

AP[®] Chemistry

Practice Exam

The questions contained in this AP® Chemistry Practice Exam are written to the content specifications of AP Exams for this subject. Taking this practice exam should provide students with an idea of their general areas of strengths and weaknesses in preparing for the actual AP Exam. Because this AP Chemistry Practice Exam has never been administered as an operational AP Exam, statistical data are not available for calculating potential raw scores or conversions into AP grades.

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AP[®] Chemistry Directions for Administration

The AP Chemistry Exam is 3 hours and 5 minutes in length and consists of a multiple-choice section and a free-response section.

- The 90-minute multiple-choice section contains 75 questions and accounts for 50 percent of the final grade.
- The 95-minute free-response section contains 6 questions and accounts for 50 percent of the final grade. Part A is timed and is 55 minutes long; Part B is 40 minutes long.

A 10-minute break should be provided after Section I is completed. Students should be given a 10-minute warning prior to the end of each of the Parts A and B in Section II of the exam.

The actual AP Exam is administered in one session. Students will have the most realistic experience if a complete morning or afternoon is available to administer this practice exam. If a schedule does not permit one time period for the entire practice exam administration, it would be acceptable to administer Section I one day and Section II on a subsequent day.

Many students wonder whether or not to guess the answers to the multiple-choice questions about which they are not certain. It is improbable that mere guessing will improve a score. However, if a student has some knowledge of the question and is able to eliminate one or more answer choices as wrong, it may be to the student's advantage to answer such a question.

- The use of a calculator* is permitted ONLY during Section II, Part A of the exam. After time is called for Part A, students must place their calculators under their chairs. The use of any other electronic devices (including a cell phone) is not permitted during any portion of the exam.
- It is suggested that Section I of the practice exam be completed using a pencil to simulate an actual administration. Students can use either a pencil or a pen for Section II.
- Teachers will need to provide paper for the students to write their free-response answers. Teachers should provide directions to the students indicating how they wish the responses to be labeled so the teacher will be able to associate the student's response with the question the student intended to answer.
- A periodic table of the elements is provided with both Section I and Section II of the exam. For Section II, a table of standard reduction potentials and tables of commonly used equations and constants are also provided.
- Remember that students are not allowed to remove any materials, including scratch work, from the testing site.

*Calculators cannot have QWERTY keyboards or be designed to communicate with other calculators (such as via infrared ports).

Section I Multiple-Choice Questions

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	Be										B	C	Z	0	<u>F</u>	Ne
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+	12										13	14	15	16	17	18
Na	Mg										Al	Si	Ь	S	C	Ar
	4.30										26.98	28.09	30.97	32.06	35.45	39.95
		22	23	24	25	56	27	28	29	30	31	32	33	34	35	36
	Ca Sc	Ţ	>	Ç	Mn	Fe	ပိ	Z	Cn	Zn	Ga	Ge	As	Se	Br	Kr
			50.94	52.00	54.94	55.85	58.93	58.69	63.55	62.39	69.72	72.59	74.92	78.96	79.90	83.80
			41	42	43	44	45	46	47	48	49	50	51	52	53	54
	Sr Y		Sp	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sp	Te	Ι	Xe
			92.91	95.94	(86)	101.1	102.91	106.42	107.87	112.41	114.82	118.71	121.75	127.60	126.91	131.29
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		58	59	09	61	62	63	64	65	99	<i>L</i> 9	89	69	70	71	
*Lanthan	*Lanthanide Series	Ce	Pr	PN	Pm	Sm	Eu	В	$\mathbf{T}\mathbf{p}$	Dy	Ho	Er	Tm	$\mathbf{A}\mathbf{P}$	Lu	
		140.12	140.91	144.24	(145)	150.4	151.97	157.25	158.93	162.50	164.93	167.26	168.93	173.04	174.97	
		06	91	92	93	94	95	96	26	86	66	100	101	102	103	
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CHEMISTRY

Section I

Time—1 hour and 30 minutes NO CALCULATOR MAY BE USED WITH SECTION I.

Note: For all questions, assume that the temperature is 298 K, the pressure is 1.00 atmosphere, and solutions are aqueous unless otherwise specified.

Throughout the test the following symbols have the definitions specified unless otherwise noted.

