Chemistry 1A03 Final Exam Dec. 13, 2010

McMaster University VERSION 1

Instructor: Drs. R. Dumont, G. Goward, J. Landry and P. Lock

Duration: 180 minutes

This test contains 28 numbered pages printed on both sides. There are **35** multiple-choice questions appearing on pages numbered 3 to 23. Pages 24, 25 and 26 are extra space for rough work. Page 27 includes some useful data and equations, and there is a periodic table on page 28. You may tear off the last page to view the periodic table and the data provided.

You must enter your name and student number on this question sheet, 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 27 are each worth 2 marks and questions 28 - 35 are worth 3 marks each; the total marks available are 78. 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.

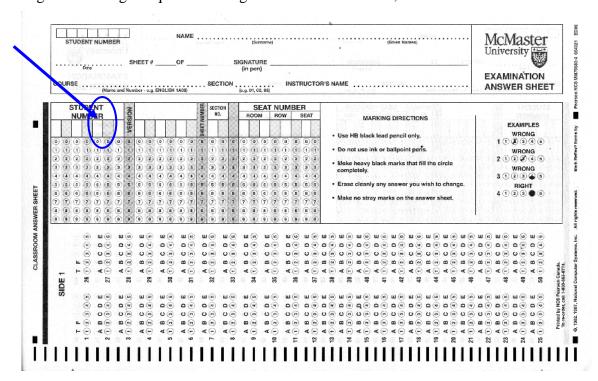
Only Casio FX 991 electronic calculators may be used; but they must NOT be transferred between students. Use of periodic tables or any aids, other than those provided, is not allowed.

OMR EXAMINATION – STUDENT INSTRUCTIONS

NOTE: IT IS YOUR RESPONSIBILITY TO ENSURE THAT THE ANSWER SHEET IS PROPERLY COMPLETED: YOUT EXAMINIATION RESULT DEPENDS UPON PROPER ATTENTION TO THESE INSTRUCTIONS.

The scanner, which reads the sheets, senses the bubble shaded areas by their non-reflection of light. A heavy mark must be made, completely filling the circular bubble, with an HB pencil. Marks made with a pen will **NOT** be sensed. Erasures must be thorough or the scanner will still sense a mark. Do **NOT** use correction fluid on the sheets. Do **NOT** put any unnecessary marks or writing on the sheet.

- 1. On SIDE 1 (**red side**) of the form, in the top box, *in pen*, print your student number, name, course name, (section number, instructor name) and the date in the spaces provided. Then you **MUST** write your signature, in the space marked SIGNATURE.
- 2. In the second box, *with a pencil*, mark your student number, **exam version number** (and course section number) in the space provided and fill in the corresponding bubble numbers underneath.
- 3. Answers: mark only **ONE** choice from the alternatives (1,2,3,4,5 or A,B,C,D,E) provided for each question. If there is a True/False question, enter response o 1 (or A) as True, and 2 (or B) as False. The question number is to the left of the bubbles. Make sure that the number of the question on the scan sheet is the same as the number on the test paper.
- 4. Pay particular attention to the Marking+ Directions on the form.
- 5. Begin answering the question using the first set of bubbles, marked "1".



Name:	Student number:
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Questions 1-27 are worth 2 marks each.

- 1. Which atomic property decreases **down** a group?
 - A) the metallic character
 - B) the first ionization energy
 - C) the atomic radius
 - D) the ionic radius
 - E) the core charge

- 2. Which **ONE** of the following pairs of reagents produces **no visible change** when they are mixed together?
 - A) Cu(s) + HCl(aq)
 - B) $KCl(aq) + AgClO_4(aq)$
 - C) Zn(s) + HCl(aq)
 - D) NaOH(aq) + phenolphthalein
 - $E) \quad Cu(s) + HNO_{3}(aq) \\$

3. **Balance** the following redox reaction in **basic** solution. Use the smallest whole number coefficients.

$$Cr(OH)_3(s) + ClO^-(aq) + OH^-(aq) \rightarrow Cr{O_4}^2 - (aq) + Cl^-(aq) + H_2O(l)$$

When this has been done correctly, the stoichiometric **coefficients** for the **reactant** species, in order from **left** to **right** are:

- A) 2, 3, 4
- B) 3, 2, 3
- C) 2, 4, 6
- D) 2, 3, 7
- E) 2, 4, 5

- 4. Which **one** species contains the greatest number of **protons**?
 - A) P⁻
 - B) Cl
 - C) S^{2+}
 - D) Si
 - E) F

5. The octanol-water partition coefficient of tetrachlorodibenzofuran (TCDF) is $K_{\rm ow} = 3.16 \times 10^7$.

 $TCDF(aq) \stackrel{?}{=} TCDF(org)$

With regard to this persistent organic pollutant (POP), choose the **one FALSE** statement.

