

AP[®] Chemistry 2010 Scoring Guidelines

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Question 1 (10 points)

Several reactions are carried out using AgBr, a cream-colored silver salt for which the value of the solubilityproduct-constant, K_{sp} , is 5.0×10^{-13} at 298 K.

(a) Write the expression for the solubility-product constant, K_{sp} , of AgBr.

$K_{sp} = [Ag^+][Br^-]$ One point is earned for the correct expression (ion charges must present; parentheses instead of square brackets not accepted).	be
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(b) Calculate the value of $[Ag^+]$ in 50.0 mL of a saturated solution of AgBr at 298 K.

Let $x =$ equilibrium concentration of Ag ⁺ (and of Br ⁻).	One point is earned for the correct
Then $K_{sp} = 5.0 \times 10^{-13} = x^2 \implies x = 7.1 \times 10^{-7} M$	value with supporting work (units not necessary).

(c) A 50.0 mL sample of distilled water is added to the solution described in part (b), which is in a beaker with some solid AgBr at the bottom. The solution is stirred and equilibrium is reestablished. Some solid AgBr remains in the beaker. Is the value of [Ag⁺] greater than, less than, or equal to the value you calculated in part (b) ? Justify your answer.

(d) Calculate the minimum volume of distilled water, in liters, necessary to completely dissolve a 5.0 g sample of AgBr(*s*) at 298 K. (The molar mass of AgBr is 188 g mol⁻¹.)

$5.0 \text{ g AgBr} \times \frac{1 \text{ mol AgBr}}{188 \text{ g AgBr}} = 0.0266 \text{ mol AgBr}$	One point is earned for the calculation of moles of dissolved AgBr.
$\frac{0.0266 \text{ mol}}{\text{V}} = 7.1 \times 10^{-7} \text{mol } \text{L}^{-1} \implies \text{V} = 3.7 \times 10^{4} \text{ L}$	One point is earned for the correct answer for the volume of water

Question 1 (continued)

(e) A student mixes 10.0 mL of $1.5 \times 10^{-4} M$ AgNO₃ with 2.0 mL of $5.0 \times 10^{-4} M$ NaBr and stirs the resulting mixture. What will the student observe? Justify your answer with calculations.

$[Ag^+] = \frac{(10.0 \text{ mL})(1.5 \times 10^{-4} M)}{12.0 \text{ mL}} = 1.3 \times 10^{-4} M$	One point is earned for calculation of concentration of ions.
$[Br^{-}] = \frac{(2.0 \text{ mL})(5.0 \times 10^{-4} M)}{12.0 \text{ mL}} = 8.3 \times 10^{-5} M$ $Q = [Ag^{+}][Br^{-}] = (1.3 \times 10^{-4} M)(8.3 \times 10^{-5} M) = 1.1 \times 10^{-8}$	One point is earned for calculation of Q and conclusion based on comparison between Q and K_{sp} .
$1.1 \times 10^{-8} > 5.0 \times 10^{-13}$, \therefore a precipitate will form.	One point is earned for indicating the precipitation of AgBr.

- (f) The color of another salt of silver, AgI(*s*), is yellow. A student adds a solution of NaI to a test tube containing a small amount of solid, cream-colored AgBr. After stirring the contents of the test tube, the student observes that the solid in the test tube changes color from cream to yellow.
 - (i) Write the chemical equation for the reaction that occurred in the test tube.

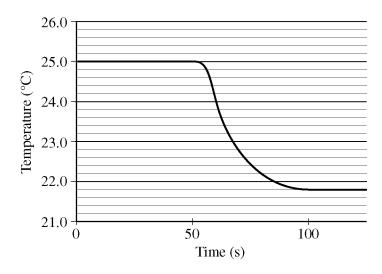
AgBr(s) + I⁻(aq) \rightarrow AgI(s) + Br⁻(aq)One point is earned for the correct equation.OROne point is earned for the correct equation.AgBr(s) + NaI(aq) \rightarrow AgI(s) + NaBr(aq)

(ii) Which salt has the greater value of K_{sp} : AgBr or AgI ? Justify your answer.

AgBr has the greater value of K_{sp} . The precipitate will consist of the less soluble salt when both I ⁻ (<i>aq</i>) and Br ⁻ (<i>aq</i>) are present. Because the color of the precipitate in the test tube turns yellow, it must be AgI(<i>s</i>) that precipitates; therefore K_{sp} for AgBr must be greater than K_{sp} for AgI. OR	One point is earned for the correct choice with justification.
K_{eq} for the displacement reaction is $\frac{K_{sp} \text{ of AgBr}}{K_{sp} \text{ of AgI}}$. Because yellow AgI forms, $K_{eq} > 1$; therefore K_{sp} of AgBr $> K_{sp}$ of AgI.	