T = temperatureL, mL = liter(s), milliliter(s)P = pressure= gram(s)V = volume= nanometer(s) nm S = entropy= atmosphere(s) atm H = enthalpymm Hg = millimeters of mercury G = Gibbs free energy J, kJ = joule(s), kilojoule(s) = volt(s)R = molar gas constantV n = number of moles= mole(s)mol M = molar

Part A

Directions: Each set of lettered choices below refers to the numbered statements immediately following it. Select the one that is best in each case and then place the letter of your choice in the corresponding box on the student answer sheet. A choice may be used once, more than once, or not at all in each set.

Questions 1-4 refer to the following chemical compounds.

m = molal

- (A) CH₄
- (B) CCl₃F
- (C) H_2S
- (D) H₂O₂
- (E) K_2CrO_4
- Commonly used as a disinfectant for minor skin wounds
- 2. A refrigerant implicated in the thinning of the stratospheric ozone layer
- 3. A major component of the fuel known as natural gas
- 4. A yellow solid at room temperature and 1 atm

Questions 5-7 refer to the following molecules.

- (A) CO
- (B) CH₄
- (C) HF
- (D) PH₃
- (E) F_2
- 5. Contains two π -bonds
- 6. Has the highest dipole moment
- 7. Has a molecular geometry that is trigonal pyramidal

Questions 8-11 refer to neutral atoms for which the atomic orbitals are represented below

$$(A) \boxed{1}$$

$$1 s$$

(B)
$$1 \downarrow 1$$
 $1 \downarrow 1$ $1 \downarrow 1$ $1 \downarrow 1$ $1 \downarrow 1$

(D)
$$\begin{array}{c|ccc}
\hline
1 & \hline
1 & \hline
2 & \hline
2 & \hline
2 & \hline
\end{array}$$

(E) [Ar]
$$\begin{array}{c|c} \hline 1 \\ \hline 4 \\ s \\ \hline \end{array}$$

$$\begin{array}{c|c} \hline 1 \\ \hline 3 \\ d \\ \hline \end{array}$$

- 8. Is in an excited state
- 9. Has exactly five valence electrons
- 10. Has the highest first ionization energy
- 11. Forms an aqueous cation that is colored

Questions 12-15 refer to the chemical reactions represented below.

(A)
$$HC_2H_3O_2(aq) + NH_3(aq) \rightarrow C_2H_3O_2^{-}(aq) + NH_4^{+}(aq)$$

(B)
$$\mathrm{Ba^{2+}}(aq) + \mathrm{SO_4^{2-}}(aq) \rightarrow \mathrm{BaSO_4}(s)$$

(C)
$$\operatorname{Zn}(\operatorname{OH})_2(s) + 2 \operatorname{OH}^-(aq) \rightarrow [\operatorname{Zn}(\operatorname{OH})_4]^{2-}(aq)$$

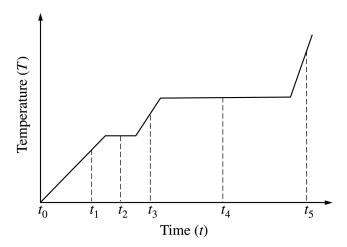
(D)
$$2 \text{ K}(s) + \text{Br}_2(l) \rightarrow 2 \text{ KBr}(s)$$

(E)
$$N_2O_4(g) \rightarrow 2 NO_2(g)$$

- 12. An oxidation-reduction reaction
- 13. A precipitation reaction
- 14. A reaction in which a coordination complex is formed
- 15. A Lewis acid-base reaction that is not a Brønsted-Lowry acid-base reaction

Questions 16-17 refer to various points in time during an experiment conducted at 1.0 atm. Heat

is added at a constant rate to a sample of a pure substance that is solid at time t_0 . The graph below shows the temperature of the sample as a function of time.



- (A) t_1
- (B) t_2
- (C) *t*₃
- (D) t_4
- (E) t_5
- 16. Time when the average distance between the particles is greatest
- 17. Time when the temperature of the substance is between its melting point and its boiling point

Directions: Each of the questions or incomplete statements below is followed by five suggested answers or completions. Select the one that is best in each case and place the letter of your choice in the corresponding box on the student answer sheet.

