

- (3 points) At a pressure of 1025 torr, the solubility of O<sub>2</sub> gas in water is 0.00175 M. If the pressure is increased to 25 atm, the solubility of O<sub>2</sub> should be **0.0324** M.
- (4 points) The heat of solution ( $\Delta H^\circ_{\text{soln}}$ ) of an ionic compound in water is approximately equal to the sum of the **lattice** energy of the crystalline solid and the **hydration** energy of the ions in solution.
- (2 points) In the **Transition State** Theory of chemical kinetics, a type of graph called a "reaction coordinate diagram" is used to illustrate Energy of Activation.
- (2 points) In water, soap molecules aggregate into small particles called **micelles**.
- (2 points) Three common examples of colloidal dispersions are: **paint, milk, soap, fog, etc.**
- (11 points) A kinetic study of the following gas-phase reaction gave the concentration vs initial rate data summarized below.  
 $\text{CH}_3\text{Cl} + 3 \text{Cl}_2 \longrightarrow \text{CCl}_4 + 3 \text{HCl}$

Expt	[CH <sub>3</sub> Cl]	[Cl <sub>2</sub> ]	initial rate (mole/L·sec)
(1)	0.250	0.150	0.0215
(2)	0.450	0.450	0.0670
(3)	1.350	0.150	0.1161
(4)	0.450	0.150	0.0387

Determine the **rate law** for this reaction. **Clearly SHOW** how you arrive at your answer. (It is not necessary to calculate the value of the rate constant.)

Compare expts 1 & 3 with constant [Cl<sub>2</sub>] to find rate  $\sim$  [CH<sub>3</sub>Cl]<sup>x</sup>

$$0.1161 / 0.0215 = (1.35 / 0.25)^x$$

$$5.4 = (5.4)^x \quad \therefore x = 1 \quad (1\text{st order in CH}_3\text{Cl})$$

Compare expts 2 and 4 with constant [CH<sub>3</sub>Cl] to find rate  $\sim$  [Cl<sub>2</sub>]<sup>y</sup>

$$0.0670 / 0.0387 = (0.450 / 0.150)^y$$

$$1.73 = 3^y \quad \therefore y = 1/2 \quad (1/2 \text{ order in Cl}_2)$$

Overall Rate Law: rate = k[CH<sub>3</sub>Cl][Cl<sub>2</sub>]<sup>1/2</sup>

7. (10 points) **SHOW ALL WORK.** A 0.725 g sample of an unknown polymer was dissolved in a suitable solvent and diluted with the solvent to make 250 mL of solution. The osmotic pressure of the solution was found to equal 1.35 torr at 20° C. Determine the molar mass of the polymer.

$$\Pi V = nRT$$

$$n = \Pi V / RT = (1.35 \text{ torr}) (1 \text{ atm} / 760 \text{ torr}) (0.250 \text{ L}) / (0.0821 \text{ L}\cdot\text{atm}/\text{mole}\cdot\text{K}) (293 \text{ K})$$

$$n = 1.846 \times 10^{-5} \text{ moles}$$

$$\text{molar mass} = (0.725 \text{ g}) / (1.846 \times 10^{-5} \text{ moles}) = 3.93 \times 10^4 \text{ g/mole}$$

8. A concentrated solution of  $\text{MgCl}_2(\text{aq})$  is 25.0 %  $\text{MgCl}_2$  by mass and has a density of 1.26 g/mL. The following problems all deal with this solution but they can be solved independently.  
[Molar masses:  $\text{H}_2\text{O} = 18.0$ ,  $\text{MgCl}_2 = 95.2$ ]

- (a) (6 points) **SHOW ALL WORK.** Determine the mole percent of  $\text{MgCl}_2$  in the solution.

