Chemistry 112-2014

Exam #2, Section 4

**Equations and Constant** 

$$\begin{split} & [A]_t = [A]_0 - kt & t_{1/2} = \frac{[A]_0}{2k} & \Delta S_{\text{phase change}} = \frac{\Delta H_{\text{phase change}}}{T_{\text{phase change}}} \\ & \ln \frac{[A]_t}{[A]_0} = -kt & t_{1/2} = \frac{0.693}{k} & \Delta H_{\text{rxn}}^\circ = \Sigma \Delta H_f^\circ (\text{products}) - \Sigma \Delta H_f^\circ (\text{reactants}) \\ & \Delta S^\circ_{\text{rxn}} = \Sigma S^\circ (\text{products}) - \Sigma S^\circ (\text{reactants}) \\ & \Delta S^\circ_{\text{rxn}} = \Sigma S^\circ (\text{products}) - \Sigma \Delta G_f^\circ (\text{reactants}) \\ & \Delta G^\circ = \Sigma \Delta G_f^\circ (\text{products}) - \Sigma \Delta G_f^\circ (\text{reactants}) \\ & \Delta G^\circ_{\text{rxn}} = \Delta H_{\text{rxn}}^\circ - T \Delta S^\circ_{\text{rxn}} \\ & \ln \frac{k_2}{k_1} = \frac{-E_a}{R} \left(\frac{1}{T_2} - \frac{1}{T_1}\right) & \Delta S_{\text{universe}} = \Delta S_{\text{system}} + \frac{-\Delta H_{\text{rxn}}}{T} \\ & k = Ae^{-E_a/RT} & \text{R} = 8.314 \text{ J/K-mol} \end{split}$$

${f 1.}$ For each pair, choose the species with the greater entropy:		NO(g)	or	NO <sub>2</sub> (g)
		H <sub>2</sub> O(g)	or	H <sub>2</sub> O(I)
a. NO and H <sub>2</sub> O(g)	b. NO and $H_2O(I)$			

c. NO<sub>2</sub>(g) and H<sub>2</sub>O(g) d. NO<sub>2</sub>(g) and H<sub>2</sub>O(l)

2. In which temperature range will the following reaction be thermodynamically most favored:

$$O=O(g) + O(g) \rightarrow O=O-O(g)$$
  
a. at all temperatures b. at no temperatures c. at high temperatures d. at low temperatures

3. What thermodynamic term controls why oil and water do not mix?

a. enthalpy change for water is disfavored	c. entropy change for water is disfavored
b. enthalpy change for oil is disfavored	d. entropy change for oil is disfavored

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

4. Consider the reaction:  $CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$ 

The reaction does not occur at room temperature. What can be said about thermodynamic control of the reaction?

- a. it is disfavored and enthalpy controlled
- b. it is disfavored and entropy controlled
- c. it is favored and enthalpy controlled
- d. it is favored and entropy controlled
- 5. For a reaction to actually occur, it must be favored by:

a. thermodynamics	c. either thermodynamics or kinetics
b. kinetics	d. both thermodynamics and kinetics

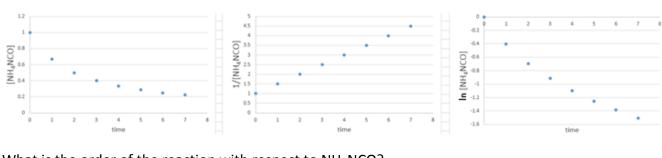
6. Initial rate data was found for the following reaction:

$$2 UO_2^+ + 4 H^+ \longrightarrow U^{4+} + UO_2^{2+} + 2 H_2O$$

Experiment	[ <b>UO</b> <sub>2</sub> <sup>+</sup> ] <sub>0</sub> , M	[ <b>H</b> <sup>+</sup> ] <sub>0</sub> , M	Initial Rate, Ms <sup>-1</sup>
1	2.85E-3	0.452	5.29
2	5.70E-3	0.452	21.1
3	2.85E-3	0.904	10.6
4	5.70E-3	0.904	42.3

