

Name: _____ Student Number: _____

Chemistry 1A03

Final Exam

February, 2010

McMaster University

VERSION 1

Instructor: P. Britz-McKibbon, G. Goward, P. Hatala, J. Landry

Duration: 3 Hours

This test contains 24 numbered pages printed on both sides. There are 35 multiple-choice questions appearing on pages numbered 3 to 21. Page 23 includes some useful data and equations. There is a periodic table on page 24. You may tear off the last page to view the periodic table and to do your rough work.

You must enter your name and student number on the question sheets, as well as on the answer sheet. Your invigilator will be checking your student card for identification.

You are responsible for ensuring that your copy of the question paper is complete. Bring any discrepancy to the attention of your invigilator.

Questions 1 to 30 are each worth 2 marks, questions 31 - 35 are each worth 3 marks; the total marks available are 75. There is no additional penalty for incorrect answers.

BE SURE TO ENTER THE CORRECT VERSION OF YOUR TEST (shown near the top of page 1), IN THE SPACE PROVIDED ON THE ANSWER SHEET.

ANSWER ALL QUESTIONS ON THE ANSWER SHEET, IN PENCIL.

Instructions for entering multiple-choice answers are given on page 2.

SELECT ONE AND ONLY ONE ANSWER FOR EACH QUESTION from the answers (A) through (E). No work written on the question sheets will be marked. The question sheets may be collected and reviewed in cases of suspected academic dishonesty.

Academic dishonesty may include, among other actions, communication of any kind (verbal, visual, etc.) between students, sharing of materials between students, copying or looking at other students' work. If you have a problem please ask the invigilator to deal with it for you. Do not make contact with other students directly. Try to keep your eyes on your own paper – looking around the room may be interpreted as an attempt to copy.

Only Casio FX 991 electronic calculators may be used; but they must NOT be transferred between students. Use of periodic tables or any aids, other than those provided, is not allowed.

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You are writing VERSION 1 of this test. Make sure you have correctly entered your version number ("1") in the correct column on your scan sheet (see p. 2 for details).

Section #1 – These questions are worth two marks each.

- _____ 1. What would be the **heat capacity** of a bomb calorimeter if the combustion of hydrogen (1 mole) with excess oxygen caused the temperature to rise from 298 K to 316 K.
($\Delta H_f^\circ \text{H}_2\text{O}(\text{l}) = -286 \text{ kJ mol}^{-1}$)
- a. 3.45 J K^{-1}
 - b. $5.02 \times 10^3 \text{ J K}^{-1}$
 - c. $9.87 \times 10^2 \text{ J K}^{-1}$
 - d. $1.59 \times 10^4 \text{ J K}^{-1}$
 - e. $6.98 \times 10^1 \text{ J K}^{-1}$
- _____ 2. A system initially has an internal energy $U = 520 \text{ J}$. It undergoes an exothermic process during which 111 J of heat is exchanged with the surroundings and 222 J of work is done on the system. What is the **final internal energy of the system?**
- a. 194 J
 - b. 431 J
 - c. 852 J
 - d. 631 J
 - e. 345 J

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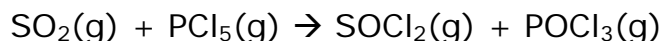
- _____ 3. Determine the **false** statement below.
- a. Converting elemental chlorine into chlorine atoms is endothermic
 - b. In a constant volume system, $\Delta U = q$
 - c. A beaker would represent an open system
 - d. Photosynthesis is an endothermic process
 - e. During the combustion of hydrogen at room temperature, work is done on the surroundings

- _____ 4. Calculate the **standard enthalpy of formation for hydrogen chloride gas** given the following data:



