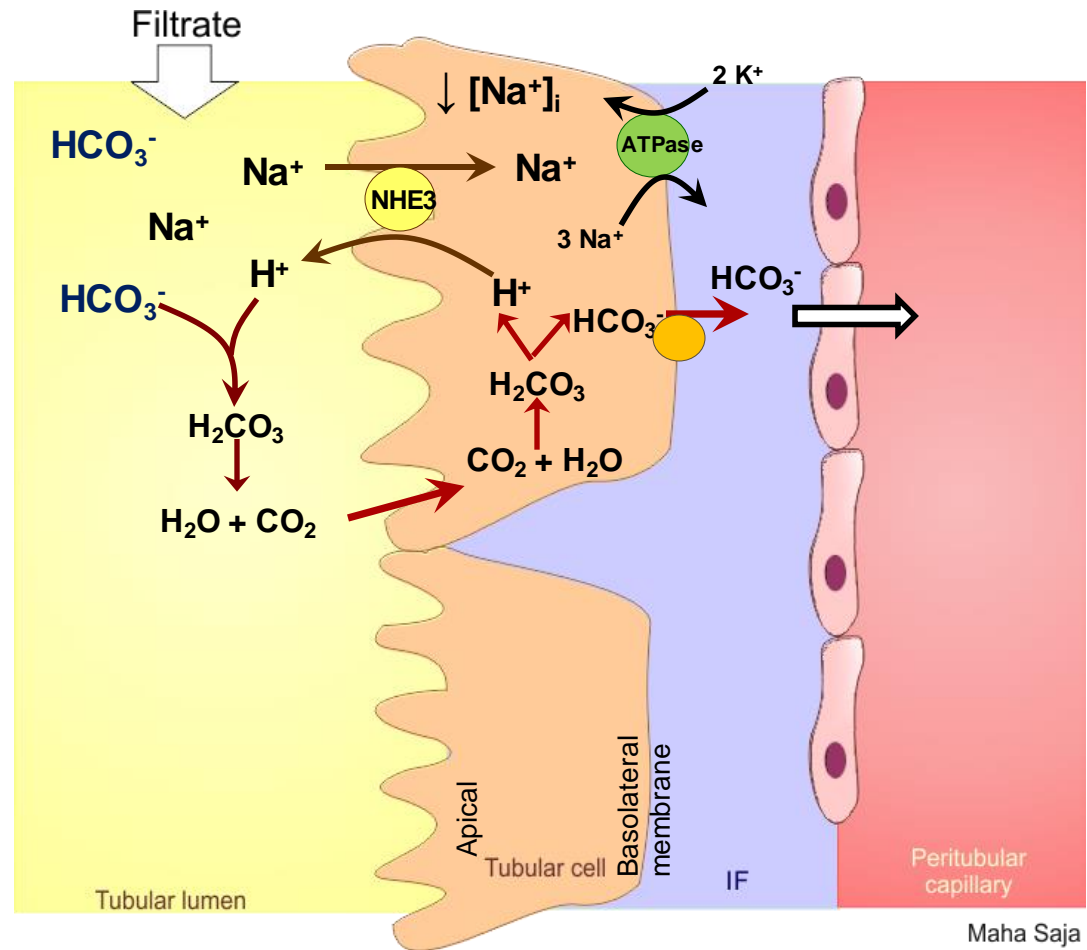
Secreted  $\text{H}^+$  binds to filtered  $\text{HCO}_3^-$  in the tubular lumen



