

Chemistry 210

Exam 2

Be sure to put your name on each page. This page can be removed from your exam so that you will have a Periodic Table handy throughout the exam, it does not need to be turned in. Show all your work for problems which require any sort of calculation, no credit will be given for answers without work shown. If you have shown a significant amount of work or multiple drawings for a problem, draw a box around what you consider your final answer.

Avogadro's Number = 6.022×10^{23} units/mol

$32.00^\circ\text{F} = 0.000^\circ\text{C} = 273.15\text{K}$

Density of Water = 1.000 g/mL

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

$PV = nRT$

$\Delta T_{\text{fp/bp}} = k_{\text{fp/bp}} \cdot m \cdot i$

For water, $k_{\text{fp}} = -1.86^\circ\text{C}/m$; $k_{\text{bp}} = 0.52^\circ\text{C}/m$

$P_1 = X_1 P_1^\circ$

$P = cRTi$

$C_1 V_1 = C_2 V_2$

Quadratic formula:

$$x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$$

Integrated Rate Laws:

0th order $[A]_t = -kt + [A]_0$

1st order $\ln[A]_t = -kt + \ln[A]_0$

2nd order $1/[A]_t = kt + 1/[A]_0$

$k = Ae^{-E_a/RT}$

$$\ln(k) = \left(\frac{-E_a}{R} \right) \left(\frac{1}{T} \right) + \ln(A)$$

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

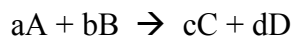
1 H 1.0079																	2 He 4.0026				
3 Li 6.941	4 Be 9.0122															5 B 10.811	6 C 12.011	7 N 14.007	8 O 15.999	9 F 18.998	10 Ne 20.180
11 Na 22.990	12 Mg 24.305															13 Al 26.982	14 Si 28.086	15 P 30.974	16 S 32.066	17 Cl 35.453	18 Ar 39.948
19 K 39.098	20 Ca 40.078	21 Sc 44.956	22 Ti 47.88	23 V 50.942	24 Cr 51.996	25 Mn 54.938	26 Fe 55.847	27 Co 58.933	28 Ni 58.69	29 Cu 63.546	30 Zn 65.39	31 Ga 69.723	32 Ge 72.61	33 As 74.922	34 Se 78.96	35 Br 79.904	36 Kr 83.80				
37 Rb 85.468	38 Sr 87.62	39 Y 88.906	40 Zr 91.224	41 Nb 92.906	42 Mo 95.94	43 Tc (98)	44 Ru 101.07	45 Rh 102.91	46 Pd 106.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.76	52 Te 127.60	53 I 126.90	54 Xe 131.29				
55 Cs 132.91	56 Ba 137.33	57 La 138.91	72 Hf 178.49	73 Ta 180.95	74 W 183.84	75 Re 186.21	76 Os 190.23	77 Ir 192.22	78 Pt 195.08	79 Au 196.97	80 Hg 200.59	81 Tl 204.38	82 Pb 207.2	83 Bi 208.98	84 Po (209)	85 At (210)	86 Rn (222)				
87 Fr (223)	88 Ra 226.03	89 Ac 227.03	104 Rf (261)	105 Db (262)	106 Sg (263)	107 Bh (262)	108 Hs (265)	109 Mt (266)	110 (269)	111 (272)	112 (277)	114 114		116 116							

58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm (145)	62 Sm 150.36	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.94	70 Yb 173.04	71 Lu 174.97
90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np 237.05	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (258)	101 Md (258)	102 No (259)	103 Lr (260)

Multiple Choice: Circle the letter of the most correct response. (6pts. per question)

1. Which of the following does **not** affect the rate of a reaction?
- The frequency of collisions between reacting particles
 - The coefficients of the reactants in the balanced equation**
 - The temperature of the system
 - The orientation of colliding particles
 - The energy of collisions between reacting particles

2. For the generic equation:

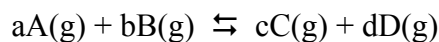


Which of the following is a correct expression of the rate of the reaction:

- $-\frac{1}{a} \frac{\Delta[A]}{\Delta t}$
- $-\frac{1}{b} \frac{\Delta[B]}{\Delta t}$
- $k[A]^a[B]^b$
- $\frac{1}{d} \frac{\Delta[D]}{\Delta t}$
- $k[C]^c[D]^d$

There should have been only 1 correct response,
this question was discarded.

