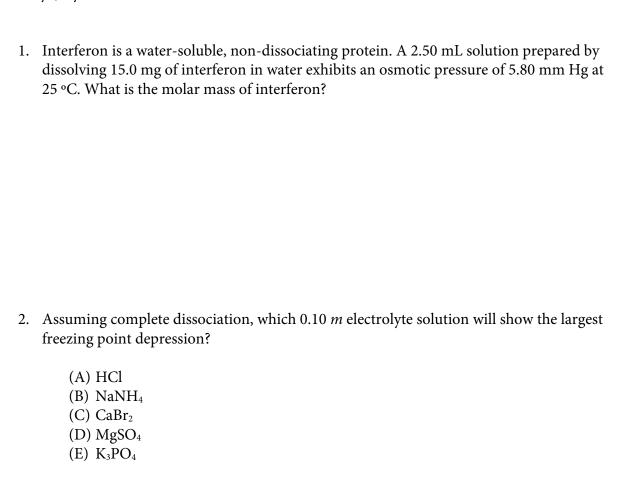
## Mock Exam I



3. The half-life of a first-order reaction is 1.5 hours. How much time is needed for 94% of

the reactant to change to product?

4. When the kinetics of the reaction,  $2A + 2B \rightarrow C$  were studied using the method of initial rates, the data in the table below were obtained.

Trial	$[A]_0(M)$	$[B]_0(M)$	Initial Rate of
			Formation of C (M/s)
1	0.060	0.040	3.6 x 10 <sup>-4</sup>
2	0.060	0.080	7.2 x 10 <sup>-4</sup>
3	0.030	0.120	5.4 x 10 <sup>-4</sup>

What is the rate law for the reaction?

5. For the reaction  $5O_{2(g)} + 4NH_{3(g)} \rightarrow 4NO_{(g)} + 6H_2O_{(g)}$ , if NH<sub>3</sub> is being consumed at a rate of 0.50 M/s, at what rate is H<sub>2</sub>O being formed?

6. A solution of 5.00 g of which ionic solid, in 1 L of solution at 25 °C, has the largest osmotic pressure?

7. A substance XY decomposes in a second-order reaction. A solution that is initially 1.00 M in XY requires 0.50 hours for its concentration to decrease to 0.50 M. How much time will it take for a solution of XY to decrease in concentration from 2.00 M to 0.25 M?

## Passage II (Questions 8-14)

A chemist interested in the reactivity of iodine concentrated his study on two reactions: the decomposition of gaseous hydrogen iodide (Reaction 1) and the reaction between iodide ions and persulfate ions (Reaction 2).

$$2HI(g)^{\frac{1}{r}}H_{2}(g) + I_{2}(g)$$

Reaction 1

$$3I^{-}(aq) + S_2O_8^{2-}(aq) \stackrel{\dashv}{\vdash} I_3^{-}(aq) + 2SO_4^{2-}(aq)$$

## Reaction 2

The value of the rate constant for Reaction 1 was studied as a function of temperature. The results are shown below.

Table 1

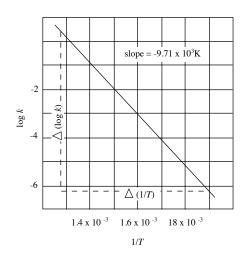
T(K)	$1/T (K^{-1})$	$k (l \cdot mol^{-1} sec^{-1})$	log k
555	1.80 ×	$3.52 \times 10^{-7}$	-6.453
575	$10^{-3}$	$1.22 \times 10^{-6}$	-5.913
645	1.74 ×	$8.59 \times 10^{-5}$	-4.066
700	$10^{-3}$	$1.16 \times 10^{-3}$	-2.936
781	1.55 ×	$3.95 \times 10^{-2}$	-1.403
	$10^{-3}$		
	1.43		
	$\times 10^{-3}$		
	1.28 ×		
	$10^{-3}$		

For any reaction, the activation energy  $(E_{\rm a})$  is related to the rate constant (k) by the Arrhenius equation (Equation 1):

$$k = A \times 10^{(-E_a / 2.303RT)}$$

## Equation 1

where R = 8.314 J mol<sup>-1</sup>K<sup>-1</sup>, T is the temperature in Kelvin, and A is a constant, called the frequency factor. Figure 1 shows a graph of log k vs. 1/T for Reaction 1.