- 18. Which of the following is the correct name for the compound with formula Ca₃P₂?
 - (A) Tricalcium diphosphorus
 - (B) Calcium phosphite
 - (C) Calcium phosphate
 - (D) Calcium diphosphate
 - (E) Calcium phosphide
- 19. What mass of KBr (molar mass 119 g mol⁻¹) is required to make 250. mL of a 0.400 *M* KBr solution?
 - (A) 0.595 g
 - (B) 1.19 g
 - (C) 2.50 g
 - (D) 11.9 g
 - (E) 47.6 g
- 20. The value of the acid-dissociation constant, K_a , for a weak monoprotic acid HA is 2.5×10^{-6} . The pH of $0.40 \, M$ HA is closest to
 - (A) 2.0
 - (B) 3.0
 - (C) 4.0
 - (D) 6.0
 - (E) 8.0

- 21. Which of the systems in equilibrium represented below will exhibit a shift to the left (toward reactants) when the pressure on the system is increased by reducing the volume of the system? (Assume that temperature is constant.)
 - (A) $2 \text{ Mg}(s) + O_2(g) \rightleftharpoons 2 \text{ MgO}(s)$
 - (B) $SF_4(g) + F_2(g) \rightleftharpoons SF_6(g)$
 - (C) $H_2(g) + Br_2(g) \rightleftharpoons 2 HBr(g)$
 - (D) $N_2(g) + 3 H_2(g) \rightleftharpoons 2 NH_3(g)$
 - (E) $SO_2Cl_2(g) \rightleftharpoons SO_2(g) + Cl_2(g)$
- 22. The standard enthalpy of formation, ΔH_f° , of $\mathrm{HI}(g)$ is +26 kJ mol^{-1} . Which of the following is the approximate mass of $\mathrm{HI}(g)$ that must decompose into $\mathrm{H_2}(g)$ and $\mathrm{I_2}(s)$ to release 500. kJ of energy?
 - (A) 250 g
 - (B) 650 g
 - (C) 1,300 g
 - (D) 2,500 g
 - (E) 13,000 g

$$CaCl_2(s) \rightarrow Ca^{2+}(aq) + 2 Cl^{-}(aq)$$

- 23. For the process of solid calcium chloride dissolving in water, represented above, the entropy change might be expected to be positive. However, ΔS for the process is actually negative. Which of the following best helps to account for the net loss of entropy?
 - (A) Cl⁻ ions are much larger in size than Ca²⁺ ions.
 - (B) The particles in solid calcium chloride are more ordered than are particles in amorphous solids.
 - (C) Water molecules in the hydration shells of Ca²⁺ and Cl⁻ ions are more ordered than they are in the pure water.
 - (D) The $Ca^{2+}(aq)$ and $Cl^{-}(aq)$ ions are more free to move around in solution than they are in $CaCl_2(s)$.
 - (E) In the solution, the average distance between $Ca^{2+}(aq)$ and $Cl^{-}(aq)$ is greater than the average distance between Ca^{2+} and Cl^{-} in $CaCl_{2}(s)$.

$$\dots$$
 CH₃OCH₃(g) + \dots O₂(g) \rightarrow \dots CO₂(g) + \dots H₂O(g)

- 24. When the equation above is balanced using the lowest whole-number coefficients, the coefficient for $O_2(g)$ is
 - (A) 6
 - (B) 4
 - (C) 3
 - (D) 2
 - (E) 1

- 25. For which of the following processes does entropy decrease ($\Delta S < 0$) ?
 - (A) $H_2O(s) \rightarrow H_2O(l)$
 - (B) $Br_2(l) \rightarrow Br_2(g)$
 - (C) Crystallization of $I_2(s)$ from an ethanol solution
 - (D) Thermal expansion of a balloon filled with $CO_2(g)$
 - (E) Mixing of equal volumes of $H_2O(l)$ and $CH_3OH(l)$
- 26. In a laboratory, a student wants to quantitatively collect the CO₂ gas generated by adding Na₂CO₃(s) to 2.5 M HCl. The student sets up the apparatus to collect the CO₂ gas over water. The volume of collected gas is much less than the expected volume because CO₂ gas
 - (A) is very soluble in water
 - (B) is produced at a low pressure
 - (C) is more dense than water vapor
 - (D) has a larger molar mass than that of N_2 gas, the major component of air
 - (E) has a slower average molecular speed than water vapor at the same temperature
- 27. What mass of Cu(s) would be produced if 0.40 mol of $Cu_2O(s)$ was reduced completely with excess $H_2(g)$?
 - (A) 13 g
 - (B) 25 g
 - (C) 38 g
 - (D) 51 g (E) 100 g

- 28. Which of the following is a formula for an ether?

