



## **AP<sup>®</sup> Chemistry 2006 Free-Response Questions**

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**DO NOT DETACH FROM BOOK.**

# PERIODIC TABLE OF THE ELEMENTS

PERIODIC TABLE OF THE ELEMENTS

\*Lanthanide Series

† Actinide Series

## STANDARD REDUCTION POTENTIALS IN AQUEOUS SOLUTION AT 25°C

| Half-reaction                                      |               |                                       | $E^\circ(\text{V})$ |
|--|---------------|---------------------------------------|---------------------|
| $\text{F}_2(\text{g}) + 2\text{e}^-$               | $\rightarrow$ | $2\text{F}^-$                         | 2.87                |
| $\text{Co}^{3+} + \text{e}^-$                      | $\rightarrow$ | $\text{Co}^{2+}$                      | 1.82                |
| $\text{Au}^{3+} + 3\text{e}^-$                     | $\rightarrow$ | $\text{Au}(\text{s})$                 | 1.50                |
| $\text{Cl}_2(\text{g}) + 2\text{e}^-$              | $\rightarrow$ | $2\text{Cl}^-$                        | 1.36                |
| $\text{O}_2(\text{g}) + 4\text{H}^+ + 4\text{e}^-$ | $\rightarrow$ | $2\text{H}_2\text{O}(\text{l})$       | 1.23                |
| $\text{Br}_2(\text{l}) + 2\text{e}^-$              | $\rightarrow$ | $2\text{Br}^-$                        | 1.07                |
| $2\text{Hg}^{2+} + 2\text{e}^-$                    | $\rightarrow$ | $\text{Hg}_2^{2+}$                    | 0.92                |
| $\text{Hg}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Hg}(\text{l})$                 | 0.85                |
| $\text{Ag}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Ag}(\text{s})$                 | 0.80                |
| $\text{Hg}_2^{2+} + 2\text{e}^-$                   | $\rightarrow$ | $2\text{Hg}(\text{l})$                | 0.79                |
| $\text{Fe}^{3+} + \text{e}^-$                      | $\rightarrow$ | $\text{Fe}^{2+}$                      | 0.77                |
| $\text{I}_2(\text{s}) + 2\text{e}^-$               | $\rightarrow$ | $2\text{I}^-$                         | 0.53                |
| $\text{Cu}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Cu}(\text{s})$                 | 0.52                |
| $\text{Cu}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Cu}(\text{s})$                 | 0.34                |
| $\text{Cu}^{2+} + \text{e}^-$                      | $\rightarrow$ | $\text{Cu}^+$                         | 0.15                |
| $\text{Sn}^{4+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Sn}^{2+}$                      | 0.15                |
| $\text{S}(\text{s}) + 2\text{H}^+ + 2\text{e}^-$   | $\rightarrow$ | $\text{H}_2\text{S}(\text{g})$        | 0.14                |
| $2\text{H}^+ + 2\text{e}^-$                        | $\rightarrow$ | $\text{H}_2(\text{g})$                | 0.00                |
| $\text{Pb}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Pb}(\text{s})$                 | -0.13               |
| $\text{Sn}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Sn}(\text{s})$                 | -0.14               |
| $\text{Ni}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Ni}(\text{s})$                 | -0.25               |
| $\text{Co}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Co}(\text{s})$                 | -0.28               |
| $\text{Cd}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Cd}(\text{s})$                 | -0.40               |
| $\text{Cr}^{3+} + \text{e}^-$                      | $\rightarrow$ | $\text{Cr}^{2+}$                      | -0.41               |
| $\text{Fe}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Fe}(\text{s})$                 | -0.44               |
| $\text{Cr}^{3+} + 3\text{e}^-$                     | $\rightarrow$ | $\text{Cr}(\text{s})$                 | -0.74               |
| $\text{Zn}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Zn}(\text{s})$                 | -0.76               |
| $2\text{H}_2\text{O}(\text{l}) + 2\text{e}^-$      | $\rightarrow$ | $\text{H}_2(\text{g}) + 2\text{OH}^-$ | -0.83               |
| $\text{Mn}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Mn}(\text{s})$                 | -1.18               |
| $\text{Al}^{3+} + 3\text{e}^-$                     | $\rightarrow$ | $\text{Al}(\text{s})$                 | -1.66               |
| $\text{Be}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Be}(\text{s})$                 | -1.70               |
| $\text{Mg}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Mg}(\text{s})$                 | -2.37               |
| $\text{Na}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Na}(\text{s})$                 | -2.71               |
| $\text{Ca}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Ca}(\text{s})$                 | -2.87               |
| $\text{Sr}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Sr}(\text{s})$                 | -2.89               |
| $\text{Ba}^{2+} + 2\text{e}^-$                     | $\rightarrow$ | $\text{Ba}(\text{s})$                 | -2.90               |
| $\text{Rb}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Rb}(\text{s})$                 | -2.92               |
| $\text{K}^+ + \text{e}^-$                          | $\rightarrow$ | $\text{K}(\text{s})$                  | -2.92               |
| $\text{Cs}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Cs}(\text{s})$                 | -2.92               |
| $\text{Li}^+ + \text{e}^-$                         | $\rightarrow$ | $\text{Li}(\text{s})$                 | -3.05               |