- A) Adverse health effects of long term exposure to POPs such as TCDF is a public health concern.
- The molecule is highly lipophilic, and therefore accumulates in fat tissue. B)
- The $\log K_{ow} = 7.500$
- D) The electron affinity of the chlorine atoms makes the molecule detectable by electron capture detection.
- The equilibrium lies far to the left due to the high hydrophilicity of the molecule. E)
- 6. Using an ice calorimeter such as that in experiment # 5, it was determined that the reaction of 0.14 g of zinc with excess HCl (aq), caused 0.68 g of ice to melt. What was the heat of reaction per mole of Zn (kJ mol⁻¹)? $[\Delta H_{\text{fus}}(\text{ice}) = 333 \text{ J g}^{-1}]$

- A) -180
- B) –95
- C) -110
- D) 240
- E) 590

- 7. For which of the following processes is work done by the system on the surroundings?
 - (i) $2 \text{ NH}_3(g) + 3 \text{ N}_2\text{O}(g) \rightarrow 4 \text{ N}_2(g) + 3 \text{ H}_2\text{O}(1)$
 - (ii) $N_2H_4(1) + H_2O(1) \rightarrow N_2O(g) + 3 H_2(g)$
 - (iii) $N_2H_4(1) + H_2O(1) \rightarrow 2 NH_3(g) + 1/2 O_2(g)$
 - (iv) $H_2(g) + 1/2 O_2(g) \rightarrow H_2O(1)$
 - (v) $N_2(g) + 2 H_2O(1) \rightarrow N_2H_4(1) + O_2(g)$

.

- A) i, iv
- B) i, ii
- C) iv, v
- D) ii, iii
- E) iii, v
- 8. How much heat (in kJ) is required to convert 36.0 g of liquid H_2O at 50 °C to gaseous H_2O at 100 °C? $\Delta H_{vaporization}$ for H_2O (l) = 44.0 kJ/mol. The specific heat of liquid H_2O is 4.184 J K⁻¹ g⁻¹.

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- A) 84.0
- B) 8.56
- C) 7.12×10^3
- D) 67.8
- E) 95.5

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9. At 1000 K, initially pure $NO_2(g)$ decomposes according to $2 NO_2(g) \implies 2 NO(g) + O_2(g)$

with an equilibrium constant, $K_p = 158$, when pressures are expressed in atm. At equilibrium, the partial pressure of $O_2 = 0.25$ atm. What are the **partial pressures** (in atm) of **NO(g)** and **NO₂(g)** (in that order) at equilibrium at 1000 K?

- A) $0.250, 2.0 \times 10^{-2}$
- B) $0.50, 4.0 \times 10^{-2}$
- C) $0.125, 2.0 \times 10^{-2}$
- D) $0.50, 2.0 \times 10^{-2}$
- E) $0.125, 4.0 \times 10^{-4}$

10. The reaction,

$$Ca(s) + 2 H_2O(1) \rightarrow Ca(OH)_2(aq) + H_2(g)$$

is carried out in a constant pressure calorimeter with heat capacity, C (J/K). The temperature was observed to increase. For this experiment, which $\bf ONE$ of the following statements is $\bf FALSE$?

- A) $\Delta H = -C\Delta T$
- B) $\Delta H < \Delta U$
- C) Work is done by the system.
- D) $\Delta H \neq \Delta U$
- E) This reaction is exothermic.

