

Name: \_\_\_\_\_

(4 points)

## Chemistry 114 Third Hour Exam

### Remember- Show all work for partial credit

All questions are worth 12 points

1.  $\text{CCl}_4$  has a normal freezing point of  $-23.0^\circ\text{C}$  and a  $K_f$  of  $30^\circ\text{C}\cdot\text{kg}/\text{mol}$ . I have dissolved 5 grams benzene (Molar mass = 78.0) in 150 grams of  $\text{CCl}_4$ . What is the freezing point of this solution?

$\Delta T = K_f m$  (Not ionic so you don't need to worry about the Van't Hoff factor)

Moles benzene =  $5 \text{ g} / (78 \text{ g}/\text{mole}) = .064 \text{ moles}$

molality = moles/ kg solvent =  $.064 \text{ moles} / .150 \text{ kg CCl}_4 = .427 \text{ m}$

$\Delta T = .427 \times 30 = 18.46^\circ\text{C}$

Freezing point =  $-23 - 18.46 = -41.46^\circ\text{C}$

2. I am studying the reaction  $\text{BrO}_3^-(\text{aq}) + 5\text{Br}^-(\text{aq}) + 6\text{H}^+(\text{aq}) \rightarrow 3\text{Br}_2(\text{l}) + 3\text{H}_2\text{O}(\text{l})$   
If the initial rate of appearance of  $\text{H}_2\text{O}$  is  $.008 \text{ mol}/\text{L}\cdot\text{sec}$ ,

A. What is the initial rate of appearance of  $\text{Br}_2$ ?

$.008 \text{ mole H}_2\text{O}/\text{L}\cdot\text{sec} \times 3 \text{ mole Br}_2/3 \text{ mole H}_2\text{O} = .008 \text{ mol Br}_2/\text{L}\cdot\text{sec}$

B. What is the initial rate of disappearance of  $\text{BrO}_3^-$ ?

$.008 \text{ mole H}_2\text{O}/\text{L}\cdot\text{sec} \times 1 \text{ mole BrO}_3^- / 3 \text{ mole H}_2\text{O} = .00267 \text{ mol BrO}_3^-/\text{L}\cdot\text{sec}$

C. What is the initial rate of disappearance of  $\text{Br}^-$ ?

$.008 \text{ mole H}_2\text{O}/\text{L}\cdot\text{sec} \times 5 \text{ mole Br}^- / 3 \text{ mole H}_2\text{O} = .00133 \text{ mol Br}^-/\text{L}\cdot\text{sec}$

3. I am studying the reaction:  $A + B \rightarrow C$

Given the following data of concentration and rates:

[A](M)	[B](M)	rate (mol/L·sec)
.06	.06	$2.76 \times 10^{-4}$
.06	0.1	$4.04 \times 10^{-4}$
0.1	0.1	$6.74 \times 10^{-4}$

Find the order parameters and the rate constant for this reaction. When you have found these numbers write the rate law for this reaction.

Dividing equation 2 by equation 1

$$\frac{4.04 \times 10^{-4}}{2.76 \times 10^{-4}} = \frac{k[A]^x[B]^y}{k[A]^x[B]^y} = \frac{k[.06]^x[.1]^y}{k[.06]^x[.06]^y}$$

$$1.46 = \left(\frac{.1}{.06}\right)^y; = 1.66^y$$

$$\ln(1.46) = y \ln(1.66)$$

$$.3784 = y(.5068); y = .3784 / .5068 = .747$$

Dividing equation 3 by equation 2

$$\frac{6.47 \times 10^{-4}}{4.04 \times 10^{-4}} = \frac{k[A]^x[B]^y}{k[A]^x[B]^y} = \frac{k[.1]^x[.1]^y}{k[.06]^x[.1]^y}$$

$$1.67 = \left(\frac{.1}{.06}\right)^x; = 1.66^x$$

$$\ln(1.67) = x \ln(1.66)$$

$$.513 = y(.507); y = .513 / .507 = 1.01 \approx 1$$

Using the last equation:

$$6.47 \times 10^{-4} = k(1.)^1(.1)^{.747}; k = 6.47 \times 10^{-4} / ((1.)^1(.1)^{.747}) = .0376$$

$$\text{rate} = .0376 [A]^1 [B]^{.747}$$

4. I have a second order reaction that has a rate constant of  $6.14 \times 10^{-2}$  L/mol·sec. If my initial reactant concentration is .01M, how long does it take until the reaction is 90% complete?