- a. -231.84 kJ
- b. -72.50 kJ
- c. -89.82 kJ
- d. -439.94 kJ
- e. -154.38 kJ

- _____ 5. Using the bond energy (BE) below to calculate the ΔH° for the reaction



$$\text{BE (S=O)} = 536 \text{ kJ} \cdot \text{mol}^{-1}$$

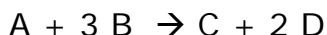
$$\text{BE (P=O)} = 460 \text{ kJ} \cdot \text{mol}^{-1}$$

$$\text{BE (P-Cl)} = 331 \text{ kJ} \cdot \text{mol}^{-1}$$

$$\text{BE (S-Cl)} = 250 \text{ kJ} \cdot \text{mol}^{-1}$$

- a. + 238 kJ
- b. + 389 kJ
- c. - 123 kJ
- d. - 348 kJ
- e. - 578 kJ

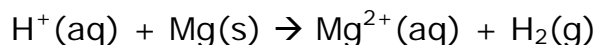
- _____ 6. Consider the following reaction where A and D are elements in their standard states.



ΔH° for the reaction would be given by

- a. $\Delta H_f^\circ (C) - \Delta H_f^\circ (A) - 3 \Delta H_f^\circ (B)$
- b. $\Delta H_f^\circ (A) + \Delta H_f^\circ (B) - \Delta H_f^\circ (C) - \Delta H_f^\circ (D)$
- c. $\Delta H_f^\circ (C) + \Delta H_f^\circ (D) - \Delta H_f^\circ (A) - \Delta H_f^\circ (B)$
- d. $\Delta H_f^\circ (C) - 3 \Delta H_f^\circ (B)$
- e. $\Delta H_f^\circ (A) + 3 \Delta H_f^\circ (B) - \Delta H_f^\circ (C) - 2 \Delta H_f^\circ (D)$

- _____ 7. In a calorimetry experiment, a respectable scientist is attempting to measure the enthalpy change for the following reaction:



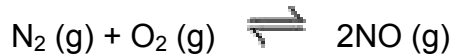
$H^+(aq)$ is the limiting reagent in the experiment and when attempting to introduce the acid into the calorimeter, a portion is spilled outside the calorimeter. **The scientist should:**

- a. Add approximately the amount of acid that was lost from the stock solution.
- b. Continue and explain the result in their discussion as a source of human error
- c. Continue with the remaining acid, estimating the acid loss and account for this in the final calculations
- d. Obtain a new, accurate amount of acid.
- e. Remove some Mg to account for the acid loss.

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8. Based on the following information:

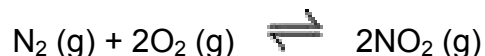


$$K_1 = 4.3 \times 10^{-25}$$



$$K_2 = 1.3 \times 10^{-5}$$

Calculate K_c for:



$$K_c = ?$$

- a. 3.1×10^{-15}
- b. 2.5×10^{-15}
- c. 7.1×10^{-25}
- d. 5.5×10^{-30}
- e. 4.8×10^{-13}

9. The partition coefficient K_{ow} describes the distribution of a persistent organic pollutant (POP) as shown by:



Select the **TRUE** statements about K_{ow} . It is:

- i. a measure of the bond energy of the POP.
- ii. related to the dipole moment of the POP.
- iii. used to assess tendency of the POP to bio-accumulate in fatty tissue.
- iv. determined by mixing the POP with water and soap.

- a. ii, iv
- b. i, iii
- c. i, ii
- d. ii, iii
- e. i, iv

- ____ 10. The following reaction takes place in a sealed steel container at 25.0°C.



The system is allowed to reach equilibrium and then 5 atm of argon gas is added. **The effect will be:**

- a. Since this is a constant volume system, the argon gas will shift the equilibrium to the right
 - b. Since this is a constant volume system, the argon gas will not affect the equilibrium
 - c. Since this is a constant pressure system, the argon gas will shift the equilibrium to the right
 - d. The argon gas will react with the hydrogen gas, lowering its partial pressure and shifting the reaction to the left
 - e. Since this is a constant pressure system, the argon gas will not affect the equilibrium
- ____ 11. **Which one of the following interactions** account for the efficacy of chemical sunscreens in preventing radiation damage to cell tissue?
- a. The quantized transitions in heavy metals like lead, upon absorbing incident light
 - b. The ability of aromatic molecules to absorb UVA/UVB light.
 - c. The absorption of electrons by low work function elements
 - d. The tendency of lipophilic drugs to bioaccumulate.
 - e. The interaction of sound waves with water.

