

Acid Base Chemistry & Metabolism



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medpathwaymcat



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Bronsted-Lowry Acids/Bases

Bronsted-Lowry Acid	Donates Protons
Bronsted-Lowry Base	Accepts Protons

Bronsted-Lowry Acids/Bases: Capable of donating and accepting protons, respectively. The system is also limited as it only focuses on protons.

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Weak Conjugate Base

$K_a = \text{Large}$

Strong

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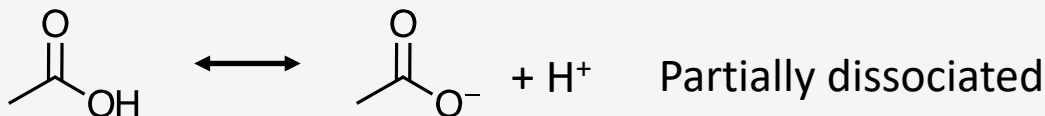
Bronsted-Lowry Acids/Bases: Capable of donating and accepting protons, respectively. The system is also limited as it only focuses on protons.



$K_a = \text{Large}$

Strong

Weak Conjugate Base



$K_a = 1.8 \times 10^{-5}$

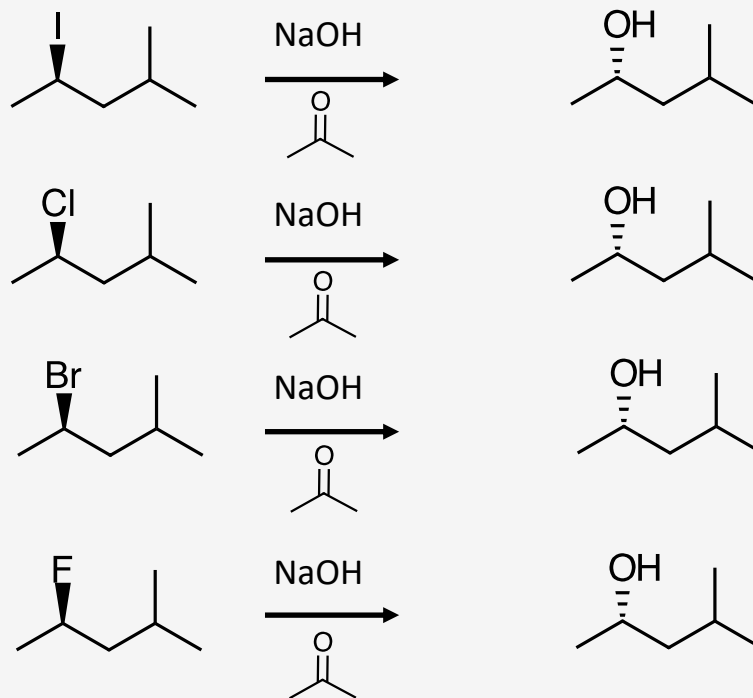
Weak

$$= \frac{[\text{CH}_3\text{COO}^-][\text{H}^+]}{\text{CH}_3\text{COOH}}$$

Strong Conjugate Base

$\text{p}K_a = 4.75$

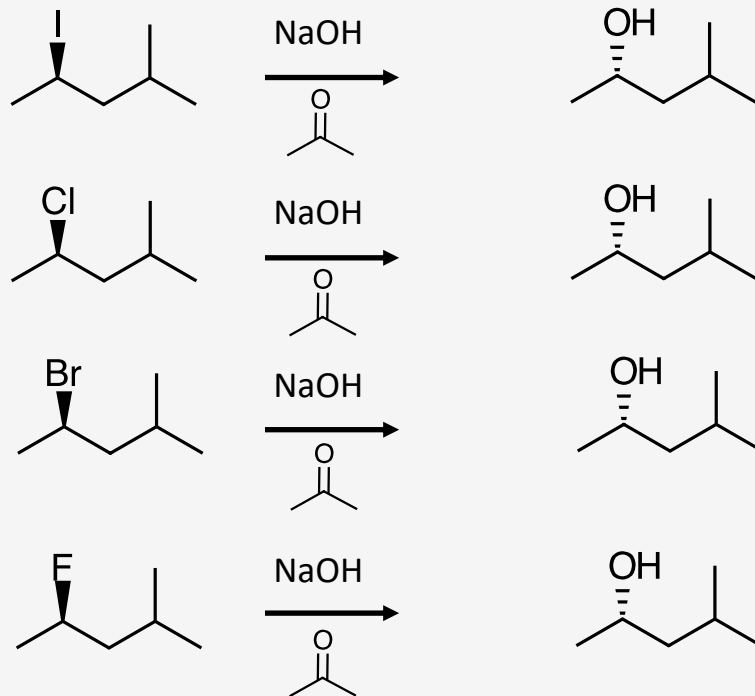
Conjugate Bases and Leaving Groups



Relative Rates of Product Formation?

Conjugate Bases and Leaving Groups

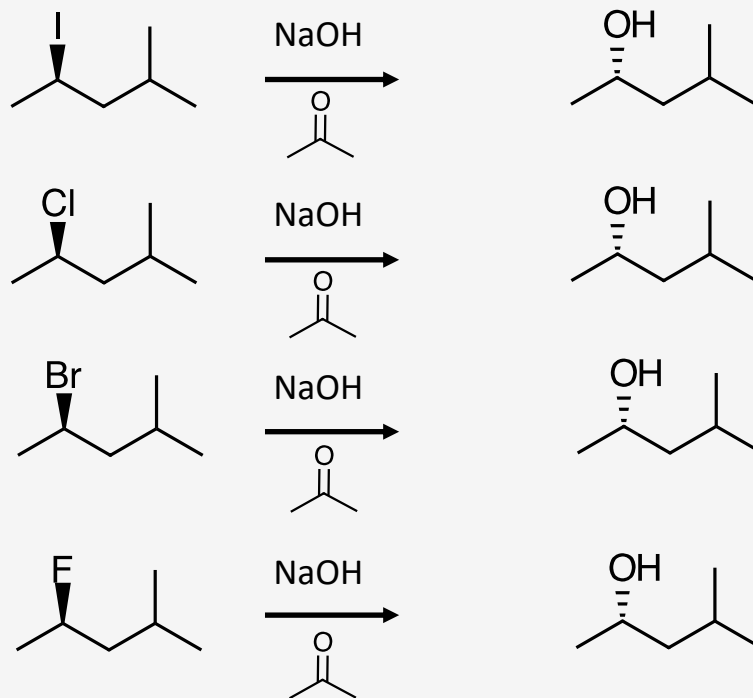
S_N2



Relative Rates of
Product Formation?

HI > HBr > HCl > HF = Relative Strength of Acids

Conjugate Bases and Leaving Groups



Relative Rates of Product Formation?

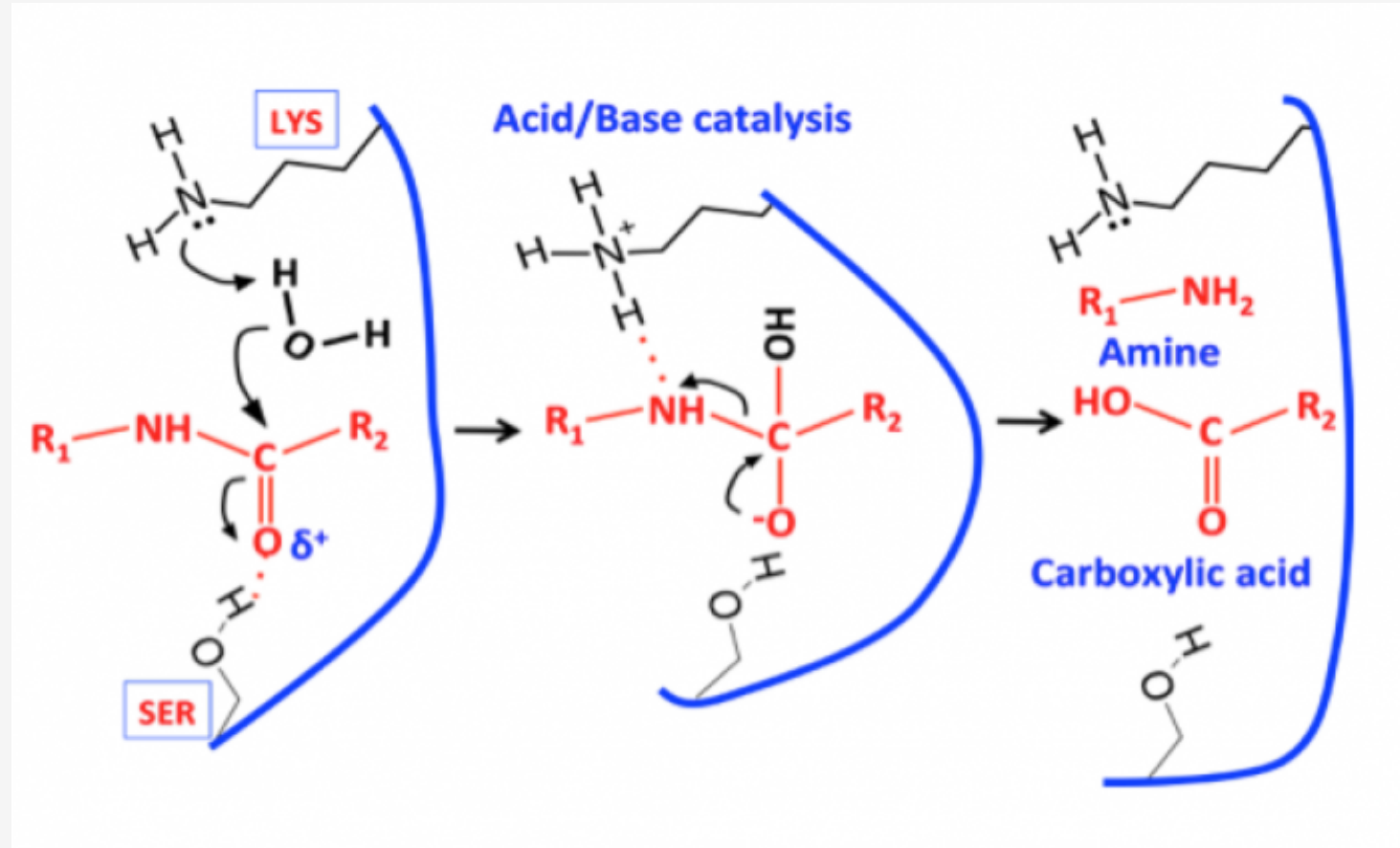
HI > HBr > HCl > HF = Relative Strength of Acids



Weakest conjugate base;
Best leaving group

Strongest conjugate base;
Worst leaving group

Acid/Base Catalysis as Leaving Groups



R-NH⁻ vs R-NH₂ leaving groups

Acid Base Problems

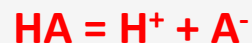
What is the pH of a 0.2 M solution of a weak acid (HA) with a $K_a = 1.8 \times 10^{-4}$?

- A. 1.3
- B. 2.2
- C. 3.4
- D. 4.1

Acid Base Problems

What is the pH of a 0.2 M solution of a weak acid (HA) with a $K_a = 1.8 \times 10^{-4}$?

Weak acids do not completely dissociate into protons and their conjugate bases. Set up an ICE table describing the initial and final changes for the components of the following equilibrium:



ICE Table:

Initial

Change

Equilibrium

Acid Base Problems

What is the pH of a 0.2 M solution of a weak acid (HA) with a $K_a = 1.8 \times 10^{-4}$?