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11. $PCl_5(s)$ can be prepared by the reaction $PCl_3(l) + Cl_2(g) \rightarrow PCl_5(s)$. Calculate the enthalpy change (in kJ) that accompanies the production of 100.0 g of $PCl_5(s)$ by the above reaction, given the following data.

$$P_4(s) + 6 Cl_2(g) \rightarrow 4 PCl_3(l)$$
 $\Delta H^o = -1280 \text{ kJ}$
 $P_4(s) + 10 Cl_2(g) \rightarrow 4 PCl_5(s)$ $\Delta H^o = -1774 \text{ kJ}$

- .
- A) +79.1
- B) -238
- C) -59.3 D) -114
- E) +164

- 12. Which **one** of the following statements is **FALSE**?
 - A) For constant pressure processes, the enthalpy change of a system ($\Delta H_{\rm sys}$) is different from its energy change ($\Delta U_{\rm sys}$) by the amount $P\Delta V$.
 - B) Enthalpy is a state function.
 - C) Work flow out of a system is considered positive when calculating $\Delta U_{\rm sys}$.
 - D) Heat flow into a system increases the energy of the system.
 - E) When neither energy nor mass can enter or escape a system, it is said to be isolated.

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- 13. When a particular gas expands against a constant external pressure of 1.00 atm, it does 5 J of work done on the surroundings. During this transformation the gas also absorbs 10 J of heat. What is the **energy change**, ΔU (in J), for the gas?
 - . A) 0
 - B) -15
 - C) +15
 - D) -5
 - E) +5

14. Predict the **approximate pH** of a solution formed when sodium formate (HCOONa) is dissolved in water.

 $K_a \text{ (HCOOH)} = 1.8 \times 10^{-4}$

- A) Neutral
- B) Weakly Acidic
- C) Strongly Acidic
- D) Weakly Basic
- E) Strongly Basic

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- 15. Which of the following statements is **TRUE** regarding the acids HClO₃ and HClO₂?
 - (i) HClO₃ is a stronger acid than HClO₂.
 - (ii) There are six nonbonding electron pairs in the charge-minimized Lewis structure of HClO₂.
 - (iii) The conjugate base of HClO₃ has three equivalent resonance structures.
 - (iv) The average formal charge on O in the conjugate base of $HClO_2$ is -1.
 - A) i, ii, iii
 - B) iii, iv
 - C) ii, iii, iv
 - D) ii, iv
 - E) i, iii

- 16. Consider all of the following molecules to act as Brønsted acids. Which one would have the **strongest conjugate base**?
 - A) NH₃
 - B) H₂O
 - C) HF
 - D) HCl
 - E) CH₄

Name:	Student number:

- 17. In the reaction of hydrofluoric acid with water to produce hydronium ion and fluoride, water is acting as?
 - i) Arrhenius Base
 - ii) Brønsted Acid
 - iii) Lewis Base
 - iv) Brønsted Base
 - v) Lewis Acid
 - vi) Arrhenius Acid
 - A) iii, iv
 - B) ii, v, vi
 - C) iv, v
 - D) v, vi
 - E) i, iii, iv
- 18. What is the **pH** of a 0.0342 M solution of HCl?
 - A) 3.492
 - B) 1.081
 - C) 1.466
 - D) 4.456
 - E) 7.568

- 19. Which **one** of the following statements is **TRUE** for a 0.1 M aqueous solution of a weak monoprotic acid?
 - A) $[OH^{-}] > [H^{+}]$
 - B) pH = 1
 - C) pH > 1
 - D) $[H^+] < [A^-]$
 - E) pH < 1

20. For which of the following reactions would you **predict** $\Delta H_{\rm rxn}^{\circ} < 0$ and $\Delta S_{\rm rxn}^{\circ} > 0$?

- (i) $N_2(g) + 3 H_2(g) \rightarrow 2 NH_3(g)$
- (ii) $Pb^{2+}(aq) + SO_4^{2-}(aq) \rightarrow PbSO_4(s)$
- (iii) $O_3(g) \rightarrow O_2(g) + O(g)$
- (iv) $5 \text{ C(s)} + 4 \text{ KNO}_3(\text{s}) \rightarrow 5 \text{ CO}_2(\text{g}) + 2 \text{ N}_2(\text{g}) + 2 \text{ K}_2\text{O(s)}$
- A) ii
- B) iv
- C) iii
- D) i
- E) none

21. You are provided with four 0.035 M metal solutions (list below) and analytical glassware for diluting samples. You are required to create an electrochemical cell that will produce a potential of 0.814 V. Which metal combination would be the best to start with?