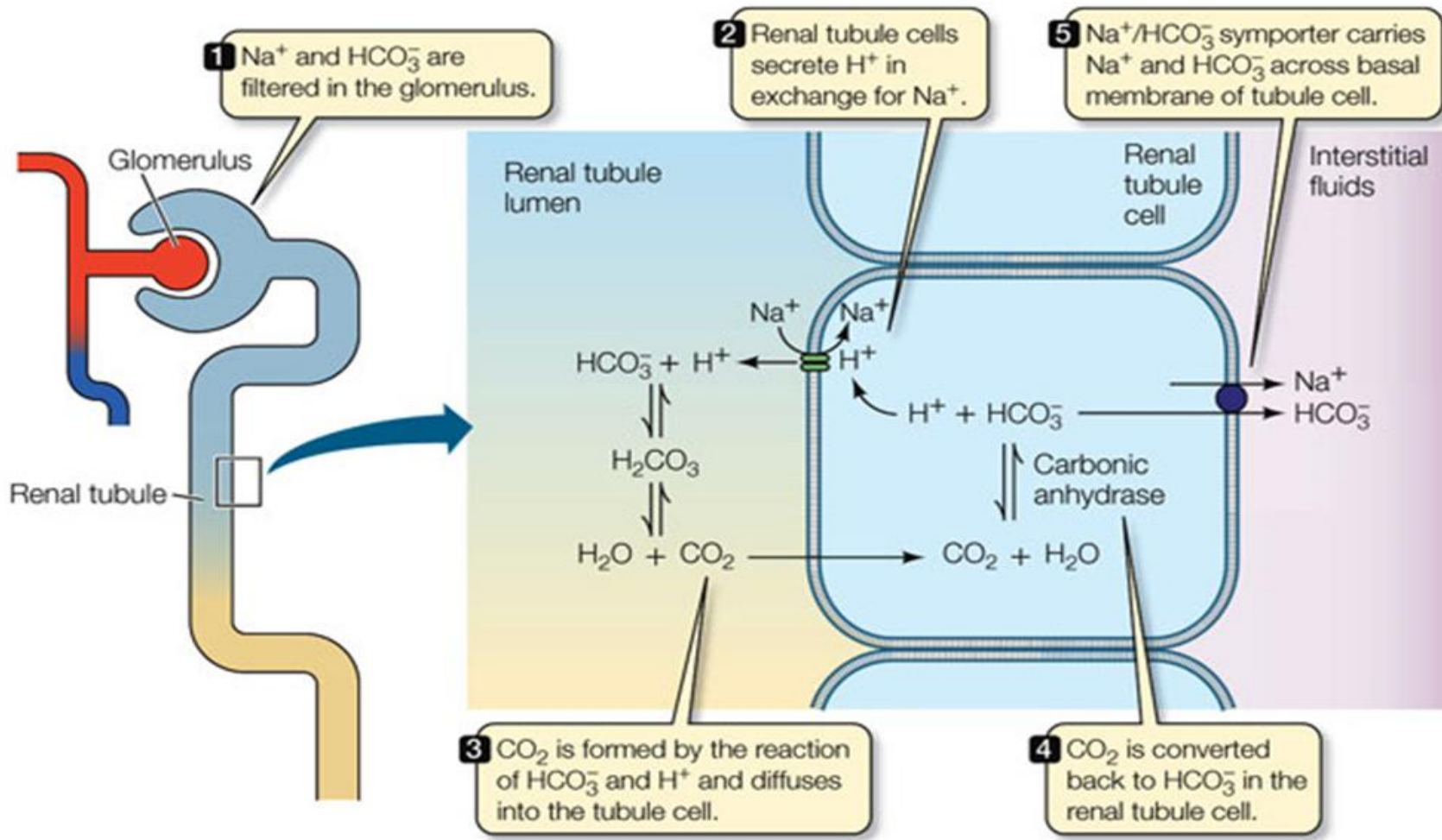
Forming  $\text{H}_2\text{CO}_3$  which dissociates into  $\text{H}_2\text{O}$  &  $\text{CO}_2$



