## Exam 2a **Chem 1142 Spring 2011**

Name:	KE	$\bigvee$

MULTIPLE CHOICE. [2 pts ea.] Choose the best response on the scantron sheet. [36 pts total.]

Q1. For the reaction: A  $\longrightarrow$  2B + C, the rate could be expressed as  $-\Delta[A]/\Delta t$ . An equivalent expression is:

a) 
$$-\frac{1}{2}\frac{\Delta[B]^2}{\Delta t}$$
 b)  $-\frac{1}{2}\frac{\Delta[B]}{\Delta t}$  c)  $+\frac{1}{2}\frac{\Delta[B]}{\Delta t}$  d)  $+\frac{\Delta[B]^2}{\Delta t}$  e)  $+\frac{\Delta[B][C]}{\Delta t}$ 

b) 
$$-\frac{1}{2}\frac{\Delta[B]}{\Delta t}$$

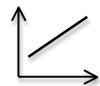
c) 
$$+\frac{1}{2}\frac{\Delta[B]}{\Delta t}$$

d) + 
$$\frac{\Delta[B]^2}{\Delta t}$$

e) + 
$$\frac{\Delta[B][C]}{\Delta t}$$

Q2.A student analyzed a second-order reaction and obtained the graph at the right, but forgot to label the axes. What should the labels be for the X and the Y coordinates respectively?

- a) time, ln [A]
- b) time, [A]
- c) temperature, [A]
- d) temperature, ln [A]
- e) time, 1/[A]



- Q3. For the overall reaction:  $2A \longrightarrow B$ , the reaction order is:
  - a) zero order

b) first order

- c) second order
- d) impossible to predict without experimental rates at various concentrations of A
- e) impossible to predict without knowing the heat of reaction
- Q4. What are the units for k, the rate constant, in a first order reaction?
  - a) M·s<sup>-1</sup>
- b) M
- c) s<sup>-1</sup>
- d) M-1 ·s-1
- e) s-1 ·M
- Q5. What is the rate law for the reaction:  $A + B \longrightarrow 2C$ , based on the following kinetic data?

Experiment #	Initial Conc of [A] / M	Initial Conc. of [B] / M	Initial rate of reaction / M/s
1	0.40	0.10	$3.6 \times 10^3$
2	0.20	0.10	$1.8 \times 10^3$
3	0.20	0.50	4.5 x 10 <sup>4</sup>

a) rate = 
$$k[A][B]^2$$

b) rate = 
$$k[A]^{1/2}[B]^5$$

c) rate = 
$$k[A]^2[B]$$

- a) rate =  $k[A][B]^2$ d) rate =  $k[A][B]^{1/5}$
- e) rate =  $k[A]^{1/2}[B]^2$
- Q6. What is unique about the half-life of any first-order reaction at 25 °C?
  - a) The units are always s<sup>-1</sup>

- b) The value only depends on the rate constant, *k*
- c) The value only depends on the initial concentration of reactant
- d)  $\Delta$ [A]/ $\Delta t = 1$
- e)  $\Delta [A]/\Delta t = \frac{1}{2}$