 - (C) H H H H H H C C C O H H H H H

Gas	Amount
Ar	0.35 mol
CH ₄	0.90 mol
N ₂	0.25 mol

- 29. Three gases in the amounts shown in the table above are added to a previously evacuated rigid tank. If the total pressure in the tank is 3.0 atm at 25°C, the partial pressure of $N_2(g)$ in the tank is closest to
 - (A) 0.75 atm
 - (B) 0.50 atm
 - (C) 0.33 atm
 - (D) 0.25 atm
 - (E) 0.17 atm

- 30. Which of the following best explains why the normal boiling point of $CCl_4(l)$ (350 K) is higher than the normal boiling point of $CF_4(l)$ (145 K)?
 - (A) The C-Cl bonds in CCl_4 are less polar than the C-F bonds in CF_4 .
 - (B) The C-Cl bonds in CCl₄ are weaker than the C-F bonds in CF₄.
 - (C) The mass of the CCl₄ molecule is greater than that of the CF₄ molecule.
 - (D) The electron cloud of the CCl₄ molecule is more polarizable than that of the CF₄ molecule.
 - (E) The bonds in the CCl₄ molecule are covalent, whereas the bonds in the CF₄ molecule are ionic.
- 31. At which of the following temperatures and pressures would a real gas be most likely to deviate from ideal behavior?

	Temperature	Pressure
	(K)	(atm)
(A)	100	50
(B)	200	5
(C)	300	0.01
(D)	500	0.01
(E)	500	1

- 32. After 195 days, a 10.0 g sample of pure ⁹⁵ Zr has decayed to the extent that only 1.25 g of the original ⁹⁵ Zr remains. The half-life of ⁹⁵ Zr is closest to
 - (A) 195 days
 - (B) 97.5 days
 - (C) 65.0 days
 - (D) 48.8 days
 - (E) 24.4 days

- 33. Which of the following would produce the LEAST mass of CO₂ if completely burned in excess oxygen gas?
 - (A) 10.0 g CH₄
 - (B) 10.0 g CH₃OH
 - (C) $10.0 \text{ g C}_2\text{H}_4$
 - (D) $10.0 \text{ g C}_2\text{H}_6$
 - (E) $10.0 \text{ g C}_4\text{H}_5\text{OH}$
- 34. Which of the following substances exhibits significant hydrogen bonding in the liquid state?
 - (A) CH_2F_2
 - (B) N_2H_4
 - (C) CH₃OCH₃
 - (D) C₂H₄
 - (E) C_2H_2
- 35. In an aqueous solution with a pH of 11.50 at 25° C, the molar concentration of $OH^{-}(aq)$ is approximately
 - (A) $3.2 \times 10^{-12} M$
 - (B) $3.2 \times 10^{-3} M$
 - (C) $2.5 \times 10^{-1} M$
 - (D) 2.5 M
 - (E) $3.2 \times 10^{11} M$
- 36. Which of the following changes to a reaction system in equilibrium would affect the value of the equilibrium constant, K_{eq} , for the reaction? (Assume in each case that all other conditions are held constant.)
 - (A) Adding more of the reactants to the system
 - (B) Adding a catalyst for the reaction to the system
 - (C) Increasing the temperature of the system
 - (D) Increasing the pressure on the system
 - (E) Removing some of the reaction products from the system

Questions 37-38 refer to a galvanic cell constructed using two half-cells and based on the two half-reactions represented below.

$$Zn^{2+}(aq) + 2e^{-} \rightarrow Zn(s)$$
 $E^{\circ} = -0.76 \text{ V}$
 $Fe^{3+}(aq) + e^{-} \rightarrow Fe^{2+}(aq)$ $E^{\circ} = 0.77 \text{ V}$

- 37. As the cell operates, ionic species that are found in the half-cell containing the cathode include which of the following?
 - I. $Zn^{2+}(aq)$
 - II. $Fe^{2+}(aq)$
 - III. $Fe^{3+}(aq)$
 - (A) I only
 - (B) II only
 - (C) III only
 - (D) I and III
 - (E) II and III
- 38. What is the standard cell potential for the galvanic cell?
 - (A) -0.01 V
 - (B) 0.01 V
 - (C) 0.78 V
 - (D) 1.53 V
 - (E) 2.31 V