Weak acids do not completely dissociate into protons and their conjugate bases. Set up an ICE table describing the initial and final changes for the components of the following equilibrium:



ICE Table:
Initial
Change
Equilibrium

	[HA]	[H ⁺]	[A ⁻]
Initial	0.2M	10 ⁻⁷ M	0 M
Change	- X	+ X	+ X
Final	0.2M - X	+ X	+ X

Autoionization of water
Is negligible



$$K_A = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]} = \frac{[\text{X}][\text{X}]}{0.2\text{M} - \text{X}}$$

$$1.8 \times 10^{-4} = \frac{[\text{X}][\text{X}]}{0.2\text{M} - \text{X}} = \frac{\text{X}^2}{0.2\text{M}}$$

$$\text{X}^2 = 3.6 \times 10^{-5}$$

$$\text{X} = .006$$

Acid Base Problems

$$\text{pH} = -\log [.006] = ??$$

Acid Base Problems

$$\text{pH} = -\log [.006] = ??$$

- A. 1.3
- B. 2.2**
- C. 3.4
- D. 4.1

[H⁺]	[pH]
0.001 M	3.0
0.006 M	2.22
0.01 M	2.0

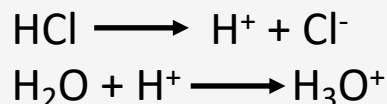
$$\text{Log} [.001] = -3.0$$

$$\text{Log} [.01] = -2.0$$

Arrhenius Acids/Bases

Arrhenius Acid	Increases $[H^+]$ in solution
Arrhenius Base	Increases $[OH^-]$ in solution

Arrhenius Acids/Bases: The Arrhenius definition of an acid is limited as it only focuses on water as the solvent.



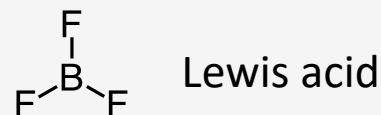
Lewis Acids/Bases

Lewis Acid	Electron Pair Acceptor
Lewis Base	Electron Pair Donor

Lewis Acids/Bases: Electron pair acceptors and donors, respectively. The Lewis acid/base definition focuses on what electrons are actually doing in a reaction.

:NH_3 = Lewis Base (Nonbonding electrons)

Nucleophile



Electrophile

3 Acid/Base Systems

Arrhenius Acid	Increases $[H^+]$ in solution
Arrhenius Base	Increases $[HO^-]$ in solution
Bronsted-Lowry Acid	Donates Protons
Bronsted-Lowry Base	Accepts Protons
Lewis Acid	Electron Pair Acceptor
Lewis Base	Electron Pair Donor

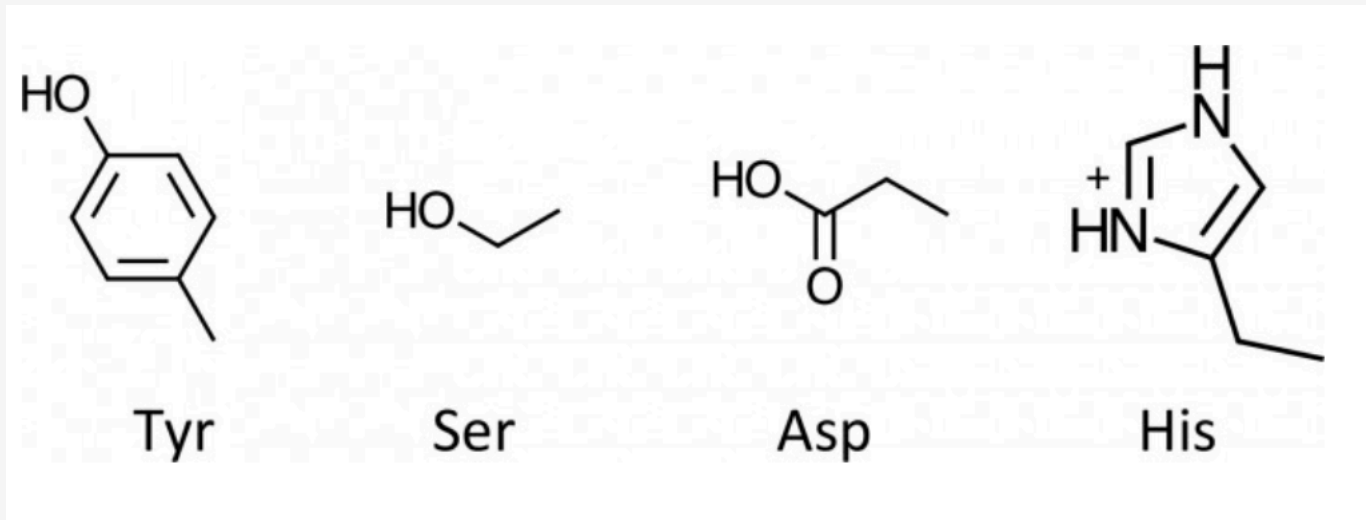
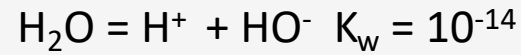
Arrhenius Acids/Bases: The Arrhenius definition of an acid is limited as it only focuses on water as the solvent.

Bronsted-Lowry Acids/Bases: Capable of donating and accepting protons, respectively. The system is also limited as it only focuses on protons.

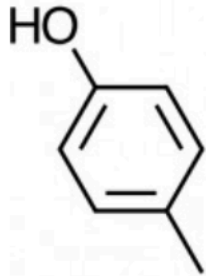
Lewis Acids/Bases: Electron pair acceptors and donors, respectively. The Lewis acid/base definition focuses on what electrons are actually doing in a reaction.

Relative Strengths of Acids

$$pK_a = -\log [K_a] \quad pK_a + pK_b = 14$$

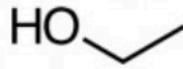


Relative pK_a values



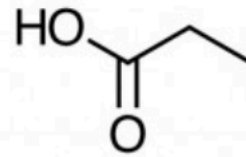
Tyr

pK_a = 10



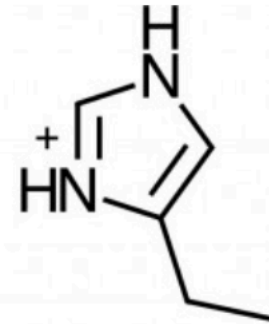
Ser

pK_a = 14



Asp

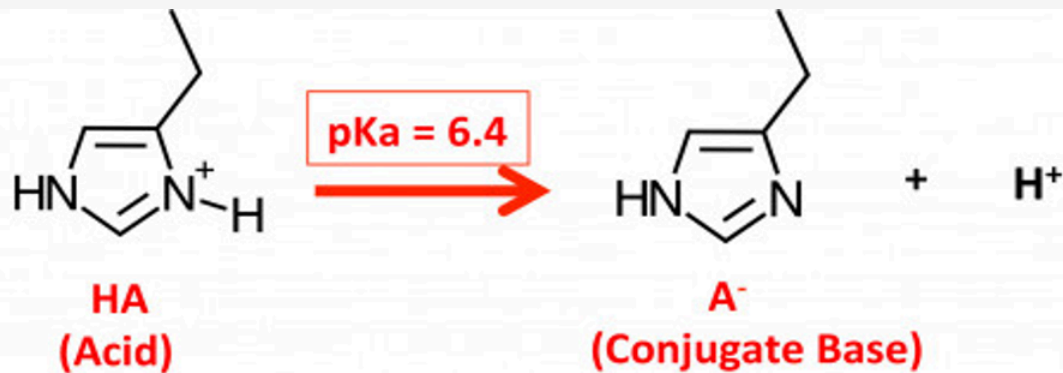
pK_a = 3



His

pK_a = 6.4

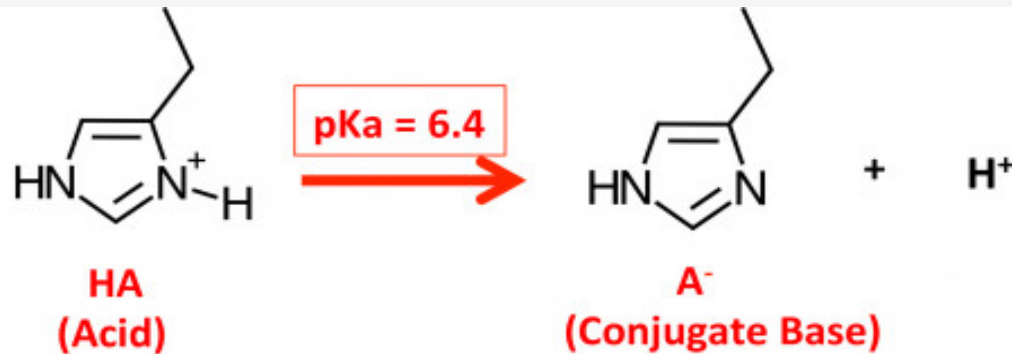
Henderson Hasselbalch



$$\text{pH} = \text{pKa} + \log\left[\frac{\text{Base}}{\text{Acid}}\right]$$

Q: What is the fraction of histidine residues that are positively charged at pH = 7.4?

Henderson Hasselbalch



$$\text{pH} = \text{pKa} + \log\left[\frac{\text{Base}}{\text{Acid}}\right]$$

Q: What is the fraction of histidine residues that are positively charged at pH = 7.4?

Solve for [Base]/[Acid] knowing that pKa = 6.4 and pH = 7.4.

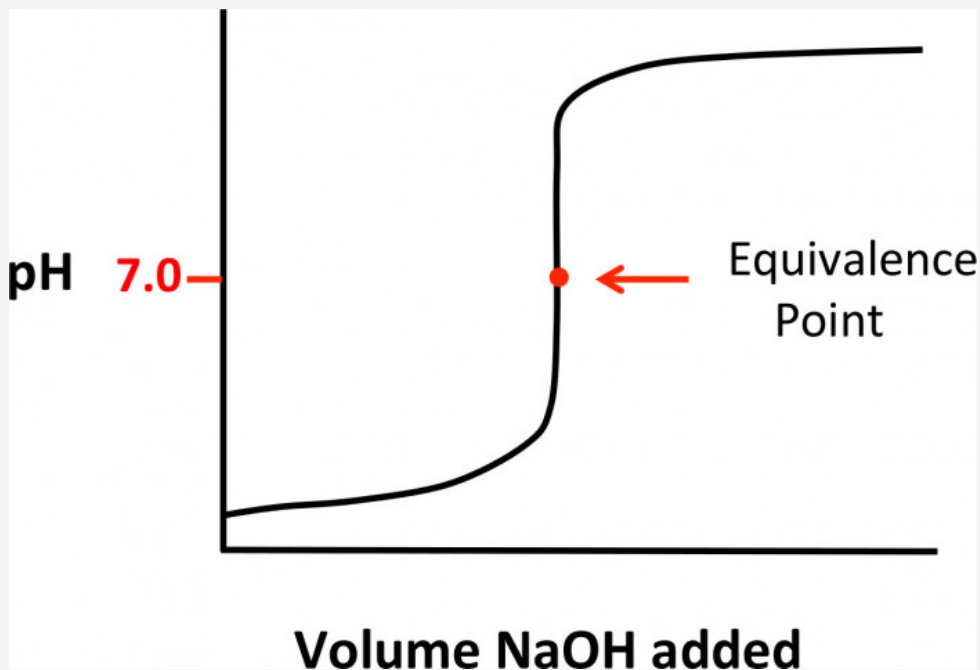
$$\text{pH} - \text{pKa} = \log(X) \quad X = \frac{[\text{A}^-]}{[\text{HA}]}$$

A: Antilog (pH-pKa) = antilog[log(X)] **Antilog 1 = X ; X = 10**

Therefore, as X = [Base]/[Acid], the ratio of [Base]/[Acid] = 10.