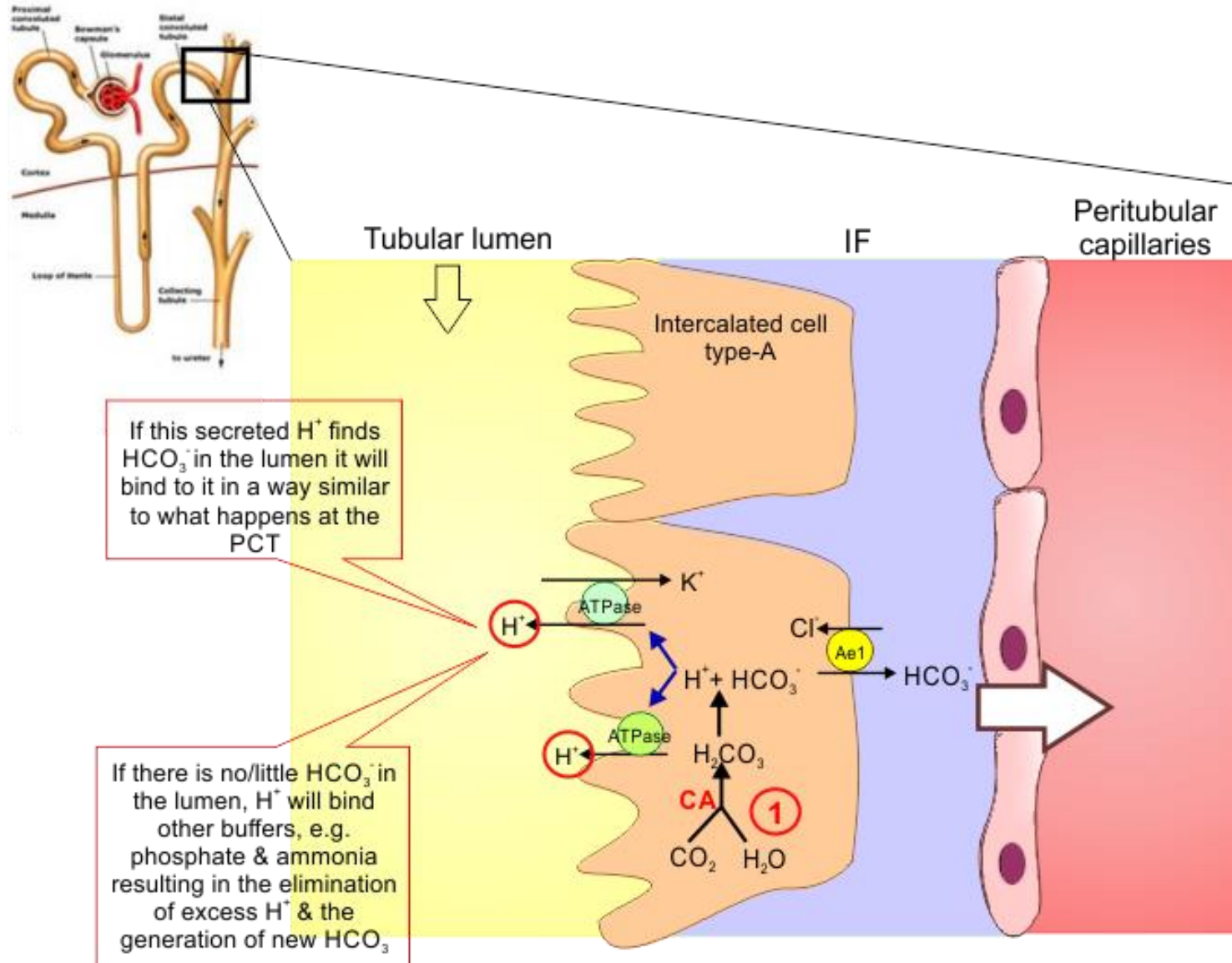
$\text{CO}_2$  diffuses through tubular cell membrane and react with water forming  $\text{H}_2\text{CO}_3$  inside the cell



# $\text{HCO}_3^-$ Reabsorption by the PCT



# What happens at the DCT & CT?



# What happens at the late DCT & CT?

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- The filtrate arriving at the DCT & CT is low in  $\text{HCO}_3^-$ .
- The distal segments of the nephron are characterised by the presence of “intercalated cells” capable of **actively secreting  $\text{H}^+$**  through  $\text{H}^+$ -ATPase and  $\text{H}^+$ - $\text{K}^+$  ATPase present on their apical membrane (**Type-A intercalated cells**).
- Only a limited number of  $\text{H}^+$  can be excreted in its free form in urine.
- Lowest possible urine  $\text{pH}=4.5 \rightarrow \approx 0.04 \text{ mmol/L}$  of free  $\text{H}^+$ .
- **How does the kidney excrete the extra  $\text{H}^+$ ?**