•	RT, may be used to	calculate the acti	vation energy from the slope of a line
a) ln k vs. 1/Temperature d) 1/k vs. 1/time	b) ln <i>k</i> vs. 1/time) ln <i>k</i> vs. e <sup>-T</sup>	ie	c) 1/k vs. Temperature
<ul><li>a) increases due to an increased</li><li>b) increases only for an endothe</li><li>c) increases due to a greater num</li></ul>	activation energy ermic reaction nber of effective c		
Q9. In basic solution, (CH <sub>3</sub> ) <sub>3</sub> CCl reacts a	according to the ed	quation:	
$(CH_3)_3CCl + OH \longrightarrow (CH_3)_3C$	COH + Cl-		
$(CH_3)_3CC1 \longrightarrow (CH_3)C^+ + Cl^-$	_	(SLOW) (FAST)	
What is the rate law expression	for the reaction?		
d) 1/k vs. 1/time e) ln k vs. e <sup>-T</sup> 3. In general, as the temperature increases, the rate of a chemical reaction a) increases due to an increased activation energy b) increases only for an endothermic reaction c) increases due to a greater number of effective collisions d) increases because bonds are weakened e) is not changed d. In basic solution, (CH <sub>3</sub> ) <sub>3</sub> CCl reacts according to the equation:  (CH <sub>3</sub> ) <sub>3</sub> CCl + OH <sup>-</sup> → (CH <sub>3</sub> ) <sub>3</sub> COH + Cl <sup>-</sup> The accepted mechanism for this reaction is:  (CH <sub>3</sub> ) <sub>3</sub> CCl → (CH <sub>3</sub> ) <sub>5</sub> C <sup>+</sup> + Cl <sup>-</sup> (CH <sub>3</sub> ) <sub>3</sub> COH (FAST)  What is the rate law expression for the reaction?  a) rate = k[(CH <sub>3</sub> ) <sub>3</sub> C <sup>+</sup> ] <sup>2</sup> [OH <sup>-</sup> ] b) rate = k[(CH <sub>3</sub> ) <sub>3</sub> CC <sup>+</sup> ][OH <sup>-</sup> ] <sup>2</sup> c) rate = k[Cl <sup>-</sup> ] d) rate = k[(CH <sub>3</sub> ) <sub>3</sub> CCl] e) rate = k[(CH <sub>3</sub> ) <sub>3</sub> CCl][OH <sup>-</sup> ] 0. What name would be used to describe an elementary reaction such as: a) NO(aq) + 2Cl <sup>-</sup> (aq) → NOCl <sub>2</sub> <sup>2</sup> -(aq) a) bimolecular b) unimolecular c) termolecular d) dimolecular e) termolecular			
-	•	reaction such as:	
a) bimolecular b) unir	molecular	c) termolecular	
$CaSO_3(s) \rightleftharpoons CaO(s) +$			
a) $\frac{\text{[CaO][SO}_2]}{\text{[CaSO}_3]}$ b) [Ca	aO][SO <sub>2</sub> ]	c) $[SO_2]$	
d) $\frac{1}{[SO_2]}$ e) $\frac{[O]}{[Ca]}$	$\frac{\text{CaSO}_3}{\text{aO}[\text{SO}_2]}$		
	of the equilibrium	constant does the	e reaction mixture consist mainly of
a) $10^5$ b) $10^3$	c) 10 <sup>0</sup>	d) 10 <sup>-3</sup>	e) 10 <sup>-5</sup>
$4NH_3(g) + 5O_2(g) \rightleftharpoons 4NO(g)$ $a) K_c = \frac{4[NH_3] + 5[O_2]}{6[HO] + 4[NO]}$	+ $6H_2O(g)$ b) $K_c = \frac{6[H_2O]}{4[NH]}$	)]+4[NO] ]+5[O]	
c) $K_c = \frac{[H_2O][NO]}{[NH_3][O_2]}$	d) $K_c = \frac{[H_2O]}{[NH_3]}$	$\begin{bmatrix} NO \end{bmatrix}^4 \\ \begin{bmatrix} A \\ O_2 \end{bmatrix}^5 $	e) $K_c = \frac{[NH_3]^4 [O_2]^3}{[H_2O]^6 [NO]^4}$

b) CO(g) + c) CO(g) + d) CO(g) +	the following equilifold $3H_2(g) \rightleftharpoons CH_4(g)$ $H_2O(g) \rightleftharpoons CO_2(g)$ $2H_2(g) \rightleftharpoons CH_3OI$ $1/2 O_2(g) \rightleftharpoons CO_2(g)$ $\rightleftharpoons N_2O_4(g)$	$+ H_2O(g)$ (x) $+ H_2(g)$ + H(g)	<i>K</i> <sub>p</sub> ?	
	$L_2(g) + Cl_2(g)$ ; $\Delta H$ am constant of 6.2	-	the following would b	e true if the temperature were
2. The cond	librium constant we centration of ICl(g) al pressure of I <sub>2</sub> we	would be increa	sed.	
a) 1 only	b) 2 only	c) 3 only	d) 1 and 2 only	e) 1 and 3 only
1. increase 2. decrease 3. increase	on system $N_2(g)$ + rium and increase the temperature the temperature the pressure the pressure			= –92 kJ/mol. In order to both
a) 1 only	b) 2 only	c) 1 and 3 or	aly d) 2 and 3 only	e) 1 and 4 only
Q17. Addition of a a TRUE	catalyst to a reaction b) FALSE	on at equilibrium	does not alter the value	e of the equilibrium constant.
is equal to 0.53, a) The reaction b) The reaction c) The reaction d) The reaction	•	to make more pr and no shift will on make more reach and causes a temp	oducts occur tants erature decrease	mperature. If the reaction quotient

