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 WEAK BASES tend to form ACIDIC solutions

$$
\mathrm{Na}_{2} \mathrm{CO}_{3}: \mathrm{Na}_{2} \mathrm{CO}_{3} \rightarrow 2 \mathrm{Na}^{+}+\mathrm{CO}_{3}^{2-}
$$

Do any of these ions have acidic or basic properties?
$\mathrm{Na}^{+}$: neutral. Not a proton donor or a proton acceptor
$\mathrm{CO}_{3}{ }^{2-}$ : BASIC, since it can accept protons to form the weak acid CARBONIC ACID in solution.

$$
\mathrm{H}_{2} \mathrm{CO}_{3}{ }^{t} 2 \mathrm{H}_{2} \mathrm{O} \rightleftharpoons 2 \mathrm{H}_{3} \mathrm{O}^{t}+\mathrm{CO}_{3}^{-2}
$$

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SALT OF A WEAK ACID
ex: $\mathrm{NaC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$


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

$$
\mathrm{A}^{-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{H}^{-} \bar{A}+\mathrm{OH}^{-} \mathrm{H}
$$

$\qquad$ 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

$$
\left.K_{b}=\frac{[H A]\left[O H^{-}\right]}{\left[A^{-}\right]} \right\rvert\, \text {This is the base ionization constant for } A^{-}
$$

Since $\bar{A}$ and HA are a conjugate pair, the ionization constants are related! You will generally not find both

$$
\begin{aligned}
& K_{w}=\left(K_{a, H A}\right)\left(K_{b, A^{-}}\right) \mid \\
& 1.0 \times 10^{-14}=p r_{a}+p r_{b} \\
& \quad L_{1}=
\end{aligned}
$$ the Ka AND Kb for a conjugate pair in the literature, since one can be easily converted to the other!

## SALT OF A WEAK BASE

ex: $\mathrm{NH}_{4} \mathrm{Cl}$

$$
\begin{aligned}
& B H C I \longrightarrow \mathrm{BH}^{+}+\mathrm{Cl}^{-} \mathrm{F}^{\text {The salt dissociates completely! }} \\
& \mathrm{BH}^{+}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{~B}+\left.\mathrm{H}_{3} \mathrm{O}^{+}\right|_{\substack{ \\
\text { EQUILIBRIUM process! } \\
\\
\left.\mathrm{K}_{a}=\frac{[\mathrm{B}]\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]}{\left[\mathrm{BH}^{+}\right]} \right\rvert\, \\
\mathrm{K}_{w}=\left(\mathrm{K}_{a, \mathrm{BH}^{+}}\right)\left(\mathrm{K}_{b, B}\right) \\
1.0 \times 10^{-14}}} ^{\text {Acid ionization constant for } \mathrm{BH}^{+}}
\end{aligned}
$$

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!
$0.100 \mathrm{M} \mathrm{NH}_{4} \mathrm{Cl}$... Find the pH of the solution

$$
\mathrm{NH}_{4} \mathrm{Cl} \rightarrow \mathrm{NH}_{4}^{+}+\mathrm{Cl}^{-}
$$

Acidic, basic, or neutral salt?
This is the WEAK BASE ammonia. Stable

$$
\begin{array}{ll}
\mathrm{NH}_{4}^{+}: & \mathrm{NH}_{4}^{+}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{+} \\
\mathrm{Cl}^{-}: & \mathrm{Cl}^{-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HCl}+\mathrm{OH}^{-} X
\end{array}
$$



This is a STRONG ACID, which does not exist as a stable molecule in water.
The conjugate of a strong acid or base is NEUTRAL - does no $\dagger$ affect pH!
$\mathrm{NH}_{4}{ }^{+}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{+} \begin{aligned} & \text { This equilibrium affects the } \mathrm{pH} \text {, so it is the } \\ & \text { equilibrium we'll need to solve to find } \mathrm{pH} \text { ! }\end{aligned}$ equilibrium well need to solve to find pH !