In order to determine the initial rate of Reaction 2, the following data were collected:

Table 2

Experiment	[I <sup>-</sup> ] (M)	$[S_2O_8^{-2}]$ (M)	Initial rate of reaction (M / sec)
1	0.21	0.15	1.14
2	0.21	0.30	2.28
3	0.42	0.15	2.28

**8.** What is the rate law for Reaction 2?

**A**. Rate = 
$$k[I^-]^2[S_2O_8^{2-}]$$

**B**. Rate = 
$$k[S_2O_8^{2-}]$$

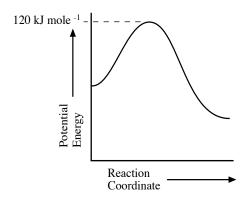
**C**. Rate = 
$$k[I^-][S_2O_8^{2-}]$$

**D**. Rate = 
$$k[I^-][S_2O_8^{2-}]^2$$

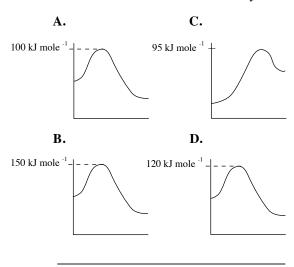
- 9. What is the numerical value of the rate constant for Reaction 2?
  - A. 7.6 mol/L•sec
  - B. 36 mol/L•sec
  - C. 172 mol/L•sec
  - D. 241 mol/L•sec

- **10.** According to Equation 1:
  - A. when the temperature is held constant, the lower the activation energy, the slower the reaction.
  - B. when the activation energy is held constant, the lower the value of A, the faster the reaction.
  - C. when the activation energy is held constant, the lower the temperature, the faster the reaction.
  - **D**. when the temperature is held constant, the lower the activation energy, the faster the reaction.
- **11.** What is the activation energy for Reaction 1?
  - 9.71 kJ/mole
  - **B**. 22.4 kJ/mole
  - C. 80.7 kJ/mole
  - **D**. 186 kJ/mole
- 12. In Figure 1, what does the intercept with the y-axis represent?
  - $\mathbf{A} \cdot \log A$
  - $\mathbf{B} \cdot \log k$
  - $\mathbf{C} \cdot -E_a$
- 13. If the rate of disappearance of  $I^-$  in Reaction 2 is 2.5  $\times$  $10^{-3}$  mol/(L•s), what is the rate of formation of  $SO_4^{2-}$ ?
  - **A** .  $1.7 \times 10^{-3}$  mol/(L•s) **B** .  $3.8 \times 10^{-3}$  mol/(L•s) **C** .  $5.0 \times 10^{-3}$  mol/(L•s) **D** .  $8.3 \times 10^{-4}$  mol/(L•s)

14. The reaction profile shown below is for an uncatalyzed reaction.



Which of the following is the reaction profile for the same reaction after the addition of a catalyst?



15. A reaction has a rate constant  $k = 8.54 \times 10^{-4} \text{ M}^{-1} \text{s}^{-1}$  at 45 °C, and an activation energy  $E_a = 90.8 \text{ kJ}$ . What is the value of k at 25 °C?

16. What is the mole fraction of CH₃OH in an aqueous solution that is 12.0 *m* in CH₃OH?

17. For the reaction,  $2A + B \rightarrow C + D$ , the rate law is: Rate = k[B].

Which of the following mechanisms is consistent with this information?