There are 10X as many His residues in conjugate base than His residues than in acidic, positively charged form.

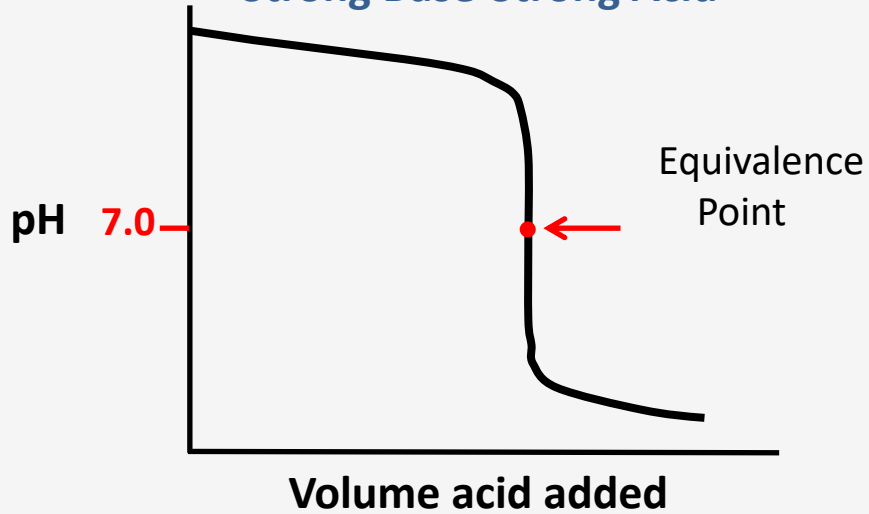
Titration of Strong Acid w/ Strong Base



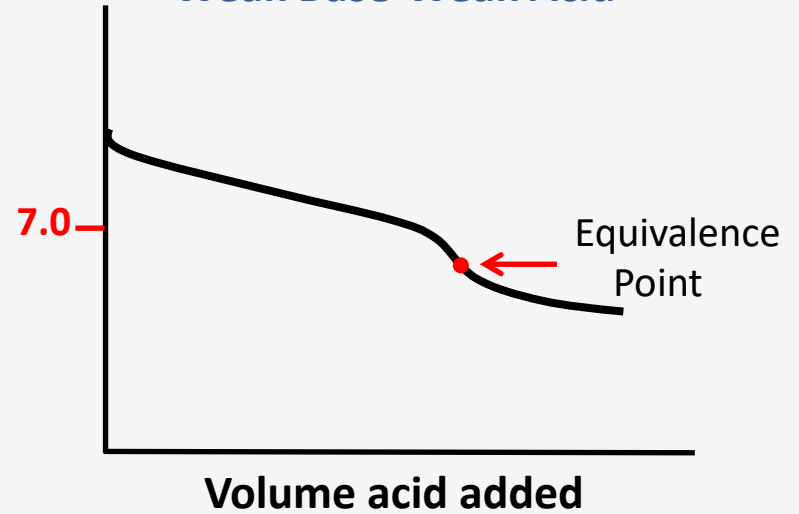
**# moles of base added = #
number of moles of acid
present in the solution**

Titration Curves

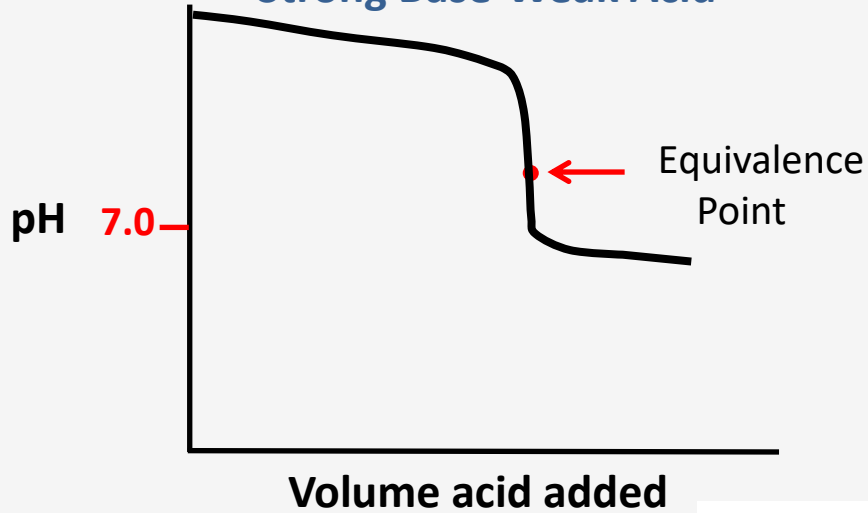
Strong Base-Strong Acid



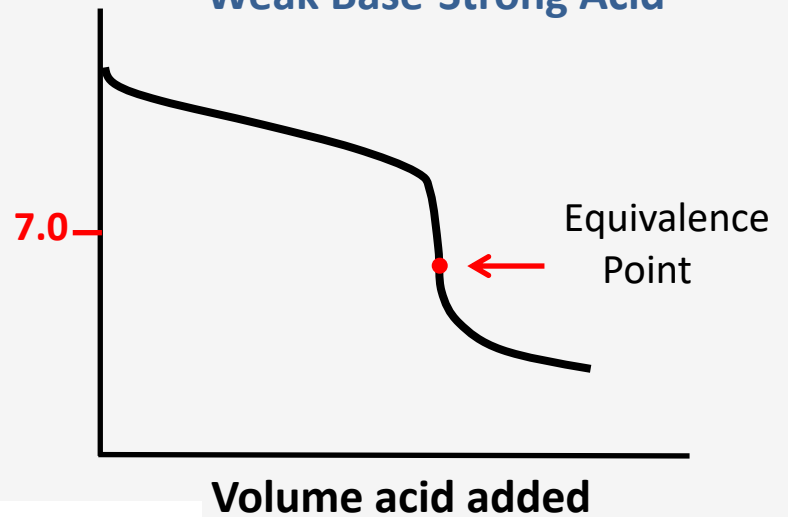
Weak Base-Weak Acid



Strong Base-Weak Acid

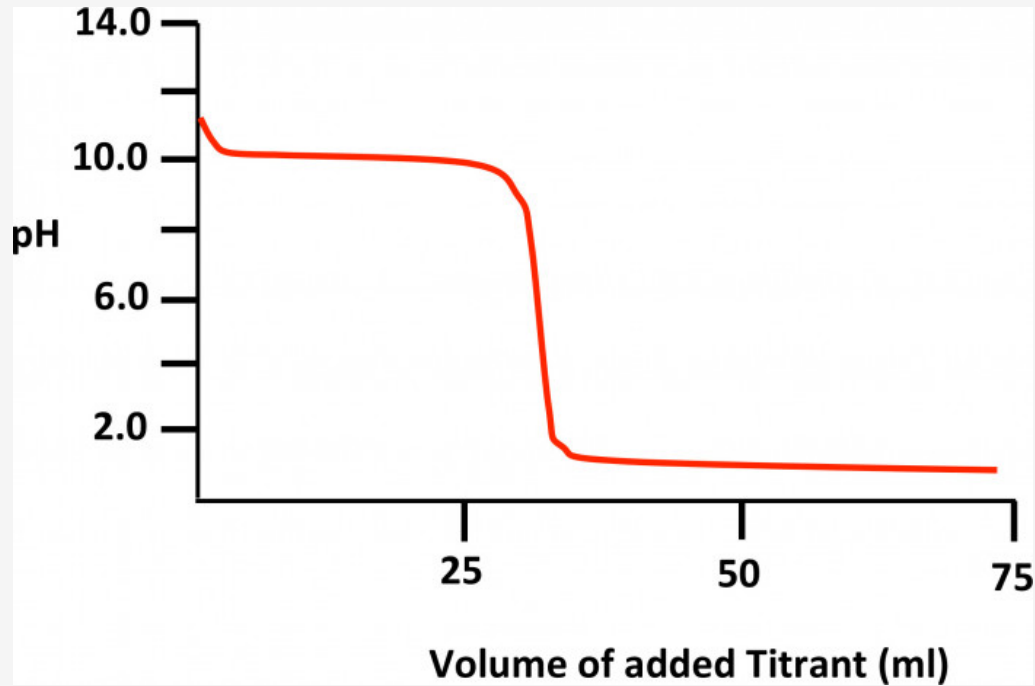


Weak Base-Strong Acid

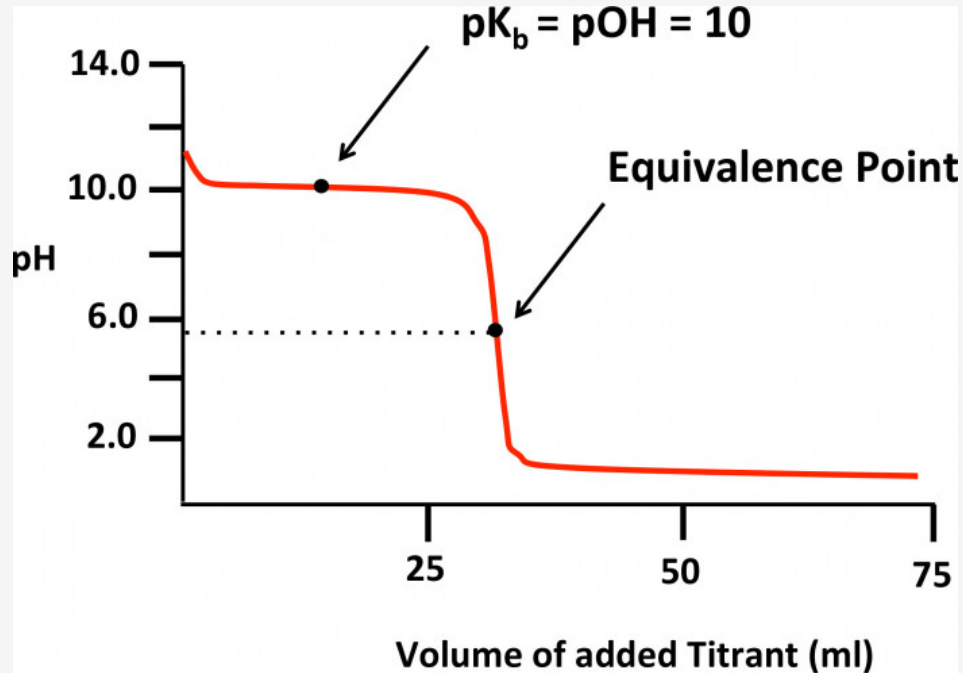


Titrations

A compound was titrated as shown. What is the K_a value of the compound?