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# ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

## ATOMIC STRUCTURE

$$E = h\nu \quad c = \lambda\nu$$

$$\lambda = \frac{h}{m\nu} \quad p = m\nu$$

$$E_n = \frac{-2.178 \times 10^{-18}}{n^2} \text{ joule}$$

## EQUILIBRIUM

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$$K_b = \frac{[\text{OH}^-][\text{HB}^+]}{[\text{B}]}$$

$$K_w = [\text{OH}^-][\text{H}^+] = 1.0 \times 10^{-14} \text{ @ } 25^\circ\text{C}$$

$$= K_a \times K_b$$

$$\text{pH} = -\log [\text{H}^+], \text{pOH} = -\log [\text{OH}^-]$$

$$14 = \text{pH} + \text{pOH}$$

$$\text{pH} = \text{p}K_a + \log \frac{[\text{A}^-]}{[\text{HA}]}$$

$$\text{pOH} = \text{p}K_b + \log \frac{[\text{HB}^+]}{[\text{B}]}$$

$$\text{p}K_a = -\log K_a, \text{p}K_b = -\log K_b$$

$$K_p = K_c(RT)^{\Delta n},$$

where  $\Delta n$  = moles product gas – moles reactant gas

## THERMOCHEMISTRY/KINETICS

$$\Delta S^\circ = \sum S^\circ \text{ products} - \sum S^\circ \text{ reactants}$$

$$\Delta H^\circ = \sum \Delta H_f^\circ \text{ products} - \sum \Delta H_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \sum \Delta G_f^\circ \text{ products} - \sum \Delta G_f^\circ \text{ reactants}$$

$$\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$$

$$= -RT \ln K = -2.303 RT \log K$$

$$= -n \mathcal{F} E^\circ$$

$$\Delta G = \Delta G^\circ + RT \ln Q = \Delta G^\circ + 2.303 RT \log Q$$

$$q = mc\Delta T$$

$$C_p = \frac{\Delta H}{\Delta T}$$

$$\ln[\text{A}]_t - \ln[\text{A}]_0 = -kt$$

$$\frac{1}{[\text{A}]_t} - \frac{1}{[\text{A}]_0} = kt$$

$$\ln k = \frac{-E_a}{R} \left( \frac{1}{T} \right) + \ln A$$

$E$  = energy

$v$  = velocity

$\nu$  = frequency

$n$  = principal quantum number

$\lambda$  = wavelength

$m$  = mass

$p$  = momentum

Speed of light,  $c = 3.0 \times 10^8 \text{ m s}^{-1}$

Planck's constant,  $h = 6.63 \times 10^{-34} \text{ J s}$

Boltzmann's constant,  $k = 1.38 \times 10^{-23} \text{ J K}^{-1}$

Avogadro's number =  $6.022 \times 10^{23} \text{ mol}^{-1}$

Electron charge,  $e = -1.602 \times 10^{-19} \text{ coulomb}$

1 electron volt per atom =  $96.5 \text{ kJ mol}^{-1}$

## Equilibrium Constants

$K_a$  (weak acid)