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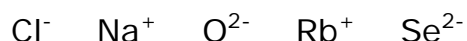
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- _____ 12. 523.0 mL of acetylene (C_2H_2) was collected over water at 23.0°C and a total barometric pressure of 0.9711 atm. If the vapour pressure of water at 23.0°C is 0.0276 atm, **how many grams of acetylene were collected?**
- a. 0.211 g
 - b. 0.318 g
 - c. 0.913 g
 - d. 0.528 g
 - e. 0.861 g
- _____ 13. **Calculate the wavelength** of a tennis ball travelling at 52 m/s if the ball weighs 70. g.
- a. 1.8×10^{-34} m
 - b. 3.1×10^{-34} m
 - c. 4.7×10^{-20} m
 - d. 9.1×10^{-25} m
 - e. 3.1×10^{-25} m
- _____ 14. Which of the following statements is **false**?
- a. Bohr's model of the atom put electrons in a random cloud around the atom
 - b. De Broglie combined Einstein's energy equation with Planck's energy equation to derive an equation that determines the wavelength of particles with mass
 - c. If there are 3 electrons in a p-subshell, the ground state configuration will have them all with the same magnetic spin
 - d. For $l=0$, there is only one subshell
 - e. An electron in hydrogen requires $-R_{\text{H}}/n^2$ joules of energy to be ionized

____ 15. Which of the following **differences in ionization energy** would you expect to be the **largest for phosphorous**?

- a. I.E. 1 \rightarrow I.E. 2
- b. I.E. 3 \rightarrow I.E. 4
- c. I.E. 2 \rightarrow I.E. 3
- d. I.E. 5 \rightarrow I.E. 6
- e. I.E. 0 \rightarrow I.E. 1

____ 16. Which of the following statements below is false about the series of ions listed:



- a. Rb^+ has a larger ionic radius than Na^+
- b. O^{2-} and Na^+ are isoelectronic
- c. Se^{2-} has the largest ionic radius
- d. Na^+ has the smallest ionic radius
- e. Rb^+ and Cl^- are isoelectronic

____ 17. Based on a formal-charge minimized Lewis structure for TeO_4^{2-} **the average Te-O bond order (BO) and O formal charge (FC) would be:**

- | | B.O. | F.C. |
|----|------|------|
| a. | 1 | 1 |
| b. | 1.5 | -0.5 |
| c. | 1 | -1.5 |
| d. | 0.7 | -0.5 |
| e. | 2.5 | -2 |

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____ 18. Which of the following salts will form a **weakly acidic solution** upon dissolution:

- i. CaI_2
- ii. NH_4Cl
- iii. NaOOCH_3
- iv. KNO_3
- v. HF

- a. i
- b. iv
- c. ii, v
- d. ii, iii
- e. i, iv

____ 19. Which of the following statements is **correct**?

- a. $\text{H}_2\text{SeO}_4 > \text{H}_2\text{SeO}_3$ - the stronger acid is listed first
- b. $\text{HCl} > \text{HBr}$ - the stronger acid is listed first
- c. $\text{PH}_3 > \text{H}_2\text{S}$ - the stronger acid is listed first
- d. $\text{CH}_3\text{COOH} > \text{CCl}_3\text{COOH}$ - the stronger acid is listed first
- e. $\text{NI}_3 > \text{NH}_3$ - the stronger base is listed first

____ 20. Which of the following statements are **TRUE**?

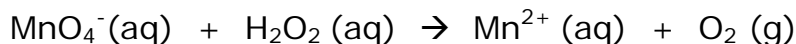
- i. For binary acids, the larger the electronegativity difference, the easier it is to remove a proton, and therefore the stronger the acid.
- ii. The position of a halogen atom relative to the carboxyl group influences the strength of an oxyacid because it weakens the O-H bond by withdrawing electron density from oxygen.
- iii. When comparing two different oxyacids, the one with the greater number of protons will be the stronger acid, because there are more protons to donate.
- iv. The presence of a halogen on an amine (for example, NH_2Cl) weakens it as a base because it makes the nitrogen more positive and therefore less likely to accept a proton.
- v. The more charge-minimized resonance structures that can be drawn for the conjugate base of an oxyacid the stronger the acid.