# Non-Bicarbonate Buffers in the Tubular Lumen?

*The extra H<sup>+</sup> secreted will need to be buffered in the tubular lumen*

2 main non-bicarbonate buffers in the tubule

Filtered

Synthesized

Phosphate buffer system

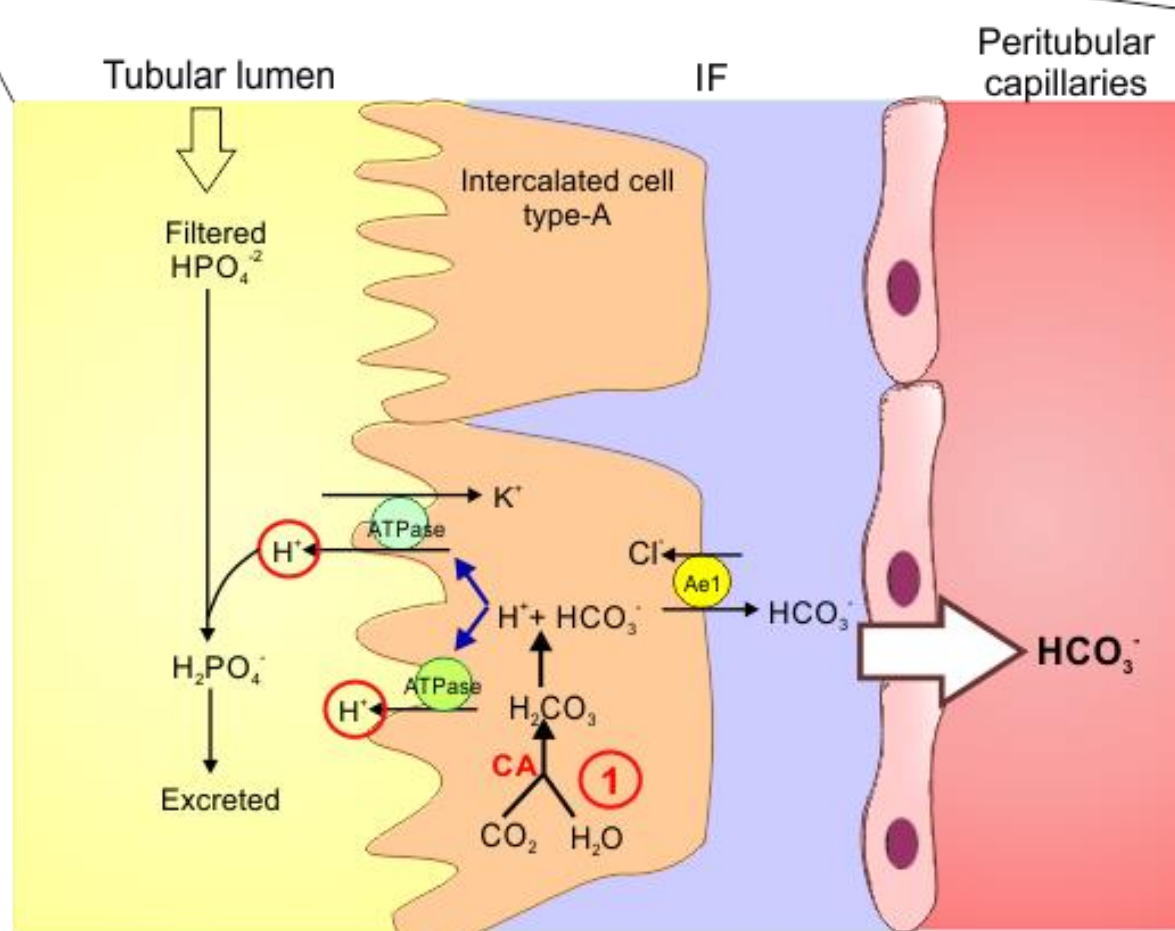
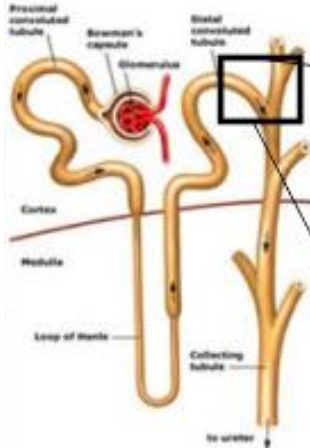


Ammonia buffer system



# Excretion of H<sup>+</sup> and Generation of New HCO<sub>3</sub><sup>-</sup>

## The phosphate buffer system



Excretion of H<sup>+</sup> as phosphate is capable of handling a limited amount of H<sup>+</sup> and will not be enough to rid the body of its daily acid load nor if there is unusually high acid production.

①  $\text{CO}_2$  combines with water within the type A intercalated cell, forming  $\text{H}_2\text{CO}_3$ .

