

Chemistry 210

Exam 1

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 = $1.000^{\text{g}}/\text{mL}$

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

$PV = nRT$

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

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 $[A]_t = -kt + [A]_o$

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

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

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

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

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

$$\text{pH} = \text{pK}_a + \log\left(\frac{[\text{conjugate base}]}{[\text{conjugate acid}]} \right)$$

$E_{\text{cell}} = E^\circ_{\text{cell}} - \frac{RT}{nF} \ln Q$

$E^\circ_{\text{cell}} = \frac{RT}{nF} \ln K^\circ$

$K^\circ = e^{(nF/RT) E^\circ_{\text{cell}}}$

$F = 96485 \text{ J}/\text{V}\cdot\text{mol of electrons}$

$\Delta G^\circ = \Delta H^\circ_{\text{system}} - T\Delta S^\circ_{\text{system}}$

$\Delta G^\circ = -nFE^\circ_{\text{cell}} = -RT \ln K^\circ$

$\Delta G = \Delta G^\circ + RT \ln Q$

$F = 96485 \text{ C}/\text{mol electrons}$

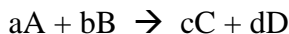
$1A = 1 \text{ C} / \text{sec}$

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		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 (5pts each): Circle the letter of the most correct response.

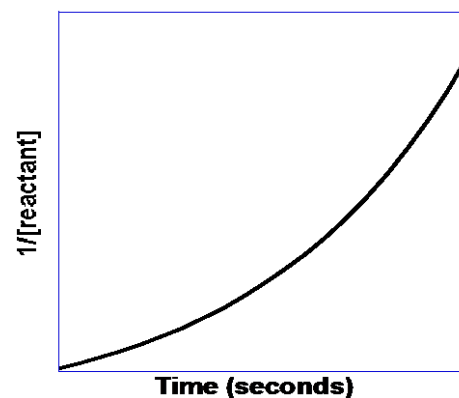
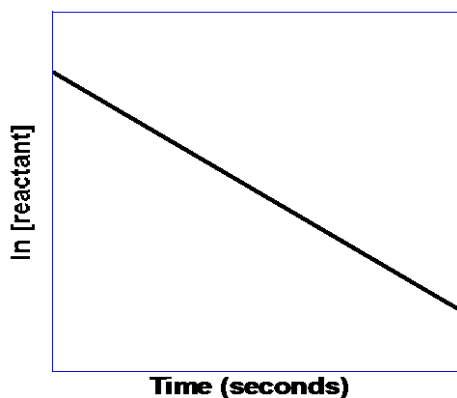
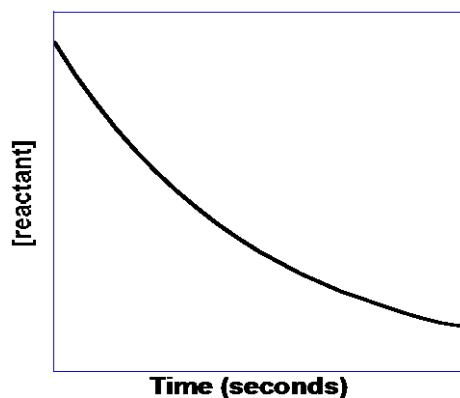
- Which of the following statements is most correct about colligative properties?
 - The presence of a solute lowers the vapor pressure of a solution.
 - The presence of a solute lowers the freezing point of a solution.
 - Colligative properties depend upon the number of solute particles, not on the identity of the solute particles.
 - The presence of a solute raises the boiling point of a solution.
 - These statements are all correct.**
- Which of the following does *not* affect the rate of a reaction?
 - The frequency of collisions between reacting particles
 - The orientation of colliding particles
 - The coefficients of the reactants in the balanced equation**
 - The temperature of the system
 - The energy of collisions between reacting particles
- If the rate of a reaction increases by a factor of 9 when the initial concentration of reactant “A” is increased by a factor of 9, the reaction must be:
 - 0th order with respect to $[A]_0$
 - 1st order with respect to $[A]_0$**
 - 2nd order overall
 - 2nd order with respect to $[A]_0$
 - The order of the reaction depends on the balanced chemical equation
- Which of the following is *true* regarding catalysts and catalyzed reactions?
 - Catalysts always have to be solids
 - 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
 - The presence of a catalyst changes the energy of the products and reactants in a reaction
- Which of the following is *false* regarding reaction mechanisms?
 - The observed rate law must agree with the rate law of the slowest step
 - 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
- For the generic equation:



