## <sup>132</sup> SALTS

- Compounds that result from the reaction of an acid and a base.

- Salts are strong electrolytes (completely dissociate in water) IF SOLUBLE (not all salts dissolve appreciably).

- Most ionic compounds are considered salts (they can be made by some reaction between the appropriate acid and base)

- Salts have acidic and basic properties! The ions that form when salts are dissolved can be acidic, basic, or neutral.

- Salts made from WEAK ACIDS tend to form BASIC solutions

- Salts made from <u>WEAK BASES</u> tend to form <u>ACIDI</u>C solutions

$$Na_2(O_3: Na_1O_3 \rightarrow 2Na^+ + CO_3^2)$$

Do any of these ions have acidic or basic properties?

 $Ma^{+}$ : neutral. Not a proton donor or a proton acceptor

 $(O_3^2 - BASIC, since it can accept protons to form the weak acid CARBONIC ACID in solution.$ 

$$H_2 (O_3 + 2H_2 O \rightleftharpoons 2H_3 O^{+} + CO_3^{-2}$$

$$ACID BASE$$

SALT OF A WEAK ACID

ex;  $NaC_2H_3O_2$  $NaA \longrightarrow Na^{+} + A^{-}$  The salt dissolves completely!

For this reaction to occur, HA MUST be stable in water. In other words, a weak acid.

+  $H_2 O \longrightarrow HA + OH^- \vdash \dots$  but the ionization of the salt's anion is an EQUILIBRIUM!

\_The anion is a BASE. It can accept a proton from water to form the weak (therefore stable as a molecule!) acid HA

$$K_b = \frac{[HA][OH^-]}{[A^-]}$$
 This is the base ionization constant for  $\overline{A}$ 

Since  $\vec{A}$  and HA are a conjugate pair, the ionization constants are related!

$$K_{W} = (K_{a,HA})(K_{b,A})$$
  
1.0 x10 14  
1.4 2 p Ka + p Kb

You will generally not find both the Ka AND Kb for a conjugate pair in the literature, since one can be easily converted to the other! xx: NH4CI  $\longrightarrow BH^+ + C [-]$  The salt dissociates completely!  $K_{a} = \frac{[B][H_{3}0^{+}]}{[R_{H}t]}$  Acid ionization constant for BH<sup>+</sup>  $Kw = (K_{a,BH^{+}})(K_{b,B})$ 1.0×10-16

Find the pH for salt solutions just like you would find pH for any other weak acid or weak base solutions. Only trick is to find out whether the salt is actually acidic or basic!  $O, IOOM NH_{4}C)$  ... Find the pH of the solution

Find the ions produced by the salt. Are they acids or bases?  $NH_4CI \longrightarrow NH_4^+ + CI^-$ 

 $NH_{4}^{4} + H_{2}O \rightleftharpoons \frac{NH_{3}}{L} + H_{3}O^{4}$ This is ammonia, a WEAK BASE. It's water-stable, so this reaction is viable.

$$CI + H_{LO} \stackrel{\rightarrow}{=} H_{CI} + OH^{-}$$
  
This is hydrochloric acid, a STRONG ACID. It's NOT water-stable, since it fully ionzes in water. This reaction won't go.

We need to solve the equilibrium of the ammonium ion.

$$NHy^{\dagger} + H_2 O \rightleftharpoons NH_3 + H_3 o^{\dagger} ; K_a = [NH_3] [H_3 o^{\dagger}]_{2}$$

$$[NH_4 f]_{2}$$

We can't find the Ka for ammonium ion in OpenStax Appendix H. We can, however, find the Kb of its conjugate (ammonia) in Appendix I:

$$K_{b_1 N H_3} = 1.8 \times 10^{-5}$$
  $K_a \times K_b = 1.0 \times 10^{-14}$   
 $K_a (1.8 \times 10^{-5}) = 1.0 \times 10^{-14}$   
 $K_{a_1 N H_4}^{+} = 5.56 \times 10^{-10}$ 

<sup>100</sup> 
$$MHy^{+} + H_20 \rightleftharpoons NH_3 + H_3 o^{+}$$
;  $K_a = [NH_3] [H_3o^{+}]_{=} = 5,56 \times 10^{-10}$   
Now that we have Ka, solve as a normal weak acid.  
Species [Jnitial]  $\Delta$  [Equilibrium]  
 $H_30^{+}$   $O$   $+X$   $X$   
 $NH_3$   $O$   $+X$   $X$   
 $NH_4^{+}$   $O,100 - X$   $O,100 - X$   
 $(x)(x) = 5.56 \times 10^{-10}$   $X = 7.45 \times 10^{-6} = [H_3o^{+}]$   
 $\downarrow \chi^{22} = 5.56 \times 10^{-10}$   $X = 7.45 \times 10^{-6} = 5.12$   
Compare:  
 $PH = -log_{10}(7.45 \times 10^{-6}) = 5.12$   
Compare:  
 $PH of 0.100 M nitrous acid: 2.17$   
 $PH of DI water: 7.00$ 

Not a B-L acid (no H+ to donate), nor is it basic ... the positive charge should REPEL protons!

