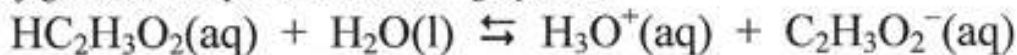


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STATION 1—CONJUGATE ACIDS & BASES

Identify the conjugate acid-base pairs in the following equation:



The conjugate acid of SO_4^{2-} is HSO_4^-

The conjugate base of HCO_3^- is CO_3^{2-}

The conjugate acid of NH_3 is NH_4^+

Which substance(s) are amphiprotic / amphoteric?

	H_3O^+	H_2O	NH_4^+	H_2CO_3	HSO_4^-
Conj acid	H_2O	OH^-	NH_3	HCO_3^-	SO_4^{2-}
Conj base	OH^-	O^{2-}	NH_2^-	CO_3^{2-}	-

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STATION 2—pH OF ACID SOLUTIONS

Calculate the pH of a 0.150 M solution of HCl. ($K_a = \text{very large}$)

STRONG ACID... $[\text{H}^+] = [\text{HCl}] = \underline{0.150 \text{ M}}$

$$\text{pH} = -\log[\text{H}^+] = -\log(0.150) = \underline{0.824}$$

Calculate the pH of a 0.150 M solution of HF. ($K_a = 7.2 \times 10^{-4}$)



i	.150	0	0
c	-x	+x	+x
e	.150 - x	x	x

ICE BOX PROBLEM
 assume $x \ll .150 \therefore (.150 - x) \approx (.150)$

$$K_a = \frac{[\text{H}^+][\text{F}^-]}{[\text{HF}]} = \frac{x^2}{.150} = 7.2 \times 10^{-4}$$

$$x = .0104 = [\text{H}^+]$$

$$\text{pH} = -\log[\text{H}^+] = -\log(.0104) = \underline{1.98}$$

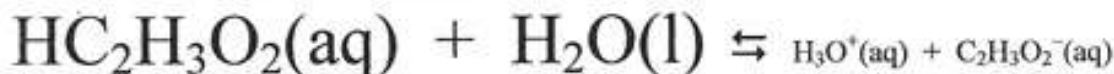
What is the % dissociation of HF in a 0.150 M solution?

$$\frac{[\text{H}^+]}{[\text{HF}]} \times 100 = \frac{.0104}{.150} \times 100 = \underline{6.9\%}$$

(I'm using 2 sig figs because $K_a = 7.2 \times 10^{-4}$)

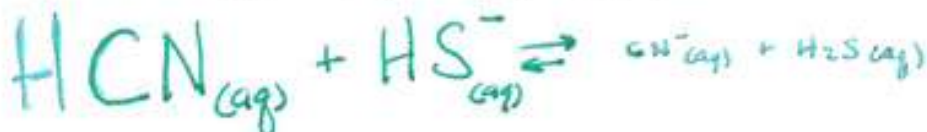
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EQUIL. FAVORS WEAKER ACID & WEAKER BASE **STATION 3—EQUILIBRIA**
 The following equation is written to represent relative concentrations in solution:



Which species is the stronger acid: $\text{HC}_2\text{H}_3\text{O}_2$ or H_3O^+ ?
 Weaker Acid: $\text{HC}_2\text{H}_3\text{O}_2$ (underlined)
 Weaker Base: H_2O (underlined)
 H_3O^+ is the BETTER DONOR
 we see it in the equation
 AFTER it HAS DONATED... H_2O

Knowing that H_2S is a stronger acid than HCN , re-write the following equilibrium to show which species are more concentrated in solution.



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STATION 4—pH & pOH

[H ⁺]	[OH ⁻]	pH	pOH	acidic or basic?
5.6×10^{-4}	1.8×10^{-11}	3.25	10.75	ACIDIC
2.5×10^{-2}	4.0×10^{-13}	1.60	12.40	ACIDIC
1.4×10^{-10}	7.1×10^{-5}	9.85	4.15	BASIC

$$[\text{H}^+] = 10^{-\text{pH}}$$

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

$$[\text{H}^+][\text{OH}^-] = 1 \times 10^{-14}$$

$$[\text{OH}^-] = 10^{-\text{pOH}}$$

$$\text{pOH} = -\log[\text{OH}^-]$$

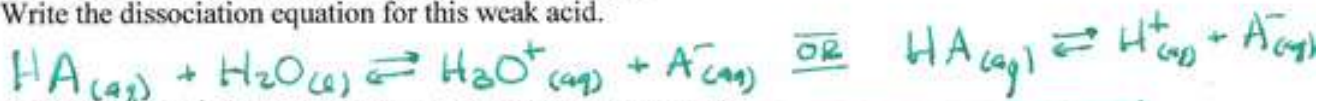
$$\text{pH} + \text{pOH} = 14$$

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STATION 5—ICE BOX

A 0.0300 M solution of the weak acid, HA, has a pH of 4.25.

a. Write the dissociation equation for this weak acid.



b. Calculate the $[H^+]$ for a 0.0300 M solution of this weak acid.

