- 9.2 a. solute=acetic acid, solvent=water b. solute=water, solvent=ethanol c. solute=oxygen, solvent=nitrogen
- 9.5 a. ionic is more soluble in water
 c. polar is more soluble in water
 d. nonpolar is more soluble in CCl₄
 d. nonpolar is more soluble in CCl₄
- 9.9 a. $KCl(aq) \rightarrow K^{+}(aq) + Cl^{-}(aq)$ b. $CaCl_{2}(aq) \rightarrow Ca^{2+}(aq) + 2Cl^{-}(aq)$ c. $K_{3}PO_{4}(aq) \rightarrow 3K^{+}(aq) + PO_{4}^{3-}(aq)$ d. $Fe(NO_{3})_{3}(aq) \rightarrow Fe^{3+}(aq) + 3NO_{3}^{-}(aq)$
- 9.12 a. mainly ions b. mainly molecules c. some ions, mainly molecules
- 9.22 a. unknown, possibly saturated b. saturated
- 9.24 You need to adjust the solute and solvent to 100 g of water to use the table.
 a. saturated (50. g KCl will not dissolve in 100. g water)
 b. saturated (200. g NaNO₃ will not dissolve in 100. g water)
 c. saturated (320. g sugar will not dissolve in 100. g water)
- 9.27 a. The solubility of sugar increases as the temperature increases.b. Gases tend to be less soluble as the temperature increases.c. Same as b.
- 9.31 5% (m/m) has 5 grams of solute per 100 g of solution while 5% (m/v) has 5 grams of solute per 100 mL of solution.

9.34 a.
$$\left(\frac{75 \text{ g}}{325 \text{ g}}\right)(100\%) = 23\% \text{ (m/m)}$$

b. $\left(\frac{2.0 \text{ g}}{22.0 \text{ g}}\right)(100\%) = 9.1\% \text{ (m/m)}$ (remember, mass of **solution**)
c. $\left(\frac{48.5 \text{ g}}{250.0 \text{ g}}\right)(100\%) = 19.4\% \text{ (m/m)}$

9.37 a.
$$\left(\frac{5.0 \text{ g}}{100 \text{ g}}\right) \left(\frac{50. \text{ g}}{1}\right) = 25 \text{ g}$$

b. $\left(\frac{4.0 \text{ g}}{100 \text{ mL}}\right) \left(\frac{1250 \text{ mL}}{1}\right) = 50. \text{ g}$

9.41 a.
$$\left(\frac{20. \text{ g}}{100 \text{ mL}}\right) \left(\frac{100. \text{ mL}}{1}\right) = 20. \text{ g}$$

b. $\left(\frac{20. \text{ g}}{100 \text{ mL}}\right) \left(\frac{100. \text{ mL}}{1 \text{ hr}}\right) \left(\frac{12 \text{ hr}}{1}\right) = 2.4 \times 10^2 \text{ g}$

9.47 a.
$$\left(\frac{2.00 \text{ mol glucose}}{4.00 \text{ L solution}}\right) = 0.500 \text{ M glucose}$$

b. $\left(\frac{4.00 \text{ g KOH}}{1}\right) \left(\frac{1 \text{ mol}}{56.1 \text{ g}}\right) \left(\frac{1}{2.00 \text{ L solution}}\right) = 0.0357 \text{ M KOH}$
c. $\left(\frac{5.85 \text{ g NaCl}}{1}\right) \left(\frac{1 \text{ mol}}{58.5 \text{ g}}\right) \left(\frac{1 \text{ ml}}{0.001 \text{ L}}\right) \left(\frac{1}{400. \text{ mL solution}}\right) = 0.250 \text{ M NaCl}$

9.55 a.
$$(6.0 \text{ M})(2.0 \text{ L}) = (C_2)(6.0 \text{ L})$$
 $C_2 = \frac{(6.0 \text{ M})(2.0 \text{ L})}{(6.0 \text{ L})} = 2.0 \text{ M}$

b.
$$(12 \text{ M})(0.50 \text{ L}) = (C_2)(3.0 \text{ L})$$
 $C_2 = \frac{(12 \text{ M})(0.50 \text{ L})}{(3.0 \text{ L})} = 2.0 \text{ M}$

c.
$$(25 \% \text{ m/v})(10.0 \text{ mL}) = (C_2)(100.0 \text{ mL})$$

 $C_2 = \frac{(25 \% \text{ m/v})(10.0 \text{ mL})}{(100.0 \text{ mL})} = 2.5 \% \text{ (m/v)}$
d. $(15 \% \text{ m/v})(50.0 \text{ mL}) = (C_2)(250 \text{ mL})$
 $C_2 = \frac{(15 \% \text{ m/v})(50.0 \text{ mL})}{(250 \text{ mL})} = 3.0 \% \text{ (m/v)}$