- a. iii, v
- b. ii, iv
- c. i, iv, v
- d. i, iii, iv
- e. ii, iv, v

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- ____ 21. Select the correct **whole-number coefficient terms** for H_2O_2 , Mn^{2+} and H_2O after balancing the following redox equation under **acidic conditions**:



- a. 3, 1, 4
- b. 1, 3, 5
- c. 2, 4, 6
- d. 4, 6, 1
- e. 5, 2, 8

- ____ 22. Which of the following statements are **FALSE**?

- a. The production of iron from iron ore requires coke, C (s) as the oxidizing agent
- b. Flue gas desulfurization technology uses a slaked lime slurry, $\text{Ca}(\text{OH})_2$ to reduce the impact of gas emissions associated with acid rain
- c. Hydroxylapatite from charred bone can be used as a filter to selectively absorb chloride from contaminated water
- d. HF is a stronger acid than HCl due to its weaker bond energy
- e. Cold packs function based on the dissolution of ammonium nitrate which is highly endothermic

- a. i, v
- b. ii, iii, v
- c. iii, iv
- d. i, iii, iv
- e. ii, iii

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____ 23. Determine the **K_{sp} of mercury (II) bromide** if the equilibrium concentration of bromide in solution is measured to be $5.00 \times 10^{-7} \text{ M}$ at 25°C :

- a. 4.37×10^{-16}
- b. 8.72×10^{-25}
- c. 1.67×10^2
- d. 2.96×10^{-44}
- e. 6.25×10^{-20}

____ 24. Predict whether the following unbalanced chemical reactions will form an **insoluble precipitate**:

- i. $\text{BaCl}_2 (\text{aq}) + \text{NaHPO}_4 (\text{aq}) + \text{K}_2\text{S} (\text{aq}) \rightarrow$
- ii. $\text{Zn} (\text{s}) + \text{H}^+ (\text{aq}) \rightarrow$
- iii. $\text{AlCl}_3 (\text{aq}) + \text{NaOH} (\text{aq}) \rightarrow$
- iv. $\text{AgNO}_3 + \text{NaCl} (\text{aq}) \rightarrow$
- v. $\text{NH}_4\text{Cl} (\text{aq}) + \text{LiBr} (\text{aq}) \rightarrow$

- a. ii, iii, v
- b. iii, v
- c. i, iii, iv
- d. ii, v
- e. i, ii

- ____ 25. The voltaic cell with the net reaction represented by the following equation has:

$$E^\circ_{\text{cell}} = -0.760 \text{ V at } 25^\circ \text{C.}$$



Calculate ΔG° at 25°C . What is the direction of spontaneous reaction in a cell at 25°C in which all species are in their standard states?

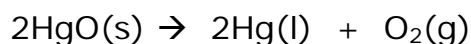
- a. 283 kJ, spontaneous in the reverse direction
 - b. 147 kJ, spontaneous in the forward direction
 - c. 147 kJ, spontaneous in the reverse direction
 - d. -84.5 kJ, spontaneous in the forward direction
 - e. -84.5 kJ, spontaneous in the reverse direction
- ____ 26. For each of the pairs listed, **which is ordered from highest entropy to lowest entropy**. Assume all substances are at 25°C , and 1atm.
- a. 1 mol of $\text{SnCl}_4(\text{g})$, 1 mol of $\text{SnCl}_4(\text{l})$
 - b. 1 mol of $\text{C}(\text{s})$ diamond, 1 mol of $\text{C}(\text{s})$ graphite
 - c. 1 mol of $\text{H}_2\text{O}(\text{l})$, 1 mol of $\text{NH}_3(\text{l})$
 - d. 1 mol of $\text{O}_2(\text{g})$, 1 mol of $\text{O}_3(\text{g})$
 - e. 1 mol of ethene (g), 1 mol of ethane (g)