What is the rate law for the reaction?

a. rate =  $k[UO_2^+][H^+]$ b. rate =  $k[UO_2^+]^2[H^+]^4$  c. rate =  $k[UO_2^+]^2[H^+]$ d. rate =  $k[UO_2^+][H^+]/[U^{4+}][UO_2^{2+}]$  7. The reaction  $NH_4NCO \rightarrow (NH_2)_2CO$  was performed and data obtained for the concentration of  $NH_4NCO$  over time. The following plots were made from the data.



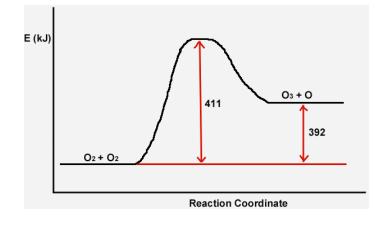
What is the order of the reaction with respect to NH<sub>4</sub>NCO?

b. first order

a. zero order

d. cannot be determined from the plots alone

8. Using the reaction coordinate diagram below, determine the activation energy and type of enthalpy change for the reaction.



c. second order

a.  $E_a = 392$  kJ, endothermicc.  $E_a = 411$  kJ, endothermicb.  $E_a = 392$  kJ, exothermicd.  $E_a = 411$  kJ, exothermic

9. The reaction 2 HI  $\rightarrow$  H<sub>2</sub> + I<sub>2</sub> is second order in [HI]. If the initial concentration of HI is

0.200 M and drops to 0.168 M in 125 seconds, what is the rate constant?

a.  $0.00760 \text{ s}^{-1}$  b.  $0.95 \text{ s}^{-1}$  c.  $1.05 \text{ s}^{-1}$  d.  $83.4 \text{ s}^{-1}$ 

10. Two mechanisms are proposed for the reaction,

$(CH_3)C=CH_2 + H_2O \rightarrow (CH_3)_3COH$				
Mechanism A:	Step 1. (CH <sub>3</sub> )C=CH <sub>2</sub> + H <sub>2</sub> O $\rightarrow$ (CH <sub>3</sub> ) <sub>3</sub> COH			
Mechanism B:	Step 1. $(CH_3)C=CH_2 + H^+ \rightarrow (CH_3)_3C^+$ (slow)			
	Step 2. $(CH_3)_3C^+ + H_2O \rightarrow (CH_3)_3COH$ (fast)			

The experimentally determined rate law is: Rate =  $k[(CH_3)C=CH_2][H^+]$ 

Which mechanism is supported by the experimental evidence?

a. Mechanism A b. Mechanism B c. I	either mechanism d. both mechanisms
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11. What is the standard entropy change for the following reaction?			S°:	
4 HCl(g) +	- O₂(g) →2 H₂O(g) + 2 C	Cl <sub>2</sub> (g)		HCI: 186 J/K O₂: 205 J/K
а. +129 J/К	b129 J/K	c. +20	d20 J/K	H <sub>2</sub> O:189 J/K Cl <sub>2</sub> : 223 J/K

12. Use the following reaction and the data given to calculate  $\Delta G^{\circ}$  at 228 °C.

2 H<sub>2</sub>S(g) + 3 O<sub>2</sub>(g) → 2 H<sub>2</sub>O(l) + 2 SO<sub>2</sub>(g)  $\Delta H^{\circ}$  = -1120 kJ  $\Delta S^{\circ}$  = -391 J/K

a. -1030 kJ b. -924 kJ c. +88000 kJ d. +195000 kJ

13. Cesium-137 is radioactive, with a half-life of 30 years. If 500 g of Cs-137 was emitted during Chernobyl nuclear disaster in the year 1986, what mass of Cs-137 will remain in the year 2046?

a. 67 g	b. 125 g	c. 250 g	d. 375 g