$K_b$  (weak base)

$K_w$  (water)

$K_p$  (gas pressure)

$K_c$  (molar concentrations)

$S^\circ$  = standard entropy

$H^\circ$  = standard enthalpy

$G^\circ$  = standard free energy

$E^\circ$  = standard reduction potential

$T$  = temperature

$n$  = moles

$m$  = mass

$q$  = heat

$c$  = specific heat capacity

$C_p$  = molar heat capacity at constant pressure

$E_a$  = activation energy

$k$  = rate constant

$A$  = frequency factor

Faraday's constant,  $\mathcal{F} = 96,500 \text{ coulombs per mole of electrons}$

Gas constant,  $R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$   
 $= 0.0821 \text{ L atm mol}^{-1} \text{ K}^{-1}$   
 $= 8.31 \text{ volt coulomb mol}^{-1} \text{ K}^{-1}$

**GO ON TO THE NEXT PAGE.**

## GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

$$\left( P + \frac{n^2 a}{V^2} \right) (V - nb) = nRT$$

$$P_A = P_{total} \times X_A, \text{ where } X_A = \frac{\text{moles A}}{\text{total moles}}$$

$$P_{total} = P_A + P_B + P_C + \dots$$

$$n = \frac{m}{M}$$

$$K = ^\circ C + 273$$

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

$$D = \frac{m}{V}$$

$$u_{rms} = \sqrt{\frac{3kT}{m}} = \sqrt{\frac{3RT}{M}}$$

$$KE \text{ per molecule} = \frac{1}{2} m v^2$$

$$KE \text{ per mole} = \frac{3}{2} RT$$

$$\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}}$$

molarity,  $M$  = moles solute per liter solution

molality = moles solute per kilogram solvent

$$\Delta T_f = iK_f \times \text{molality}$$

$$\Delta T_b = iK_b \times \text{molality}$$

$$\pi = iMRT$$

$$A = abc$$

$P$  = pressure

$V$  = volume

$T$  = temperature

$n$  = number of moles

$D$  = density

$m$  = mass

$v$  = velocity

$u_{rms}$  = root-mean-square speed

$KE$  = kinetic energy

$r$  = rate of effusion

$M$  = molar mass

$\pi$  = osmotic pressure

$i$  = van't Hoff factor

$K_f$  = molal freezing-point depression constant

$K_b$  = molal boiling-point elevation constant

$A$  = absorbance

$a$  = molar absorptivity

$b$  = path length

$c$  = concentration

$Q$  = reaction quotient

$I$  = current (amperes)

$q$  = charge (coulombs)

$t$  = time (seconds)

$E^\circ$  = standard reduction potential

$K$  = equilibrium constant

## OXIDATION-REDUCTION; ELECTROCHEMISTRY

$$Q = \frac{[C]^c [D]^d}{[A]^a [B]^b}, \text{ where } a A + b B \rightarrow c C + d D$$

$$I = \frac{q}{t}$$

$$E_{\text{cell}} = E_{\text{cell}}^\circ - \frac{RT}{n\mathcal{F}} \ln Q = E_{\text{cell}}^\circ - \frac{0.0592}{n} \log Q @ 25^\circ C$$

$$\log K = \frac{nE^\circ}{0.0592}$$

$$\text{Gas constant, } R = 8.31 \text{ J mol}^{-1} \text{ K}^{-1}$$

$$= 0.0821 \text{ L atm mol}^{-1} \text{ K}^{-1}$$

$$= 8.31 \text{ volt coulomb mol}^{-1} \text{ K}^{-1}$$

$$\text{Boltzmann's constant, } k = 1.38 \times 10^{-23} \text{ J K}^{-1}$$

$$K_f \text{ for H}_2\text{O} = 1.86 \text{ K kg mol}^{-1}$$

$$K_b \text{ for H}_2\text{O} = 0.512 \text{ K kg mol}^{-1}$$

$$1 \text{ atm} = 760 \text{ mm Hg}$$

$$= 760 \text{ torr}$$

$$\text{STP} = 0.000^\circ C \text{ and } 1.000 \text{ atm}$$

$$\text{Faraday's constant, } \mathcal{F} = 96,500 \text{ coulombs per mole of electrons}$$