Question 2 (10 points)

A student performs an experiment to determine the molar enthalpy of solution of urea, H_2NCONH_2 . The student places 91.95 g of water at 25°C into a coffee cup calorimeter and immerses a thermometer in the water. After 50 s, the student adds 5.13 g of solid urea, also at 25°C, to the water and measures the temperature of the solution as the urea dissolves. A plot of the temperature data is shown in the graph below.



(a) Determine the change in temperature of the solution that results from the dissolution of the urea.

$\Delta T = 21.8 - 25.0 = -3.2 \text{ Celsius degrees}$	One point is earned for the correct temperature change.
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(b) According to the data, is the dissolution of urea in water an endothermic process or an exothermic process? Justify your answer.

The process is endothermic. The decrease in temperature indicates that the process for the dissolution of urea in water requires energy.	One point is earned for the correct choice with justification.
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Question 2 (continued)

- (c) Assume that the specific heat capacity of the calorimeter is negligible and that the specific heat capacity of the solution of urea and water is 4.2 J g^{-1} °C⁻¹ throughout the experiment.
 - (i) Calculate the heat of dissolution of the urea in joules.

Assuming that no heat energy is lost from the calorimeter and given that the calorimeter has a negligible heat capacity, the sum of the heat of dissolution, q_{soln} and the change in heat energy of the urea-water mixture must equal zero. $q_{soln} + mc\Delta T = 0 \implies q_{soln} = -mc\Delta T$ $m_{soln} = 5.13 \text{ g} + 91.95 \text{ g} = 97.08 \text{ g}$ $q_{soln} = -(97.08 \text{ g})(4.2 \text{ J g}^{-1\circ}\text{C}^{-1})(-3.2^{\circ}\text{C}) = 1.3 \times 10^{3} \text{ J}$	One point is earned for the correct setup. One point is earned for the correct numerical result for the heat of dissolution.
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(ii) Calculate the molar enthalpy of solution, ΔH_{soln}° , of urea in kJ mol⁻¹.

$\Delta H_{soln}^{\circ} = \frac{q_{soln}}{\text{mol solute}}$ molar mass of urea = 4(1.0) + 2(14.0) + 12.0 + 16.0 = 60.0 g mol ⁻¹	One point is earned for the calculation of moles of urea.
moles of urea = 5.13 g urea $\times \frac{1 \text{ mol urea}}{60.0 \text{ g urea}} = 0.0855 \text{ mol}$	One point is earned for the correct numerical result with
$\Delta H_{soln}^{\circ} = \frac{1.3 \times 10^3 \mathrm{J}}{0.0855 \mathrm{mol}} = 1.5 \times 10^4 \mathrm{J mol}^{-1} = 15 \mathrm{kJ mol}^{-1}$	correct algebraic sign.

(d) Using the information in the table below, calculate the value of the molar entropy of solution, ΔS_{soln}° , of urea at 298 K. Include units with your answer.

	Accepted Value
ΔH_{soln}° of urea	14.0 kJ mol ⁻¹
ΔG_{soln}° of urea	-6.9 kJ mol^{-1}

$\Delta G^{\circ} = \Delta H^{\circ} - T \Delta S^{\circ}$	One point is earned for the correct setup.
- 6.9 kJ mol ⁻¹ = 14.0 kJ mol ⁻¹ - (298 K)(ΔS°) $\Delta S_{soln}^{\circ} = 0.0701$ kJ mol ⁻¹ K ⁻¹ = 70.1 J mol ⁻¹ K ⁻¹	One point is earned for the correct numerical result with correct units.

Question 2 (continued)

(e) The student repeats the experiment and this time obtains a result for ΔH_{soln}° of urea that is 11 percent below the accepted value. Calculate the value of ΔH_{soln}° that the student obtained in this second trial.

Error = $(0.11)(14.0 \text{ kJ mol}^{-1}) = 1.54 \text{ kJ mol}^{-1}$	One point is earned for the correct
$14.0 \text{ kJ mol}^{-1} - 1.54 \text{ kJ mol}^{-1} = 12.5 \text{ kJ mol}^{-1}$	numerical result.

(f) The student performs a third trial of the experiment but this time adds urea that has been taken directly from a refrigerator at 5°C. What effect, if any, would using the cold urea instead of urea at 25°C have on the experimentally obtained value of ΔH_{soln}° ? Justify your answer.