3. For the generic equation



The value of the equilibrium constant, K_c :

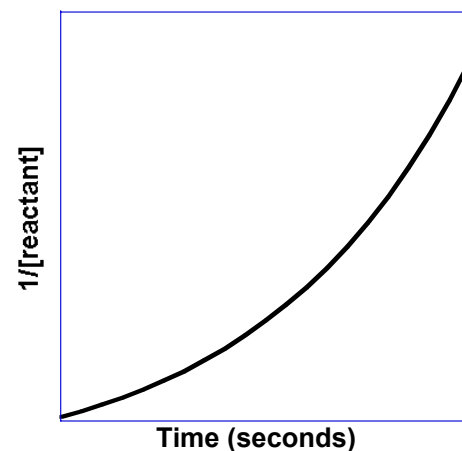
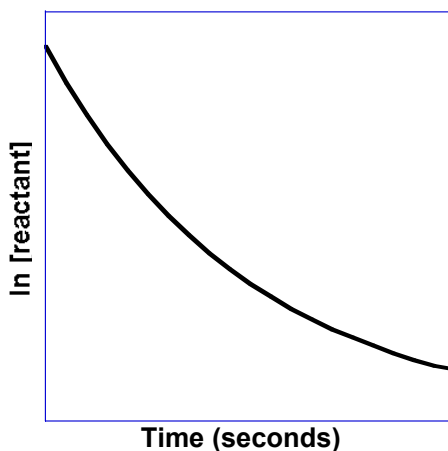
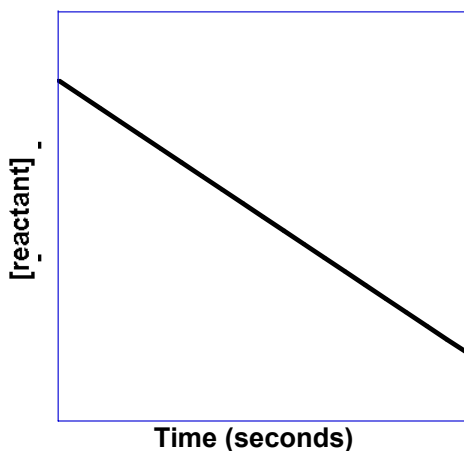
- Must be measured, it cannot be derived from the balanced equation
 - Is equal to $k[A]^a[B]^b$
 - Is equal to $([A]^a[B]^b)/([C]^c[D]^d)$
 - Is not affected by temperature
 - Is equal to $([C]^c[D]^d)/([A]^a[B]^b)$**
4. If the rate of a reaction increases by a factor of 4 when the initial concentration of reactant "A" is increased by a factor of 4, the reaction must be:
- 2nd order overall
 - 4th order with respect to $[A]_0$
 - 1st order with respect to $[A]_0$**
 - 2nd order with respect to $[A]_0$
 - The order of the reaction depends on the balanced chemical equation

5. For a first order reaction:

- The slope of the integrated rate law plot is equal to k
- The intercept of the integrated rate law plot is equal to the initial concentration
- The slope of the integrated rate law plot is equal to $(-E_a/R)$
- The intercept of the integrated rate law is equal to the \ln of the initial concentration**
- The slope of the integrated rate law is equal to the frequency factor, A .

6. Which of the following is **true** regarding catalysts and catalyzed reactions?

- The presence of a catalyst changes the equilibrium constant for a reaction
- The presence of a catalyst changes the energy of the products and reactants in a reaction
- The presence of a catalyst changes the activation energy for a reaction**
- The presence of a catalyst does not change the mechanism of a reaction
- The concentration of a catalyst cannot appear in the rate law for a reaction