For the reaction to be 90 % complete, only 10 % of the reactant can remain so  
 $[ ] = .01(.1) = .001$

Plugging these values into the second order rate equation:

$$\frac{1}{.001} = 6.14 \times 10^{-2} t + \frac{1}{.01}$$

$$1000 = 6.14 \times 10^{-2} t + 100$$

$$900 = 6.14 \times 10^{-2} t; t = 900 / 6.14 \times 10^{-2} = 14,658 \text{ seconds}$$

5. Give the equations for the half-life of a zero order reaction

$$t_{1/2} = \frac{[A_0]}{2k}$$

first order reaction

$$t_{1/2} = \frac{.693}{k}$$

second order reaction

$$t_{1/2} = \frac{1}{k[A_0]}$$

6. Define the following terms

Homogeneous catalyst

A catalyst that is in the same phase as the reactants and products.

steric factor

The steric factor is  $p$  in the equation  $k = zpe^{-E_a/RT}$  and reflects the fraction of collisions that are actually in the correct orientation to result in a chemical reaction.

elementary step

In a reaction mechanism an elementary step is a reaction whose rate law can be written directly from the molecularly of the reaction.

half-life of a reaction

The time it takes for one half of the reactant to change to product in a given reaction.

rate constant (in a differential rate law equation)

$k$  in the equation  $\text{rate} = k[A]^n$ . It is a proportionality constant that reflects the how a rate is proportional to the concentration term.

7. If a reaction with an activation energy of 20 kJ/mol has a k of  $3 \times 10^{-5} \text{ s}^{-1}$  at  $20^\circ \text{C}$ , what is the k of the reaction at  $100^\circ \text{C}$ ?

$$20^\circ \text{C} = 293 \text{ K}$$

$$100^\circ \text{C} = 373 \text{ K}$$

$$\ln\left(\frac{K_1}{K_2}\right) = \frac{E_a}{R} \left(\frac{1}{T_2} - \frac{1}{T_1}\right)$$

$$\ln\left(\frac{3 \times 10^{-5}}{X}\right) = \frac{20000}{8.314} \left(\frac{1}{373} - \frac{1}{293}\right)$$

$$\ln\left(\frac{3 \times 10^{-5}}{X}\right) = 2406(.002681 - .003413)$$

$$\ln\left(\frac{3 \times 10^{-5}}{X}\right) = -1.761$$

$$\frac{3 \times 10^{-5}}{X} = e^{-1.761} = 1.72$$

$$X = 3 \times 10^{-5} / 1.72 = 1.74 \times 10^{-4}$$

8. I am studying the reaction  $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$

A. (For Bryan L.) What is the equilibrium expression for this reaction?

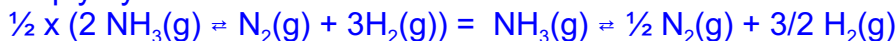
$$K = \frac{[\text{NH}_3]^2}{[\text{N}_2][\text{H}_2]^3}$$

B. (For me) If the  $K_c$  for the above reaction is  $6.0 \times 10^{-2}$ , what is the  $K_c$  for the reaction  $\text{NH}_3(\text{g}) \rightleftharpoons \frac{1}{2}\text{N}_2(\text{g}) + \frac{3}{2}\text{H}_2(\text{g})$

First, reverse the above reaction to  $2\text{NH}_3(\text{g}) \rightleftharpoons \text{N}_2(\text{g}) + 3\text{H}_2(\text{g})$

$$K_{\text{reverse}} = 1/K_{\text{forward}} = 1/6.0 \times 10^{-2} = 16.67$$

Now multiply by  $\frac{1}{2}$



$$K_{\text{new}} = K_{\text{old}}^{\frac{1}{2}} = \text{sqrt}(16.67) = 4.076$$

C. (Also for me)

What is  $K_p$  for the reaction  $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$  if  $T = 200\text{K}$ ?

$$K_p = K_c(RT)^{\Delta n}; \Delta n = 2 - 4 = -2$$

$$K_p = 6.0 \times 10^{-2} (.08206 \times 200)^{-2} = 2.23 \times 10^{-4}$$