Ionization Energies for Element X

	1st	2nd	3rd	4th	5th	6th	7th
Ionization Energy (kJ mol ⁻¹)	787	1,580	3,200	4,400	16,000	20,000	24,000

- 39. The first seven ionization energies of element X are shown in the table above. On the basis of these data, element X is most likely a member of which of the following groups (families) of elements?
 - (A) Alkaline earth metals
 - (B) Boron group
 - (C) Carbon group
 - (D) Nitrogen group
 - (E) Halogen group
- 40. Which of the following particles is emitted by an atom of ³⁹Ca when it decays to produce an atom of ³⁹K?
 - (A) $^{4}_{2}$ He
 - (B) ${}_{0}^{1}$ n
 - (C) ${}_{1}^{1}H$
 - (D) β⁻
 - (E) β⁺
- 41. At approximately what temperature will 40. g of argon gas at 2.0 atm occupy a volume of 22.4 L?
 - (A) 1,200 K
 - (B) 600 K
 - (C) 550 K
 - (D) 270 K
 - (E) 140 K
- 42. Which of the following aqueous solutions has the highest boiling point at 1.0 atm?
 - (A) 0.20 *M* CaCl₂
 - (B) $0.25 \, M \, \text{Na}_2 \text{SO}_4$
 - (C) 0.30 M NaCl
 - (D) 0.30 *M* KBr
 - (E) $0.40 M C_6 H_{12} O_6$

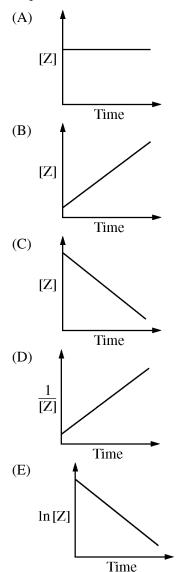
- 43. A certain reaction is spontaneous at temperatures below 400. K but is not spontaneous at temperatures above 400. K. If ΔH° for the reaction is -20. kJ mol⁻¹ and it is assumed that ΔH° and ΔS° do not change appreciably with temperature, then the value of ΔS° for the reaction is
 - (A) $-50. \text{ J mol}^{-1} \text{ K}^{-1}$
 - (B) $-20. \text{ J mol}^{-1} \text{ K}^{-1}$
 - (C) $-0.050 \text{ J mol}^{-1} \text{ K}^{-1}$
 - (D) $20. \text{ J mol}^{-1} \text{ K}^{-1}$
 - (E) 8,000 J mol⁻¹ K⁻¹
- 44. A sample of a solution of RbCl (molar mass 121 g mol⁻¹) contains 11.0 percent RbCl by mass. From the following information, what is needed to determine the molarity of RbCl in the solution?
 - I. Mass of the sample
 - II. Volume of the sample
 - III. Temperature of the sample
 - (A) I only
 - (B) II only
 - (C) I and II only
 - (D) II and III only
 - (E) I, II, and III

...
$$\operatorname{Au}^{3+}(aq) + ... \operatorname{I}^{-}(aq) \rightarrow ... \operatorname{Au}(s) + ... \operatorname{I}_{2}(s)$$

- 45. When the equation above is balanced using the lowest whole-number coefficients, the coefficient for $I_2(s)$ is
 - (A) 8
 - (B) 6
 - (C) 4
 - (D) 3
 - (E) 2
- 46. A closed rigid container contains distilled water and $N_2(g)$ at equilibrium. Actions that would increase the concentration of $N_2(g)$ in the water include which of the following?
 - I. Shaking the container vigorously
 - II. Raising the temperature of the water
 - III. Injecting more $N_2(g)$ into the container
 - (A) I only
 - (B) II only
 - (C) III only
 - (D) I and II only
 - (E) I, II, and III

$$Z \rightarrow X + Y$$

47. A pure substance Z decomposes into two products, X and Y, as shown by the equation above. Which of the following graphs of the concentration of Z versus time is consistent with the rate of the reaction being first order with respect to Z?



$$CS_2(l) + 3 O_2(g) \rightarrow CO_2(g) + 2 SO_2(g)$$

- 48. When 0.60 mol of $CS_2(l)$ reacts as completely as possible with 1.5 mol of $O_2(g)$ according to the equation above, the total number of moles of reaction products is
 - (A) 2.4 mol
 - (B) 2.1 mol
 - (C) 1.8 mol
 - (D) 1.5 mol
 - (E) 0.75 mol

Questions 49-50 refer to an experiment to determine the value of the heat of fusion of ice. A student used a calorimeter consisting of a polystyrene cup and a thermometer. The cup was weighed, then filled halfway with warm water, then weighed again. The temperature of the water was measured, and some ice cubes from a 0°C ice bath were added to the cup. The mixture was gently stirred as the ice melted, and the lowest temperature reached by the water in the cup was recorded. The cup and its contents were weighed again.