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27. Oxygen was first made by Joseph Priestley (1733-1804) by heating HgO(s). Use the thermodynamic data below, that apply at 25 °C, to **estimate the temperature** at which HgO(s) in a sealed vessel decomposes sufficiently to be in equilibrium with 1 atm pressure of O₂(g).

(Hint: Finding the temperature at which $\Delta G_r^\circ = 0$)



$$S^\circ(\text{Hg,l}) = 76.02 \text{ JK}^{-1}\text{mol}^{-1}$$

$$\Delta H_f^\circ(\text{HgO,s}) = -90.83 \text{ kJmol}^{-1}$$

$$S^\circ(\text{O}_2\text{,g}) = 205.138 \text{ JK}^{-1}\text{mol}^{-1}$$

$$S^\circ(\text{HgO,s}) = 70.29 \text{ JK}^{-1}\text{mol}^{-1}$$

- a. T = 735.8 K
- b. T = 911.5 K
- c. T = 553.4 K
- d. T = 838.7 K
- e. T = 445.2 K

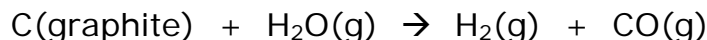
28. Which of the following correctly represent the thermodynamics of the given reactions:

Reaction	ΔH° kJ	ΔS° J/K	Reaction Type
i. $\text{CH}_4\text{(g)} + 2\text{O}_2\text{(g)} \rightarrow 2\text{H}_2\text{O(l)} + \text{CO}_2\text{(g)}$	-890.6	-242.8	spontaneous at low T, NOT spontaneous at high T
ii. $2\text{Fe}_2\text{O}_3\text{(s)} + 3\text{C(graphite)} \rightarrow 4\text{Fe(s)} + 3\text{CO}_2\text{(g)}$	+467.9	+560.7	NOT spontaneous at low T, spontaneous at high T
iii. $\text{C(graphite)} + \text{O}_2\text{(g)} \rightarrow \text{CO}_2\text{(g)}$	-393.5	+3.1	spontaneous at all T
iv. $\text{N}_2\text{(g)} + 3\text{F}_2\text{(g)} \rightarrow 2\text{NF}_3\text{(g)}$	-264.2	-277.8	non spontaneous at all T

- a. i, ii, iii
- b. ii, iv
- c. iii, iv
- d. ii, iii
- e. i, iv

- ____ 29. A crucial reaction for the production of synthetic fuels is the conversion of coal to H₂ with steam. **Calculate the value of K** for the reaction at 25 °C.

$$\Delta G_f^0(\text{H}_2\text{O}, \text{g}) = -228.572 \text{ kJmol}^{-1} \quad \Delta G_f^0(\text{CO}, \text{g}) = -137.168 \text{ kJmol}^{-1}$$



- a. 9.50×10^{-17}
 - b. 4.43×10^{-3}
 - c. 6.55×10^{18}
 - d. 1.10×10^{-8}
 - e. 3.87×10^{-12}
- ____ 30. Choose the **false** statement from the following:
- a. For a spontaneous reaction, the sum of the entropy change of the system and the surroundings must be greater than zero.
 - b. The spontaneous reaction of sodium azide upon detonation of an airbag is driven by the formation of the stable nitrogen triple bond.
 - c. The spontaneous reaction in a galvanic cell provides a means of capturing electrical energy with high efficiency.
 - d. The usefulness of the Gibbs Free Energy is that the spontaneity of a reaction may be determined through consideration of the system alone.
 - e. Phase transitions occur spontaneously because ΔG at the transition is less than zero, and the equilibrium constant is greater than 1.

