

LECTURE TOPICS:

TODAY'S CLASS: continue Ch.17

NEXT CLASS: continue Ch.17

(1)

17.4 Equilibrium constants for acids & bases

For eqm involving reaction of an ACID with H_2O : " K_a "

$$K = \frac{[A^-][H_3O^+]}{[HA][H_2O]}$$

$[H_2O]$ constant since dilute solution \Rightarrow not included..
(in advanced treatment, value $\rightarrow 1$ & disappears...)

$$K_a = \frac{[A^-][H_3O^+]}{[HA]}$$

K_a = "Acid
Dissociation
Constant"
Larger K_a for
stronger acids

$$pK_a = -\log(K_a)$$

Smaller pK_a for
stronger acids

Acids with $K_a > 1$ described as STRONG acids
Acids with $K_a < 1$ described as WEAK acids

(2)

K_a & pK_a : a way to quantify acid strength...

See K&T
huge Table 17.3
Zumdahl's table ↓

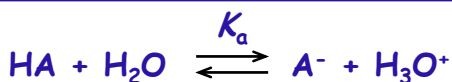


TABLE 14.2 Values of K_a for Some Common Monoprotic Acids

Formula	Name	Value of K_a^*	pK_a	
HSO_4^-	Hydrogen sulfate ion	1.2×10^{-2}	1.92	↑ Increasing acid strength
$HClO_2$	Chlorous acid	1.2×10^{-2}	1.92	
$HC_2H_2ClO_2$	Monochloroacetic acid	1.35×10^{-3}	2.87	
HF	Hydrofluoric acid	7.2×10^{-4}	3.14	
HNO_2	Nitrous acid	4.0×10^{-4}	3.40	
$HC_2H_3O_2$	Acetic acid	1.8×10^{-5}	4.74	
$[Al(H_2O)_6]^{3+}$	Hydrated aluminum(III) ion	1.4×10^{-5}	4.85	
HOCl	Hypochlorous acid	3.5×10^{-8}	7.46	
HCN	Hydrocyanic acid	6.2×10^{-10}	9.21	
NH_4^+	Ammonium ion	5.6×10^{-10}	9.25	
HOC_6H_5	Phenol	1.6×10^{-10}	9.80	

NOTE: K_a 's of "strong" acids too large to measure in H_2O
...because they ionize ~fully, we can't measure all []'s!

17.7 Calculations with equilibrium constants

Some typical scenarios:

1.) Use solution pH to find K_a (or K_b).

Q: When a solution of weak acid HA (or base B) of known initial concentration is prepared, the pH of the solution is _____. Find K_a (or K_b).

2.) Determine the pH of a solution of a weak acid (or base).

Q: When a solution of known initial concentration of a certain weak base (known K_b) is prepared, what is the pH?

3.) Determine the pH of a solution after an acid/base rxn.

Q: When a known volume of solution of acid HA is mixed with a known volume of strong base, what is the pH of the resulting solution?

(4)

APPLYING OUR EQM CALCULATIONS: (17.7)
Finding the pH of a weak acid solution...

Q: What is the pH of a typical vinegar solution?

→ vinegar = 5% v/v acetic acid

→ acetic acid: MM = 60.05 g/mol; d = 1.049 g/mL

APPROACH:

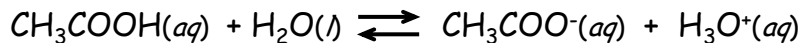
- LOOK UP: K_a of CH_3COOH ...
 - USE EQM ATTACK PLAN:
 - balance chemical equation
 - write eq'm expression for K_a
 - set up an ICE table
 - calculate $[\text{CH}_3\text{COOH}]_0$ in molarity & put into ICE table
 - solve for $[\text{H}_3\text{O}^+]$
- (5) ...& use $\text{pH} = -\log[\text{H}_3\text{O}^+]$

Q: What is the pH of a typical vinegar solution?

• vinegar = 5% v/v acetic acid → 5mL HA in approx. 100mL total

• acetic acid: $K_a = 1.82 \times 10^{-5}$; MM = 60.05 g/mol; d = 1.049 g/mL

→ $[\text{HA}]_0 = [(5\text{mL} \cdot 1.049\text{g/mL}) / 60.05\text{g/mol}] / (0.100\text{L}) \approx 0.873\text{ M}$