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$$
\begin{aligned}
& \mathrm{NH}_{4}^{+}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{NH}_{3}+\mathrm{H}_{3} \mathrm{O}^{+} \\
& \mathrm{Ka}_{1} \mathrm{NH}_{4}{ }^{+}=\frac{\left[\mathrm{NH}_{3}\right]\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]}{\left[\mathrm{NH}_{4}^{+}\right]}
\end{aligned}
$$

We can get Ka for the acid from Kb for the base!

$$
\begin{aligned}
& K_{0, N H_{3}}=1.8 \times 10^{-s} \quad \overline{p A-14} \\
& K_{a} \times h_{b}=1.0 \times 10^{-14} \\
& \text { So, } h_{a}=5.56 \times 10^{-10} \quad(\text { forNHy+ })
\end{aligned}
$$

| Species | [Initial] | $\triangle$ | [Equilibrium] |
| :--- | :---: | :---: | :---: |
| $\mathrm{H}_{3}{ }^{+}$ | 0 | $+X$ | $x$ |
| $\mathrm{NH}_{3}$ | 0 | $+X$ | $x$ |
| $\mathrm{NH}^{+}$ | 0.100 | $-x$ | $0.100-x$ |

$$
\begin{aligned}
{\left[430^{+}\right] } & =7.46 \times 10^{-6} \\
p H & =5.13
\end{aligned}
$$

Compare:
$\mathrm{pH}=1.00$ for 0.100 M strong acid $\mathrm{pH}=2.16$ for 0.100 M nitrous acid $\mathrm{pH}=7.00$ for distilled water
$\mathrm{O} .100 \mathrm{M} \mathrm{NaC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$, Find pH

$$
\mathrm{NaC}_{2} \mathrm{H}_{3} \mathrm{O}_{2} \rightarrow \mathrm{Na}^{+}+\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}
$$

Check the ions formed to see if they have acidic or basic properties:
$\mathrm{Na}^{+}$: Cannot be B-L acid (no H), and not likely to be B-L base, since it's a simple positively charged metal ion.
$\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{2}$ : Has protons, but also a negative charge - so it might be a more likely proton acceptor rather than a donor.

$$
\begin{aligned}
& \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+\mathrm{OH}^{-} \\
& \text {Acetic acid is a WEAK ACID and stable } \\
& \text { in water, so the acetate ion CAN function } \\
& \text { as a base. } \\
& \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+\mathrm{OH}^{-}
\end{aligned}
$$

$$
\begin{aligned}
& K_{a_{1} H_{2} H_{3} O_{2}}=1.7 \times 10^{-5} ; K_{a} \times K_{b}=1.0 \times 10^{-14}, 50 k_{b}=5.88 \times 10^{-10}
\end{aligned}
$$

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$$
\begin{aligned}
& \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}+\mathrm{OH}^{-} \\
& \mathrm{K}_{\mathrm{b}_{1} \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}}=\frac{\left[\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}\right]\left[\mathrm{OH}^{-}\right]}{\left[\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}\right]}=5.88 \times 10^{-10}
\end{aligned}
$$

| Species | [Initial] | $\Delta$ | [Equilibrium] |
| :--- | :---: | :---: | :---: |
| $\mathrm{OH}^{-}$ | 0 | $+X$ | $X$ |
| $\mathrm{HC}_{2} \mathrm{H}_{3} \mathrm{O}_{2}$ | 0 | $+x$ | $x$ |
| $\mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}$ | 0.100 | $-X$ | $0.100-x$ |

$$
\left.\frac{x^{2}}{0.100-x}=5.88 \times 10^{-10} \right\rvert\, x=7.67 \times 10^{-6}=\left[\mathrm{OH}^{-}\right]
$$

Calculate POH first...

$$
\begin{aligned}
P O H & =-\log _{10}\left(7.67 \times 10^{-6}\right) \\
& =5.12
\end{aligned}
$$