# 2006 AP<sup>®</sup> CHEMISTRY FREE-RESPONSE QUESTIONS

## CHEMISTRY

### Section II

(Total time—90 minutes)

#### Part A

Time—40 minutes

**YOU MAY USE YOUR CALCULATOR FOR PART A.**

CLEARLY SHOW THE METHOD USED AND THE STEPS INVOLVED IN ARRIVING AT YOUR ANSWERS. It is to your advantage to do this, since you may obtain partial credit if you do and you will receive little or no credit if you do not. Attention should be paid to significant figures.

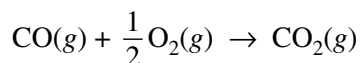
Be sure to write all your answers to the questions on the lined pages following each question in the booklet with the pink cover. Do NOT write your answers on the green insert.

Answer Question 1 below. The Section II score weighting for this question is 20 percent.

1. Answer the following questions that relate to solubility of salts of lead and barium.
  - (a) A saturated solution is prepared by adding excess  $\text{PbI}_2(s)$  to distilled water to form 1.0 L of solution at 25°C. The concentration of  $\text{Pb}^{2+}(aq)$  in the saturated solution is found to be  $1.3 \times 10^{-3} M$ . The chemical equation for the dissolution of  $\text{PbI}_2(s)$  in water is shown below.
$$\text{PbI}_2(s) \rightleftharpoons \text{Pb}^{2+}(aq) + 2 \text{I}^{-}(aq)$$
    - (i) Write the equilibrium-constant expression for the equation.
    - (ii) Calculate the molar concentration of  $\text{I}^{-}(aq)$  in the solution.
    - (iii) Calculate the value of the equilibrium constant,  $K_{sp}$ .
  - (b) A saturated solution is prepared by adding  $\text{PbI}_2(s)$  to distilled water to form 2.0 L of solution at 25°C. What are the molar concentrations of  $\text{Pb}^{2+}(aq)$  and  $\text{I}^{-}(aq)$  in the solution? Justify your answer.
  - (c) Solid  $\text{NaI}$  is added to a saturated solution of  $\text{PbI}_2$  at 25°C. Assuming that the volume of the solution does not change, does the molar concentration of  $\text{Pb}^{2+}(aq)$  in the solution increase, decrease, or remain the same? Justify your answer.
  - (d) The value of  $K_{sp}$  for the salt  $\text{BaCrO}_4$  is  $1.2 \times 10^{-10}$ . When a 500. mL sample of  $8.2 \times 10^{-6} M$   $\text{Ba}(\text{NO}_3)_2$  is added to 500. mL of  $8.2 \times 10^{-6} M$   $\text{Na}_2\text{CrO}_4$ , no precipitate is observed.
    - (i) Assuming that volumes are additive, calculate the molar concentrations of  $\text{Ba}^{2+}(aq)$  and  $\text{CrO}_4^{2-}(aq)$  in the 1.00 L of solution.
    - (ii) Use the molar concentrations of  $\text{Ba}^{2+}(aq)$  ions and  $\text{CrO}_4^{2-}(aq)$  ions as determined above to show why a precipitate does not form. You must include a calculation as part of your answer.

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Answer EITHER Question 2 below OR Question 3 printed on page 8. Only one of these two questions will be graded. If you start both questions, be sure to cross out the question you do not want graded. The Section II score weighting for the question you choose is 20 percent.



2. The combustion of carbon monoxide is represented by the equation above.

- (a) Determine the value of the standard enthalpy change,  $\Delta H_{\text{rxn}}^{\circ}$ , for the combustion of  $\text{CO}(g)$  at 298 K using the following information.



