Chem 104 - Section 1 Hour Examination II April 5, 2006

This test consists of seven (7) pages, including this cover page. Be sure your copy is complete before beginning your work. If this test packet is defective, ask for another one.

Useful Information

R = 0.08206 L·atm/K·mol = 8.314 J/K·mol $K = {}^{\circ}\text{C} + 273$ $K_p = K_c(RT)^{\Delta n}$ $t_{\gamma_2} = 0.693/k$

DO NOT WRITE BELOW THIS LINE

1.

2.

3.

4.

5.

TOTAL

α	Name								
1.	(20 points; 4 points each part) Circle the correct answer to each of the following.								
a.	Consider the reaction $CH_4(g) + 2H_2S(g) - CS_2(g) + 4H_2(g)$, for which the rate is defined as $Rate = -d[CH_4]/dt$. At any time in the course of this reaction, the rate of disappearance of $H_2(g)$ is equal to								
	Rate $4 \times Rate$ $1/4 \times Rate$ $2 \times Rate$ $1/2 \times Rate$								
b.	The reaction $AB(g) - A(g) + B(g)$ has the rate law $Rate = k[AB]$ and has a half-life of 9.6 s. If the initial concentration of AB is 1.00 mol/L, what will be the concentration of AB after 24.0 s?								
	0.000 mol/L								
c.	For the reaction described in part b, what is the value of the rate constant, k , with the appropriate units?								
	0.0289 L/mol/s 0.0433 s^{-1} 0.0722 s^{-1} 1.00 L/mol/s 2.50 s^{-1}								
d.	Kinetic data are collected for the reaction described in part b at several temperatures, and a value of k is determined at each temperature. Which of the following plots of these data would yield the indicated information?								
	A plot of k vs. T should be a straight line with a slope equal to E_a .								
	A plot of k vs. $1/T$ should be a straight line with a slope equal to $-E_d/R$.								
	A plot of $1/k$ vs. T should be a straight line with a slope equal to E_d/R .								
	A plot of $\ln k$ vs. $1/T$ should be a straight line with a slope equal to E_d/R .								
	A plot of $\ln k$ vs. $1/T$ should be a straight line with a slope equal to $-E_d/R$.								
e.	For the reaction described in part b, at any temperature a plot of ln [AB] vs. t (time) should give a straight line whose slope is								
	$\ln [AB]_o$ E_a $t_{1/2}$ $\ln A$								

 $N_2O(g)$

 $NO_{s}(g)$

2. (16 points) Consider the reaction equilibrium

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$$3NO(g) \rightleftharpoons NO_2(g) + N_2O(g)$$

for which $\Delta H^0 = -155.7$ kJ/mol in the forward direction.

3NO(a)

a. (4 points) An empty one-liter vessel is charged with 0.619 mol NO(g). When equilibrium is established at 300 K it contains 0.124 mol of $N_2O(g)$. Fill in the following table with the missing information.

	5110(g)	1102(8)	1.20(8)
Initial (mol/L)	0.619	. 0	0
Change (mol/L)	-0.372	+0.124	+0.124
Equilibrium (mol/L)	0.247	0.124	0.124

b. (6 points) In the space below, write the expression for the equilibrium constant K_c for the reaction, and calculate its numerical value.

$$K_c = \frac{[NO_2][N_2O]}{[NO]^3} = \frac{(0.124)(0.124)}{(0.247)^3} = 1.02$$

c. (3 points) If additional $N_2O(g)$ were added to the equilibrium mixture, would the reaction form more products, more reactant, or stay the same?

Answer More reactant

d. (3 points) If the temperature were increased, would the new equilibrium mixture contain more products, more reactant, or stay the same?

Answer More reactant

Name Key

3. (24 points) Consider the equilibrium

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$$H_2(g) + I_2(g) \Rightarrow 2HI(g)$$

for which K = 47.9 at 458 °C. A one-liter vessel at 458 °C is found to contain 0.800 mol $H_2(g)$, 0.800 mol $I_2(g)$, and 0.800 mol HI(g).

a. (4 points) Will the reaction proceed to the right or to the left to achieve equilibrium?

