Constants and Equations: R = 8.31 J/mol-K **Beer-Lambert Law:** $A = log\left(\frac{I_o}{I}\right) = \varepsilon bc$ **Michaelis-Menten Equation:** $v_0 = \frac{V_m[S]}{K_M + [S]}$



CHEM 1423 - Exam 1 – February 11, 2016

Name_____

(60) **PART I. MULTIPLE CHOICE (Circle the ONE correct answer)**

- 1. Consider the hypothetical reaction, $2A + B \rightarrow 3C$. If the rate of change of [C] is Δ [C]/dt = +0.60 M hr⁻¹. What is the rate of change of [A], Δ [A]/dt ?
 - (A) -0.20 M hr^{-1} (B) -0.40 M hr^{-1}
 - (C) -0.30 M hr^{-r} (D) $+0.40 \text{ M hr}^{-1}$
- 2. The rate of the chemical reaction involving two substances, A and B, is measured. It is found that if the initial concentration of A used is tripled, keeping the B concentration the same, the rate increases by a factor of 27 (relative to the first experiment). If the concentrations of both A and B are doubled, the rate increases by a factor of 32 (relative to the first experiment). The rate law for this reaction is: Rate =
 - (A) $k[A]^{2}[B]^{3}$ (B) $k[A]^{2}[B]^{2}$ (C) $k[A]^{3}[B]$ (D) $k[A]^{3}[B]^{2}$

For #3 - #4: Consider a reaction, $A \rightarrow$ Products, which is of order "n"; i.e. Rate = k[A]ⁿ. For this reaction, the following initial rate data was obtained.

When $[A]_{o} = 0.20$ M, the initial rate is 0.50 M/s

When $[A]_{\circ} = 0.60$ M, the initial rate is 13.5 M/s

- 3. The order of this reaction (i.e. "n") is:
 - (A) +3 (B) +2 (C) +1 (D) -1
- 4. The rate constant for this reaction (i.e. "k") is:

(A) $313 \text{ M}^{-3}\text{s}^{-1}$ (B) $12.5 \text{ M}^{-1}\text{s}^{-1}$ (C) $62.5 \text{ M}^{-2}\text{s}^{-1}$ (D) 2.5 s^{-1}

- 5. Radioactive decay follows first order kinetics. Thallium-201 is a radioisotope which is used for myocardial imaging tests. Its half-life is 73 hours. If 300 mg of ²⁰¹TI is injected into a patient, how many mg will remain 185 hours after the injection ?
 - (A) 124 mg (B) 52 mg (C) 34 mg (D) 83 mg.
- For the zeroth order reaction, A → Products, when the initial concentration of A is 0.90 M, it takes 25 s for the concentration to decrease to 0.30 M. Therefore, the rate constant for this reaction is approximately:

```
(A) 0.044 Ms<sup>-1</sup> (B) 0.089 Ms<sup>-1</sup> (C) 0.015 Ms<sup>-1</sup> (D) 0.024 Ms<sup>-1</sup>
```

For #7 - #8: Consider a reaction, $A \rightarrow$ Products, which is of **second** order; i.e. Rate = k[A]². For this reaction, the rate constant is 8.5x10⁻³ M⁻¹s⁻¹.

7. For this reaction, a plot of _____ vs. time is a straight line with a _____ slope.

(A)	In([A] _t) , negative	(B) $[A]_t$, negative
(C)	1/(A]t), negative	(D) 1/[A]t, positive

8. If the initial concentration of A is 0.75 M, what will be the approximate concentration of A 600 seconds after the start of the reaction.

(A) 0.20 M (B) 0.16 M (C) 0.28 M (D) 0.046 M

9. Which **one** statement below is correct? The frequency factor in the Arrhenius equation can best be described as:

(A) A numerical description of the amount of energy released by colliding reactant molecules when they form products.

(B) The probability that a collision between molecules will have the correct relative orientation for reaction to occur.

(C) A numerical description of how often molecules collide with the proper orientation to react at a specific concentration.

(D) A numerical description of the amount of energy needed by colliding reactant molecules in order to form products.

- 10. Which of the following statements is/are NOT true?
 - (1) The mechanism for a catalyzed reaction is different from the mechanism of the same reaction without the catalyst.
 - (2) The enthalpy change of the reaction is lower for the catalyzed reaction than for the uncatalyzed chemical reaction.
 - (3) The catalyst in a reaction is present at the start of the reaction and is regenerated during the reaction.
 - (4) The Rate Determining Step in a catalyzed reaction mechanism has a higher activation energy and, therefore, is faster than the Rate Determining Step for the uncatalyzed reaction.
 - (A) 2 & 4 (B) 2 & 3 & 4 (C) 1 & 3 (D) 2 & 4