 $(_2H_3O_2^-; Does have hydrogens, but also has a negative charge. Let's see if this is basic:$  $<math>(_2H_3O_2^- + H_2O \longrightarrow H(_2H_3O_2^- + OH^-)$ 

Acetic acid (weak!)

Solve equilibrium 
$$(2H_3O_7 + H_2O \Longrightarrow H(2H_3O_2 + OH^-;K_b = \frac{[H(2H_3O_2]Con-]}{[C_2H_3O_7]})$$

Look at OpenStax appendicies: Chart doesn't have acetate ion, but DOES have acetic acid!  $K_{A}, H(_{2}H_{3}O_{2} = 1, & \chi | O - S', (1, & \chi | O - S) \\ K_{b} = \leq , \leq 6 \times 10^{-10} \\ K_{b} = \leq , \leq 6 \times 10^{-10} \\ \frac{16}{16} \\ \frac{16}{16}$ 

$$\frac{[135]}{[C_{2}H_{3}O_{2}][0H^{-}]} = 5.56 \times 10^{-10} \quad (2H_{3}O_{2}^{-} + H_{2}O \Longrightarrow H(2H_{3}O_{2} + OH^{-})$$
Solve like any other base!
$$\frac{[Species]}{[OH^{-}]} \quad (2H_{1}H_{1H}] \land [Ce_{qui} | ib_{rium}]$$
Let "x" equal the change in hydroxide ion concentration.
$$\frac{H(2H_{3}O_{2})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + X) \times X$$

$$\frac{H(2H_{3}O_{2})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + X) \times X$$

$$\frac{(2H_{3}O_{2})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + X) \times X$$

$$\frac{(2H_{3}O_{2})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + 2H_{3}) \times X$$

$$\frac{H(2H_{3}O_{2})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + 2H_{3}) \times X$$

$$\frac{H(2H_{3}O_{2} + 2H_{3})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + 2H_{3}) \times X$$

$$\frac{H(2H_{3}O_{2} + 2H_{3})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + 2H_{3}) \times X$$

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$$\frac{H(2H_{3}O_{2} + 2H_{3})}{[CH_{3}O_{2}]} \quad (2H_{3}O_{2} + 2H_{3}) \times X$$

$$\frac{H(2H_{3}$$

For comparison:

0.100 M sodium acetate, pH = 8.87

0.100 M ammonia, pH = 11.13

0.100 M NaOH (strong base), pH = 13.00

The acetate ion is basic, but it's a very weak base!

$$O.100 M NaCl, Find pH$$
  
 $NaCl \rightarrow Na^+ + Cl^-$ 

 $N_{A}$  +  $\frac{1}{2}$  Not a B-L acid (no H+ to donate), nor is it basic ... the positive charge should REPEL protons!

CIT ; Not a B-L acid (no H+ to donate). Base? CIT + H20 Strong base, so this is NOT stable in water. CI- cannot hold onto the proton! Since neither ion acts as an acid or base, then water's own ionization controls pH. The pH is 7.00.

## <sup>141</sup> POLYPROTIC ACIDS

... what's special about phosphoric acid?

 $K_{a1} = 7.5 \times 10^{-3}$   $K_{a2} = 6.2 \times 10^{-8}$  $K_{a3} = 4.2 \times 10^{-13}$ 

(1) 
$$H_3 PO_4 + H_2 O \rightleftharpoons H_2 PO_4^- + H_3 O^+$$
  
(2)  $H_2 PO_4^- + H_2 O \rightleftharpoons H PO_4^{2-} + H_3 O^+$   
(3)  $H PO_4^{2-} + H_2 O \rightleftharpoons PO_4^{3-} + H_3 O^+$ 

Phosphoric acid has THREE acidic protons!

The first dissocation is dominant here, and for simple calculations of phosphoric acid in water, we will simply use the first ionization and ignore the other two.

Remember: This is a weak acid. It exists in water mostly as undissociated phosphoric acid molecules.

Solve the equilibrium of phosphoric acid's FIRST proton:  
H<sub>3</sub> PO<sub>44</sub> + H<sub>2</sub>O 
$$\rightleftharpoons$$
 H<sub>2</sub> PO<sub>47</sub> + H<sub>3</sub>O<sup>+</sup>; K<sub>6</sub> = 7.5 × 10<sup>-3</sup>  
K<sub>6</sub> =  $\frac{1}{2}$  H<sub>2</sub> PO<sub>47</sub> = 7.5 × 10<sup>-3</sup>  
K<sub>6</sub> =  $\frac{1}{2}$  H<sub>3</sub> PO<sub>4</sub> = 7.5 × 10<sup>-3</sup>  
Let "x' equal the change in hydronium ion concentration  
H<sub>3</sub>O<sup>+</sup> O + X ×  
H<sub>2</sub> PO<sub>4</sub> O + X ×  
H<sub>2</sub> PO<sub>4</sub> O + X ×  
(X)(X) = 7.5 × 10<sup>-3</sup>  
(V, 1)(X) = 7.5 × 10<sup>-3</sup>  
 $\frac{1}{2}$  × 2.5 × 10<sup>-3</sup>