25.0 % by mass: 25.0 g  $\text{MgCl}_2$  mixed with 75.0 g  $\text{H}_2\text{O}$

$$(25.0 \text{ g MgCl}_2) (1 \text{ mole} / 95.2 \text{ g}) = 0.263 \text{ mole MgCl}_2$$

$$(75.0 \text{ g H}_2\text{O}) (1 \text{ mole} / 18.0 \text{ g}) = 4.167 \text{ mole H}_2\text{O}$$

$$\text{mole \%} = (0.263) / (0.263 + 4.167) \times 100 \% = 5.94 \text{ mole \%}$$

- (b) (8 points) **SHOW ALL WORK.** Assuming complete dissociation, determine the freezing point (in °C) of this  $\text{MgCl}_2$  solution. (*Note:* For water,  $K_f = 1.86 \text{ }^\circ\text{C}/\text{m}$ )

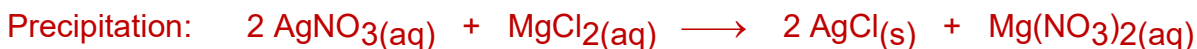
$$\text{molality of MgCl}_2 = (0.263 \text{ mole MgCl}_2) / (0.075 \text{ kg H}_2\text{O}) = 3.51 \text{ m}$$

1 mole of  $\text{MgCl}_2$  yields 3 moles of ions in solution

$$\therefore \text{effective molality is } 3 \times 3.51 = 10.5 \text{ m}$$

$$\Delta T = K_f m = (1.86 \text{ }^\circ\text{C}/\text{m}) (10.5 \text{ m}) = 19.5 \text{ }^\circ\text{C} \quad \text{freezing point} = -19.5 \text{ }^\circ\text{C}$$

- (c) (10 points) **SHOW ALL WORK.** Determine the volume (in mL) of this  $\text{MgCl}_2$  solution that is required to react exactly with 1.50 L of 0.250 M  $\text{AgNO}_3(\text{aq})$  to precipitate all of the  $\text{Ag}^+$  as  $\text{AgCl}(\text{s})$ .



0.263 moles  $\text{MgCl}_2$  is contained in 100 g of solution

$$(100 \text{ g}) (1 \text{ mL} / 1.26 \text{ g}) = 79.37 \text{ mL} = 0.07937 \text{ L}$$

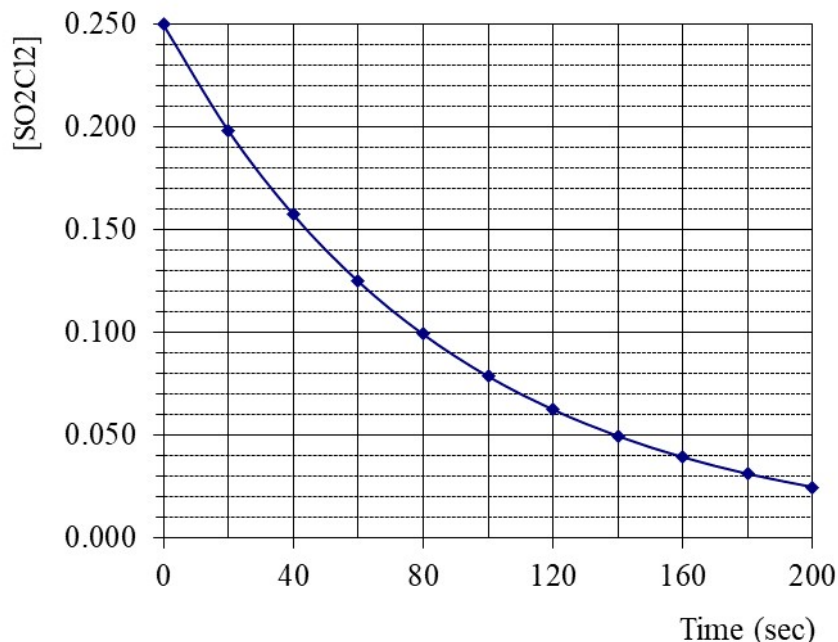
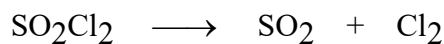
$$(0.263 \text{ mole}) / (0.07937 \text{ L}) = 3.314 \text{ M MgCl}_2$$

$$(1.50 \text{ L}) (0.25 \text{ mole Ag}^+/\text{L}) = 0.375 \text{ mole Ag}^+$$

$$(0.375 \text{ mole Ag}^+) (1 \text{ mole MgCl}_2 / 2 \text{ mole Ag}^+) = 0.1875 \text{ mole MgCl}_2$$

$$(0.1875 \text{ mole MgCl}_2) (1000 \text{ mL} / 3.314 \text{ mole}) = 56.6 \text{ mL}$$