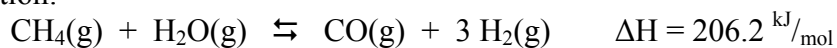
7. The reaction represented by the plots above:
- Is zero order**
 - Is first order
 - Is second order
 - Is third order
 - The order can't be determined by these graphs
8. Which of the following is **false** regarding equilibrium?
- The concentrations of products and reactants does not change once the reaction has reached equilibrium
 - Equilibrium can be shifted by changing pressure or temperature
 - The forward and reverse reactions stop when a system reaches equilibrium**
 - The rates of the forward and reverse reactions are equal
 - Equilibrium concentrations do not depend upon whether you approach equilibrium from the left or the right
9. Which of the following is **false** regarding reaction mechanisms?
- The steps of the mechanism can contain chemical species that do not appear in the overall correctly balanced chemical equation
 - The observed rate law is equal to the sum of the rate laws from all steps**
 - Catalysts can appear in the steps of a mechanism
 - A mechanism must be composed of elementary reactions
 - The observed rate law must agree with the rate law of the slowest step
10. Which of the following statements is **false** regarding the reaction quotient, Q ?
- It tells the direction that the reaction must shift to reach equilibrium
 - If $Q > K_c$, the system needs to shift toward the reactants to reach equilibrium
 - If $Q = K_c$, the system is at equilibrium
 - If $Q < K_c$, the system needs to shift toward the reactants to reach equilibrium**
 - It has the same mathematical form as the equilibrium constant

Multiple Choice Calculations: (12pts each)

11. A reaction is found to be first order with respect to reactant A and second order with respect to reactant B. If $[A]_0 = 0.318\text{M}$, $[B]_0 = 0.934\text{M}$ and $k = 5.49 \times 10^{-5} \text{M}^{-2}\text{sec}^{-1}$, what is the initial rate of the reaction?

- a. $1.98 \times 10^{-4} \text{M}/\text{sec}$
- b. $1.85 \times 10^{-4} \text{M}/\text{sec}$
- c. $1.63 \times 10^{-5} \text{M}/\text{sec}$
- d. $1.52 \times 10^{-5} \text{M}/\text{sec}$
- e. $5.19 \times 10^{-6} \text{M}/\text{sec}$

12. For the reaction:



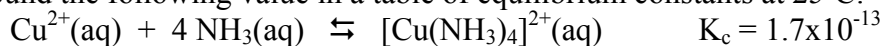
The equilibrium concentrations have been found to be $[\text{CO}]_{\text{eq}} = 2.22\text{M}$, $[\text{H}_2]_{\text{eq}} = 1.81\text{M}$, $[\text{CH}_4]_{\text{eq}} = 8.81 \times 10^{-5} \text{M}$, $[\text{H}_2\text{O}]_{\text{eq}} = 1.53 \times 10^{-2} \text{M}$. What is the equilibrium constant?

- a. 1.02×10^{-7}
- b. 388
- c. 2.98×10^6
- d. 8.94×10^6
- e. 9.77×10^6

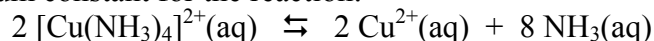
13. A reaction is found to be second order with respect to fluoride ion, a reactant. If $[\text{F}^-]_0 = 2.19\text{M}$ and $k = 1.16 \times 10^{-2} \text{M}^{-1}\text{sec}^{-1}$, what will the concentration be after 12 minutes have passed?

- a. 8.81 M
- b. 1.68 M
- c. 0.596 M
- d. 0.127 M
- e. **0.114 M**

14. You have found the following value in a table of equilibrium constants at 25°C:



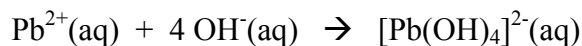
What is the equilibrium constant for the reaction:



- a. **3.5×10^{25}**
- b. 5.9×10^{12}
- c. 2.9×10^{12}
- d. 3.4×10^{-13}
- e. 2.9×10^{-26}

Problems: (21pts each)

15. For the reaction:



You have collected the following data at 16.93°C:

Experiment	$[\text{Pb}^{2+}]_0$	$[\text{OH}^{-}]_0$	Rate _{observed}
1	0.267 M	1.621 M	$1.16 \times 10^{-2} \text{ M/sec}$
2	0.267 M	3.242 M	$2.32 \times 10^{-2} \text{ M/sec}$
3	0.801 M	1.621 M	$3.48 \times 10^{-2} \text{ M/sec}$

What are the rate law and the value of the rate law constant, k, for this reaction?