- 49. The purpose of weighing the cup and its contents again at the end of the experiment was to
 - (A) determine the mass of ice that was added
 - (B) determine the mass of the thermometer
 - (C) determine the mass of water that evaporated
 - (D) verify the mass of water that was cooled
 - (E) verify the mass of the calorimeter cup
- 50. Suppose that during the experiment, a significant amount of water from the ice bath adhered to the ice cubes. How does this affect the calculated value for the heat of fusion of ice?
 - (A) The calculated value is too large because less warm water had to be cooled.
 - (B) The calculated value is too large because more cold water had to be heated.
 - (C) The calculated value is too small because less ice was added than the student assumed.
 - (D) The calculated value is too small because the total mass of the calorimeter contents was too large.
 - (E) There is no effect on the calculated value because the water adhered to the ice cubes was at 0°C.

- 51. Which of the following molecules contains bonds that have a bond order of 1.5 ?
 - (A) N₂
 - (B) O_3
 - (C) NH₃
 - (D) CO₂
 - (E) CH₂CH₂
- 52. Of the following metals, which reacts violently with water at 298 K?
 - (A) Au
 - (B) Ag
 - (C) Cu
 - (D) Mg
 - (E) Rb
- 53. Heat energy is added slowly to a pure solid covalent compound at its melting point. About half of the solid melts to become a liquid. Which of the following <u>must</u> be true about this process?
 - (A) Covalent bonds are broken as the solid melts.
 - (B) The temperature of the solid/liquid mixture remains the same while heat is being added.
 - (C) The intermolecular forces present among molecules become zero as the solid melts.
 - (D) The volume of the compound increases as the solid melts to become a liquid.
 - (E) The average kinetic energy of the molecules becomes greater as the molecules leave the solid state and enter the liquid state.
- 54. A steady electric current is passed through molten MgCl₂ for exactly 1.00 hour, producing 243 g of Mg metal. If the same current is passed through molten AlCl₃ for 1.00 hour, the mass of Al metal produced is closest to
 - (A) 27.0 g
 - (B) 54.0 g
 - (C) 120. g
 - (D) 180. g
 - (E) 270. g

$$F^{-}(aq) + H_2O(l) \rightleftharpoons HF(aq) + OH^{-}(aq)$$

- 55. Which of the following species, if any, acts as a Brønsted-Lowry base in the reversible reaction represented above?
 - (A) HF(aq)
 - (B) $H_2O(l)$
 - (C) $F^-(aq)$ only
 - (D) Both $F^-(aq)$ and $OH^-(aq)$ act as Brønsted-Lowry bases.
 - (E) No species acts as a Brønsted-Lowry base.
- 56. What is the empirical formula of a hydrocarbon that is 10.0 percent hydrogen by mass?
 - (A) CH₃
 - (B) C_2H_5
 - (C) C_3H_4
 - (D) C_4H_9
 - (E) C_9H_{10}

$$Pb(s) \rightleftarrows Pb(l)$$

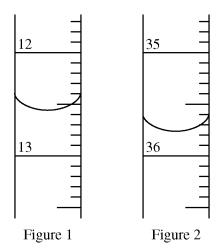
- 57. Which of the following is true for the process represented above at 327°C and 1 atm? (The normal melting point of Pb(s) is 327°C.)
 - (A) $\Delta H = 0$
 - (B) $T\Delta S = 0$
 - (C) $\Delta S < 0$
 - (D) $\Delta H = T\Delta G$
 - (E) $\Delta H = T\Delta S$

$$N_2(g) + O_2(g) + Cl_2(g) \rightleftharpoons 2 \text{ NOCl}(g)$$
 $\Delta H^{\circ} = +104 \text{ kJ mol}^{-1}$