Answer right

b. (20 points) Calculate the concentrations $[H_2]$, $[I_2]$, and [HI] for the system when equilibrium is established at 458 °C. (Show work in the space provided.) [Note: This calculation does not require solving a quadratic equation.]

Equilibrium 0.800-x 0.800-x 0.800+2x
$$K = \frac{LHIJ^{2}}{LH_{2}ILI_{2}} = 47.9 = \frac{(0.800+2x)^{2}}{(0.800-x)^{2}}$$

$$K = \frac{[HI]^2}{[H_2][I_2]} = 47.9 = \frac{(0.800 + 2x)^2}{(0.800 - x)^2}$$

$$6.92 = \frac{0.800 + 2x}{0.800 - x}$$

$$6.53_{68} - 6.92x = 0.800 + 2x$$

 $9.92x = 4.73_{1}$

[H2]=[I2]=0.800-0.531=0.269 ml/L [HI] = 0.800 + (2)(0,531) = 1,86 mol/L

Check:

$$Q = \frac{(1.96)^2}{(0.269)^2} = 47.8 \implies 0 \times$$

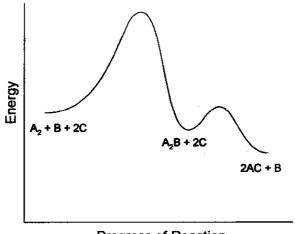
Name Key

4. (20 points) The catalyzed reaction $A_2(g) + 2C(g) \rightarrow 2AC(g)$ is thought to proceed by the following mechanism:

Step 1: $A_2(g) + B(g) \xrightarrow{k_1} A_2B(g)$

Step 2: $A_2B(g) + 2C(g)$ $\xrightarrow{k_2}$ 2AC(g) + B(g)

The reaction profile for this two-step mechanism is shown below.



Progress of Reaction

- a. (3 points) Identify any reaction intermediate species. AB
- b. (3 points) Identify the catalyst species.
- c. (4 points) Which step (Step 1 or Step 2) is rate determining? Step 1
- d. (4 points) Write the rate law expression for each step.

 $rate_1 = \frac{1}{2} \left[\frac{1}{2} \left[\frac{1}{2} \right] \right]$ $rate_2 = \frac{1}{2} \left[\frac{1}{2} \right] \left[\frac{1}{2} \right]$

- e. (4 points) Is the proposed mechanism plausible if the observed rate is $Rate = k[A_2][B]$?
- f. (2 points) Is the overall reaction exothermic or endothermic? exothermic

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Name Key

5. (20 points) All parts of this question and the bonus question on the next page refer to the hypothetical reaction

$$A_2 + B + C \rightarrow AB + AC$$

a. (15 points) The following data were collected for the rate of the reaction.

Exp	[A ₂] (M)	[B] (M)	[C] (M)	Initial Rate (M/s)	
1	0.200	0.150	0.120	1.60 x 10 ⁻²	
2	0.800	0.150	0.120	3.20 x 10 ⁻²	>Ritca[A] 1/2
3	0.200	0.600	0.120	6.40 x 10 ⁻²	>Pate = [B]
4	0.200	0.150	0.360	4.80 x 10 ⁻²	⇒ Extca Cc]

What is the rate law for the reaction? $Rate = k LA_1 J^2 LB J LC J$

b. (5 points) What is the value of the rate constant, k, with the appropriate units, based on the data from Experiment #1? (Show work in the space below)

Name Key

BONUS (5 points) The following mechanism has been proposed for this reaction.

$$A_2 \xrightarrow{k_1} 2A$$
 fast
 $A + B + C \xrightarrow{k_2} ABC$ slow
 $A + ABC \xrightarrow{k_3} AB + AC$ fast

Prove that this mechanism has a rate law consistent with the correct observed rate law in part a.

$$Rxtc = rxt_{2} = e_{2} CAT[B]CC]$$

$$e_{1} CA_{2}T = e_{1} CAT^{2} \Rightarrow CAT = \left(\frac{R_{1}}{R_{-1}}\right)^{2} CA_{2}T^{2}$$

$$\Rightarrow Rxt = e_{2} \left(\frac{R_{1}}{R_{-1}}\right)^{2} CA_{3}T^{2} CBTCCT$$