Titrations

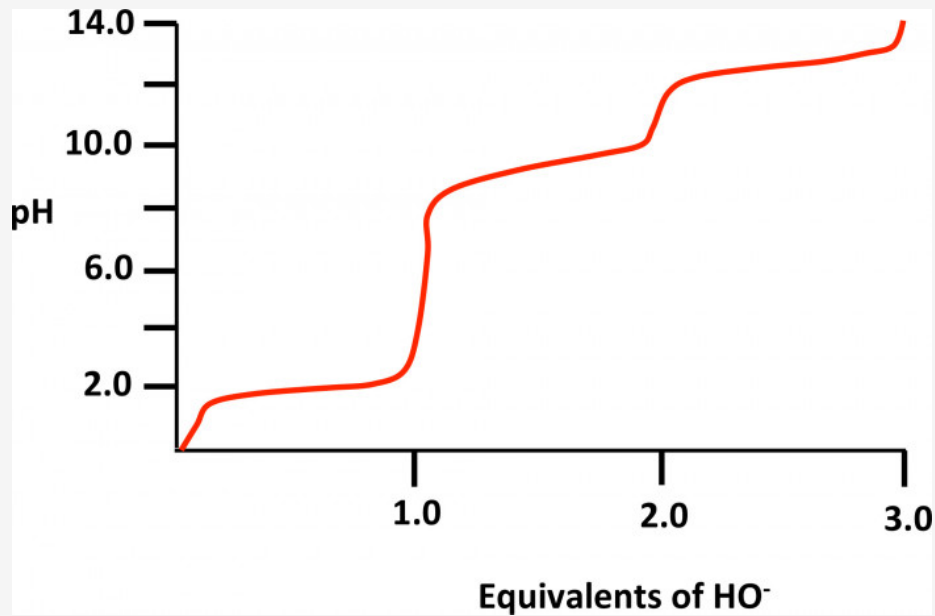


The titration is between a weak base and an acid as the pH decreases with the addition of titrant.

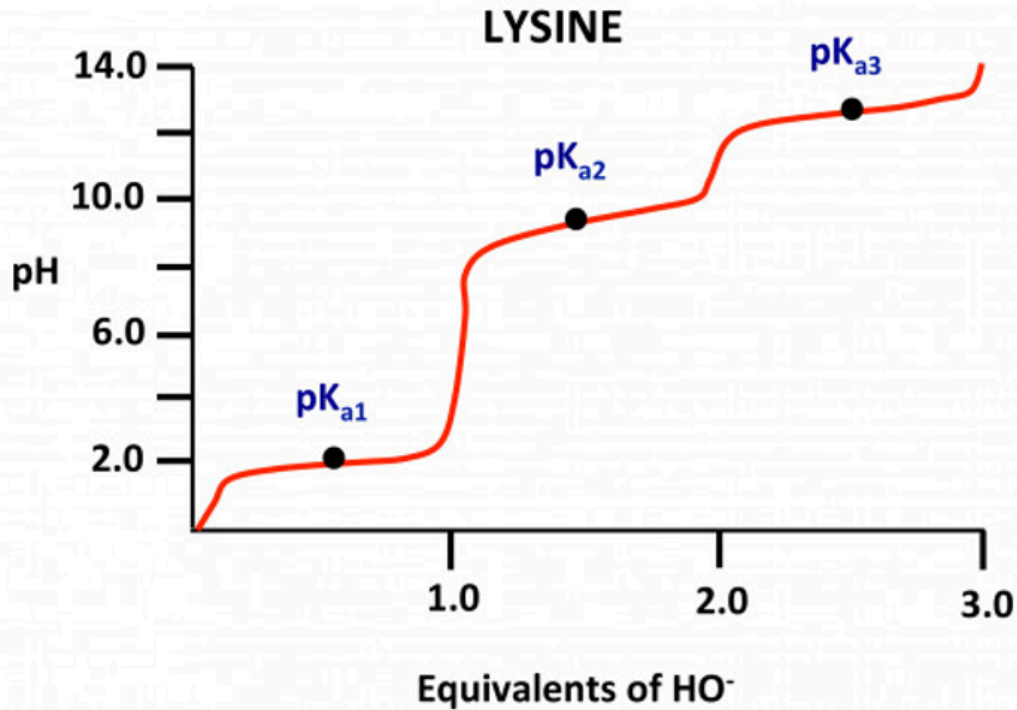
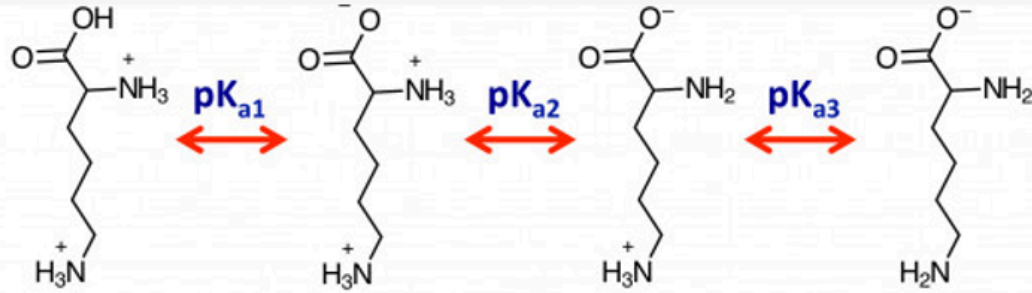
Observe from the titration that the inflection point (where the slope = 0), represents the pK_b . Therefore, $pOH = pK_b = 10$. Recall that $K_b = 10^{-pK_b} = 10^{-10}$ and $K_a K_b = K_w = 10^{-14}$. Therefore, $K_a = 10^{-14}/10^{-10} = 10^{-4}$.

Amino Acid Titrations

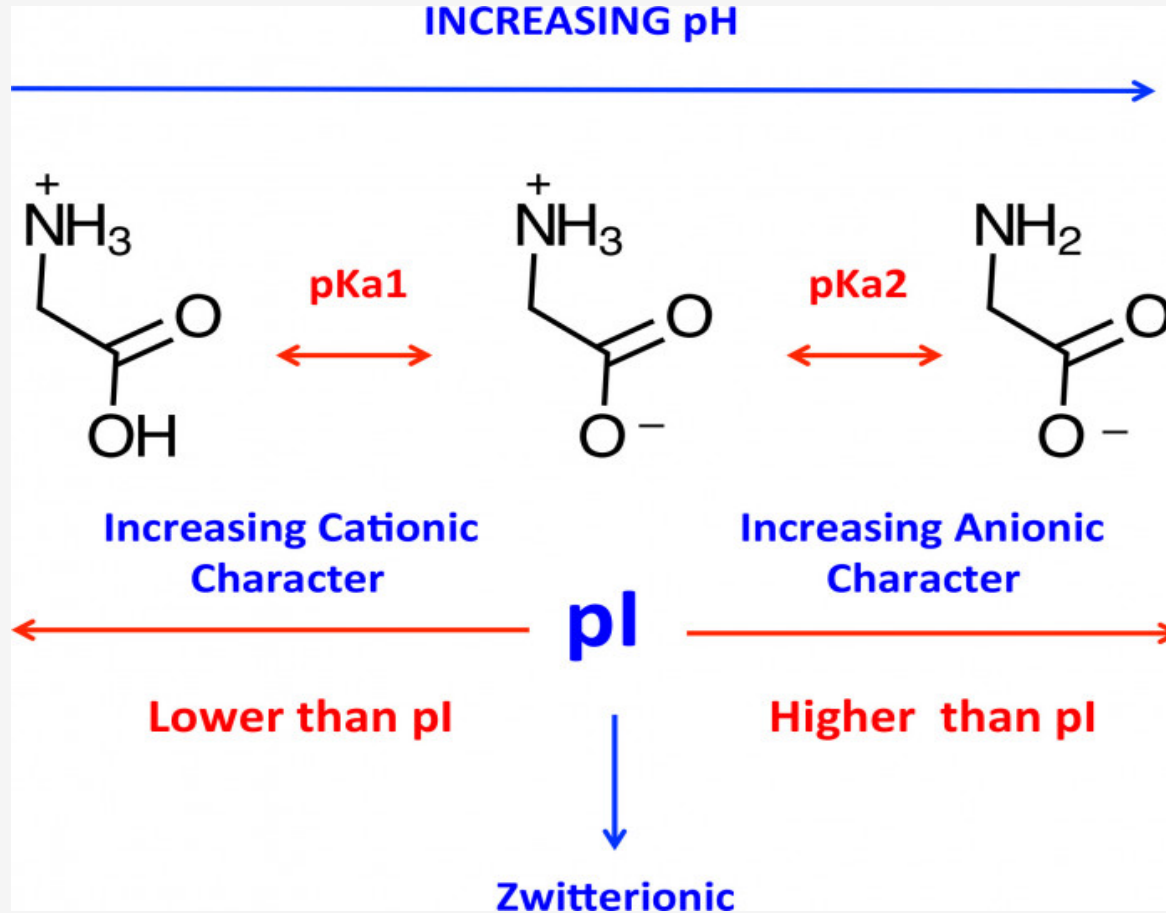
Which of the following amino acids best fits the titration profile shown below?