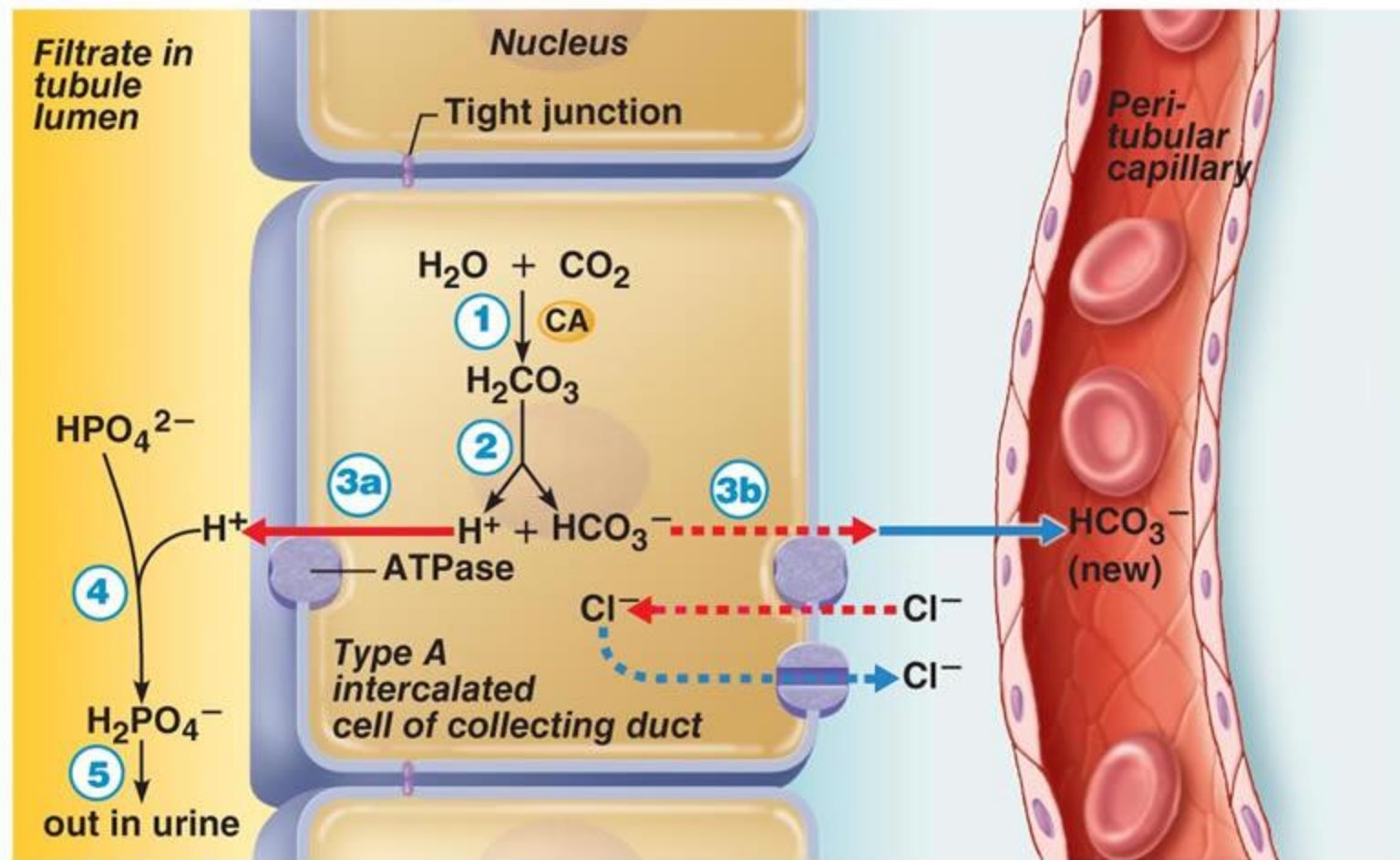
②  $\text{H}_2\text{CO}_3$  is quickly split, forming  $\text{H}^+$  and bicarbonate ion ( $\text{HCO}_3^-$ ).

③a  $\text{H}^+$  is secreted into the filtrate by a  $\text{H}^+$  ATPase pump.

③b For each  $\text{H}^+$  secreted, a  $\text{HCO}_3^-$  enters the peritubular capillary blood via an antiport carrier in a  $\text{HCO}_3^-$ - $\text{Cl}^-$  exchange process.