9. (3 points) The vapor pressure of pure hexane ( $C_6H_{14}$ ) is 120 torr at 20 °C. If 1.0 mole of a non-volatile solute is dissolved in 5.0 moles of hexane at that temperature, the vapor pressure of the resulting solution should be **100** torr.
- 10 At a certain temperature, a kinetic study of the decomposition reaction of  $SO_2Cl_2$  gave the following concentration vs time data. (**Do not assume the reaction is first order!**)



- (a) (2 points) The half-life for this reaction is **60** sec.
- (b) (4 points) **SHOW ALL WORK.** Determine the *initial rate* of this reaction in units of mole/L·sec.
- $$\text{rate} = \Delta \text{conc} / \Delta \text{time} = (0.250 \text{ M} - 0.200 \text{ M}) / 20 \text{ sec} = 2.5 \times 10^{-3} \text{ M/sec}$$
- (c) (4 points) **SHOW ALL WORK.** Determine the *instantaneous* rate of this reaction when time = 120 sec.
- $$\text{rate} = (0.080 \text{ M} - 0.050 \text{ M}) / 40 \text{ sec} = 7.5 \times 10^{-4} \text{ M/sec}$$
- (d) (5 points) **In 50 words or less, describe** how you would re-plot the above data in order to prove that this reaction is first order? Also, explain how to determine the rate constant (k) from your new graph.

For a first order process, the integrated rate law is:  $\ln[SO_2Cl_2]_t = -kt + \ln[SO_2Cl_2]_0$

Therefore, a plot of  $\ln[SO_2Cl_2]$  vs t should be a straight line with slope = -k.

11. (2 points) Two important factors that affect the rate of a reaction are: **concentration**, **temperature**, or **nature of reactants**. (Do **not** write "time" as one of your answers!)
12. A kinetic study of the gas-phase decomposition reaction of  $\text{N}_2\text{O}_5$  shows it to be a first-order process. In one experiment at  $25\text{ }^\circ\text{C}$ , the reaction vessel initially contained pure  $\text{N}_2\text{O}_5$  at a pressure of 250 torr. After the reaction occurred for 10.0 hours, the **total** pressure in the reaction vessel was 484 torr.



- (a) (2 points) Write the rate law for this reaction.

$$\text{rate} = k[\text{N}_2\text{O}_5]$$

- (b) (10 points) **SHOW ALL WORK.** Determine the rate constant ( $k$ ) for this reaction at  $25\text{ }^\circ\text{C}$  in units of  $\text{sec}^{-1}$ .

Since the Rx is first order:  $\ln P(\text{N}_2\text{O}_5)_t = -kt + \ln P(\text{N}_2\text{O}_5)_0$

As the reaction occurs, based on the balanced equation above,  $2x$  torr of  $\text{N}_2\text{O}_5$  reacts to form  $4x$  torr of  $\text{NO}_2$  and  $x$  torr of  $\text{O}_2$ .

$$P(\text{total}) = P(\text{N}_2\text{O}_5) + P(\text{NO}_2) + P(\text{O}_2)$$

$$484 \text{ torr} = (250 \text{ torr} - 2x) + 4x + x = 250 \text{ torr} + 3x \quad \text{solve for } x = 78 \text{ torr}$$

After 10 hrs has elapsed:  $P(\text{N}_2\text{O}_5) = 250 \text{ torr} - 2(78 \text{ torr}) = 94 \text{ torr}$

$$\ln(94) = -k(10.0 \text{ hr})(3600 \text{ sec/hr}) + \ln(250)$$

$$k = 2.72 \times 10^{-5} \text{ sec}^{-1}$$

13. (10 points) **SHOW ALL WORK.** A certain chemical reaction has an Activation Energy of 105 kJ/mole. Determine the temperature (in  $^\circ\text{C}$ ) at which the reaction would occur  $10^5$  times faster than it does at  $0\text{ }^\circ\text{C}$ .

$$\ln(k_2 / k_1) = -E_a/RT [1/T_2 - 1/T_1]$$

$$\ln(10^5) = (-105 \text{ kJ/mole}) / (8.314 \times 10^{-3} \text{ kJ/mole}\cdot\text{K}) [1/T_2 - 1/273 \text{ K}]$$

$$11.513 = -(12629) [1/T_2 - 1/273]$$

$$1/T_2 = 2.751 \times 10^{-3}$$

$$T_2 = 363 \text{ K} = 90.5\text{ }^\circ\text{C}$$