$$\Delta H^{\circ} = +104 \text{ kJ mol}^{-1}$$

- 58. The equilibrium system represented above is contained in a sealed, rigid vessel. Which of the following will increase if the temperature of the mixture is raised?
 - (A) $[N_2(g)]$
 - (B) The rate of the forward reaction only
 - (C) The rate of the reverse reaction only
 - (D) The rates of both the forward and reverse reactions
 - (E) The total number of moles of gas in the vessel
- 59. If a metal X forms an ionic chloride with the formula XCl₃, then which of the following formulas is most likely to be that of a stable sulfide of X?
 - (A) XS₂
 - (B) X_2S_3
 - (C) XS₆
 - (D) $X(SO_3)_3$
 - (E) $X_2(SO_3)_3$

Questions 60-61 refer to the figures below. The figures show portions of a buret used in a titration of an acid solution of known concentration with a saturated solution of Ba(OH)₂. Figures 1 and 2 show the level of the Ba(OH)₂ solution at the start and at the endpoint of the titration, respectively. Phenolphthalein was used as the indicator for the titration.



- 60. What is the evidence that the endpoint of the titration has been reached?
 - (A) The color of the solution in the buret changes from pink to colorless.
 - (B) The color of the solution in the buret changes from blue to red.
 - (C) The color of the contents of the flask below the buret changes from colorless to pink.
 - (D) The color of the contents of the flask below the buret changes from blue to red.
 - (E) The contents of the flask below the buret change from clear to cloudy.
- 61. The volume of saturated $Ba(OH)_2$ used to neutralize the acid was closest to
 - (A) 6.60 mL
 - (B) 22.80 mL
 - (C) 23.02 mL
 - (D) 23.20 mL
 - (E) 29.80 mL

$C(diamond) \rightarrow C(graphite)$

- 62. For the reaction represented above, the standard Gibbs free energy change, $\Delta G_{298}^{\,\circ}$, has a value of $-2.90 \, \mathrm{kJ \ mol^{-1}}$. Which of the following best accounts for the observation that the reaction does NOT occur (i.e., diamond is stable) at 298 K and 1.00 atm?
 - (A) ΔS° for the reaction is positive.
 - (B) The activation energy, E_a , for the reaction is very large.
 - (C) The reaction is slightly exothermic ($\Delta H^{\circ} < 0$).
 - (D) Diamond has a density greater than that of graphite.
 - (E) Diamond has a heat capacity lower than that of graphite.

$$8 H_2(g) + S_8(s) \rightarrow 8 H_2S(g)$$

- 63. When 25.6 g of $S_8(s)$ (molar mass 256 g mol⁻¹) reacts completely with an excess of $H_2(g)$ according to the equation above, the volume of $H_2S(g)$, measured at 0°C and 1.00 atm, produced is closest to
 - (A) 30 L
 - (B) 20 L
 - (C) 10 L
 - (D) 5 L
 - (E) 2 L

$$2 \text{ HClO} + 3 \text{ O}_2 \rightarrow 2 \text{ HClO}_4$$

- 64. As the reaction represented above proceeds to the right, the oxidation number of chlorine changes from
 - (A) -1 to +3
 - (B) -1 to +5
 - (C) +1 to +5
 - (D) +1 to +7
 - (E) +3 to +7

- 65. By mixing only 0.15 *M* HCl and 0.25 *M* HCl, it is possible to create all of the following solutions EXCEPT
 - (A) 0.23 M HCl
 - (B) 0.21 M HCl
 - (C) 0.18 M HCl
 - (D) 0.16 M HCl
 - (E) 0.14 M HCl
- 66. At 25°C a saturated solution of a metal hydroxide, $M(OH)_2$, has a pH of 9.0. What is the value of the solubility-product constant, K_{sp} , of $M(OH)_2(s)$ at 25°C?
 - (A) 5.0×10^{-28}
 - (B) 1.0×10^{-27}
 - (C) 5.0×10^{-19}
 - (D) 5.0×10^{-16}
 - (E) 1.0×10^{-15}