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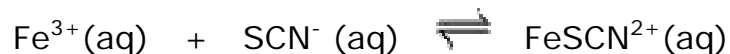
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Section #2 – These questions are worth three marks each.

- ____ 31. Two beakers of ethylene glycol are mixed in a closed system. Beaker A is twice the volume and at 4 times the temperature of Beaker B, which is at 10°C . What is the **final temperature** of the combination.
- a. 30°C
 - b. 80°C
 - c. 15°C
 - d. 33.33°C
 - e. 20°C

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___ 32. Consider the following chemical equilibrium:



If absorbance = $540 \times [\text{FeSCN}^{2+}]$ (the only species that absorbs visible light), **determine the equilibrium constant** when equal volumes of 2.5×10^{-3} M solutions of $\text{Fe}(\text{NO}_3)_3$ and KSCN are mixed together and produce an absorbance of 0.34 at 447 nm.

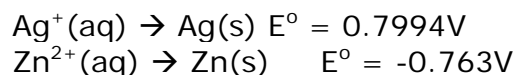
- a. 1600
- b. 40
- c. 560
- d. 1.5×10^{-3}
- e. 3400

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- _____ 33. Calculate the **pH** of a saturated solution containing the **sparingly soluble solid**, Ca(OH)_2 given that its $K_{\text{sp}} = 5.02 \times 10^{-6}$ at 25°C :
- a. 4.652
 - b. 7.254
 - c. 12.334
 - d. 2.527
 - e. 9.843

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- ____ 34. One half-cell in a voltaic cell at 25 °C is constructed from a silver wire dipped into a AgNO_3 solution of unknown concentration. The other half-cell consists of a zinc electrode in a 1.0 M $\text{Zn(NO}_3)_2$ solution. The cell emf is measured to be 1.48 V. **What is the concentration of $\text{Ag}^+(\text{aq})$ ions?** The reduction potentials involved are:



- a. 0.84 M
- b. 0.018 M
- c. 0.22 M
- d. 0.35 M
- e. 0.041 M

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- ____ 35. The thermodynamic data relevant to the vapour pressure of ethanol (at 25°C) is $\Delta H_{\text{vap}}^\circ$ is 39.33 kJ mol⁻¹, and $\Delta G_{\text{vap}}^\circ$ is 6.29 kJ mol⁻¹. **Estimate the boiling point of ethanol** at 1.0 atm pressure (the temperature at which the equilibrium vapour pressure is equal to 1.0 atm).
- a. 76
 - b. 82
 - c. 90
 - d. 94
 - e. 73

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ROUGH WORK

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Some general data are provided on this page and the next page. Other data appear with the questions.

A periodic table is provided on the next page.

$$\text{STP} = 273.15 \text{ K}, 1 \text{ atm} \quad F = 96485 \text{ C/mol}$$

$$R = 8.3145 \text{ J/K}\cdot\text{mol} = 0.08206 \text{ L}\cdot\text{atm/K}\cdot\text{mol} \quad N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$$

$$1 \text{ atm} = 760 \text{ mm Hg} = 101.325 \text{ kPa} \quad 0^\circ\text{C} = 273.15 \text{ K}$$

$$1 \text{ J} = 1 \text{ kg m}^2 \text{ s}^{-2} = 1 \text{ kPa}\cdot\text{L} = 1 \text{ Pa}\cdot\text{m}^3 \quad 1 \text{ m} = 10^9 \text{ nm} = 10^{10} \text{ \AA}$$

$$1 \text{ cm}^3 = 1 \text{ mL} \quad 1 \text{ g} = 10^3 \text{ mg} = 10^{-3} \text{ kg}$$

$$1 \text{ Hz} = 1 \text{ cycle/s} \quad c = 2.9979 \times 10^8 \text{ m/s}$$

$$h = 6.6256 \times 10^{-34} \text{ J}\cdot\text{s} \quad m_e = 9.10 \times 10^{-31} \text{ kg}$$

$$\lambda = h / \mu = h / p$$

$$E_n = -R_H / n^2 = -2.179 \times 10^{-18} \text{ J} / n^2 \text{ (} R_H \text{ is the energy form of the Rydberg constant for H)}$$

$$K_w = 1.0 \times 10^{-14} \text{ (} 25^\circ\text{C)} \quad C_{\text{water}} = 4.184 \text{ J/g}\cdot\text{K}$$