$$Au^{3+} + 3e^{-} \rightarrow Au (s);$$
 $E^{\circ}_{red} = 1.52 \text{ V}$
 $Ag^{+} + e^{-} \rightarrow Ag (s);$ $E^{\circ}_{red} = 0.800 \text{ V}$
 $Cd^{2+} + 2e^{-} \rightarrow Cd (s);$ $E^{\circ}_{red} = -0.403 \text{ V}$
 $Mn^{2+} + 2e^{-} \rightarrow Mn (s);$ $E^{\circ}_{red} = -1.18 \text{ V}$

- .
- A) Ag^+/Ag^+
- B) Cd^{2+}/Au^{3+}
- C) Mn^{2+}/Cd^{2+}
- Cd^{2+}/Ag^{+}
- E) Mn^{2+}/Au^{3+}

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22. What is the value of ΔG° (in kJ) at 298K for $4 \text{ Cl}_2(g) + \text{CH}_4(g) \rightarrow \text{CCl}_4(l) + 4 \text{ HCl}(g)$? Use the following ΔG_f° data at 298K:

$$\Delta G_f^{\circ}(CH_4(g)) = -50.8 \text{ kJ/mol}$$

 $\Delta G_f^{\circ}(CCl_4(l)) = -65.3 \text{ kJ/mol}$

$$\Delta G_f^{\circ}(HCl(g)) = -95.3 \text{ kJ/mol}$$

.

- A) -260.8
- B) -433.6
- C) -371.9
- D) -395.7
- E) +290.3

23. Acid rain is produced through the reaction of sulfur dioxide with water to give an acidic solution. The reaction is $SO_2(g) + H_2O(l) \rightarrow H_2SO_3(aq)$. Use the thermochemical information given below to calculate the **standard Gibbs free energy** (in kJ) of the combination reaction at 298K.

Data:

$$\begin{array}{cccc} & H_2O(l) & SO_2(g) & H_2SO_3(aq) \\ \Delta H_f^{\circ} \, / \, (kJ \; mol^{-1}) & -285.83 & -296.83 & -627.98 \\ S^{\circ} \, / \, (J \; mol^{-1}K^{-1}) & 69.91 & 248.22 & 132.38 \end{array}$$

.

- A) +129
- B) +10.0
- C) -254
- D) +55.3
- E) -162

Name:	Student number:

24. The reaction,

$$2 \text{ SO}_3(g) \rightarrow 2 \text{ SO}_2(g) + \text{O}_2(g)$$
 carried out at 298K, where $\Delta H = 198$ kJ, will

- A) be spontaneous only in the reverse direction
- B) be spontaneous at any temperature
- C) not be spontaneous at any temperature
- D) be spontaneous only at low temperature
- E) be spontaneous only at high temperature

- 25. Choose the **FALSE** statement regarding entropy.
 - A) Entropy is a property of state; its changes do not depend on the path taken by the system.
 - B) At T = 0 K, a pure, perfect crystal has zero entropy.
 - C) A spontaneous process always implies an increase in the entropy of the universe.
 - D) Entropy is an intensive property, in other words, it does not depend on the amount of substance present.
 - E) The processes of melting and boiling are accompanied by positive changes of entropy of the substance.

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26. Calculate the **boiling point** (in °C) of bromine given the following data:

$$\Delta H_{\rm f}^{\circ}[{\rm Br}_2({\rm g})] = 30.907 \text{ kJ mol}^{-1}$$

 $S^{\circ}[{\rm Br}_2({\rm l})] = 152.2 \text{ J K}^{-1} \text{ mol}^{-1}$
 $S^{\circ}[{\rm Br}_2({\rm g})] = 245.463 \text{ J K}^{-1} \text{ mol}^{-1}$

- .
- A) 19
- B) 311
- C) 123
- D) 4
- E) 58

- 27. The interior and exterior of a nerve cell behave as a concentration cell. If the concentration of K^+ outside the cell is 0.030 M and the concentration inside is 0.30 M, what is the **potential difference** across the cell membrane, in volts V? Assume normal body temperature, $T = 37^{\circ}C$ and a one-electron transfer (z = 1).
 - .
 - A) 0.00517
 - B) 0.132
 - C) 0.0267
 - D) 0.0615
 - E) 0.198

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Questions 28-35 are worth 3 marks each.

