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.

$$\mathsf{C_4H_8}_{(g)} \rightleftharpoons \mathsf{2C_2H_4}_{(g)}$$

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^{-1} ; rate = $k[C_2H_4]^2$
- b. 88 s^{-1} ; rate = $k[C_4H_8]$
- c. 92 s^{-1} ; rate = $k[C_4H_8]$
- d. 88 s^{-1} ; rate = $k[C_2H_4]^2$
- e. 65 s^{-1} ; rate = $k[C_4H_8]$
- f. 92 s^{-1} ; rate = $k[C_2H_4]^2$

16.2 The decomposition of hydrogen bromide is shown below.

$$2\mathsf{HBr}_{(g)} \rightleftarrows \mathsf{H}_{2(g)} + \mathsf{Br}_{2(g)}$$

At 600 K, the rate constant is 1.50×10^{-5} L/mol s. At 800 K, the rate constant is 6.80×10^{-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. 1.22×10^2 kJ/mol
- e. $2.02 \times 10^2 \text{ kJ/mol}$

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. 626 K
- 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]²[B][C]

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

- 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		Shift
a)	Increase in Concentration of X	I.	No shift
b)	Decrease in Concentration of X	II.	Away from heat
c)	Increase in Pressure	III.	Away from substance X
d)	Decrease in Pressure	IV.	Toward Heat
e)	Increase in Temperature	V.	Towards fewer moles of gas
f)	Decrease in Temperature	VI.	Towards substance X
g)	Catalyst	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 ₃]	
Decrease [N ₂]	•
Decrease [NH ₃]	
Increase Pressure	•
Increase Temperature	•
Catalyst	
Increase [H ₂]	

17.3 A fictional reaction is shown below.

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

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.

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

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.

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

Calculate the equilibrium partial pressures of COCl₂, CO, and Cl₂ when the initial partial pressure of COCl₂ is 0.28 atm.

- a. COCl₂=0.5 atm, CO=0.24 atm, Cl₂=0.014 atm
- b. COCl₂=0.071 atm, CO=Cl₂=0.85 atm
- c. COCl₂=0.26 atm, CO= Cl₂=0.024 atm
- d. COCl₂=0.29 atm, CO=0.21 atm, Cl₂=0.79 atm
- e. COCl₂=0.21 atm, CO= Cl₂=0.073 atm

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

- 18.1 Calculate the [H₃O⁺], pH, [OH⁻], and pOH, of 0.5 M of H₂SO₄.
- a. $[H_3O^+]=0.5 \text{ M}$, pH=0.30, $[OH^-]=2\times10^{-14} \text{ M}$, pOH=13.70
- b. $[H_3O^+]=0.25M$, pH=0.60, $[OH^-]=5\times10^{-10}M$, pOH=10.5
- c. $[H_3O^+]=0.15 \text{ M}$, pH=0.90, $[OH^-]=0.27 \text{ M}$, and pOH=13.2
- d. $[H_3O^+]=0.90 \text{ M}$, pH=3.1, $[OH^-]=8.1\times10^{-5} \text{ M}$, 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 NH₃.

A Lewis acid is an electron pair acceptor. An example is BF₃.

A Lewis base is an electron pair donor. An example is H₂O.

- 18.3 Methylamine (CH₃NH₂) has a K_b of 4.4×10^{-4} . What is the pH of 5 M CH₃NH₂?
- a. 10.94
- b. 7.62
- c. 12.67
- d. 13.63

18.4 Potassium propionate (KCH ₃ CH ₂ 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 ₃ CH ₂ COO at 298 K? The K_a of propionic acid is 1.34×10^{-5} .
a. 10.13
b. 8.34
c. 9.86
d. 9.23
18.5 Which of the following is a potential Lewis acid?
a. Cl ⁻
b. OH-
c. BCl ₃
d. NH ₄ ⁺
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, Cyclohexane e. Cyclodecane, Cyclononane, Cyclohexane
15.2 Draw an example of each functional group.
a. Alkane
b. Alkene
c. Alkyne
d. Alcohol

e. Ether
f. Halide
g. Aldehyde
h. Ketone
i. Carboxylic Acid
j. Ester
k. Amine
$15.3\ C_6H_{14}$ has 5 different constitutional isomers. Draw each and name according to IUPAC naming rules.
a.
b.
b.
b. c.
b.c.d.