# Different professors are covering different material in their classes. Therefore, the questions in this review are separated by chapter.

### **Chapter 16: Kinetics, Rate Laws**

16.1 Cyclobutane decomposes in a first order reaction shown below.

$$C_4H_{8(g)} \rightleftharpoons 2C_2H_{4(g)}$$

Given that the initial concentration of  $C_4H_8$  is 5M and the final concentration is 0.06M after 0.05 seconds, what is the rate constant and the expected rate law?

- a. 65 s<sup>-1</sup>; rate =  $k[C_2H_4]^2$
- b. <u>88 s<sup>-1</sup></u>; rate =  $k[C_4H_8]$
- c. 92 s<sup>-1</sup>; rate =  $k[C_4H_8]$
- d. 88 s<sup>-1</sup>; rate =  $k[C_2H_4]^2$
- e. 65 s<sup>-1</sup>; rate =  $k[C_4H_8]$
- f. 92 s<sup>-1</sup>; rate =  $k[C_2H_4]^2$

16.2 The decomposition of hydrogen bromide is shown below.

$$2\text{HBr}_{(g)} \rightleftharpoons \text{H}_{2(g)} + \text{Br}_{2(g)}$$

At 600 K, the rate constant is  $1.50 \times 10^{-5}$  L/mol s. At 800 K, the rate constant is  $6.80 \times 10^{-3}$  L/mol s. What is the activation energy for this reaction?

- a.  $1.82 \times 10^2 \text{ kJ/mol}$
- b.  $1.39 \times 10^2 \text{ kJ/mol}$
- c.  $1.58 \times 10^2 \text{ kJ/mol}$
- d. <u>1.22×10<sup>2</sup> kJ/mol</u>
- e. 2.02×10<sup>2</sup> kJ/mol

16.3 Using the information in the previous question (16.2), determine what temperature should be used if you wanted the rate to be three times as fast as it is at 600K?

- a. 576 K
- b. 582 K
- c. <u>626 K</u>
- d. 654 K
- e. 730 K

16.4 A fictional reaction is shown below with its rate law.

$$2A + B + C \rightarrow 3D$$
, rate = k[A]<sup>2</sup>[B][C]

Which of the following mechanisms can be considered possible mechanisms for this reaction and rate law?

I: 
$$A + B \rightleftharpoons X$$
 (fast)II:  $2A + B + C \rightarrow 3D$  (one step)III:  $B + C \rightleftharpoons X$  (slow) $X + A + C \rightarrow 3D$  (slow) $M + 2A \rightarrow 3D$  (fast)

- a. I only
- b. II only
- c. III only
- d. I and II
- e. I, and III
- f. I, II, and III

## Chapter 17: Chemical Equilibrium, K & Q, Le Chatelier's Principle

17.1 Using Le Chatelier's Principle, match the correct shift for the given change.

## Change

- a) Increase in Concentration of X
- b) Decrease in Concentration of X
- c) Increase in Pressure
- d) Decrease in Pressure
- e) Increase in Temperature
- f) Decrease in Temperature
- g) Catalyst

### Shift

- I. No shift
- II. Away from heat
- III. Away from substance X
- IV. Toward Heat
- V. Towards fewer moles of gas
- VI. Towards substance X
- VII. Towards more moles of gas

17.2 Ammonia is created on an industrial scale using the Haber-Bosch process shown below.

$$N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$$
  $\Delta H = -92.4 \text{ kJ} * \text{mol}^{-1}$ 

Predict the shift (left, right, or none) for each given change.

Change	Shift
Increase [NH <sub>3</sub> ]	Left
Decrease [N <sub>2</sub> ]	Left
Decrease [NH <sub>3</sub> ]	Right
Increase Pressure	Right
Increase Temperature	Left
Catalyst	No shift
Increase [H <sub>2</sub> ]	Right

17.3 A fictional reaction is shown below.

 $A_{(q)} + B_{(q)} \rightleftharpoons 3C_{(q)}$   $\Delta H = +36 \text{ kJ mol}^{-1}$ 

To favor the production of the product (C), under what conditions should this reaction be carried out?

a. High pressure, Low temperature

b. High pressure, High temperature

c. Low pressure, High temperature

d. Low pressure, Low temperature

17.4 Calculate Kc given the following reaction.

$$PCl_{3(g)} + Cl_{2(g)} \rightleftharpoons PCl_{5(g)}$$
  $K_p = 4.07 \times 10^{-1} (at 226.85^{\circ}C)$ 

If the Qc was 20.5, is the reaction at equilibrium? Or will it proceed to the left or to the right?

a. Kc=16.7, proceed left until Qc=Kc

b. Kc=38.5, proceed right until Qc=Kc

c. Kc=20.5, the reaction is at equilibrium

d. Kc=30.2, proceed right until Qc=Kc

17.5 The decomposition of phosgene is shown in the reaction below.

 $\text{COCl}_{2(g)} \rightleftharpoons \text{CO}_{(g)} + \text{Cl}_{2(g)}$   $K_c = 6.3 \times 10^{-4} \text{ (at 500K)}$ 