Initial	0.873 M	constant...	0	0
Change	-x		+ x	+ x
Eqm	0.873 - x		x	x

$$K_a = \frac{[\text{CH}_3\text{COO}^-][\text{H}_3\text{O}^+]}{[\text{CH}_3\text{COOH}]} \quad \rightarrow \quad 1.82 \times 10^{-5} = \frac{x^2}{0.873 - x}$$

$$\approx \frac{x^2}{0.873}$$

Check approximation:

$$\frac{x}{[\text{HA}]_{\text{initial}}} = \frac{0.00399}{0.873}$$

$$= 0.0047 < 5\% \therefore \text{ok}$$



$$x^2 = (0.873)(1.82 \times 10^{-5})$$

$$x = 3.99 \times 10^{-3} \text{ M} = [\text{H}_3\text{O}^+]$$

$$\text{pH} = -\log[\text{H}_3\text{O}^+]$$

$$= 2.4 \quad (1 \text{ SF})$$

Thus: % dissociation = ~0.5 %

Summary of acid strengths



PROPERTY	STRONG acid	WEAK acid
K_a value	very large	small
$\text{p}K_a$ value	very small	large
Equilibrium position	right	left
$[\text{H}^+]$ compared to $[\text{HA}]_0$	$[\text{H}^+] \approx [\text{HA}]_0$	$[\text{H}^+] \ll [\text{HA}]_0$
Strength of conjugate base A^- compared to H_2O	A^- is very WEAK	A^- is much STRONGER

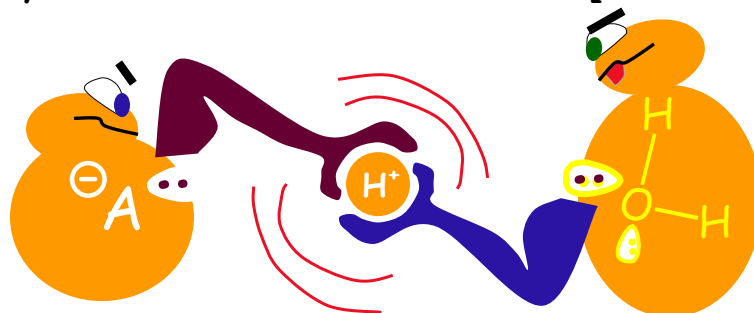
(7)

Where does the equilibrium lie? *i.e.*, How strong is HA?



IT'S ALL ABOUT THE BASE...

Really a **COMPETITION** between A^- & H_2O for the H^+ :



The stronger base wins!

Many molecules...distribution of energies...fwd & reverse rxn...

At EQM: more molecules of the stronger base are bonded to H^+

RELATING STRENGTHS OF ACIDS & THEIR CONJUGATE BASES

The numerical values shown refer to:

pK_a values of conj. acids (○) & pK_b values of conj. bases (●).

The labels

strong, weak & very weak

refer to the

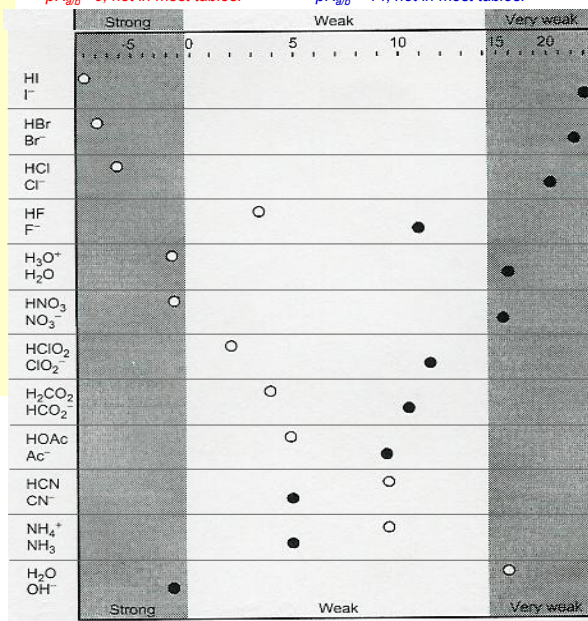
acid (○) OR base (●)

that appears

in that region of the figure.

STRONG acids & bases
react 100% with H₂O.
 $pK_{a/b} < 0$, not in most tables.

Very weak conj. bases & acids
don't react noticeably with H₂O.
 $pK_{a/b} > 14$, not in most tables.



Weak acids have weak conj. bases.
Both react noticeably with H₂O.

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ASSIGNED READINGS:

BEFORE NEXT CLASS:

Read: Ch.16 (all),
Ch.17 up to section 17.4 (to 6th Ed. p.809),
& 17.7 (to 6th Ed. p.824)

+ **WORK ON** Problems from Ch.16, Ch.17
including finding the pH of weak acid solutions
e.g., section 17.7 problems, #47-50

Extra information:

% DISSOCIATION: extent of ionization depends on [initial]

More concentrated More dilute

← Acid concentration

→ Percent dissociation

← H⁺ concentration

$$\% \text{ diss'n} = \frac{[A^-]}{[HA]_0} \times 100\%$$

For 5% acetic acid example:

$$\begin{aligned} \% \text{ diss'n} &= \frac{[3.99 \times 10^{-3} \text{ M}]}{[0.873 \text{ M}]} \times 100\% \\ &= 0.457 \% \\ &= 0.5 \% \text{ (1SF in data)} \end{aligned}$$

PROVE IT TO YOURSELF: On your own, use calculations to show that a **lower %** of the acetic acid molecules are ionized in 2.5 M CH₃COOH than in a 2.5 × 10⁻³ M solution, even though the more concentrated acid solution indeed has a lower pH.