If you redo Experiment 2 at 34.26°C, the rate is $7.53 \times 10^{-2} \text{ M/sec}$. What is the activation energy for this reaction?

Comparing runs 1 and 2, $[\text{OH}^{-}]_0$ doubles, rate doubles, the rxn is 1st order with respect to $[\text{OH}^{-}]_0$
Comparing runs 1 and 3, $[\text{Pb}^{2+}]_0$ triples, rate triples, the rxn is 1st order with respect to $[\text{Pb}^{2+}]_0$

$$\text{Rate}_0 = k [\text{Pb}^{2+}]_0 [\text{OH}^{-}]_0$$

Plugging in values to solve for k:

$$k = \frac{\text{Rate}_0}{[\text{Pb}^{2+}]_0 [\text{OH}^{-}]_0} = \frac{1.16 \times 10^{-2} \frac{\text{M}}{\text{sec}}}{(0.267 \text{ M})(1.621 \text{ M})} = 2.68 \times 10^{-2} \frac{1}{\text{M} \cdot \text{sec}}$$

At 34.26°C, the rate law *expression* remains the same, but the value of k changes. Plugging in the new conditions:

$$k = \frac{\text{Rate}_0}{[\text{Pb}^{2+}]_0 [\text{OH}^{-}]_0} = \frac{7.53 \times 10^{-2} \frac{\text{M}}{\text{sec}}}{(0.267 \text{ M})(3.242 \text{ M})} = 8.70 \times 10^{-2} \frac{1}{\text{M} \cdot \text{sec}}$$

Plugging in to the comparative form of the Arrhenius equation:

$$\ln \left(\frac{2.68 \times 10^{-2} \frac{1}{\text{M} \cdot \text{sec}}}{8.70 \times 10^{-2} \frac{1}{\text{M} \cdot \text{sec}}} \right) = \frac{E_a}{8.314 \text{ J/mol} \cdot \text{K}} \left(\frac{1}{307.41 \text{ K}} - \frac{1}{290.08 \text{ K}} \right) \rightarrow E_a = 50.4 \text{ kJ/mol}$$

16. When 3.512g of nitrogen dioxide $\{\text{NO}_2(\text{g})\}$ and 4.913g of chlorine gas $\{\text{Cl}_2(\text{g})\}$ are sealed together in a 2.000L vessel, they reach equilibrium with nitrogen trichloride $\{\text{NCl}_3(\text{g})\}$ and oxygen $\{\text{O}_2(\text{g})\}$. If the equilibrium concentration of $\text{Cl}_2(\text{g})$ is found to be $3.124 \times 10^{-4} \text{ M}$:

- What are the equilibrium concentrations of all products and reactants?
- What is the value of K_c ?
- Is the reaction Product-favored or reactant-favored?

	$2 \text{NO}_2(\text{g})$	$+ 3 \text{Cl}_2(\text{g}) \leftrightarrow$	$2 \text{NCl}_3(\text{g})$	$+ 2 \text{O}_2(\text{g})$
Initial	$3.512 \text{ g} / 46.005 \frac{\text{g}}{\text{mol}} / 2.000 \text{ L}$	$4.913 \text{ g} / 70.906 \frac{\text{g}}{\text{mol}} / 2.000 \text{ L}$		
	$3.817 \times 10^{-2} \text{ M}$	$3.464 \times 10^{-2} \text{ M}$	0	0
Δ	-2x	-3x	+2x	+2x
@ Eq	$(3.817 \times 10^{-2} - 2x) \text{ M}$	$(3.464 \times 10^{-2} - 3x) \text{ M}$	2x M	2x M
	$1.523 \times 10^{-2} \text{ M}$	$3.124 \times 10^{-4} \text{ M}$	$2.289 \times 10^{-2} \text{ M}$	$2.289 \times 10^{-2} \text{ M}$

$$K_c = \frac{[\text{NCl}_3]_{\text{eq}}^2 [\text{O}_2]_{\text{eq}}^2}{[\text{NO}_2]_{\text{eq}}^2 [\text{Cl}_2]_{\text{eq}}^3}$$

$$K_c = 3.881 \times 10^7, \text{ product-favored}$$