Calculate the equilibrium partial pressures of  $COCl_2$ , CO, and  $Cl_2$  when the initial partial pressure of  $COCl_2$  is 0.28 atm.

a.  $COCl_2=0.5$  atm, CO=0.24 atm,  $Cl_2=0.014$  atm

- b. COCl<sub>2</sub>=0.071 atm, CO=Cl<sub>2</sub>=0.85 atm
- c. COCl<sub>2</sub>=0.26 atm, CO= Cl<sub>2</sub>=0.024 atm
- d.  $COCl_2=0.29$  atm, CO=0.21 atm,  $Cl_2=0.79$  atm
- e. <u>COCl<sub>2</sub>=0.21 atm, CO= Cl<sub>2</sub>=0.073 atm</u>

#### Chapter 18: Acid-Base Equilibrium (Not all professors are covering Chapter 18)

18.1 Calculate the  $[H_3O^+]$ , pH,  $[OH^-]$ , and pOH, of 0.5 M of  $H_2SO_4$ .

a. [H<sub>3</sub>O<sup>+</sup>]= 0.5 M, pH=0.30, [OH<sup>-</sup>]=2×10<sup>-14</sup> M, pOH=13.70

b.  $[H_3O^+]=0.25M$ , pH=0.60,  $[OH^-]=5\times10^{-10}$  M, pOH=10.5

c. [H<sub>3</sub>O<sup>+</sup>]=0.15 M, pH=0.90, [OH<sup>-</sup>]=0.27 M, and pOH=13.2

d. [H<sub>3</sub>O<sup>+</sup>]=0.90 M, pH=3.1, [OH<sup>-</sup>]=8.1×10<sup>-5</sup> M, and pOH=10.9

18.2 Define each of the following and provide an example.

A Brønsted-Lowry acid is a proton donor. An example is HCl.

A Brønsted-Lowry base is a proton acceptor. An example is <u>NH<sub>3</sub></u>.

A Lewis acid is an <u>electron pair acceptor</u>. An example is <u>BF<sub>3</sub></u>.

A Lewis base is an <u>electron pair donor</u>. An example is  $H_2O$ .

18.3 Methylamine (CH<sub>3</sub>NH<sub>2</sub>) has a K<sub>b</sub> of  $4.4 \times 10^{-4}$ . What is the pH of 5 M CH<sub>3</sub>NH<sub>2</sub>?

- a. 10.94
- b. 7.62
- <u>c. 12.67</u>
- d. 13.63

18.4 Potassium propionate (KCH<sub>3</sub>CH<sub>2</sub>COO) is a food preservative that is most commonly used as an antimicrobial agent in bread. What is the pH of 0.38 M KCH<sub>3</sub>CH<sub>2</sub>COO at 298K? The K<sub>a</sub> of propionic acid is  $1.34 \times 10^{-5}$ .

- a. 10.13
- b. 8.34
- c. 9.86
- <u>d. 9.23</u>

18.5 Which of the following is a potential Lewis acid?

a. Cl<sup>-</sup>

b. OH-

c. BCl<sub>3</sub>

d. NH4<sup>+</sup>

## Chapter 15: Organic Chemistry (Not all professors are covering Chapter 15)

15.1 What are the names of these four structures, in order (left to right)?



a. Cyclohexane, Cyclobutane, Cycloheptane, Cyclooctane

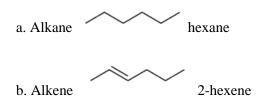
b. Cyclopropane, Cyclobutane, Cyclopentane, Cyclohexane

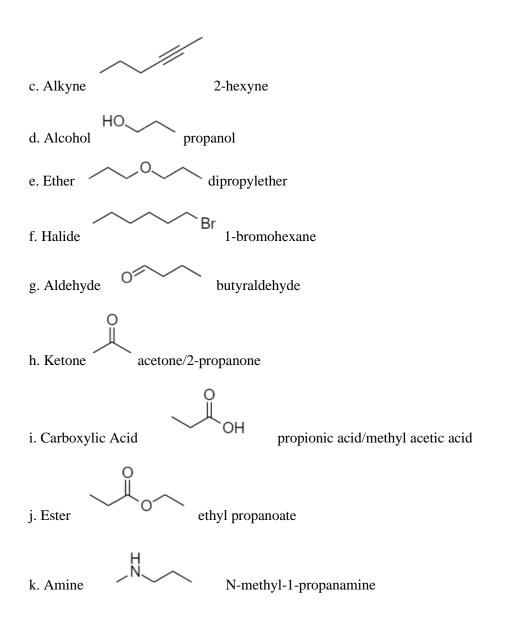
c. Cyclopropane, Cyclooctane, Cyclononane, Cyclohexane

d. Cyclobutane, Cyclopropane, Cyclohexane, Cyclononane

e. Cyclodecane, Cyclononane, Cyclopropane, Cyclohexane

15.2 Draw an example of each functional group.





15.3 C<sub>6</sub>H<sub>14</sub> has 5 different constitutional isomers. Draw each and name according to IUPAC naming rules.

Hexane

a.

b. 2-methylpentane

3-methylpentane



2,2-dimethylbutane

d.

c.

2,3-dimethylbutane

e.