11. For the reaction, $Cd_2^{2+}(aq) + Zr^{3+}(aq) \rightarrow 2 Cd^{2+}(aq) + Zl^{+}(aq)$, the accepted reaction mechanism is:

$$Cd_2^{2+} \xrightarrow{\kappa} Cd + Cd^{2+}$$
 Fas

Fast Pre-Equilibrium

 $Zr^{3+} + Cd \xrightarrow{k_2} Zr^+ + Cd^{2+}$ Slow Rate Determining Step An acceptable rate law for the rate, Rate = $\Delta[Zr^+]/\Delta t$, is:

(A)
$$Rate = k' \frac{[Zr^{3+}][Cd_2^{2+}]}{[Cd]}$$
 (B) $Rate = k' \frac{[Zr^{3+}][Cd^{+}]^2}{[Cd^{2+}]}$
(C) $Rate = k'[Zr^{3+}][Cd_2^{2+}][Cd^{2+}]$ (D) $Rate = k' \frac{[Zr^{3+}][Cd_2^{2+}]}{[Cd^{2+}]}$

12. Consider a reaction, R → P (i.e. Reactants → Products). If the activation energy for the reverse reaction is 85 kJ/mol and the overall energy (aka enthalpy) change for the reaction is -30 kJ/mol. Therefore, the activation energy for the forward reaction is:

(A) +115 kJ/mol (B) +55 kJ/mol (C) +105 kJ/mol (D) -55 kJ/mol

13. In an enzyme catalyzed reaction, when the Substrate Concentration is much greater than the Michaelis-Constant, K_M, then the rate of the reaction is _____ order with respect to [S]

(A) Second order (B) First Order (C) Zeroth Order (D) Indeterminate without know the values of
$$K_M$$
 and [S].

- 14. In an enzyme catalyzed reaction, for what ratio, [S]/K_M, is the initial velocity, $v_0 = 0.70 \text{ V}_m$?
 - (A) 1.75 (B) 0.43 (C) 2.33 (D) 3.41

15. Consider the gas phase equilibrium, $A(g) \Rightarrow B(g) + 3C(g)$,

 $K_c = 1.0 \times 10^{-5}$. 3.0 mol of A(g) is placed in a 2.0 L container and the mixture is allowed to come to equilibrium. Calculate the approximate concentration of C(g) at equilibrium.

NOTE: You can assume that very little A(g) reacts to form B(g) and C(g)

(A) $8.2x10^{-2}$ M (B) $1.1x10^{-1}$ M (C) $2.7x10^{-2}$ M (D) $9.7x10^{-2}$ M

For #16 - #20 For the gas phase reaction, $2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$, $K_c = 120$. at 600 K. The Enthalpy change for this reaction is +95 kJ/mol

16. If a mixture is prepared with [SO₂] = 0.4 M, [O₂] = 0.3 M and [SO₃] = 1.5 M, the reaction quotient is _____ and the reaction will proceed towards the _____.

(A) 47., Right	(B) 188., Left
----------------	----------------

(C) 63., Right (D) 188., Right



PART II. THERE ARE FOUR (4) PROBLEMS BELOW.

(10) 1. The measured value of the rate constant for a first order reaction is 1.50x10³ min⁻¹ at a temperature of 120 °C, and 1.35x10⁵ min⁻¹ at 180 °C. Calculate the Arrhenius Activation Energy (in kJ/mol) and the Frequency Factor (in min⁻¹) for this reaction.

(10) 2. Consider a molecule, B, which reacts to form Products via First Order kinetics:

$$A \rightarrow P$$
 $R = -\frac{d[A]}{dt} = k[A]$

The molecule, B, absorbs visible light at 480 nm. The Molar Absorptivity for this absorption is given by $\varepsilon = 800 \text{ M}^{-1} \text{ cm}^{-1}$.

When a sample of B is prepared in a 0.5 cm cell, the Percent Transmission at the beginning of the reaction was 25%. After the reaction has run for 50 minutes, the percent transmission is 70%. Calculate the rate constant for this reaction (give the units)

(10) 3. Consider the reaction: $Cl_2(g) + 2 NO_2(g) \Rightarrow 2 NOCI(g)$.

If 1.50 mol of $Cl_2(g)$ and 1.50 mol of $NO_2(g)$ are placed in a 5.0 Liter container, which is heated to 200 $^{\circ}$ C, the equilibrium concentration of $NO_2(g)$ is 0.120 M. Calculate the equilibrium constant, K_c, for this reaction at 200 $^{\circ}$ C.

(10) 4. Consider the reaction: $H_2(g) + I_2(g) \xleftarrow{K_c} 2HI(g)$ K_c = 9.0 (at 550 K) 8.0 mol of HI(g) is placed in a 20.0 L container and heated to 550 K, where equilibrium is established. Calculate the equilibrium concentrations (**in M**) of H₂(g), I₂(g) and HI(g) ?