There would be an increase in the obtained value for ΔH_{soln}° because the colder urea would have caused a larger negative temperature change.	One point is earned for the correct prediction with justification.
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Question 3 (9 points)

 $8 \operatorname{H}^{+}(aq) + 4 \operatorname{Cl}^{-}(aq) + \operatorname{MnO}_{4}^{-}(aq) \rightarrow 2 \operatorname{Cl}_{2}(g) + \operatorname{Mn}^{3+}(aq) + 4 \operatorname{H}_{2}O(l)$

 $Cl_2(g)$ can be generated in the laboratory by reacting potassium permanganate with an acidified solution of sodium chloride. The net-ionic equation for the reaction is given above.

- (a) A 25.00 mL sample of 0.250 *M* NaCl reacts completely with excess KMnO₄(*aq*). The Cl₂(*g*) produced is dried and stored in a sealed container. At 22°C the pressure of the Cl₂(*g*) in the container is 0.950 atm.
 - (i) Calculate the number of moles of $Cl^{-}(aq)$ present before any reaction occurs.

mol Cl ⁻ = $(0.02500 \text{ L})(0.250 \text{ M}) = 6.25 \times 10^{-3} \text{ mol}$	One point is earned for the correct numerical value.
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(ii) Calculate the volume, in L, of the $Cl_2(g)$ in the sealed container.

$\operatorname{mol} \operatorname{Cl}_{2} = \frac{\operatorname{mol} \operatorname{Cl}^{-}}{2} = \frac{6.25 \times 10^{-3} \operatorname{mol}}{2} = 3.125 \times 10^{-3} \operatorname{mol} \operatorname{Cl}_{2}$ $V = \frac{nRT}{P} = \frac{(3.125 \times 10^{-3} \operatorname{mol} \operatorname{Cl}_{2})(0.0821 \operatorname{L} \operatorname{atm} \operatorname{mol}^{-1} \operatorname{K}^{-1})(295 \operatorname{K})}{0.950 \operatorname{atm}}$	One point is earned for the correct number of moles of Cl_2 based on stoichiometry.
P = 0.950 atm	_
	substitution into ideal gas law
$= 0.0797 \text{ L Cl}_2$	and correct numerical result.

An initial-rate study was performed on the reaction system. Data for the experiment are given in the table below.

Trial	[Cl ⁻]	$[MnO_4^{-}]$	$[\mathrm{H}^+]$	Rate of Disappearance of MnO_4^- in $M s^{-1}$
1	0.0104	0.00400	3.00	2.25×10^{-8}
2	0.0312	0.00400	3.00	2.03×10^{-7}
3	0.0312	0.00200	3.00	1.02×10^{-7}

Question 3 (continued)

- (b) Using the information in the table, determine the order of the reaction with respect to each of the following. Justify your answers.
 - (i) Cl⁻

The reaction is second order. Tripling [Cl ⁻] between trials 1 and 2 with no change in $[MnO_4^-]$ results in a nine-fold increase in the rate:	
$\left(\frac{0.0312 \ M}{0.0104 \ M}\right)^{x} = \frac{2.03 \times 10^{-7}}{2.25 \times 10^{-8}}$	One point is earned for the correct order of reaction
$3^x = 9$	with justification.
x = 2	
Thus the order of the reaction must be 2 with respect to Cl ⁻ .	

(ii) MnO₄⁻

The reaction is first order. Doubling $[MnO_4^-]$ between trials 3 and 2 with no change in $[Cl^-]$ results in a doubling of the rate:	
$\left(\frac{0.00400 M}{0.00200 M}\right)^{y} = \frac{2.03 \times 10^{-7}}{1.02 \times 10^{-7}}$ $2^{y} = 2$ $y = 1$	One point is earned for the correct order of reaction with justification.
Thus the order of the reaction must be 1 with respect to MnO_4^{-} .	

- (c) The reaction is known to be third order with respect to H⁺. Using this information and your answers to part (b) above, complete both of the following:
 - (i) Write the rate law for the reaction.

rate = $k[Cl^{-}]^{2}[MnO_{4}^{-}][H^{+}]^{3}$	One point is earned for the correct rate law.
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Question 3 (continued)

(ii) Calculate the value of the rate constant, k, for the reaction, including appropriate units.

Using data from trial 1: $2.25 \times 10^{-8} M s^{-1} = k(0.0104 M)^2 (0.00400 M) (3.00 M)^3$	One point is earned for the correct numerical result.
$k = 1.93 \times 10^{-3} M^{-5} s^{-1}$	One point is earned for the correct units.

(d) Is it likely that the reaction occurs in a single elementary step? Justify your answer.

