

AP[®] Chemistry 2006 Free-Response Questions

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INFORMATION IN THE TABLE BELOW AND IN THE TABLES ON PAGES 3-5 MAY BE USEFUL IN ANSWERING THE QUESTIONS IN THIS SECTION OF THE EXAMINATION.

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				6	Ŧ	19.00	17	\Box	35.453	35	\mathbf{Br}	79.90	53	Ι	126.91	85	At	(210)			
				8	0	16.00	16	S	32.06	34	Se	78.96	52	Te	127.60	84	P_0	(506)			
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	140.12	140.91	144.24	(145)	150.4	151.97	157.25	158.93	162.50	164.93	167.26		173.04	174.97
	06	91	92	93	94	95	96	26	86	66	100		102	103
†Actinide Series	Th	Pa	Ω	Np	Pu	Am	Cm	Bk	$\mathbf{C}\mathbf{f}$	$\mathbf{E}\mathbf{s}$	Fm		No	Γ
	232.04	232.04 231.04 238.03	238.03	237.05	(244)	(243)	(247)	(247)	(251)	(252)	(257)		(259)	(262)

STANDARD REDUCTION POTENTIALS IN AQUEOUS SOLUTION AT 25°C

STANDARD REDUCTION	Half-reaction		$E^{\circ}(V)$
F () : 2 =		2 E=	2.87
$F_2(g) + 2e^-$	\rightarrow	2 F ⁻	1.82
$Co^{3+} + e^{-}$	\rightarrow	Co ²⁺	
$Au^{3+} + 3e^{-}$	\rightarrow	Au(s)	1.50
$\operatorname{Cl}_2(g) + 2e^-$	\rightarrow	2 C1	1.36
$O_2(g) + 4 H^+ +$		$2 H_2O(l)$	1.23
$Br_2(l) + 2e^-$	\rightarrow	2 Br	1.07
$2 \text{ Hg}^{2+} + 2 e^{-}$	\rightarrow	Hg_2^{2+}	0.92
$Hg^{2+} + 2e^{-}$	\rightarrow	Hg(l)	0.85
$Ag^+ + e^-$	\rightarrow	Ag(s)	0.80
$Hg_2^{2+} + 2e^-$	\rightarrow	$2 \operatorname{Hg}(l)$	0.79
$Fe^{3+} + e^{-}$	\rightarrow	Fe ²⁺	0.77
$I_2(s) + 2e^-$	\rightarrow	2 I ⁻	0.53
$Cu^+ + e^-$	\rightarrow	Cu(s)	0.52
$Cu^{2+} + 2e^{-}$	\rightarrow	Cu(s)	0.34
$Cu^{2+} + e^{-}$	\rightarrow	Cu ⁺	0.15
$\mathrm{Sn}^{4+} + 2 e^{-}$	\rightarrow	Sn ²⁺	0.15
$S(s) + 2H^+ + 2$	$e^- \longrightarrow$	$H_2S(g)$	0.14
$2 \mathrm{H^+} + 2 e^-$	\rightarrow	$H_2(g)$	0.00
$Pb^{2+} + 2e^{-}$	\rightarrow	Pb(s)	-0.13
$\mathrm{Sn}^{2+} + 2 e^{-}$	\rightarrow	Sn(s)	-0.14
$Ni^{2+} + 2e^{-}$	\rightarrow	Ni(s)	-0.25
$Co^{2+} + 2e^{-}$	\rightarrow	Co(s)	-0.28
$Cd^{2+} + 2e^{-}$	\rightarrow	Cd(s)	-0.40
$Cr^{3+} + e^{-}$	\rightarrow	Cr ²⁺	-0.41
$Fe^{2+} + 2e^{-}$	\rightarrow	Fe(s)	-0.44
$Cr^{3+} + 3e^{-}$	\rightarrow	Cr(s)	-0.74
$Zn^{2+} + 2e^{-}$	\rightarrow	Zn(s)	-0.76
$2 H_2 O(l) + 2 e^-$	\rightarrow	$H_2(g) + 2 OH^-$	-0.83
$Mn^{2+} + 2e^{-}$	\rightarrow	Mn(s)	-1.18
$Al^{3+} + 3e^{-}$	\rightarrow	Al(s)	-1.66
$Be^{2+} + 2e^{-}$	\rightarrow	Be(s)	-1.70
$Mg^{2+} + 2e^{-}$	\rightarrow	Mg(s)	-2.37
$Na^+ + e^-$	\rightarrow	Na(s)	-2.71
$Ca^{2+} + 2e^{-}$	\rightarrow	Ca(s)	-2.87
$Sr^{2+} + 2e^{-}$	\rightarrow	Sr(s)	-2.89
$Ba^{2+} + 2e^{-}$	\rightarrow	Ba(s)	-2.90
$Rb^{+} + e^{-}$	\rightarrow	Rb(s)	-2.92
$K^+ + e^-$	\rightarrow	K(s)	-2.92
$Cs^+ + e^-$	\rightarrow	Cs(s)	-2.92
$Li^{+} + e^{-}$	\rightarrow	Li(s)	-3.05
Di 10	·	LI(3)	5.05