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

$$E_{\text{cell}} = E^\circ_{\text{cell}} - \frac{RT}{nF} \ln Q = E^\circ_{\text{cell}} - \frac{0.0592}{n} \log_{10} Q$$

TABLE 5.1 Solubility Guidelines for Common Ionic Solids

Follow the lower-numbered guideline when two guidelines are in conflict. This leads to the correct prediction in most cases.

1. Salts of group 1 cations (with some exceptions for Li^+) and the NH_4^+ cation are soluble.
2. Nitrates, acetates, and perchlorates are soluble.
3. Salts of silver, lead, and mercury(I) are insoluble.
4. Chlorides, bromides, and iodides are soluble.
5. Carbonates, phosphates, sulfides, oxides, and hydroxides are insoluble (sulfides of group 2 cations and hydroxides of Ca^{2+} , Sr^{2+} , and Ba^{2+} are slightly soluble).
6. Sulfates are soluble except for those of calcium, strontium, and barium.

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PERIODIC TABLE OF THE ELEMENTS															
ALDRICH®															
Transition Metals															
I	II	III	IV	V	VI	VII	VIII	IX	X	XI	XII	III	IV	V	VI
1 H 1.0079	2 He 4.0026	3 Li 6.941	4 Be 9.0122	5 B 10.811	6 C 12.011	7 N 14.007	8 O 15.999	9 F 18.998	10 Ne 20.180	11 Na 22.990	12 Mg 24.305	13 Al 26.982	14 Si 28.086	15 P 30.974	16 S 32.066
19 K 39.098	20 Ca 40.078	21 Sc 44.956	22 Ti 47.88	23 V 50.942	24 Cr 51.996	25 Mn 54.938	26 Fe 55.847	27 Co 58.933	28 Ni 58.69	29 Cu 63.546	30 Zn 65.39	31 Ga 69.723	32 Ge 72.61	33 As 74.922	34 Se 78.96
37 Rb 85.468	38 Sr 87.62	39 Y 88.906	40 Zr 91.224	41 Nb 92.906	42 Mo 95.94	43 Tc [98]	44 Ru 101.07	45 Rh 102.91	46 Pd 105.42	47 Ag 107.87	48 Cd 112.41	49 In 114.82	50 Sn 118.71	51 Sb 121.75	52 Te 127.60
55 Cs 132.91	56 Ba 137.33	57 *La 138.91	58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm [145]	62 Sm 150.36	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04
87 Fr [223]	88 Ra 226.03	89 **Ac 227.03	90 Th 232.04	91 Pa 231.04	92 U 238.03	93 Np 237.05	94 Pu [244]	95 Am [243]	96 Cm [247]	97 Bk [247]	98 Cf [251]	99 Es [252]	100 Fm [257]	101 Md [258]	102 No [259]
Atomic weights are based on $^{12}\text{C} = 12$ and conform to the 1987 IUPAC report values rounded to 5 significant digits. Numbers in [] indicate the most stable isotope.															

* Lanthanides													
58 Ce 140.12	59 Pr 140.91	60 Nd 144.24	61 Pm [145]	62 Sm 150.36	63 Eu 151.97	64 Gd 157.25	65 Tb 158.93	66 Dy 162.50	67 Ho 164.93	68 Er 167.26	69 Tm 168.93	70 Yb 173.04	71 Lu 174.97

** Actinides													
90	91	92	93	94	95	96	97	98	99	100	101	102	103
Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	Lr
232.04	231.04	238.03	237.05	[244]	[243]	[247]	[247]	[251]	[252]	[257]	[258]	[259]	[262]