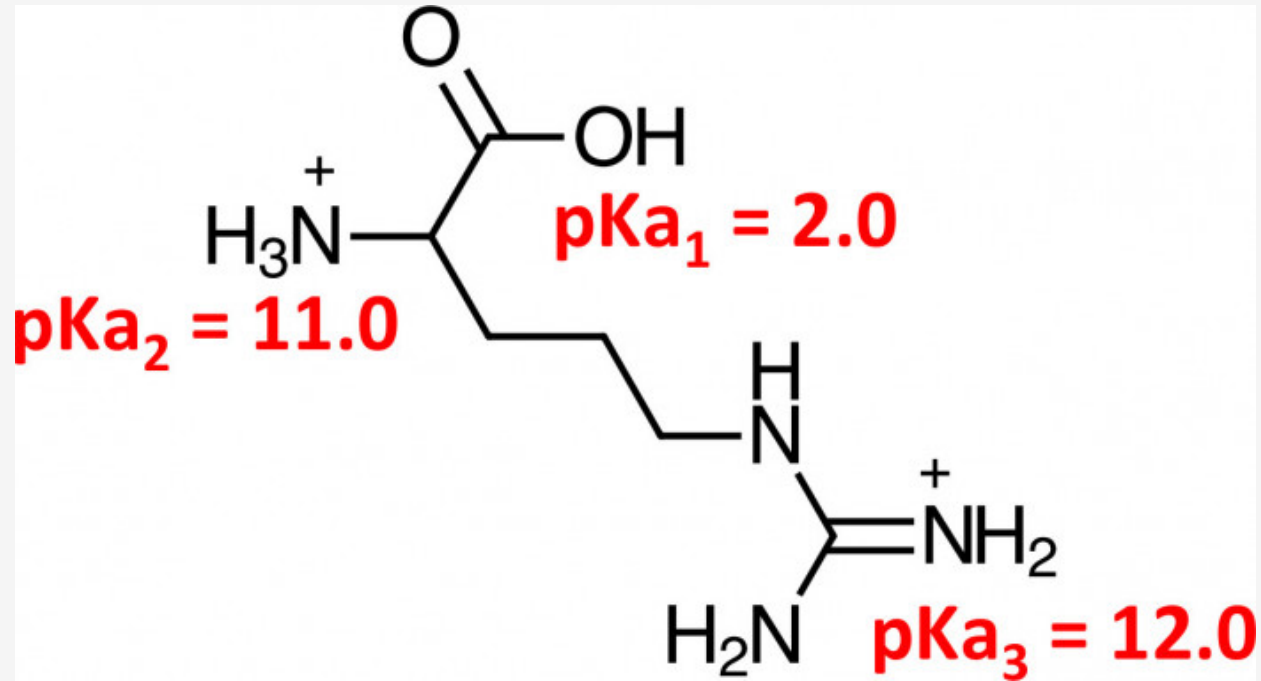
Lysine Titration



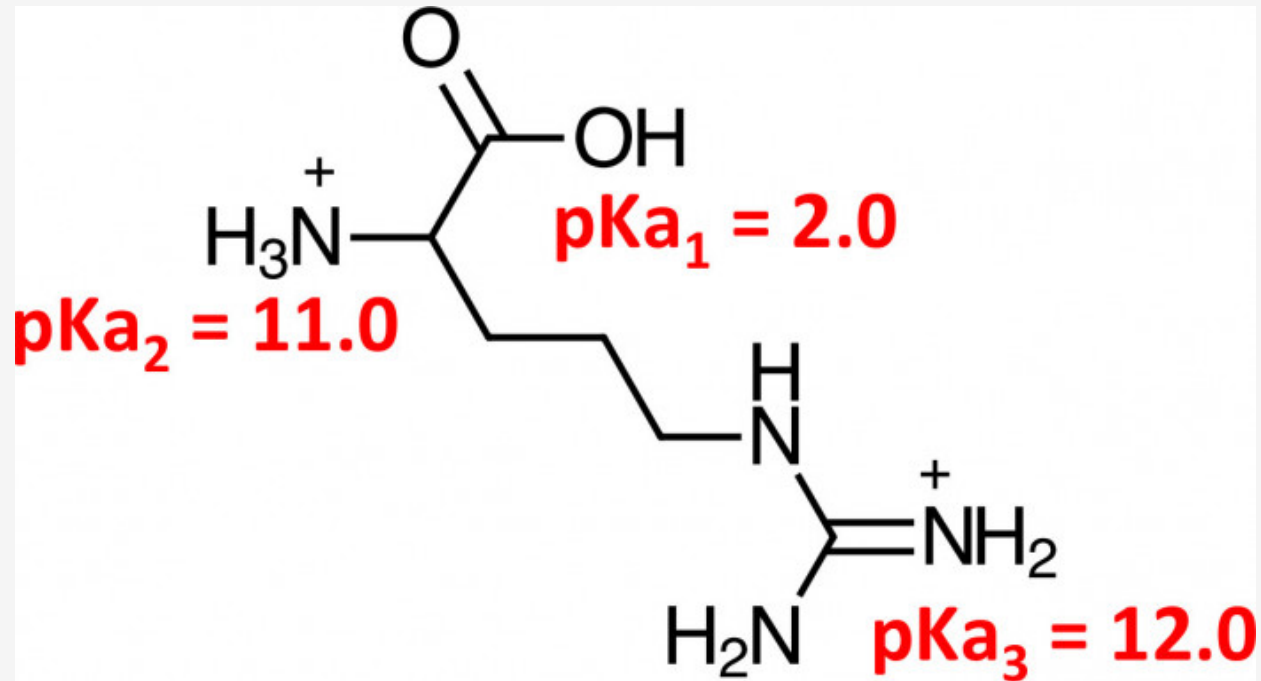
Isoelectric Point



What is the isoelectric point of arginine?



Isoelectric point of arginine?

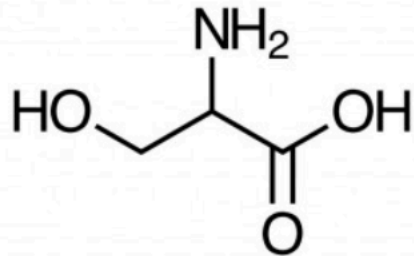


$$\text{pI} = (11 + 12) \times 0.5 = 11.5$$

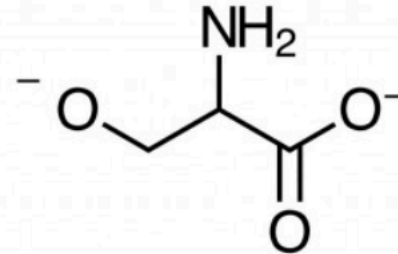
Relative pK_a Values

Which of the following forms of serine would fail to exist at any pH?

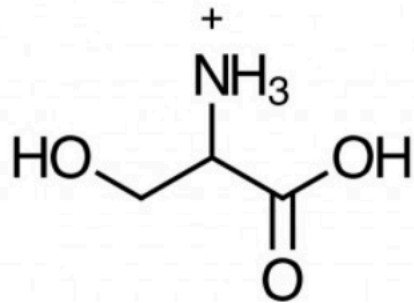
A



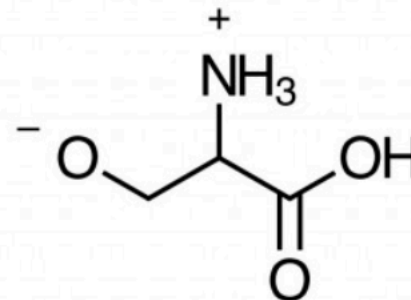
B



C



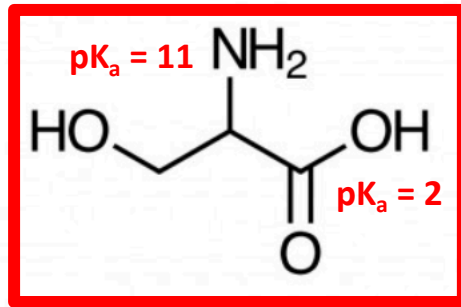
D



Relative pK_a Values

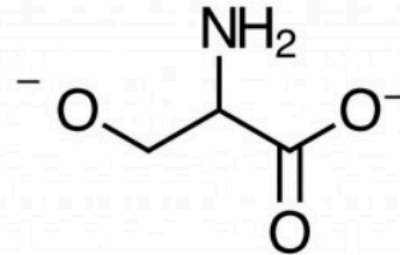
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A



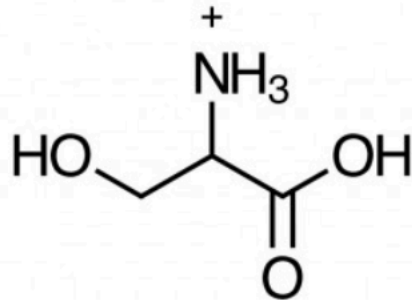
Loss of amino proton
before Carboxyl Proton

B



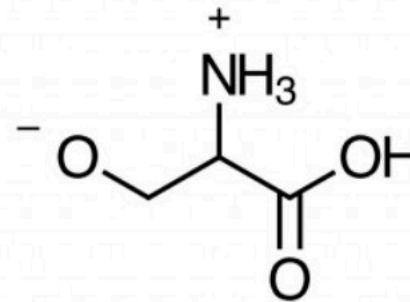
$pH > pI$

C



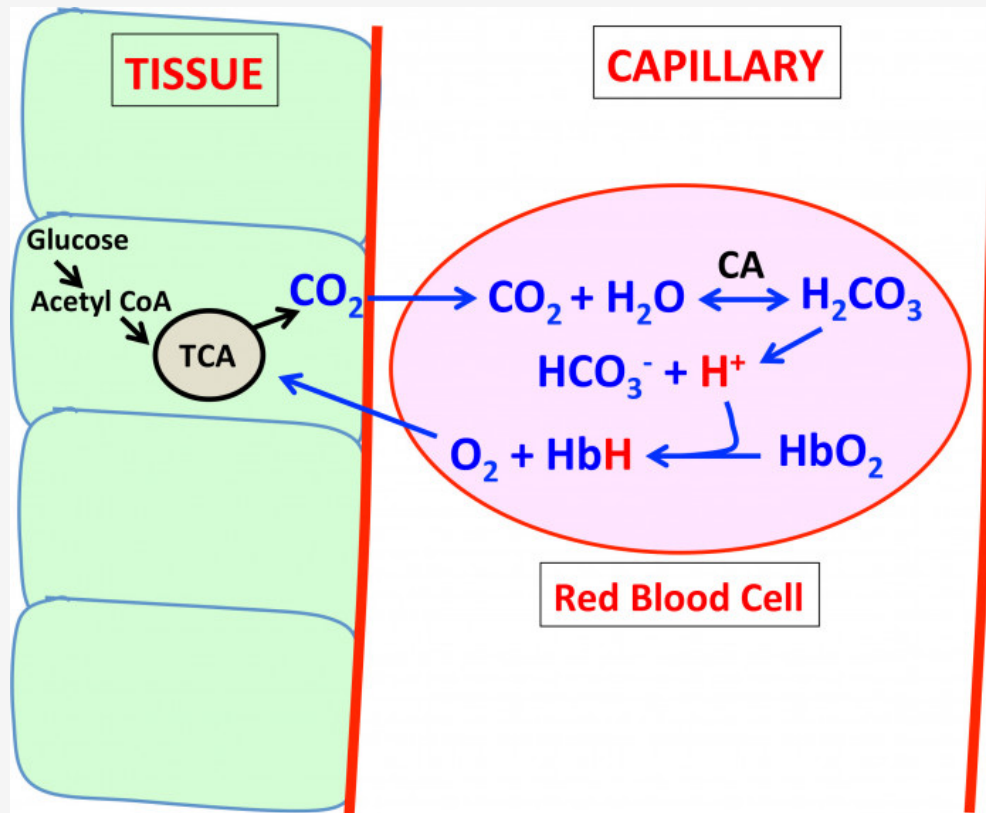
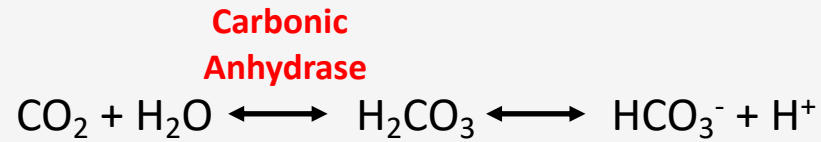
$pH < pI$