It is not likely that the reaction occurs in a single step because the orders of the reaction with respect to the reactants do not correspond to the coefficients in the balanced equation <i>OR</i>	One point is earned for the correct
It is not likely that the reaction occurs in a single step because the reaction requires the collision of many (13) reactant particles and the frequency of a 13-particle collision is negligible.	answer with justification.

Question 4 (15 points)

For each of the following three reactions, write a balanced equation for the reaction in part (i) and answer the question about the reaction in part (ii). In part (i), coefficients should be in terms of lowest whole numbers. Assume that solutions are aqueous unless otherwise indicated. Represent substances in solutions as ions if the substances are extensively ionized. Omit formulas for any ions or molecules that are unchanged by the reaction. You may use the empty space at the bottom of the next page for scratch work, but only equations that are written in the answer boxes provided will be scored.

EXAMPLE: A strip of magnesium metal is added to a solution of silver(I) nitrate.	
(i) Balanced equation: $Mg + 2 Ag^{+} \longrightarrow Mg^{2+} + 2 Ag^{-}$	
(ii) Which substance is oxidized in the reaction? Mg is optidized.	

(a) A 0.2 *M* potassium hydroxide solution is titrated with a 0.1 *M* nitric acid solution.

(i) Balanced equation:	One point is earned for each correct reactant.
$H_3O^+ + OH^- \rightarrow 2 H_2O$	One point is earned for the correct product.
OR	One point is earned for correctly balancing (mass
$\rm H^{+} + OH^{-} \rightarrow \rm H_{2}O$	and charge) the equation.

(ii) What would be observed if the solution was titrated well past the equivalence point using bromthymol blue as the indicator? (Bromthymol blue is yellow in acidic solution and blue in basic solution.)

The solution would appear yellow.	One point is earned for the correct description of the solution.
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Question 4 (continued)

(b) Propane is burned completely in excess oxygen gas.

(i) Balanced equation: $C_3H_8 + 5 O_2 \rightarrow 3 CO_2 + 4 H_2O$	One point is earned for both correct reactants. Two points are earned for the correct products. One point is earned for correctly balancing the equation.	

(ii) When the products of the reaction are bubbled through distilled water, is the resulting solution neutral, acidic, or basic? Explain.

The resulting solution would be acidic because CO_2	One point is earned for the correct choice with
reacts with water as a weak acid.	justification.

(c) A solution of hydrogen peroxide is heated, and a gas is produced.

(i) Polonard equation:	One point is earned for the correct reactant.
(i) Balanced equation: $2 H_2O_2 \rightarrow 2 H_2O + O_2$	Two points are earned for the correct products.
$2 \Pi_2 O_2 \rightarrow 2 \Pi_2 O + O_2$	One point is earned for correctly balancing the equation.

(ii) Identify the oxidation state of oxygen in hydrogen peroxide.

The oxidation state of O in H_2O_2 is -1.	One point is earned for the correct oxidation state.
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Question 5 (8 points)

Use the information in the table below to respond to the statements and questions that follow. Your answers should be in terms of principles of molecular structure and intermolecular forces.

Compound	Formula	Lewis Electron-Dot Diagram
Ethanethiol	CH ₃ CH ₂ SH	нн н:С:С:Ї:ін нн
Ethane	CH ₃ CH ₃	нн н:С:С:н йй
Ethanol	CH ₃ CH ₂ OH	нн н:ё:ё:ё:ю:н нйн
Ethyne	C ₂ H ₂	H:C∷C:H or H−C≡C−H

(a) Draw the complete Lewis electron-dot diagram for ethyne in the appropriate cell in the table above.

See the lower right cell in the table above.	One point is earned for the correct Lewis structure.	

(b)Which of the four molecules contains the shortest carbon-to-carbon bond? Explain.

Ethyne, which contains a triple bond, has the shortest C-to-C bond. The other molecules have single C-to-C bonds, and triple bonds are shorter than single bonds.	One point is earned for the correct choice. One point is earned for the correct explanation.
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Question 5 (continued)

(c) A Lewis electron-dot diagram of a molecule of ethanoic acid is given below. The carbon atoms in the molecule are labeled x and y, respectively.

Identify the geometry of the arrangement of atoms bonded to each of the following.

(i) Carbon *x*

Trigonal planar One point is earned for the correct geometry.

(ii) Carbon y

Distorted tetrahedral, tetrahedral or trigonal pyramidal	One point is earned for the correct geometry.
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(d) Energy is required to boil ethanol. Consider the statement "As ethanol boils, energy goes into breaking C-C bonds, C-H bonds, C-O bonds, and O-H bonds." Is the statement true or false? Justify your answer.