- (b) Determine the value of the standard entropy change,  $\Delta S_{\text{rxn}}^{\circ}$ , for the combustion of  $\text{CO}(g)$  at 298 K using the information in the following table.

| Substance           | $S_{298}^{\circ}$<br>(J mol <sup>-1</sup> K <sup>-1</sup> ) |
|---------------------|---|
| CO(g)               | 197.7   |
| CO <sub>2</sub> (g) | 213.7   |
| O <sub>2</sub> (g)  | 205.1   |

- (c) Determine the standard free energy change,  $\Delta G_{\text{rxn}}^{\circ}$ , for the reaction at 298 K. Include units with your answer.
- (d) Is the reaction spontaneous under standard conditions at 298 K? Justify your answer.
- (e) Calculate the value of the equilibrium constant,  $K_{\text{eq}}$ , for the reaction at 298 K.

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3. Answer the following questions that relate to the analysis of chemical compounds.
- (a) A compound containing the elements C, H, N, and O is analyzed. When a 1.2359 g sample is burned in excess oxygen, 2.241 g of  $\text{CO}_2(\text{g})$  is formed. The combustion analysis also showed that the sample contained 0.0648 g of H.
- (i) Determine the mass, in grams, of C in the 1.2359 g sample of the compound.
  - (ii) When the compound is analyzed for N content only, the mass percent of N is found to be 28.84 percent. Determine the mass, in grams, of N in the original 1.2359 g sample of the compound.
  - (iii) Determine the mass, in grams, of O in the original 1.2359 g sample of the compound.
  - (iv) Determine the empirical formula of the compound.
- (b) A different compound, which has the empirical formula  $\text{CH}_2\text{Br}$ , has a vapor density of  $6.00 \text{ g L}^{-1}$  at 375 K and 0.983 atm. Using these data, determine the following.
- (i) The molar mass of the compound
  - (ii) The molecular formula of the compound

**STOP**

**If you finish before time is called, you may check your work on this part only.  
Do not turn to the other part of the test until you are told to do so.**



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## CHEMISTRY

### Part B

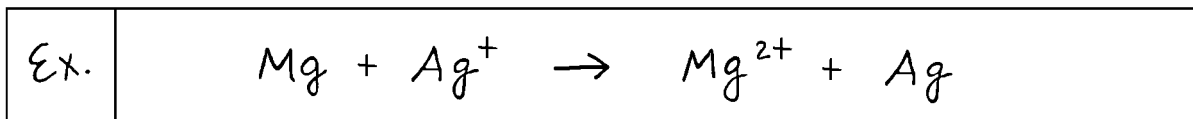
Time—50 minutes

**NO CALCULATORS MAY BE USED FOR PART B.**

Answer Question 4 below. The Section II score weighting for this question is 15 percent.

4. Write the formulas to show the reactants and the products for any FIVE of the laboratory situations described below. No more than five choices will be graded. In all cases, a reaction occurs. Assume that solutions are aqueous unless otherwise indicated. Represent substances in solution as ions if the substances are extensively ionized. Omit formulas for any ions or molecules that are unchanged by the reaction. You need not balance the equations.

Example: A strip of magnesium is added to a solution of silver nitrate.



- (a) Solid potassium chlorate is strongly heated.
- (b) Solid silver chloride is added to a solution of concentrated hydrochloric acid.
- (c) A solution of ethanoic (acetic) acid is added to a solution of barium hydroxide.
- (d) Ammonia gas is bubbled into a solution of hydrofluoric acid.
- (e) Zinc metal is placed in a solution of copper(II) sulfate.
- (f) Hydrogen phosphide (phosphine) gas is added to boron trichloride gas.
- (g) A solution of nickel(II) bromide is added to a solution of potassium hydroxide.
- (h) Hexane is combusted in air.

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Your responses to the rest of the questions in this part of the examination will be graded on the basis of the accuracy and relevance of the information cited. Explanations should be clear and well organized. Examples and equations may be included in your responses where appropriate. Specific answers are preferable to broad, diffuse responses.

Answer BOTH Question 5 below AND Question 6 printed on pages 11-12. Both of these questions will be graded. The Section II score weighting for these questions is 30 percent (15 percent each).