D



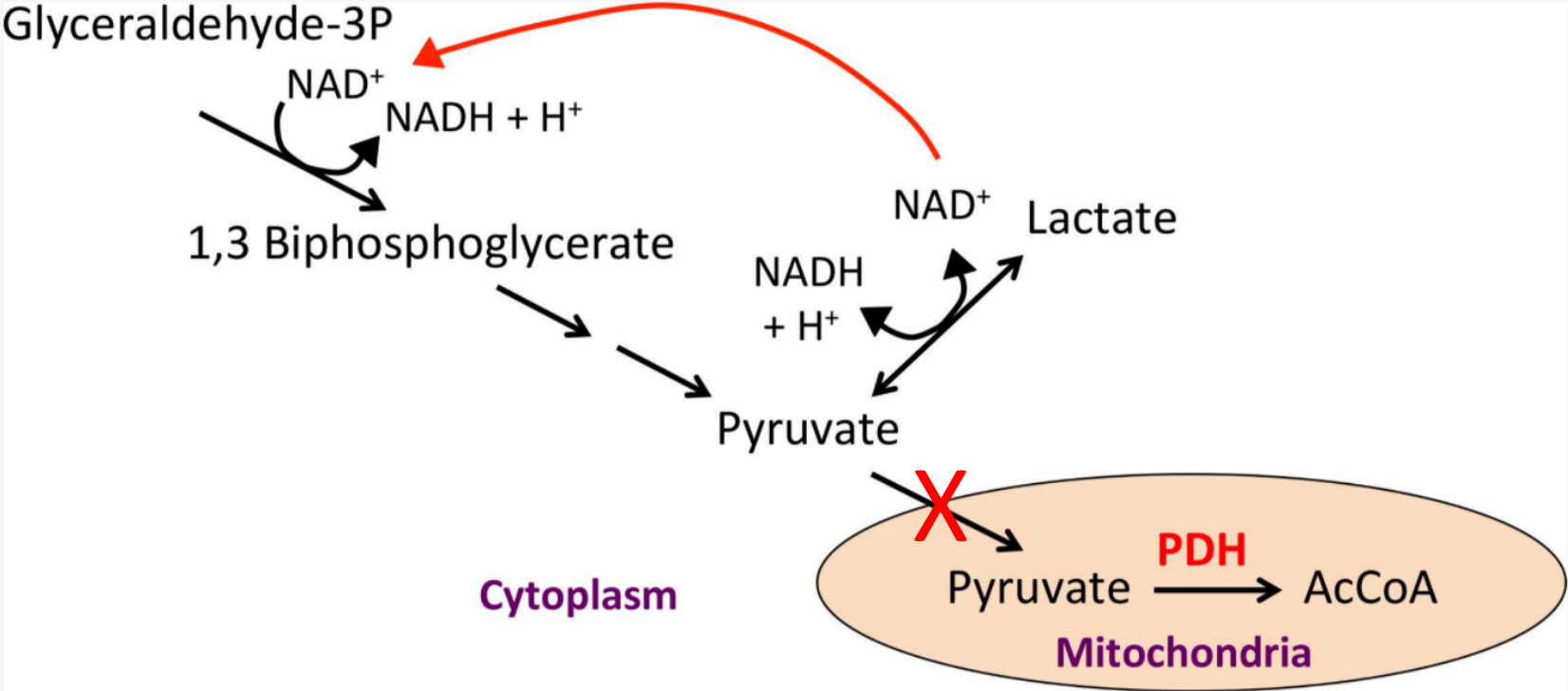
Isoelectric Point

Carbonic Acid Buffering System

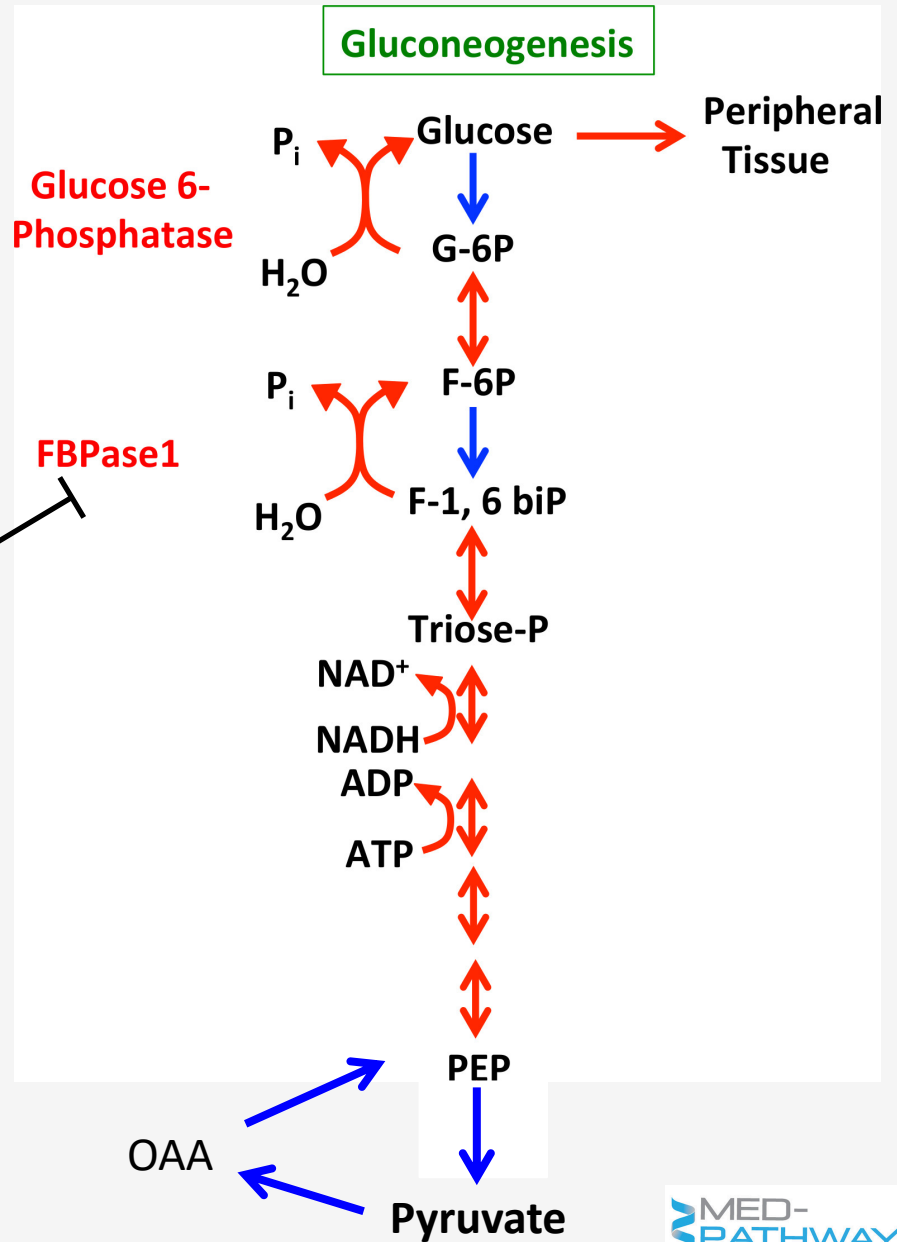


Metabolic Acidosis

Lactic Acidosis

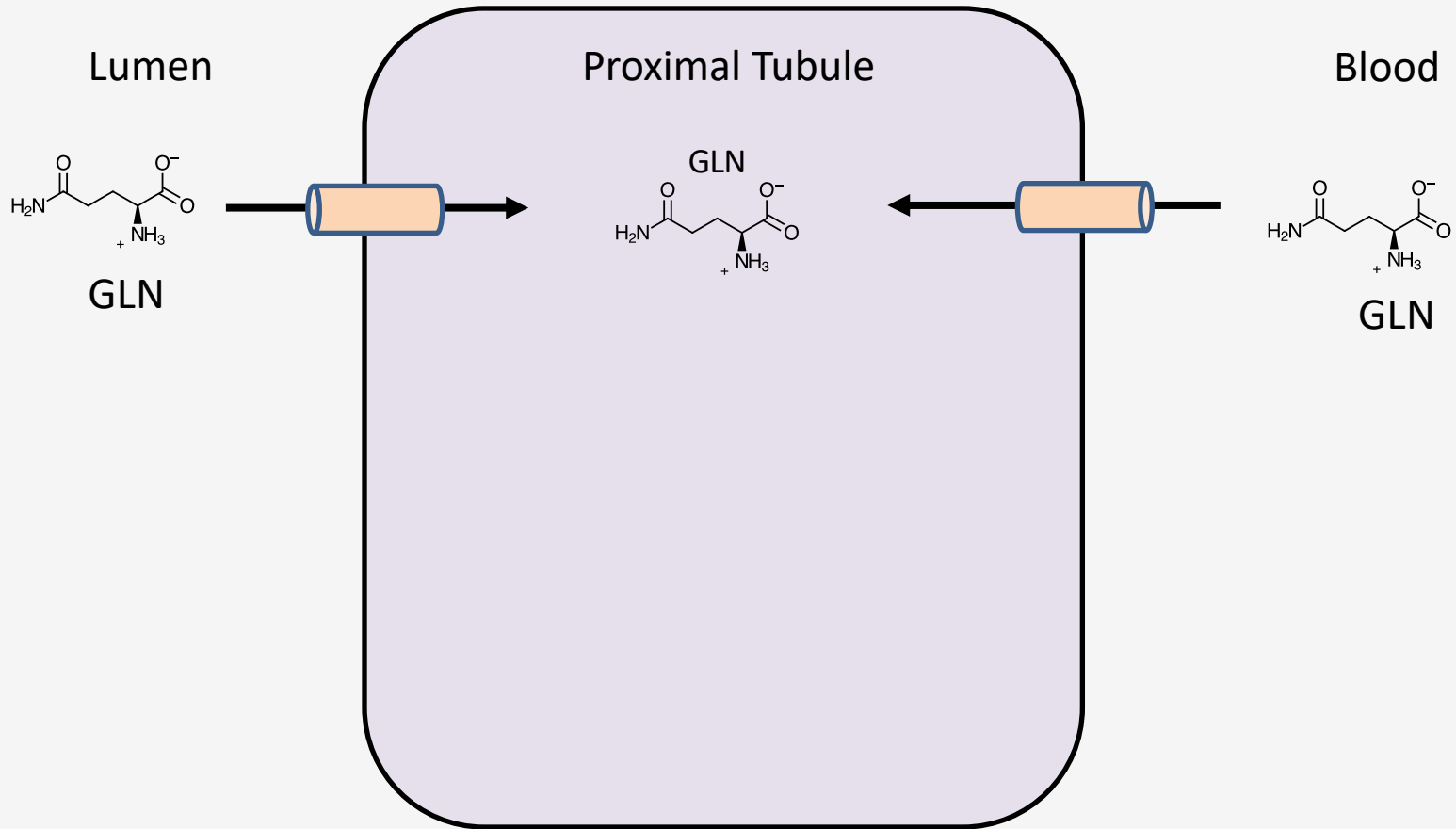