- 28. The standard enthalpies of formation of $SO_3(g)$, S(g) and O(g) are -359.7, +278.8 and +249.0 kJ mol^{-1} , respectively. Use these data to calculate the **average bond energy** (in kJ mol^{-1}) of the **S=O** double bond.
 - A) +277.9
 - B) +461.8
 - C) -277.9
 - D) -461.8
 - E) +307.8

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- 29. Which of the following statements is(are) **FALSE**? Be sure to minimize formal charges.
 - IBr₃ is T-shaped. (i)
 - The iodine atom in IBr_3 carries two nonbonded electrons pairs. The average bond order in SO_3^{2-} is 5/3. SO_3^{2-} is triangular pyramidal. The formal charge on S in SO_3^{2-} is +1. (ii)
 - (iii)
 - (iv)
 - (v)
 - A) v
 - B) iii, v
 - C) i, iii
 - D) iii, iv
 - E) i, ii

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- 30. A 1.0 M solution of HCN(aq) has a pH of 4.60. What is the K_b of CN⁻?
 - A) 3.4×10^{-6}

 - B) 2.9×10⁻⁶ C) 9.1×10⁻⁵ D) 1.6×10⁻⁵ E) 7.3×10⁻⁴

31. Consider a standard electrochemical cell built from the two half-cells below and identify the **TRUE** statement(s).

$$Cr^{3+}(aq) + 3 e^{-} \rightarrow Cr(s)$$
 $E^{\circ}_{red} = -0.740 \text{ V}$
 $NO_{3}^{-}(aq) + 4 \text{ H}^{+}(aq) + 3 e^{-} \rightarrow NO(g) + 2 \text{ H}_{2}O(l)$ $E^{\circ}_{red} = +0.956 \text{ V}$

(i) The spontaneous cell reaction is

$$Cr^{3+}(aq) + NO(g) + 2 H_2O(l) \rightarrow Cr(s) + NO_3^-(aq) + 4 H^+(aq).$$

- (ii) A schematic cell diagram could be $Cr(s) \mid Cr^{3+}(aq) \parallel HNO_3(aq) \mid NO(g) \mid Pt(s)$.
- (iii) The standard cell potential $E^{\circ}_{\text{cell}} = +0.216 \text{ V}$.
- (iv) Nitrate ions are reduced at the cathode.
- (v) ΔG° for the spontaneous cell reaction (with the smallest integer stoichiometric coefficients) is -490.9 kJ.
- A) i, ii, iii
- B) ii, iv
- C) v, iii
- D) ii, iv, v
- E) i, ii

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32. Calculate the standard **entropy change of the universe,** ΔS_{univ}° (in J K⁻¹), at 25.00°C for

$$CO(g) + 2 H_2(g) \rightarrow CH_3OH(l)$$

from the following data:

$$\begin{array}{cccc} & \Delta H_{\rm f}^{\circ} \ ({\rm kJ\ mol}^{-1}) & S^{\circ} \ ({\rm J\ K}^{-1} \ mol}^{-1}) \\ H_2(g) & 0 & 130.684 \\ CO(g) & -110.525 & 197.674 \\ CH_3OH(l) & -238.66 & 126.8 \end{array}$$

- .
- A) +818.8
- B) +97.52
- C) +1121
- D) -312.2
- E) +423.8

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33. The maximum wavelength of light sufficient to break a Cl-Cl bond is 489 nm. Given this information, and the data below, determine the **bond energy** (kJ mol⁻¹) for the H-Cl bond.

$$BE(H-H) = 436 \text{ kJ mol}^{-1}$$

 $\Delta H_1^{\circ}(HCl, (g)) = -92.31 \text{ kJ mol}^{-1}$

- A) 386
- B) -433
- C) -386
- D) 433
- E) 288

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- 34. A 1.25 L vessel contains 1.00 L of HF(g) at 3.00 atm and 250. mL of an HF(aq) solution at 298K. 23.5% of the total HF in the vessel is in the aqueous solution. What is the **pH** of the solution? K_a (HF) = 6.6×10^{-4}
 - . A) 2.75
 - B) 2.50
 - C) 1.75
 - D) 2.25
 - E) 2.00

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35. The K_{sp} value for SrSO₄ is 7.6×10^{-7} , and the K_{sp} value for SrF₂ is 7.9×10^{-10} . Sr(NO₃)₂(s) is added to 1.00 L of solution containing 0.020 mol F and 0.10 mol of SO₄²⁻ with constant volume.

