

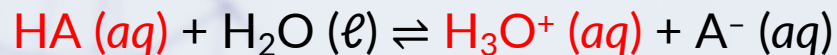


EXPERIMENT 8 (DAY 1)

Acids, Bases, & Buffers

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ACIDS
PROTON
DONORS

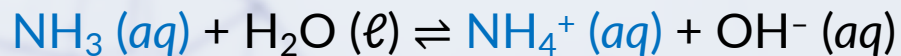


If $K_a > 1$, the acid is **strong**
examples: HCl , HNO_3 , ...

If $K_a < 1$, the acid is **weak**

If $K_a \ll 1$, the acid is **very weak**
examples: CH_3COOH , HCOOH , ...

BASES
PROTON
ACCEPTORS



If $K_b > 1$, the base is **strong**
examples: NaOH, KOH, Ca(OH)₂ ...

If $K_b < 1$, the base is **weak**

If $K_b \ll 1$, the base is **very weak**
examples: CH₃COO⁻, any conjugate base

AUTOIONIZATION OF WATER

When an **ACID** or **BASE** or **BOTH** are added to water...



Three simultaneous equilibria will automatically establish.



In many situations, only one of the three equilibria will dominate over the other two.

CONJUGATE ACID-BASE PAIR

$$K_a \times K_b = K_w = 1 \times 10^{-14}$$

If HA is strong, then A⁻ is very weak.

If HA is (medium) weak, then A⁻ is (medium) weak.

If HA is very weak, then A⁻ is strong.

ACID-BASE TITRATIONS

pH = $-\log [H^+]$ vs. volume NaOH added

Steep rise around the EQUIVALENCE POINT – the point where the moles of NaOH added equals the initial moles of (a monoprotic) acid.

Indicator chosen must change color within this steep rise.

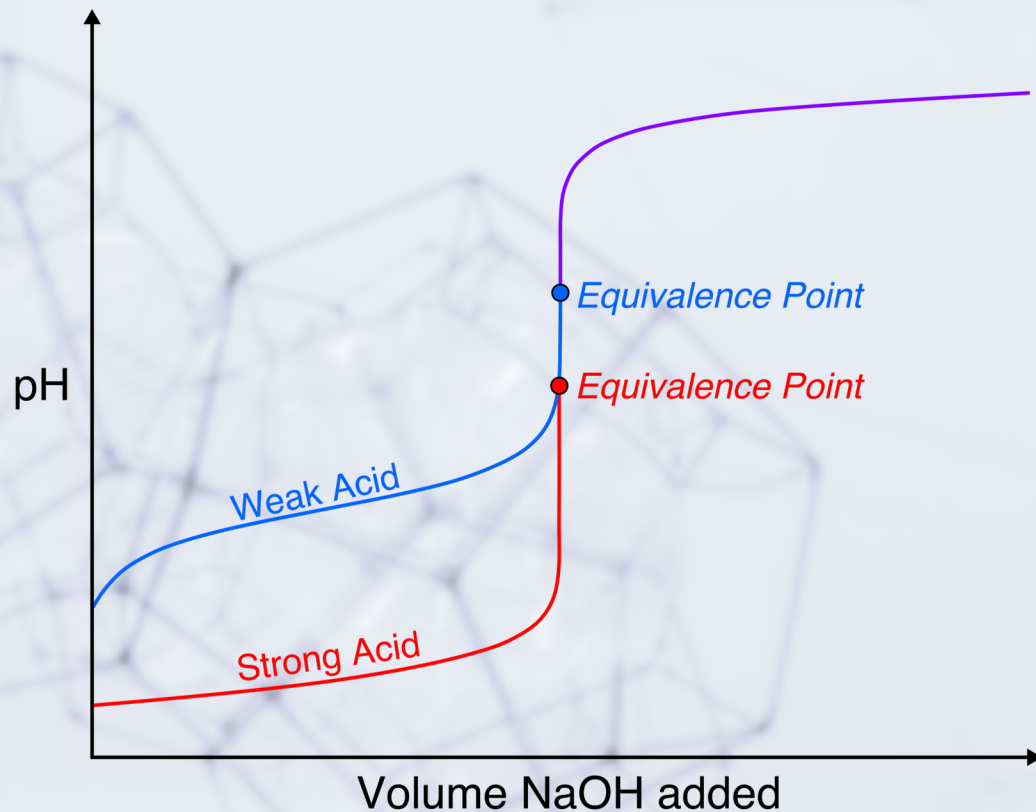
END POINT is the point where there is a distinct visual change in color.

What would the pH be at the EQUIVALENCE POINT in a strong **ACID**-strong **BASE** titration?

pH = 7

What would the pH be at the EQUIVALENCE POINT in a weak **ACID**-strong **BASE** titration?

pH > 7



BUFFERS

Consists of a weak **ACID**-conjugate **BASE** pair.
They are essential in all biochemical processes.

Buffers have the ability to *resist* change in pH when a small amount of a strong **ACID** or **BASE** is added.

CALCULATING THE pH OF A BUFFER

$$[\text{H}^+]_{\text{buffer}} = K_a \times \frac{[\text{HA}]_{\text{eq}}}{[\text{A}^-]_{\text{eq}}} \approx K_a \times \frac{[\text{HA}]_{\text{initial}}}{[\text{A}^-]_{\text{initial}}}$$

NOTES

1. The pH electrode is a delicate item, so handle it with care.
To work well, it needs to be well-immersed in the solution.
2. Plot both pH and its first derivative ($\partial\text{pH}/\partial V$).
→ Use the latter to judiciously choose NaOH increments.
3. In the vicinity of the EQUIVALENCE POINT, you must use very small increments of base (one drop at a time).
If not, you will not be able to determine the EQUIVALENCE POINT accurately.
4. Part 3: Either do it today or next week.
5. No lab report due next week.