The statement is false. All of the bonds described are intramolecular; these bonds are not broken during vaporization. When ethanol boils, the added energy overcomes <u>inter</u> molecular, not <u>intra</u> molecular, forces.	One point is earned for the correct choice with justification.
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(e) Identify a compound from the table above that is nonpolar. Justify your answer.

Either ethane or ethyne may be identified as nonpolar. The ethane/ethyne molecule is nonpolar because all of the bond dipoles in the molecule cancel. <i>OR</i>	
The ethane/ethyne molecule is nonpolar because the molecule is symmetric.	One point is earned for a correct choice with justification.
<u>Note:</u> Explanation must refer to the shape of the molecule. Statements such as: "all hydrocarbons are nonpolar', "the carbons are surrounded by hydrogens" or "there are no lone pairs" do not earn this point.	

Question 5 (continued)

(f) Ethanol is completely soluble in water, whereas ethanethiol has limited solubility in water. Account for the difference in solubilities between the two compounds in terms of intermolecular forces.

Ethanol is able to form strong hydrogen bonds with water whereas ethanethiol does not have similar capability. The formation of hydrogen bonds increases the attraction between molecules of ethanol and molecules of water, making them more soluble in each other.	One point is earned for the correct explanation.
<u>Note:</u> The answer must clearly focus on the solute- solvent interaction. Just the mention of hydrogen bonding does not earn the point.	

Ouestion 6 (9 points)

 $2 \operatorname{Al}(s) + 3 \operatorname{Zn}^{2+}(aq) \rightarrow 2 \operatorname{Al}^{3+}(aq) + 3 \operatorname{Zn}(s)$

Respond to the following statements and questions that relate to the species and the reaction represented above.

(a) Write the complete electron configuration (e.g., $1s^2 2s^2 \dots$) for Zn^{2+} .

$1s^2 2s^2 2p^0 3s^2 3p^0 3d^{10}$ One point is earlied for the conject configuration.	$1s^2 2s^2 2p^6 3s^2 3p^6 3d^{10}$	One point is earned for the correct configuration.
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(b) Which species, Zn or Zn^{2+} , has the greater ionization energy? Justify your answer.

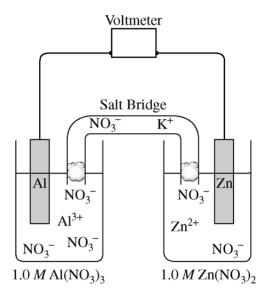
Zn^{2+} has the greater ionization energy. The electron being removed from Zn^{2+} experiences a larger effective nuclear charge than the electron being removed from Zn because Zn^{2+} has two fewer electrons shielding the nucleus. <i>OR</i>	One point is earned for identifying Zn ²⁺ with justification.
It takes more energy to remove a negatively charged electron from a positive ion than from a neutral atom.	

(c) Identify the species that is oxidized in the reaction.

Al(s)	One point is earned for identifying Al.
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Question 6 (continued)

The diagram below shows a galvanic cell based on the reaction. Assume that the temperature is 25°C.



(d) The diagram includes a salt bridge that is filled with a saturated solution of KNO_3 . Describe what happens in the salt bridge as the cell operates.

As the cell operates, NO_3^- ions flow toward the Al half-cell	One point is earned for correctly
and K^+ ions flow toward the Zn half-cell.	indicating the direction of ion flow.

(e) Determine the value of the standard voltage, E° , for the cell.

$E^{\circ} = (-0.76 \text{ V}) - (-1.66 \text{ V}) = 0.90 \text{ V}$	One point is earned for the correct E° .
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(f) Indicate whether the value of the standard free-energy change, ΔG° , for the cell reaction is positive, negative, or zero. Justify your answer.

ΔG° is negative since E° is positive and $\Delta G^{\circ} = -n \mathcal{F} E^{\circ}$. <i>OR</i>	One point is earned for indicating that ΔG° is negative.
ΔG° must be negative because the reaction is spontaneous under standard conditions.	One point is earned for a correct justification.

Question 6 (continued)

(g) If the concentration of Al(NO₃)₃ in the Al(s)/Al³⁺(aq) half-cell is lowered from 1.0 *M* to 0.01 *M* at 25°C, does the cell voltage increase, decrease, or remain the same? Justify your answer.

Lowering [Al ³⁺] causes an increase in the cell voltage.	
The value of Q will fall below 1.0 and the log term in the Nernst	One point is earned for indicating
equation will become negative. This causes the value of E_{cell} to	that E_{cell} increases.
become more positive.	
OR	One point is earned for the correct
A decrease in a product concentration will increase the spontaneity	justification.
of the reaction, increasing the value of E_{cell} .	