Which salt **precipitates first**, and what is the $[Sr^{2+}]$ in solution when the precipitate forms?

- A) SrSO₄ precipitates first, when $[Sr^{2+}] = 5.3 \times 10^{-9} \text{ M}$
- B) SrF₂ precipitates first, when $[Sr^{2+}] = 2.0 \times 10^{-6} \text{ M}$
- C) SrF₂ precipitates first, when $[Sr^{2+}] = 1.6 \times 10^{-7} \text{ M}$
- D) SrF₂ and SrSO₄ precipitate together when $[Sr^{2+}] = 3.8 \times 10^{-4} \text{ M}$
- E) SrSO₄ precipitates first, when $[Sr^{2+}] = 7.7 \times 10^{-8} \text{ M}$

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Extra space for rough work:

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- Some general data are provided on this page.
- A Periodic Table with atomic weights is provided on the next page.

STP = 273.15 K, 1 atm F = 96485 C/mol $R = 8.3145 \text{ J K}^{-1} \text{ mol}^{-1} = 0.08206 \text{ L atm K}^{-1} \text{ mol}^{-1}$ $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$ $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$ $N_A = 6.6256 \times 10^{-34} \text{ Js}$ $N_A = 6.6256 \times 10^{-34} \text{ Js}$

1 atm = 101.325 kPa = 760 mm Hg 1 J = 1 kg m² s⁻² = 1 kPa L = 1 Pa m³ 1 cm³ = 1 mL 1 Hz = 1 cycle/s 0°C = 273.15 K 1 m = 10⁹ nm = 10^{10} Å 1 g = 10^3 mg

De Broglie wavelength:

 $\Delta H^{o}_{vap}[H_2O] = 44.0 \text{ kJ mol}^{-1}$

$$\lambda = h / mv = h / p$$

Hydrogen atom energy levels:

$$E_n = -R_H / n^2 = -2.178 \times 10^{-18} \text{ J} / n^2$$

Nernst Equation:

$$E = E^{\circ} - \frac{RT}{zF} \ln Q = E^{\circ} - \frac{0.0257 \text{ V}}{z} \ln Q = E^{\circ} - \frac{0.0592 \text{ V}}{z} \log_{10} Q$$

Entropy change: $\Delta S = \frac{q_{\text{rev}}}{T}$

Solubility Guidelines for Common Ionic Solids

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 ${\rm Li}^+$) and the ${\rm NH_4}^+$ 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 Ca²⁺, Sr²⁺, and Ba²⁺ are slightly soluble).
- 6. Sulfates are soluble except for those of calcium, strontium, and barium.

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PERIODIC TABLE	5				9	24	ပ်	51.996	42	ğ	95.94	74	≥	183.85	106	Unh	[263]
RICH	3				5	23	>	50.942	41	운	95.906	23	Ħ	180.95	105	Unp	[262]
O O	,				4	25	F	47.88	40	Ž	91.224	72	Ξ	178.49	104	Unq	[261]
					3	21	သွ	44.956	39	>	88.906	22	۳ پ	138.91	68	**AcUn	227.03
= 0	⁴ A	9.0122	12	Mg	24.305	20	င္မ	40.078	38	ഗ്	87.62	26	Ba	137.33	88	Ra	226.03
T 0.0.1	e	6.941	= '	Na	22.990	19	¥	39.098	37	윤	85.468	25	Cs	132.91	87	Ţ	[223]

	58	26	09	19	62	63	49	99	99	29	88	69	70	71
Lanthanides	ပ္ပ	4	ž	Pm	Sm	Eu	В	4	٥	웃	ш	Ę	Ϋ́	3
	140.12	140.91	144.24	[145]	150.36	151.97	157.25	158.93	162.50	164.93	167.26	168.93	173.04	174.97
						3								
	06	91	85	93	94	95	96	26	86	66	901	101	102	103
Actinides	두	Pa	-	Š	Pu	Am	Cm	쓢	ర	Es	ᇤ	ğ	ž	۲
	232.04	231.04	238.03	237.05	[244]	[243]	[247]	[247]	[251]	[252]	[257]	[258]	[259]	[262]