5. Three pure, solid compounds labeled  $X$ ,  $Y$ , and  $Z$  are placed on a lab bench with the objective of identifying each one. It is known that the compounds (listed in random order) are  $\text{KCl}$ ,  $\text{Na}_2\text{CO}_3$ , and  $\text{MgSO}_4$ . A student performs several tests on the compounds; the results are summarized in the table below.

| Compound | pH of an Aqueous Solution of the Compound | Result of Adding 1.0 $M$ $\text{NaOH}$ to a Solution of the Compound | Result of Adding 1.0 $M$ $\text{HCl}$ Dropwise to the Solid Compound |
|----------|---|--|--|
| $X$      | $> 7$                                     | No observed reaction   | Evolution of a gas   |
| $Y$      | 7   | No observed reaction   | No observed reaction   |
| $Z$      | 7   | Formation of a white precipitate                                     | No observed reaction   |

- (a) Identify each compound based on the observations recorded in the table.

Compound  $X$  \_\_\_\_\_

Compound  $Y$  \_\_\_\_\_

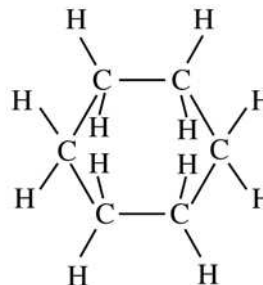
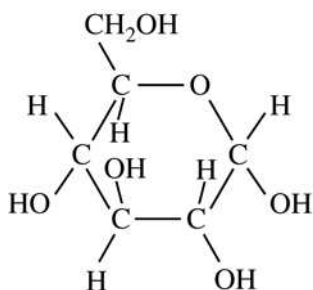
Compound  $Z$  \_\_\_\_\_

- (b) Write the chemical formula for the precipitate produced when 1.0  $M$   $\text{NaOH}$  is added to a solution of compound  $Z$ .
- (c) Explain why an aqueous solution of compound  $X$  has a pH value greater than 7. Write an equation as part of your explanation.
- (d) One of the testing solutions used was 1.0  $M$   $\text{NaOH}$ . Describe the steps for preparing 100. mL of 1.0  $M$   $\text{NaOH}$  from a stock solution of 3.0  $M$   $\text{NaOH}$  using a 50 mL buret, a 100 mL volumetric flask, distilled water, and a small dropper.
- (e) Describe a simple laboratory test that you could use to distinguish between  $\text{Na}_2\text{CO}_3(s)$  and  $\text{CaCO}_3(s)$ . In your description, specify how the results of the test would enable you to determine which compound was  $\text{Na}_2\text{CO}_3(s)$  and which compound was  $\text{CaCO}_3(s)$ .

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6. Answer each of the following in terms of principles of molecular behavior and chemical concepts.

(a) The structures for glucose,  $C_6H_{12}O_6$ , and cyclohexane,  $C_6H_{12}$ , are shown below.

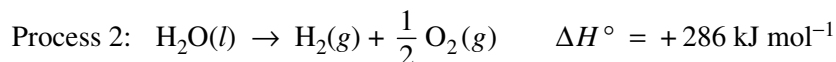


Identify the type(s) of intermolecular attractive forces in

- (i) pure glucose
- (ii) pure cyclohexane

(b) Glucose is soluble in water but cyclohexane is not soluble in water. Explain.

(c) Consider the two processes represented below.



- (i) For each of the two processes, identify the type(s) of intermolecular or intramolecular attractive forces that must be overcome for the process to occur.
- (ii) Indicate whether you agree or disagree with the statement in the box below. Support your answer with a short explanation.

When water boils,  $H_2O$  molecules break apart to form hydrogen molecules and oxygen molecules.

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(d) Consider the four reaction-energy profile diagrams shown below.