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

ATOMIC STRUCTURE

$$E = hv$$
 $c = \lambda v$
 $\lambda = \frac{h}{mv}$ $p = mv$
 $E_n = \frac{-2.178 \times 10^{-18}}{n^2}$ joule

EQUILIBRIUM

$$K_{a} = \frac{[H^{+}][A^{-}]}{[HA]}$$

$$K_{b} = \frac{[OH^{-}][HB^{+}]}{[B]}$$

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

$$= K_{a} \times K_{b}$$

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

$$14 = pH + pOH$$

$$pH = pK_{a} + \log \frac{[A^{-}]}{[HA]}$$

$$pOH = pK_{b} + \log \frac{[HB^{+}]}{[B]}$$

$$pK_{a} = -\log K_{a}, pK_{b} = -\log K_{b}$$

$$K_{p} = K_{c}(RT)^{\Delta n},$$

where $\Delta n = \text{moles product gas} - \text{moles reactant gas}$

THERMOCHEMISTRY/KINETICS

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

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

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

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

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

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

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

$$q = mc\Delta T$$

$$C_{p} = \frac{\Delta H}{\Delta T}$$

$$\ln[A]_{t} - \ln[A]_{0} = -kt$$

$$\frac{1}{[A]_{t}} - \frac{1}{[A]_{0}} = kt$$

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

$$E = \text{energy}$$
 $v = \text{velocity}$

$$v = \text{frequency}$$
 $n = \text{principal quantum number}$

$$\lambda$$
 = wavelength m = mass

$$p = momentum$$

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

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,
$$e = -1.602 \times 10^{-19}$$
 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)

$$K_n$$
 (gas pressure)

$$K_c$$
 (molar concentrations)

$$S^{\circ}$$
 = standard entropy

$$H^{\circ}$$
 = standard enthalpy

$$G^{\circ}$$
 = standard free energy

$$E^{\circ}$$
 = standard reduction potential

$$T = temperature$$

$$n = moles$$

$$m = \text{mass}$$

$$q = \text{heat}$$

$$c =$$
specific heat capacity

$$C_p$$
 = molar heat capacity at constant pressure

$$E_a$$
 = activation energy

$$k = \text{rate constant}$$

$$A =$$
frequency factor

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

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

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

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

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

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

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

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

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

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

$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$

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

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

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

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

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

$$molarity, M = \text{ moles solute per liter solution}$$

$$molality = \text{ moles solute per kilogram solvent}$$

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

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

$$\pi = iMRT$$

OXIDATION-REDUCTION; ELECTROCHEMISTRY

A = abc

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

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

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

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

P = pressure V = volume T = temperature n = number of moles D = density m = mass

 u_{rms} = root-mean-square speed

KE = kinetic energy r = rate of effusion

v = velocity

r = rate of effusion

M = molar mass

 π = 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° = standard reduction potential

K = equilibrium constant

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

 $= 0.0821 \text{ L atm mol}^{-1} \text{ K}^{-1}$
 $= 8.31 \text{ volt coulomb mol}^{-1} \text{ K}^{-1}$
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 atm = 760 mm Hg
= 760 torr
STP = 0.000°C and 1.000 atm
Faraday's constant, $\mathcal{F} = 96,500$ coulombs per mole

of electrons

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 $PbI_2(s)$ to distilled water to form 1.0 L of solution at 25°C. The concentration of $Pb^{2+}(aq)$ in the saturated solution is found to be $1.3 \times 10^{-3} M$. The chemical equation for the dissolution of $PbI_2(s)$ in water is shown below.

$$PbI_2(s) \rightleftharpoons Pb^{2+}(aq) + 2 I^{-}(aq)$$

- (i) Write the equilibrium-constant expression for the equation.
- (ii) Calculate the molar concentration of $I^-(aq)$ in the solution.
- (iii) Calculate the value of the equilibrium constant, K_{sn} .
- (b) A saturated solution is prepared by adding $PbI_2(s)$ to distilled water to form 2.0 L of solution at 25°C. What are the molar concentrations of $Pb^{2+}(aq)$ and $I^{-}(aq)$ in the solution? Justify your answer.
- (c) Solid NaI is added to a saturated solution of PbI_2 at 25°C. Assuming that the volume of the solution does not change, does the molar concentration of $Pb^{2+}(aq)$ in the solution increase, decrease, or remain the same? Justify your answer.
- (d) The value of K_{sp} for the salt BaCrO₄ is 1.2×10^{-10} . When a 500. mL sample of 8.2×10^{-6} M Ba(NO₃)₂ is added to 500. mL of 8.2×10^{-6} M Na₂CrO₄, no precipitate is observed.
 - (i) Assuming that volumes are additive, calculate the molar concentrations of $Ba^{2+}(aq)$ and $CrO_4^{2-}(aq)$ in the 1.00 L of solution.
 - (ii) Use the molar concentrations of $Ba^{2+}(aq)$ ions and $CrO_4^{2-}(aq)$ 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.