④ Secreted  $\text{H}^+$  combines with  $\text{HPO}_4^{2-}$  in the tubular filtrate, forming  $\text{H}_2\text{PO}_4^-$ .

⑤ The  $\text{H}_2\text{PO}_4^-$  is excreted in the urine.



→ Primary active transport  
- - - Secondary active transport  
→ Simple diffusion  
- - - Facilitated diffusion

● Transport protein  
● Ion channel  
● CA Carbonic anhydrase

# Excretion of H<sup>+</sup> and Generation of New HCO<sub>3</sub><sup>-</sup>

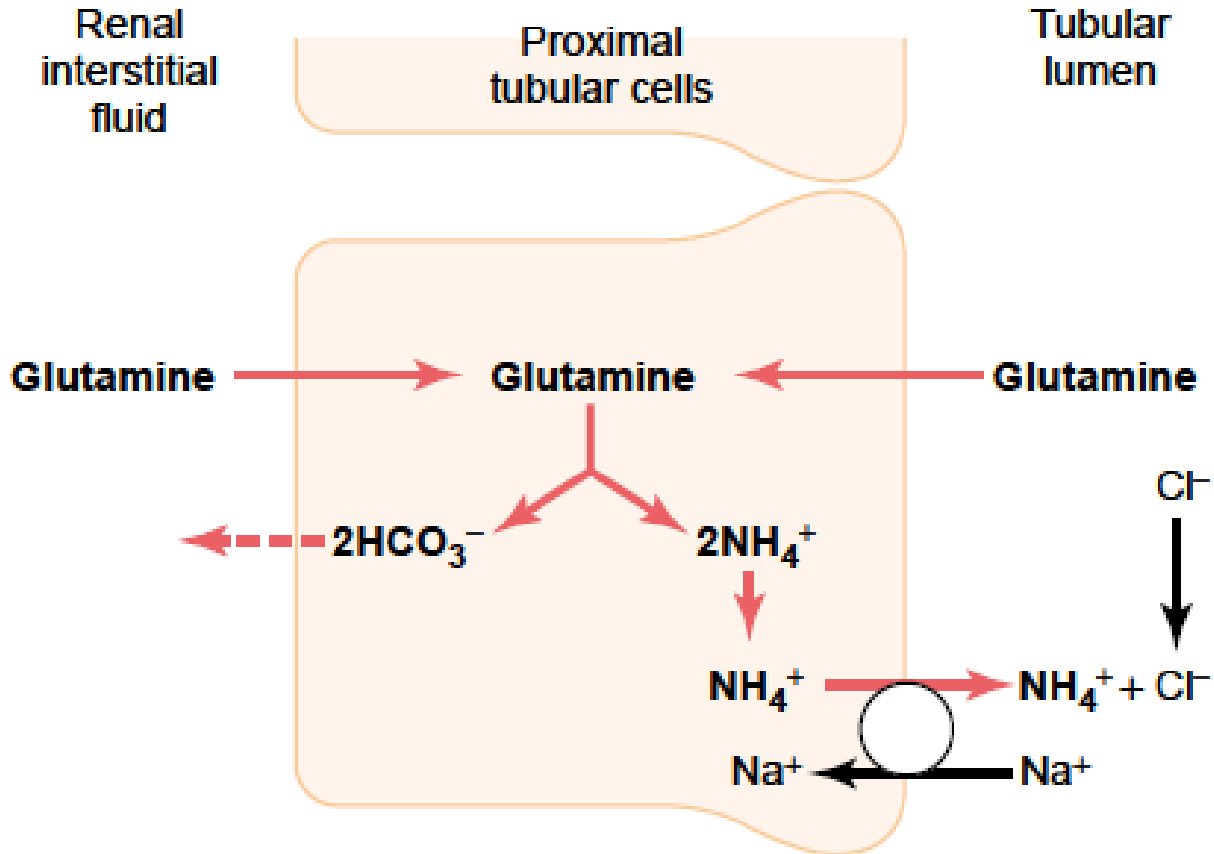
## The Ammonia Buffer System

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- Renal tubular cells, especially PCT, are capable of generating ammonium (NH<sub>4</sub><sup>+</sup>) “**ammoniogenesis**” which is then excreted in urine carrying with it H<sup>+</sup>.
- The rate of ammoniogenesis can be modified according to the needs of the body.
- Quantitatively, the ammonia buffer system is more important than the phosphate buffer system for H<sup>+</sup> excretion in urine.
- It is the most important system in case of acidosis.