Metabolic Acidosis

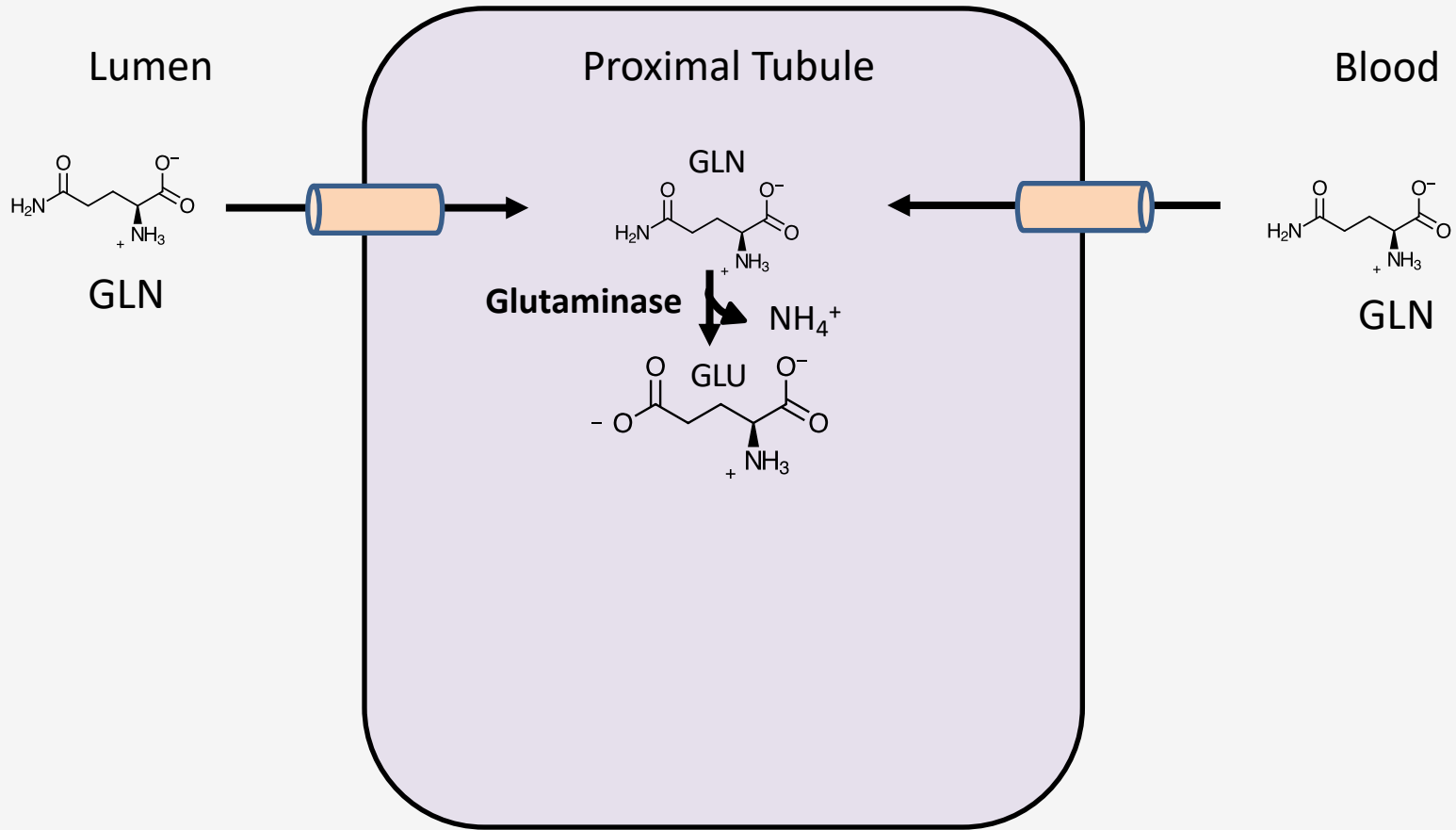


Metabolic Acidosis: The Kidney

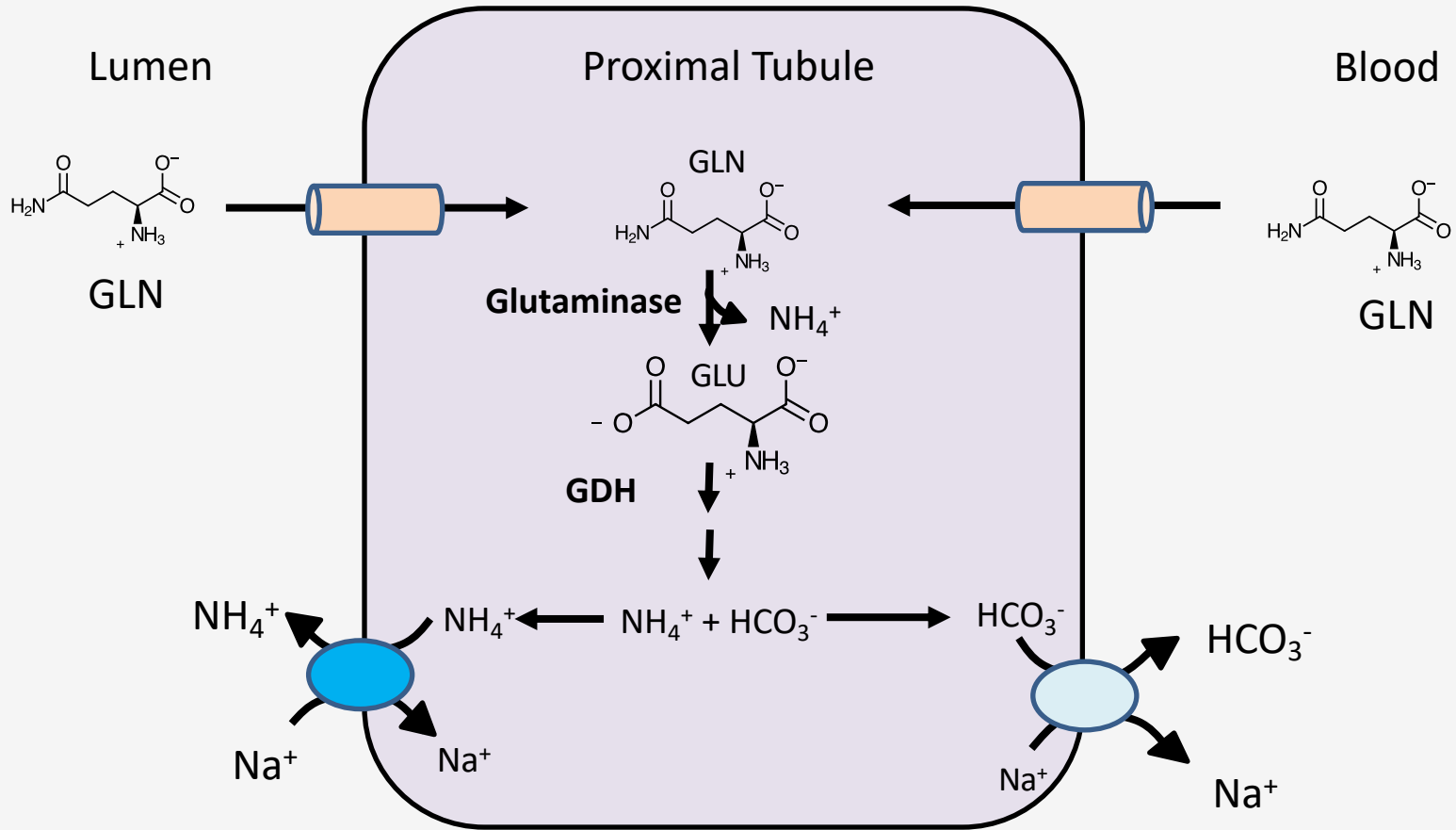
Physiological Goal: To Raise Blood pH levels