Convert to pH

$$
\begin{aligned}
& p H+p O H=14.00 \\
& p H+5.12=14.00 \\
& p H=8.88
\end{aligned}
$$

Compare:
$\mathrm{pH}=7.00$ for pure distilled water
$\mathrm{pH}=11.13$ for 0.100 M ammonia
$\mathrm{pH}=13.00$ for 0.100 M strong base
0.100 M NaCl , Find PH

$$
\mathrm{NaCl} \rightarrow \mathrm{Na}^{+}+\mathrm{Cl}^{-}
$$

Check the ions formed to see if they have acidic or basic properties:
$\mathrm{Na}^{+}$; Cannot be B-L acid (no H), and not likely to be B-L base, since it's a simple positively charged metal ion.
$\mathrm{Cl}^{-}$: Cannot be B-L acid (no H), but can it accept protons?
 does not exist readily in water in the molecular form.
... so chloride ion will not function as a base and will be NEUTRAL.

Since neither sodium ion nor chloride ion affect the water equilibrium, the pH of the solution will be the same as that of pure distilled water: 7.00

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Find pH of $\mathrm{O}, 10 \mathrm{M} \mathrm{H}_{3} \mathrm{PO}_{4}$
... what's special about phosphoric acid?
(1) $\mathrm{H}_{3} \mathrm{PO}_{4}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{H}_{2} \mathrm{PO}_{4}^{-}+\mathrm{H}_{3} \mathrm{O}^{+}$

Phosphoric acid has THREE acidic protons!
(2)
(3)

$$
\left.\mathrm{HPO}_{4}^{2-}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{PO}_{4}^{3-}+\mathrm{H}_{3} \mathrm{O}^{+}\right]
$$

$$
\begin{aligned}
& K_{a_{1}}=6,9 \times 10^{-3} \\
& K_{a_{2}}=6.2 \times 10^{-8} \\
& K_{a_{3}}=4.8 \times 10^{-13}
\end{aligned}
$$

The first dissocation is dominant here, and for
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.

Solving the equilibrium of phosphoric acid's first proton:

$$
\begin{aligned}
& \mathrm{H}_{3} \mathrm{PO}_{4}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{H}_{2} \mathrm{PO}_{4}^{-}+\mathrm{H}_{3} \mathrm{O}^{+} \\
& \mathrm{Ka}=6.9 \times 10^{-3}=\frac{\left[\mathrm{H}_{2} \mathrm{PO}_{4}^{-}\right]\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]}{\left[\mathrm{H}_{3} \mathrm{PO}_{4}\right]}
\end{aligned}
$$

| Species | [Initial] | $\Delta$ | [Equilibrium] |
| :--- | :---: | :---: | :---: |
| $\mathrm{H}_{30^{+}}$ | 0 | $+x$ | $x$ |
| $\mathrm{H}_{2} \mathrm{PO}_{4}^{-}$ | 0 | $+x$ | $x$ |
| $\mathrm{H}_{3} \mathrm{PO}_{4}$ | 0.10 | $-x$ | $0.10-x$ |

$\frac{x^{2}}{0.10-x}=6.9 \times 10^{-3} \quad \begin{aligned} & \text { This time, let's use the quadratic. Were not as confident } \\ & x \ll 0.10 \text { as we were in the previous }\end{aligned}$ $x \ll 0.10$ as we were in the previous few examples.

$$
\begin{aligned}
& x^{2}=6.4 \times 10^{-4}-6.9 \times 10^{-3} x \\
& x^{2}+6.9 \times 10^{-3} x-6.9 \times 10^{-4}=0 \\
& a=1, b=6.9 \times 10^{-3}, c=-6.9 \times 10^{-4} \\
& x=2.3043 \times 10^{-2} \text { or }-2.9943 \times 10^{-2} \\
& \\
& =[430+] \\
& \rho H=1.64
\end{aligned}
$$