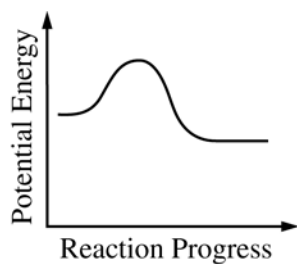


Diagram 1

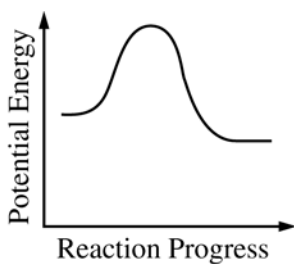


Diagram 2

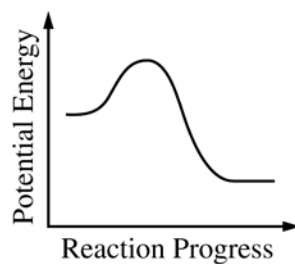


Diagram 3

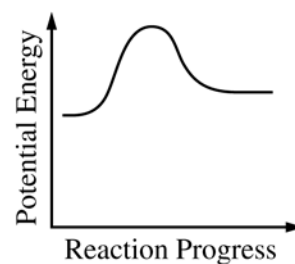


Diagram 4

- (i) Identify the two diagrams that could represent a catalyzed and an uncatalyzed reaction pathway for the same reaction. Indicate which of the two diagrams represents the catalyzed reaction pathway for the reaction.
- (ii) Indicate whether you agree or disagree with the statement in the box below. Support your answer with a short explanation.

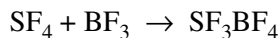
Adding a catalyst to a reaction mixture adds energy that causes the reaction to proceed more quickly.

## 2006 AP<sup>®</sup> CHEMISTRY FREE-RESPONSE QUESTIONS

Answer EITHER Question 7 below OR Question 8 printed on page 14. Only one of these two questions will be graded. If you start both questions, be sure to cross out the question you do not want graded. The Section II score weighting for the question you choose is 15 percent.

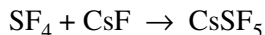
7. Answer the following questions about the structures of ions that contain only sulfur and fluorine.

(a) The compounds  $\text{SF}_4$  and  $\text{BF}_3$  react to form an ionic compound according to the following equation.



- (i) Draw a complete Lewis structure for the  $\text{SF}_3^+$  cation in  $\text{SF}_3\text{BF}_4$ .
- (ii) Identify the type of hybridization exhibited by sulfur in the  $\text{SF}_3^+$  cation.
- (iii) Identify the geometry of the  $\text{SF}_3^+$  cation that is consistent with the Lewis structure drawn in part (a)(i).
- (iv) Predict whether the F–S–F bond angle in the  $\text{SF}_3^+$  cation is larger than, equal to, or smaller than  $109.5^\circ$ . Justify your answer.

(b) The compounds  $\text{SF}_4$  and  $\text{CsF}$  react to form an ionic compound according to the following equation.



- (i) Draw a complete Lewis structure for the  $\text{SF}_5^-$  anion in  $\text{CsSF}_5$ .
- (ii) Identify the type of hybridization exhibited by sulfur in the  $\text{SF}_5^-$  anion.
- (iii) Identify the geometry of the  $\text{SF}_5^-$  anion that is consistent with the Lewis structure drawn in part (b)(i).
- (iv) Identify the oxidation number of sulfur in the compound  $\text{CsSF}_5$ .

## 2006 AP<sup>®</sup> CHEMISTRY FREE-RESPONSE QUESTIONS

8. Suppose that a stable element with atomic number 119, symbol Q, has been discovered.
- (a) Write the ground-state electron configuration for Q, showing only the valence-shell electrons.
  - (b) Would Q be a metal or a nonmetal? Explain in terms of electron configuration.
  - (c) On the basis of periodic trends, would Q have the largest atomic radius in its group or would it have the smallest? Explain in terms of electronic structure.
  - (d) What would be the most likely charge of the Q ion in stable ionic compounds?
  - (e) Write a balanced equation that would represent the reaction of Q with water.
  - (f) Assume that Q reacts to form a carbonate compound.
    - (i) Write the formula for the compound formed between Q and the carbonate ion,  $\text{CO}_3^{2-}$ .
    - (ii) Predict whether or not the compound would be soluble in water. Explain your reasoning.

**STOP**

**END OF EXAM**