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

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

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



8. 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

Problems:

9. What is the freezing point of a solution made by dissolving 22.618g of sodium sulfate in 150.0g of water? (12pts)

$$\Delta T_{fp} = (1.86 \text{ } ^\circ\text{C}/m) (22.618\text{g} / 142.042\text{g/mol} / 0.1500\text{kg}) (3 \text{ particles/mol}) = 5.92^\circ\text{C}$$

$$\{m = 1.06 \text{ mol/kg}\}$$

$$T_{fp} = -5.92^\circ\text{C}$$

10. A reaction is found to be first order with respect to reactant A and second order with respect to reactant B. If $[A]_0 = 0.318\text{M}$, $[B]_0 = 0.196\text{M}$ and $k = 6.27 \times 10^{-3} \text{ M}^{-2}\text{sec}^{-1}$, what is the initial rate of the reaction? (12pts)

$$\text{Rate}_0 = k [A]_0^1 [B]_0^2 = (6.27 \times 10^{-3} \text{ M}^{-2}\text{sec}^{-1})(0.318\text{M})(0.196\text{M})^2 = 7.66 \times 10^{-5} \text{ M}/\text{sec}$$

11. A reaction is found to be zero order with respect to methane, a reactant. If $[\text{CH}_4]_0 = 1.61\text{M}$ and $k = 0.103\text{ M}\cdot\text{min}^{-1}$, what will the concentration of methane be after 7.43 minutes have passed? (13pts)

Zeroth order integrated rate law...

$$[\text{CH}_4]_t = -kt + [\text{CH}_4]_0 = -(0.103\text{M}/\text{min})(7.43\text{min}) + (1.61\text{M}) = 0.84\text{M}$$

12. A 3.00kg sample of ice is initially at -14.23°C and 1 atm pressure. This sample is heated to 28.19°C at constant pressure. How much energy was required to heat the water? (13pts)

$$\{C_s(\text{ice}) = 2.09\text{ J}/\text{g}\cdot\text{K}; C_s(\text{water}) = 4.184\text{ J}/\text{g}\cdot\text{K}; C_s(\text{steam}) = 2.01\text{ J}/\text{g}\cdot\text{K}; \Delta H_{\text{fusion}}(\text{water}) = 6.02\text{ kJ}/\text{mol}; \Delta H_{\text{vaporization}}(\text{water}) = 40.7\text{ kJ}/\text{mol}\}$$

3 parts: heat capacity of ice -14.23°C to 0°C , ΔH_{fusion} , heat capacity of water 0°C to 28.19°C

$$(2.09\text{ J}/\text{g}\cdot\text{K})(3000\text{g})(14.23^\circ\text{C}) = 89200\text{ J} = 89.2\text{ kJ}$$

$$(6.02\text{ kJ}/\text{mol})(3000\text{g} / 18.015\text{ g}/\text{mol}) = 1000\text{ kJ}$$

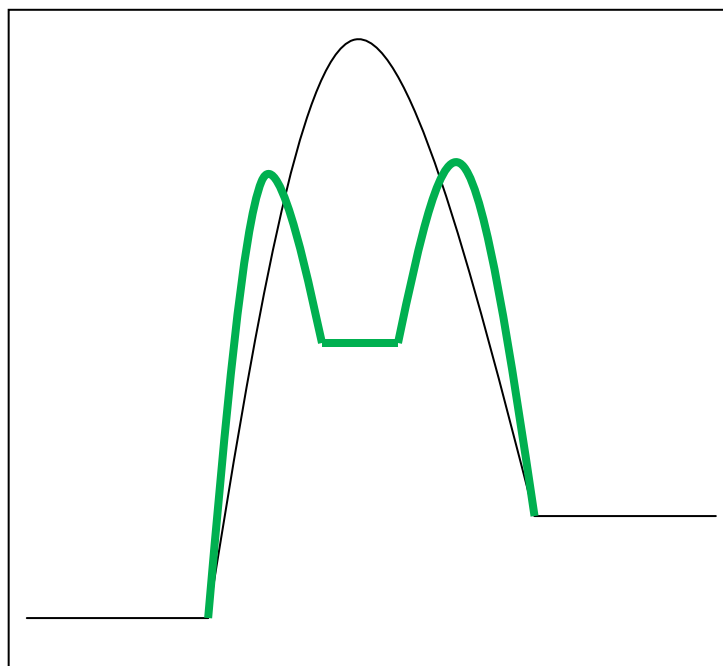
$$(4.184\text{ J}/\text{g}\cdot\text{K})(3000\text{g})(28.19^\circ\text{C}) = 3.54 \times 10^5\text{ J} = 354\text{ kJ}$$

$$\text{Total} = 1440\text{ kJ}$$

13. The reaction coordinate diagram at the right represents an uncatalyzed reaction. Modify the diagram to represent the same reaction in the presence of a catalyst and describe why you made any changes. (10pts)

