

- **Blood Buffering**

- Our blood-buffering system is much more effective at buffering against added H^+ than added OH^- because there is more of the conjugate base (bicarbonate: carbonic acid = 20:1)

- **Picking a Buffer**

- **Q1: You need to make a buffer solution at pH = 7.40. In order to make the best buffer possible (i.e., effectively buffers against both H^+ and OH^-), which would be the best choice?**

When $[A^-] = [HA]$, the buffer is optimal.

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$pH = pK_a + \log (1)$$

So pH = pK_a

- When challenge with 0.5 mL of 1M HCl, 1M, 0.1M, 0.01M Tris all had the same initial pH ($[HA] / [A^-]$ was 1:1 for all pH=8)
 - Initial pH is a result of $[A^-]/[HA]$ ratio
 - Buffer capacity decreased with concentrations of A^- and HA
- A good buffer has a pK_a close to the desired pH.
- Excess acid can be exhaled as CO₂; excess base excreted as bicarb.
- Rule of thumb: Choose a buffer with a pK_a within 1 pH unit of the desired pH.
 - Buffer at pH = 9.0, choose boric acid, pK_a = 9.24
 1. Need conjugate base, borate (sodium borate)
 2. Add NaOH (strong base) to boric acid to make borate
$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$9.0 = 9.24 + \log \frac{[A^-]}{[HA]}$$

Solve for $[A^-] : [HA]$

- **Henderson–Hasselbalch equation**

- **Calculating pH**

- What is the pH of a buffer that is 0.12 M in lactic acid and 0.10 M in sodium lactate? K_a for lactic acid is 1.4×10^{-4} .

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$= -\log (1.4 \times 10^{-4}) + \log \frac{0.10 M}{0.12 M}$$

$$= 3.85 + (-0.08) = 3.77$$

- **Calculating ratio of $[A^-]/[HA]$**

- To make a buffer solution at pH = 7.40, what ratio of $[A^-]/[HA]$ do you need in your phosphate buffer? pK_a = 7.21.

$$pH = pK_a + \log \frac{[A^-]}{[HA]}$$

$$7.40 = 7.21 + \log \frac{[A^-]}{[HA]}$$

$$0.19 = \log \frac{[A^-]}{[HA]} \quad \text{so } \frac{[A^-]}{[HA]} = 1.5$$

○ **Calculating pK_a**

- A solution of 0.225 M NaNO₂ and 1.0 M HNO₂ has a pH = 2.51. What is the pK_a of HNO₂?

$$\text{pH} = \text{pK}_a + \log \frac{[A^-]}{[HA]}$$

$$2.51 = \text{pK}_a + \log \frac{(0.225)}{(1.0)}$$

$$2.51 = \text{pK}_a - 0.65 \quad \text{so } \text{pK}_a = 3.16$$

● **Adding a Strong Acid or Base to a Buffer**

- Adding a strong acid/base converts the weak base/acid to its conjugate acid/base.
- Find [HA] and [A⁻] when the added acid/base reacts.
- Use the Henderson–Hasselbalch equation to find pH.