(A) 
$$A + B \rightarrow M$$
 (slow)  
 $A + M \rightarrow C + D$  (fast)

(B) 
$$A + A \rightarrow M$$
 (fast)  
  $B \rightarrow C + D$  (slow)

(C) 
$$B \rightarrow M$$
 (slow)  
 $A + M \rightarrow N$  (fast)  
 $N + A \rightarrow C + D$  (fast)

(D) 
$$B \rightarrow M$$
 (fast)  
 $A + M \rightarrow N$  (slow)  
 $N + A \rightarrow C + D$  (fast)

(E) 
$$A + A \rightarrow M$$
 (slow)  
 $M + B \rightarrow C + D$  (fast)

- 18. Which of the following decreases with increasing intermolecular forces?
  - (A) Boiling point
  - (B) Molar enthalpy of vaporization ( $\Delta_{vap}H^o$ )
  - (C) Vapor pressure
  - (D) Viscosity
  - (E) Surface tension
- 19. The Henry's Law constant for oxygen gas in water at 25 °C  $k_{O2} = 1.3 \times 10^{-3}$  M/atm. What is the partial pressure of  $O_2$  above a solution at 25 °C with an  $O_2$  concentration of 2.3 x  $10^{-4}$  M at equilibrium?

- 20. Which mixture of water and H<sub>2</sub>SO<sub>4</sub> represents a solution with a concentration that is closest to 30% by mass H<sub>2</sub>SO<sub>4</sub>?
  - (A)  $30 g H_2SO_4 + 100 g H_2O$
  - (B)  $1 \text{ mol } H_2SO_4 + 200 \text{ g } H_2O$
  - (C)  $30 \text{ mol } H_2SO_4 + 0.70 \text{ kg } H_2O$
  - (D)  $0.30 \text{ mol } H_2SO_4 + 0.70 \text{ mol } H_2O$
  - (E)  $0.30 \text{ mol } H_2SO_4 + 100 \text{ mol } H_2O$
- 21. The vapor pressure of water at 31 °C is 33.7 mmHg. When you dissolve 931.0 g of acetone  $(CH_3COCH_3)$  in 32.50 kg of water, what is the vapor pressure of water over the solution at 31 °C?

22. The data below were collected for this reaction at 500 °C: CH3CN (g)  $\rightarrow$  CH3NC (g)

Time (hr)	[CH <sub>3</sub> CN] (M)
0.0	1.000
5.0	0.794
10.0	0.631
15.0	0.501
20.0	0.393
25.0	0.316

a) What is the order of the reaction? Please briefly explain your reasoning.

b) What is the value of the rate constant at this temperature?

c) What is the half-life for this reaction (at the initial concentration)?

d) How long will it take for 90% of the CH3CN to convert to CH3NC?

23. During the winter, much of the salt you see melting ice is calcium chloride, which has	s a
solubility of 74.5 g per 100.0g of cold water. What is the lowest temperature ice that	
calcium chloride salt can melt assuming complete dissociation?	

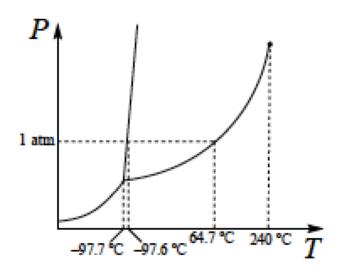
24. The data shown below were collected for the second-order reaction:

$$Cl_2(\mathbf{g}) + 2 H_2(\mathbf{g}) \rightarrow 2 HCl(\mathbf{g}) + H_2(\mathbf{g})$$

$\operatorname{Cl}_{2}\left(\mathbf{g}\right)+2\operatorname{H}_{2}\left(\mathbf{g}\right) \rightarrow 2\operatorname{HCl}\left(\mathbf{g}\right)+\operatorname{H}_{2}\left(\mathbf{g}\right)$		
Temp. (K)	Rate Constant (M <sup>-1</sup> s <sup>-1</sup> )	
90	0.00357	
100	0.0773	
110	0.956	

Please determine the activation energy and frequency factor for the reaction.

25. Below is the phase diagram of methanol:



- a) True or False:
  - i. \_\_\_\_\_ Solid methanol is denser than liquid methanol.
  - ii. \_\_\_\_\_ Solid methanol sublimes at atmospheric pressure.
- iii. Solid, liquid, and gaseous methanol cannot coexist at atmospheric pressure.
- iv. \_\_\_\_ It is possible to have liquid methanol at -100 °C.
- v. \_\_\_\_ It is possible to have gaseous methanol at 1 atm.
- b) What is the boiling point of methanol at atmospheric pressure? \_\_\_\_\_
- c) What is the freezing point of methanol at atmospheric pressure? \_\_\_\_\_