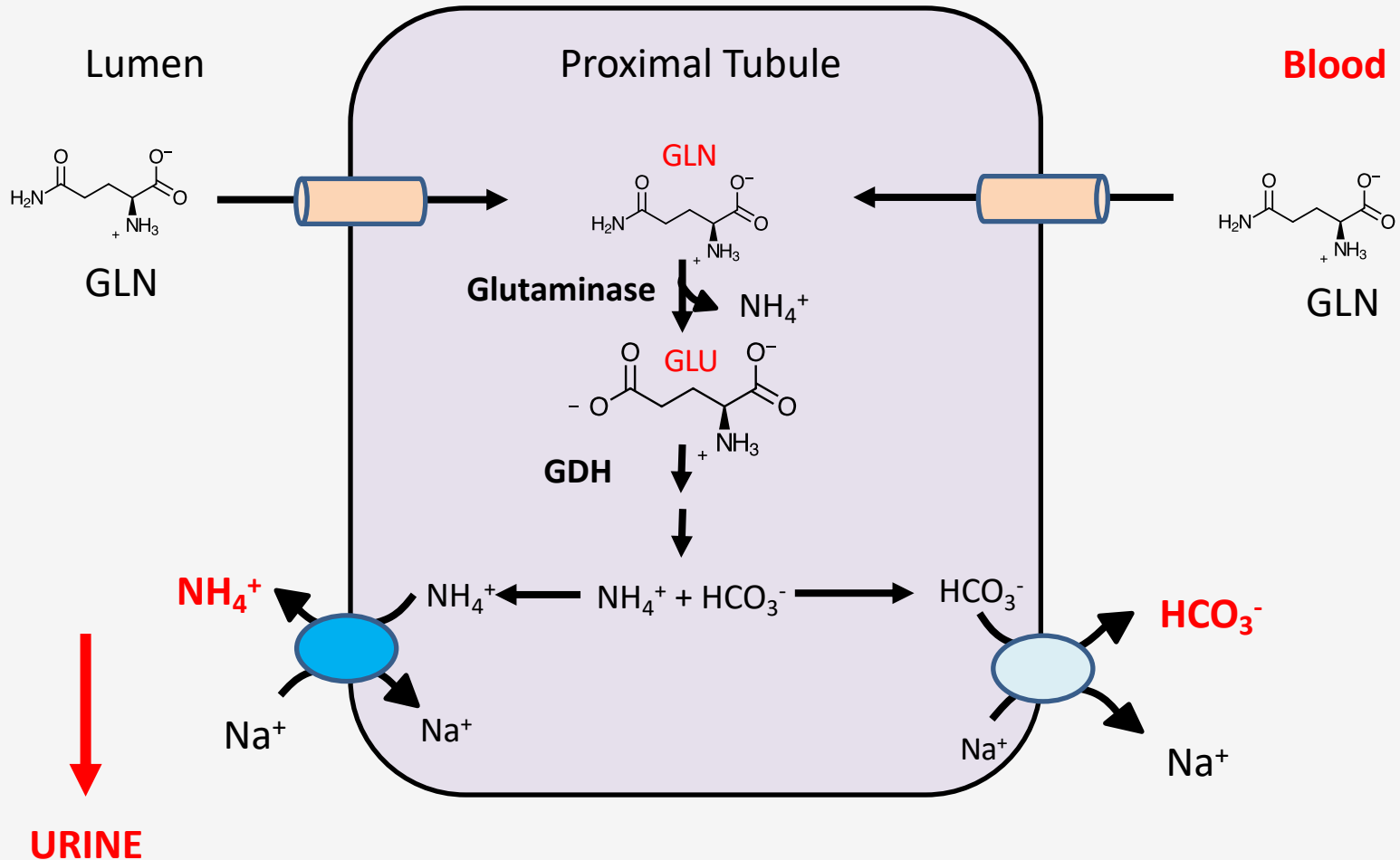
Metabolic Acidosis



Metabolic Acidosis

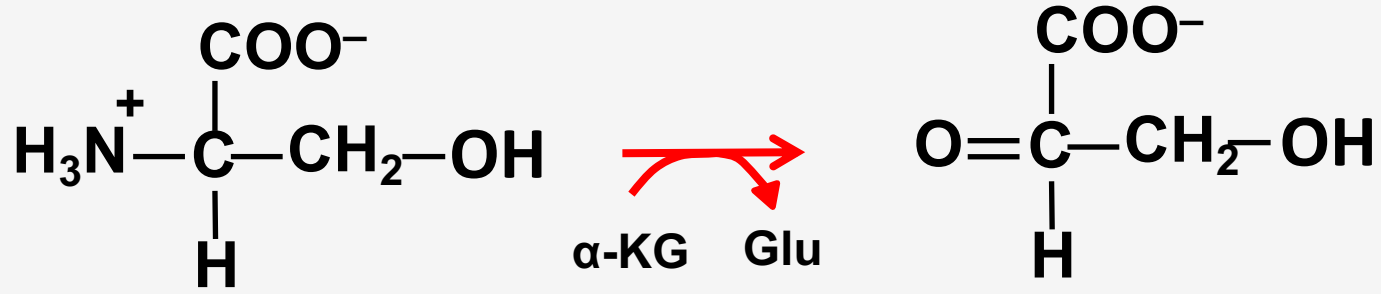


Metabolic Acidosis

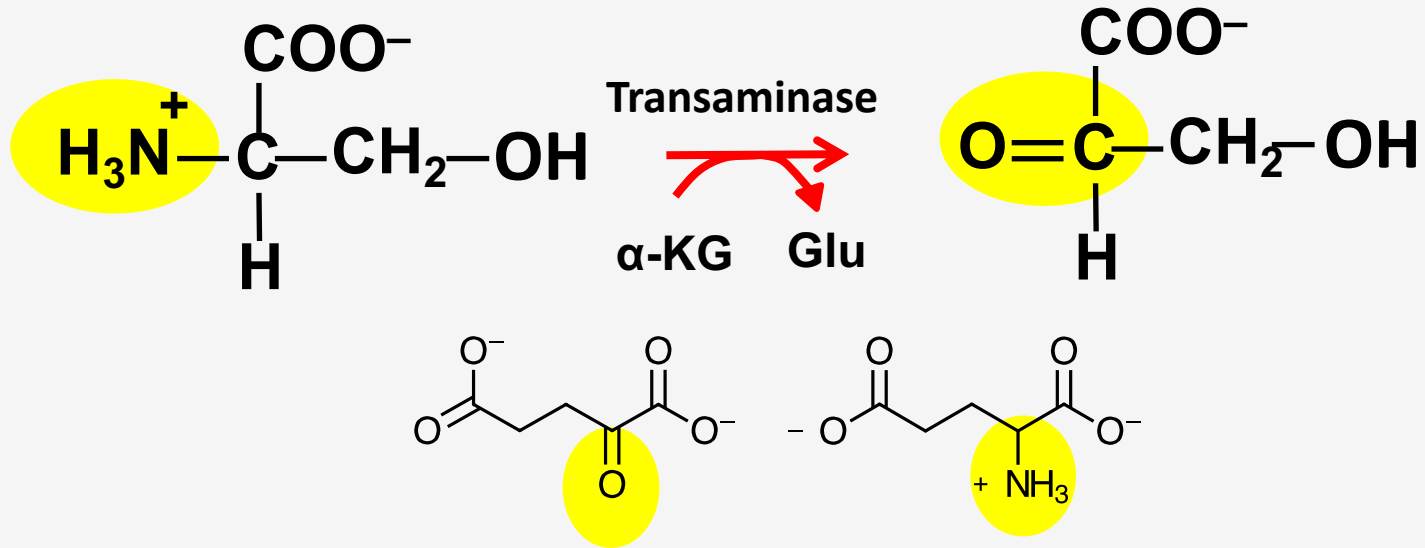


pH is linked to nitrogen balance

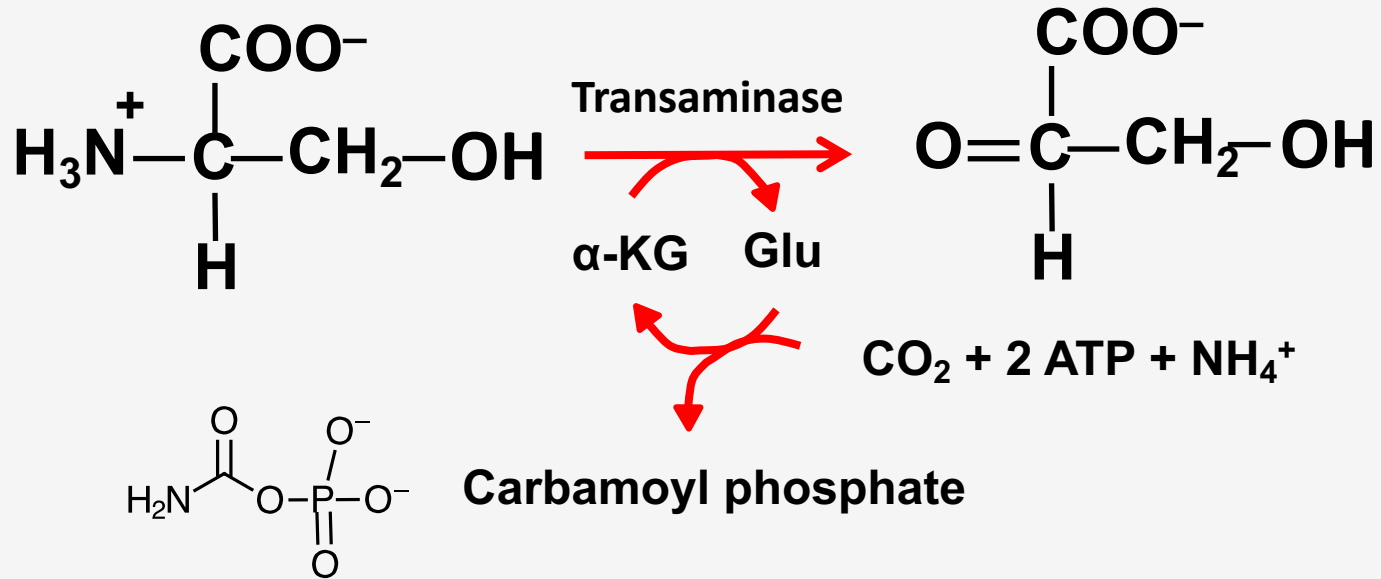
Nitrogen Balance & pH



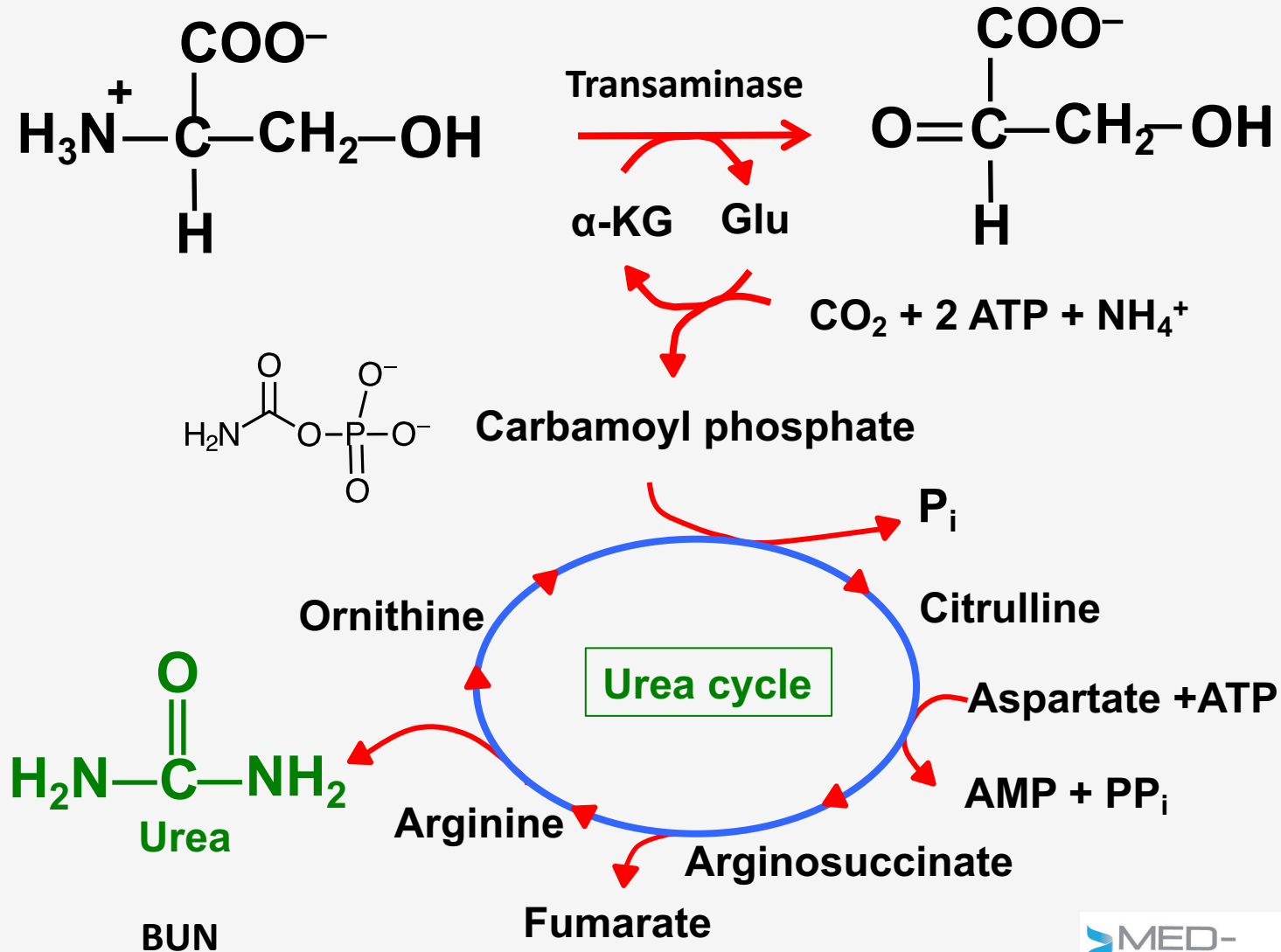
Nitrogen Balance & pH



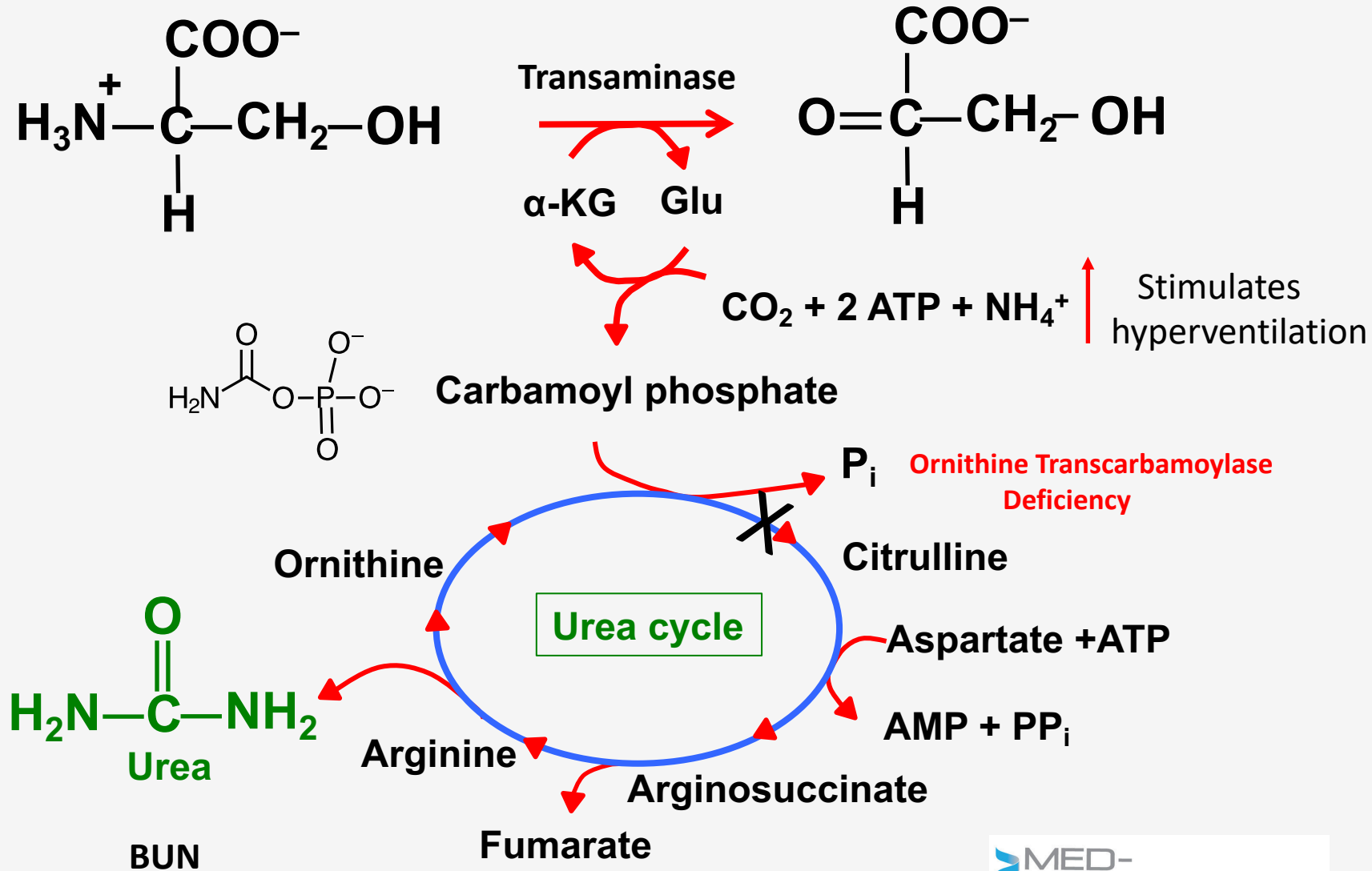
Nitrogen Balance & pH



Nitrogen Balance & pH



Nitrogen Balance & pH



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