USE THE pH

$$pH = 4.25 \quad [H^+] = 10^{-pH} = \boxed{5.6 \times 10^{-5} M}$$

c. Calculate the K_a of this weak acid.

$$HA + H_2O \rightleftharpoons H_3O^+ + A^-$$

I	0.0300 M	0	0
C	-5.6×10^{-5}	$+5.6 \times 10^{-5}$	$+5.6 \times 10^{-5}$
E	$0.0300 - 5.6 \times 10^{-5}$	5.6×10^{-5}	5.6×10^{-5}

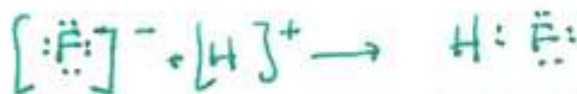
NOTE: DON'T USE "K" ... YOU KNOW THE VALUES.

$$K_a = \frac{[H_3O^+][A^-]}{[HA]} = \frac{(5.6 \times 10^{-5})^2}{(0.0300 - 5.6 \times 10^{-5})} = 1.056 \times 10^{-7} = \boxed{1.1 \times 10^{-7}}$$

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STATION 6—LEWIS & BRONSTED-LOWRY DEFINITIONS

Consider the reaction: $F^- + H_2O \rightleftharpoons HF + OH^-$



F⁻ would be a (circle all answers that apply):

Brønsted-Lowry acid

Brønsted-Lowry base

Lewis acid

Lewis base

F⁻ accepts an H⁺

F⁻ DONATES e⁻ pair to make bond

Consider the reaction: $BF_3 + NH_3 \rightarrow BF_3NH_3$

BF₃ would be a (circle all answers that apply):

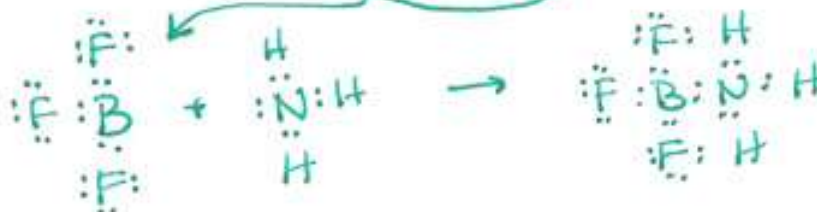
Brønsted-Lowry acid

Brønsted-Lowry base

Lewis acid

Lewis base

e⁻ pair acceptor

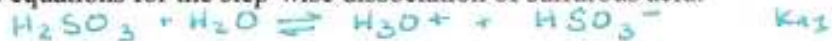


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STATION 7—DIPROTIC ACIDS

Consider the diprotic acid, sulfurous acid: H_2SO_3 $K_{a1} = 1.2 \times 10^{-2}$ $K_{a2} = 6.6 \times 10^{-8}$

a. Write the equations for the step-wise dissociation of sulfurous acid.



b. Calculate the pH of a 0.0200 M solution of H_2SO_3 .



I	.0200	0	0
C	-x	+x	+x
E	.0200 - x	x	x

In a solution of H_2SO_3 , all of the H_3O^+ 's come from the 1st step.

(b) ICE box using K_{a1}

oops! You must USE QUADRATIC
 $\boxed{\text{pH} = 1.97}$

c. What is the $[\text{SO}_3^{2-}]$ in a 0.0125 M solution of H_2SO_3 ?

$$K_{a2} = \frac{[\text{H}_3\text{O}^+][\text{SO}_3^{2-}]}{[\text{HSO}_3^-]} = 6.6 \times 10^{-8}$$

BECAUSE OF STEP 1, $[\text{H}_3\text{O}^+] \approx [\text{HSO}_3^-]$

$$\text{so } \frac{[\text{H}_3\text{O}^+][\text{SO}_3^{2-}]}{[\text{HSO}_3^-]} = \boxed{6.6 \times 10^{-8}}$$

$[\text{SO}_3^{2-}] = K_{a2}$
 MEMORIZE!

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STATION 8—SALT SOLUTIONS

For each solution, state whether it would be Acidic, Basic, or Neutral.

A NH_4Cl (NH_3)

N Na_2SO_4

B $\text{KC}_2\text{H}_3\text{O}_2$ ($\text{HC}_2\text{H}_3\text{O}_2$)

B CaF_2 (HF)

B MgSO_3 (H_2SO_3)

N KI

NOTE: THE ONLY conjugate acid we usually see is NH_4^+ (NH_3) or CH_3NH_3^+ (CH_3NH_2)

a. Write the equation for the equilibrium that exists in a solution of KCN.



$\text{CN}^- \dots \text{K}^+$ is a spectator (conjugate base of HCN)

b. Calculate the K_b for CN^- .

$$K_b = \frac{K_w}{K_a} = \frac{1 \times 10^{-14}}{6.2 \times 10^{-10}} = \boxed{1.6 \times 10^{-5}}$$

c. Calculate the pH of a 0.0200 M solution of KCN.

I	.0200	0	0
C	-x	+x	+x
E	.0200 - x	x	x

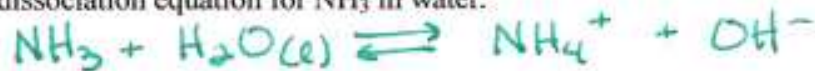
assume $x \ll .02$ etc.
 $x = [\text{OH}^-] = \sqrt{(0.0200)(1.6 \times 10^{-5})}$
 $= 5.68 \times 10^{-4}$
 $\text{pOH} = 3.245$
 $\text{pH} = 14 - \text{pOH} = \boxed{10.75}$

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STATION 9—WEAK BASES

NH_3 is a weak base with a $K_b = 1.8 \times 10^{-5}$.

- a. Write the dissociation equation for NH_3 in water.



- b. Calculate the pH of a 0.100 M solution of NH_3 . ICE BOX ... SHORT CUT?

$$x = [\text{OH}^-] = \sqrt{(0.100)(1.8 \times 10^{-5})} = 1.34 \times 10^{-3}$$

$$\text{pOH} = -\log [\text{OH}^-] = 2.87$$

$$\text{pH} = 14 - \text{pOH} = \boxed{11.13}$$

- c. Calculate the K_a for the conjugate acid of NH_3 .

$$K_a = \frac{K_w}{K_b} = \frac{1.0 \times 10^{-14}}{1.8 \times 10^{-5}} = \boxed{5.56 \times 10^{-10}}$$

- d. Write the equation for the equilibrium that exists in a solution of NH_4Cl .