$$CO(g) + \frac{1}{2}O_2(g) \rightarrow CO_2(g)$$

- 2. The combustion of carbon monoxide is represented by the equation above.
 - (a) Determine the value of the standard enthalpy change, ΔH_{rxn}° , for the combustion of CO(g) at 298 K using the following information.

$$C(s) + \frac{1}{2}O_2(g) \rightarrow CO(g)$$
 $\Delta H_{298}^{\circ} = -110.5 \text{ kJ mol}^{-1}$

$$C(s) + O_2(g) \rightarrow CO_2(g)$$
 $\Delta H_{298}^{\circ} = -393.5 \text{ kJ mol}^{-1}$

(b) Determine the value of the standard entropy change, ΔS_{rxn}° , for the combustion of CO(g) at 298 K using the information in the following table.

Substance	S_{298}° (J mol ⁻¹ K ⁻¹)
CO(g)	197.7
$CO_2(g)$	213.7
$O_2(g)$	205.1

- (c) Determine the standard free energy change, ΔG_{rxn}° , 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_{eq} , for the reaction at 298 K.

- 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 $CO_2(g)$ 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 CH_2Br , has a vapor density of 6.00 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.

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.

$$Ex.$$
 $Mg + Ag^+ \rightarrow Mg^{2+} + Ag$

- (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.

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 KCl, Na₂CO₃, and MgSO₄. 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 NaOH to a Solution of the Compound	Result of Adding 1.0 <i>M</i> 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 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 $1.0\,M$ NaOH. Describe the steps for preparing $100.\,\text{mL}$ of $1.0\,M$ NaOH from a stock solution of $3.0\,M$ NaOH using a $50\,\text{mL}$ buret, a $100\,\text{mL}$ volumetric flask, distilled water, and a small dropper.
- (e) Describe a simple laboratory test that you could use to distinguish between $Na_2CO_3(s)$ and $CaCO_3(s)$. In your description, specify how the results of the test would enable you to determine which compound was $Na_2CO_3(s)$ and which compound was $CaCO_3(s)$.

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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

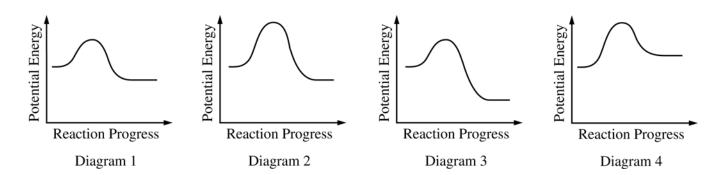
- (i) pure glucose
- (ii) pure cyclohexane
- (b) Glucose is soluble in water but cyclohexane is not soluble in water. Explain.
- (c) Consider the two processes represented below.

Process 1:
$$H_2O(l) \to H_2O(g)$$
 $\Delta H^{\circ} = +44.0 \text{ kJ mol}^{-1}$
Process 2: $H_2O(l) \to H_2(g) + \frac{1}{2} O_2(g)$ $\Delta H^{\circ} = +286 \text{ kJ mol}^{-1}$

- (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, H₂O molecules break apart to form hydrogen molecules and oxygen molecules.

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



- (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.

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 SF₄ and BF₃ react to form an ionic compound according to the following equation.

$$SF_4 + BF_3 \rightarrow SF_3BF_4$$

- (i) Draw a complete Lewis structure for the SF_3^+ cation in SF_3BF_4 .
- (ii) Identify the type of hybridization exhibited by sulfur in the SF₃⁺ cation.
- (iii) Identify the geometry of the 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 SF₃⁺ cation is larger than, equal to, or smaller than 109.5°. Justify your answer.
- (b) The compounds SF₄ and CsF react to form an ionic compound according to the following equation.

$$SF_4 + CsF \rightarrow CsSF_5$$

- (i) Draw a complete Lewis structure for the SF₅⁻ anion in CsSF₅.
- (ii) Identify the type of hybridization exhibited by sulfur in the SF_5^- anion.
- (iii) Identify the geometry of the SF₅⁻ anion that is consistent with the Lewis structure drawn in part (b)(i).
- (iv) Identify the oxidation number of sulfur in the compound CsSF₅.

- 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, CO_3^{2-} .
 - (ii) Predict whether or not the compound would be soluble in water. Explain your reasoning.

STOP

END OF EXAM