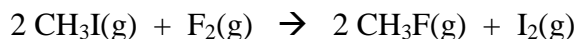
Addition of a catalyst lowers the activation energy of a reaction, usually by changing the mechanism by adding intermediates

↑
E



Reaction Coordinate →

14. For the reaction:



You have collected the following data at 38.42°C:

Experiment	$[\text{CH}_3\text{I}]_0$	$[\text{F}_2]_0$	Initial Rate
1	0.412	0.397	$4.91 \times 10^{-3} \text{ M/sec}$
2	0.824	0.397	$9.82 \times 10^{-3} \text{ M/sec}$
3	0.412	0.794	$4.91 \times 10^{-3} \text{ M/sec}$

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

If you redo Experiment 3 at 22.68°C, the rate slows to $3.83 \times 10^{-3} \text{ M/sec}$. What is the activation energy for this reaction? (25pts)

Comparing Expt 1 & 2, the concentration of methane doubles and the initial rate doubles, 1st order w.r.t. methane
Comparing Expt 1 & 3, the concentration of fluorine doubles and the initial rate is unchanged, 0th order w.r.t. fluorine

$$\text{Rate}_o = k [\text{CH}_4]_o$$

$$k = \text{Rate}_o / [\text{CH}_4]_o = (4.91 \times 10^{-3} \text{ M/sec}) / (0.412 \text{ M}) = 1.19 \times 10^{-2} \text{ sec}^{-1}$$

$$\text{Plugging in the new conditions for experiment 3, } k = (3.83 \times 10^{-3} \text{ M/sec}) / (0.412 \text{ M}) = 9.30 \times 10^{-3} \text{ sec}^{-1}$$

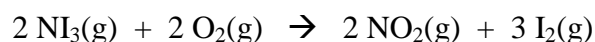
Plugging in to the comparative form of the Arrhenius equation:

$$\ln (1.19 \times 10^{-2} \text{ sec}^{-1} / 9.30 \times 10^{-3} \text{ sec}^{-1}) = (E_a / 8.314 \text{ J/mol.K}) (1/295.83 \text{ K} - 1/311.57 \text{ K})$$

$$E_a = 1.20 \times 10^4 \text{ J/mol} = 12.0 \text{ kJ/mol}$$

15. Triiodoamine (NI_3) reacts with oxygen to form nitrogen dioxide and iodine. Under some set of conditions at some point in time, you find that 0.137 mols of oxygen react every second in a 500.0 mL vessel. (25pts)

- What is the rate of oxygen consumption?
- What is the rate of NI_3 consumption?
- What is the rate of nitrogen dioxide production?
- What is the rate of iodine production?
- What is the rate of the *reaction*?



$$\text{Rate}_{\text{O}_2} = (0.137 \text{ mols} / 0.5000 \text{ L}) / 1 \text{ sec} = 0.274 \text{ M/sec}$$

$$\text{Rate}_{\text{NI}_3} = (2 \text{ mol NI}_3 / 2 \text{ mol O}_2) (\text{Rate}_{\text{O}_2}) = 0.274 \text{ M/sec}$$

$$\text{Rate}_{\text{NO}_2} = (2 \text{ mol NO}_2 / 2 \text{ mol O}_2) (\text{Rate}_{\text{O}_2}) = 0.274 \text{ M/sec}$$

$$\text{Rate}_{\text{I}_2} = (3 \text{ mol I}_2 / 2 \text{ mol O}_2) (\text{Rate}_{\text{O}_2}) = 0.411 \text{ M/sec}$$

$$\text{Rate}_{\text{rxn}} = (1 \text{ mol "rxn"} / 2 \text{ mol O}_2) (\text{Rate}_{\text{O}_2}) = 0.137 \text{ M/sec}$$