9.66 a. A b. A c. B

10.1	a. acid	b. acid	c. acid	d. base	e. both
10.8	a. $\text{CO}_3^{2-}(\text{aq})$ base	,	HCO ₃ ⁻ (aq)+ acid	-OH⁻(aq) base	
	b. $H_2SO_4(aq) + H_2O(\ell) \rightarrow HSO_4^{-}(aq) + H_3O^{+}(aq)$ acid base base acid				
10.9	a. F ⁻	b. OH⁻	c. HCO ₃ ⁻	d. SO_4^{2-}	
10.12	a. HSO ₄ -	b. HCN	c. H ₂ O	d. HClO ₂	
10.13	a. $H_2CO_3(aq) + H_2O(\ell) \rightleftharpoons HCO_3^-(aq) + H_3O^+(aq)$ acid base				
	conjugate base of $H_2CO_3 = HCO_3^-$ conjugate acid of $H_2O = H_3O^+$				
	b. $NH_4^+(aq) + H_2O(\ell) \rightleftharpoons NH_3(aq) + H_3O^+(aq)$ acid base				
	conjugate base of $NH_4^+ = NH_3$ conjugate acid of $H_2O = H_3O^+$				
	c. $HCN(aq) + NO_2^{-}(aq) \rightleftharpoons CN^{-}(aq) + HNO_2(aq)$ acid base				
	conjugate base of HCN = CN^{-} conjugate acid of NO_{2}^{-} = HNO_{2}				
10.18	a. HCO ₃ ⁻	b. H ₂ O	c. H ₂ CO ₃		

- 10.25 In pure water, any water molecules that dissociate will produce **one** hydronium and **one** hydroxide ion.
- 10.26 It's the ion-product of hydronium and hydroxide ions concentrations in the solution. At 25 °C it's 1.0×10^{-14} .

- 10.30 a. basic, $[H_3O^{1+}] < 1.0 \times 10^{-7} M$ b. acidic, $[H_3O^{1+}] > 1.0 \times 10^{-7} M$ c. acidic, $[H_3O^{1+}] = 2.0 \times 10^{-3} M$, $[H_3O^{1+}] > 1.0 \times 10^{-7} M$ d. basic, $[H_3O^{1+}] = 2.2 \times 10^{-13} M$, $[H_3O^{1+}] < 1.0 \times 10^{-7} M$
- 10.32 Plug into $K_w = [H_3O^+][OH^-]$ and solve for $[H_3O^+]$ a. $[H_3O^+] = 1.0 \times 10^{-8}$ M b. $[H_3O^+] = 2.0 \times 10^{-4}$ M c. $[H_3O^+] = 5.0 \times 10^{-7}$ M d. $[H_3O^+] = 4.8 \times 10^{-12}$ M
- 10.34 Plug into $K_w = [H_3O^+][OH^-]$ and solve for $[OH^-]$ a. $[OH^-] = 1.0 \times 10^{-2}$ Mb. $[OH^-] = 1.7 \times 10^{-11}$ Mc. $[OH^-] = 1.0 \times 10^{-5}$ Md. $[OH^-] = 1.9 \times 10^{-13}$ M
- 10.39 a. pH=4.0 b. 8.5 c. pH=9.0 d. pH=3.4 e. pH=7.2 f. pH=10.9

10.54 a. No. It's just an acid.

- b. No. It's just a neutral salt.
- c. Yes. Acetic acid (HC₂H₃O₂, acid) and sodium acetate (C₂H₃O₂⁻, acetate is the conjugate base of acetic acid)
- d. No. Both are strong and you need a **weak** acid and its conjugate base or a **weak** base and it's conjugate acid.
- 10.55 a. (3) A buffer solution maintains the pH. So if a small amount of an acidic or basic solution is added to the buffer, the pH is not expected to change much.
 - b. (1, 2) An acid added to this buffer solution will convert the base, F⁻, to its conjugate acid, HF. A base added to this buffer solution will convert the acid, HF, to its conjugate base, F¹⁻. You need to have **both** present because the product is the conjugate of either the acid or the base.
 - c. The best answer for the given choices is (3). However, HF is also present.