#### Short Response. Show all work.

Q19. [7 pts.] The rate law for the reaction

$$NH_4^+(aq) + NO_2^-(aq) \longrightarrow N_2(g) + 2H_2O(l)$$

is given by rate= $k[NH_4^+][NO_2^-]$ . At 25 °C, the rate constant is 3.0 x 10<sup>-4</sup> M<sup>-1</sup> 's<sup>-1</sup>. Calculate the rate of the reaction at this temperature if  $[NH_4^+] = 0.26$  M, and  $[NO_2^-] = 0.080$  M.

Rate = 
$$3.0 \times 10^{-4} \, \text{M}^{-1} \cdot \text{S}^{-1} \times 0.26 \, \text{M} \times 0.080 \, \text{M}$$
  
=  $6.2 \times 10^{-6} \, \frac{\text{M}}{\text{s}}$ 

Q20. [15 pts.] The rate at which tree crickets chirp is 2.0 x 10<sup>2</sup> per minute at 27 °C, but only 39.6 per minute at 5 °C. From these data, calculate the "energy of activation" for the chirping process. (*Hint:* The ratio of rates is equal to the ratio of rate constants.)

$$\lim_{x \to \infty} \left(\frac{K_2}{K_1}\right) = \frac{E_A}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right)$$

$$= \lim_{x \to \infty} \left(\frac{E_2}{K_1}\right) \cdot \frac{R}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right)$$

$$= \lim_{x \to \infty} \left(\frac{2 \cdot 0 \times 10^2 / m_{in}}{39 \cdot 6 / m_{in}}\right) \times 83145$$

$$= \lim_{x \to \infty} \left(\frac{1}{278K} - \frac{1}{300 \cdot K}\right)$$

$$= 51,000$$

$$= \lim_{x \to \infty} \int_{m_0}^{\infty} dx$$

Q21. [5 pts.] How does a catalyst increase the rate of a reaction?

### It lowers the activation energy by providing an alternate mechanism.

Q22. [6 pts.] Write equilibrium constant expressions for  $K_c$  and  $K_p$  for the following reactions:

a) 
$$2CO_2(g) \rightleftharpoons 2CO(g) + O_2(g)$$

$$k_{c} = \frac{[co]^{2}[o_{2}]}{[co_{2}]^{2}}$$
,  $k_{p} = \frac{\rho_{co}^{2} \cdot \rho_{o_{2}}}{\rho_{co_{2}}^{2}}$ 

b) 
$$2HgO(s) \rightleftharpoons 2Hg(l) + O_2(g)$$

$$K_c = [0_2]$$
,  $K_p = P_{02}$ 

Q23. [10 pts.] Write out electron configurations for the following ATOMS or IONS:

a) Li 
$$|s^2/s|^6$$

#### Q24. [11 pts.] The equilibrium constant $K_c$ for the reaction

$$H_2(g) + CO_2(g) \rightleftharpoons H_2O(g) + CO(g)$$

is 4.2 at 1650 °C. Initially 0.80 mol H<sub>2</sub> and 0.80 mol CO<sub>2</sub> are injected into a 5.0-L flask. Calculate the concentration of each species at equilibrium.