# The Ammonia Buffer System

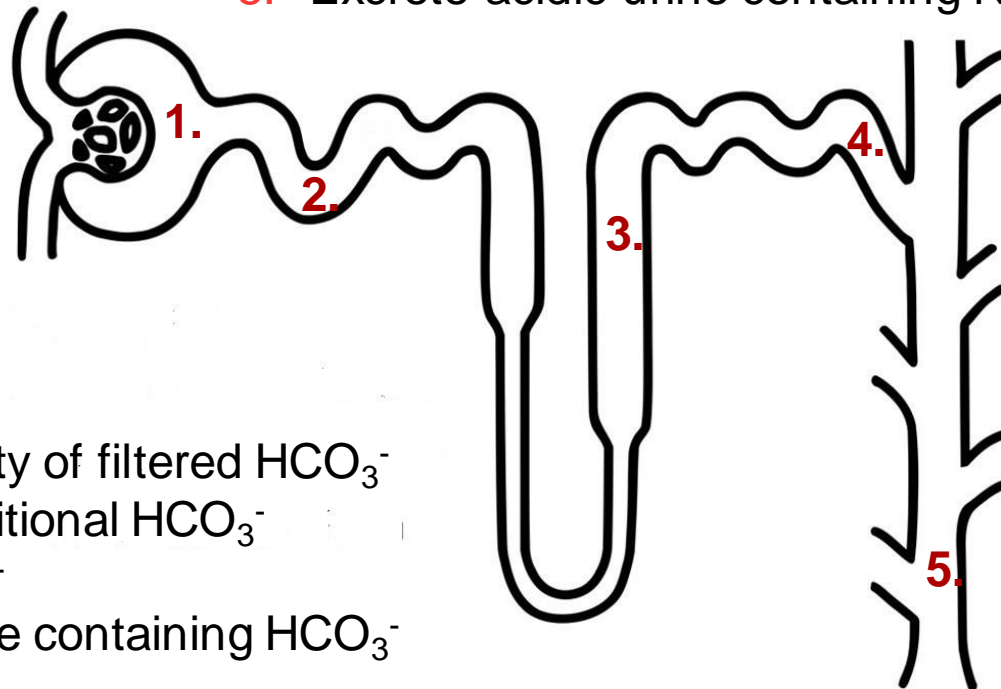




# The Overall Scheme of Renal Excretion of Acids & Bases

## *To excrete acid:*

1. Freely filter  $\text{HCO}_3^-$
2. Reabsorb the majority of filtered  $\text{HCO}_3^-$
3. Reabsorb some additional  $\text{HCO}_3^-$
4. Secrete  $\text{H}^+$  (titrate filtered bases, i.e.  $\text{HPO}_4^{-2}$ ) and secrete  $\text{NH}_4^+$
5. Excrete acidic urine containing  $\text{NH}_4^+$



## *To excrete base:*

1. Freely filter  $\text{HCO}_3^-$
2. Reabsorb the majority of filtered  $\text{HCO}_3^-$
3. Reabsorb some additional  $\text{HCO}_3^-$
4. Secrete some  $\text{HCO}_3^-$
5. Excrete alkaline urine containing  $\text{HCO}_3^-$

# Factors Affecting H<sup>+</sup> Secretion and HCO<sub>3</sub><sup>-</sup> Reabsorption

Table 30-2

## Factors That Increase or Decrease H<sup>+</sup> Secretion and HCO<sub>3</sub><sup>-</sup> Reabsorption by the Renal Tubules

### Increase H<sup>+</sup> Secretion and HCO<sub>3</sub><sup>-</sup> Reabsorption

↑ PCO<sub>2</sub>

↑ H<sup>+</sup>, ↓ HCO<sub>3</sub><sup>-</sup>

↓ Extracellular fluid volume

↑ Angiotensin II

↑ Aldosterone

Hypokalemia

### Decrease H<sup>+</sup> Secretion and HCO<sub>3</sub><sup>-</sup> Reabsorption

↓ PCO<sub>2</sub>

↓ H<sup>+</sup>, ↑ HCO<sub>3</sub><sup>-</sup>

↑ Extracellular fluid volume

↓ Angiotensin II

↓ Aldosterone

Hyperkalemia

**THANK YOU**

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