$$[H_{2}]_{i} = [C0_{2}]_{i} = \underbrace{0.80 \, \text{mol}}_{\text{5.0 L}} = 0.16 \, \text{M}$$

$$= 0.328 = 3.05 \times \\
= 0.328 \\
3.05 = 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.106$$

$$= 0.108 \, \text{M}$$

$$= (H_{2})[C0]_{ea} = (C0)_{ea} = 0.052 \, \text{M}$$

$$= (W)(X)$$

Q25. [10 pts.] Write formulas for the following compounds:

a) ammonium sulfate 
$$(NH_4)_2 SO_4$$
  $NH_4$   $SO_4$ 
b) copper(I) acetate  $(L_1L_3O_2)_2$   $C_4$   $C_4$   $C_4$   $C_5$ 
c) iron(III) nitride  $(L_1L_3O_2)_2$   $C_4$   $C_4$   $C_4$   $C_5$ 
d) heptanitrogen disulfide  $(L_1L_3O_2)_2$   $(L_2L_3O_2)_3$ 
e) nitric acid  $(L_1L_3O_3)_3$   $(L_2L_3O_4)_4$   $(L_1L_3O_4)_5$   $(L_1L_$ 

#### **BONUS QUESTIONS**

Predict the molecular geometry of XeF4.

Draw a diagram showing the formation of hydrogen bonds between molecules of CH<sub>3</sub>OH. Clearly label the location of the hydrogen bonds in your diagram!

# Useful Information

#### Periodic Table of the Flements

			Perio	oaic i	able (	ot the	Elem	ients									
IA 1	IIA											IIIA	IVA	VA	VIA	VIIA	VIIIA
1	Ī																2
Н																	He
1.01	2											13	14	15	16	17	4.00
3	4											5	6	7	8	9	10
Li	Be											В	С	N	0	F	Ne
6.94	9.01											10.81	12.01	14.01	16.00	19.00	20.18
11	12											13	14	15	16	17	18
Na	Mg											Al	Si	P	S	CI	Ar
22.99	24.31	3	4	5	6	7	8	9	10	11	12	26.98	28.09	30.97	32.07	35.45	39.95
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
39.10	40.08	44.96	47.87	50.94	52.00	54.94	55.85	58.93	58.69	63.55	65.39	69.72	72.61	74.92160	78.96	79.90	83.80
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr	Υ	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	ı	Xe
85.47	87.62	88.91	91.22	92.91	95.94	[98]	101.07	102.91	106.42	107.87	112.41	114.82	118.71	121.76	127.60	126.90	131.29
55	56	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba*	Lu	Hf	Ta	W	Re	Os	lr	Pt	Au	Hg	TI	Pb	Bi	Po	At	Rn
132.91	137.33	174.97	178.49	180.95	183.84	186.21	190.23	192.22	195.08	196.97	200.59	204.38	207.20	208.98	[210]	[210]	[222]
87	88	103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118
Fr	Ra**	Lr	Rf	Db	Sg	Bh	Hs	Mt									
[223]	[226]	[262]	[261]	[262]	[266]	[264]	[265]	[268]	[269]	[272]	[277]		[285]		[289]		[293]
																,	
		57	58	59	60	61	62	63	64	65	66	67	68	69	70		
	*	La	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb		
		138.91	140.12	140.91	144.24	[145]	150.36	151.96	157.25	158.93	162.50	164.93	167.26	168.93	173.04		

Cf

Es

Fm

Md

No [259]

Bk

 $R = 8.314 \text{ J/mol \cdot K} = 0.08206 \text{ (L ·atm)/(mol · K)}$ 

Th

Αc

Рa

$$k = \mathcal{A}e^{-Ea/RT} \qquad \ln k = (-E_a/R)(1/T) + \ln \mathcal{A}$$

$$\ln \left(\frac{k_2}{k_1}\right) = \frac{E_a}{R} \left(\frac{1}{T_1} - \frac{1}{T_2}\right)$$

U

•1-order: 
$$\ln[A]_t = -kt + \ln[A]_0$$
  $\ln\left(\frac{[A]_t}{[A]_0}\right) = -kt$   $t_{1/2} = 0.693 / k$   
•2-order:  $1/[A]_t = kt + 1/[A]_0$   $t_{1/2} = 1 / ([A]_0 \cdot k)$ 

Pu

Αm

Cm

Np

$$K_{\rm p} = K_{\rm c}({\rm R}T)^{\rm ang}$$

Given: 
$$ax^2 + bx + c = 0$$
, then  $x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$