- 67. A student weighs out 0.0154 mol of pure, dry NaCl in order to prepare a 0.154 *M* NaCl solution. Of the following pieces of laboratory equipment, which would be most essential for preparing the solution?
 - (A) Large crucible with lid
 - (B) 50 mL volumetric pipet
 - (C) 100 mL Erlenmeyer flask
 - (D) 100 mL graduated beaker
 - (E) 100 mL volumetric flask
- 68. In which of the following are the chemical species correctly ordered from smallest radius to largest radius?
 - (A) B < C < N
 - (B) Ar < Xe < Kr
 - (C) $Cl < S < S^{2-}$
 - (D) $Na < Na^+ < K$
 - (E) $K^+ < Ca^{2+} < K$
- 69. A large piece of wood can burn slowly, but wood in the form of sawdust can combust explosively. The primary reason for the difference is that compared with a large piece of wood, sawdust
 - (A) has a greater surface area per kilogram
 - (B) has a greater carbon content per kilogram
 - (C) absorbs more atmospheric moisture per kilogram
 - (D) contains more compounds that act as catalysts for combustion
 - (E) contains more compounds that have higher heats of combustion
- 70. Of the following elements, which would be expected to have chemical properties most similar to those of sulfur, S?
 - (A) Br
 - (B) Cl
 - (C) N
 - (D) P
 - (E) Se

- 71. When a solution is formed by adding some methanol, CH₃OH, to water, processes that are endothermic include which of the following?
 - I. Methanol molecules move water molecules apart as the methanol goes into solution.
 - II. Water molecules move methanol molecules apart as the methanol goes into solution.
 - III. Intermolecular attractions form between molecules of water and methanol as the methanol goes into solution.
 - (A) I only
 - (B) III only
 - (C) I and II only
 - (D) II and III only
 - (E) I, II, and III
- 72. Of the following gases, which has the greatest average molecular speed at 298 K?
 - (A) $Cl_2(g)$
 - (B) NO(*g*)
 - (C) $H_2S(g)$
 - (D) HCN(g)
 - (E) $PH_3(g)$
- 73. Types of hybridization exhibited by carbon atoms in a molecule of propyne, CH₃CCH, include which of the following?
 - I. sp
 - II. sp^2
 - III. sp^3
 - (A) I only
 - (B) III only
 - (C) I and III only
 - (D) II and III only
 - (E) I, II, and III

$$\mathrm{XY}_2(aq) \ \rightleftarrows \ \mathrm{X}^{2+}(aq) + 2 \ \mathrm{Y}^-(aq)$$

- 74. A soluble compound XY_2 dissociates in water according to the equation above. In a 0.050 m solution of the compound, the $XY_2(aq)$ species is 40.0 percent dissociated. In the solution, the number of moles of particles of solute per 1.0 kg of water is closest to
 - (A) 0.15
 - (B) 0.090
 - (C) 0.070
 - (D) 0.040
 - (E) 0.020

- 75. In which of the following processes are covalent bonds broken?
 - (A) Solid silver melts.
 - (B) Solid potassium chloride melts.
 - (C) Solid carbon (graphite) sublimes.
 - (D) Solid iodine sublimes.
 - (E) Glucose dissolves in water.

END OF SECTION I

IF YOU FINISH BEFORE TIME IS CALLED, YOU MAY CHECK YOUR WORK ON THIS SECTION.

DO NOT GO ON TO SECTION II UNTIL YOU ARE TOLD TO DO SO.

Section II Free-Response Questions

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_				工区	PEKIODIC IABLE OF THE ELEMENTS		IAB	LE	OF	III	(L)	LINIE L		^			2
Н																	He
1.008												s			2		4.00
3	4											5	9	7	8	6	10
Li	Be											B	C	Z	0	\	Ne
6.94	9.01											10.81	12.01	14.01	16.00	19.00	20.18
11	12										•	13	14	15	16	17	18
Na	Mg											Al	Si	Ь	S	C	Ar
22.99	24.30											26.98	28.09	30.97	32.06	35.45	39.95
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
X	Ca	Sc	Ti	>	Cr	Mn	Fe	Co	Z	Cu	Zn	Ga	Ge	As	Se	Br	Kr
39.10	40.08	44.96	47.90	50.94	52.00	54.94	55.85	58.93	58.69	63.55	65.39	69.72	72.59	74.92	78.96	79.90	83.80
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	\mathbf{Sr}	Y	\mathbf{Zr}	Sp	Mo	Tc	Ru	Rh	Pd	Ag	Cq	In	Sn	Sb	Te	Ι	Xe
85.47	87.62	88.91	91.22	92.91	95.94	(86)	101.1	102.91	106.42	107.87	112.41	114.82	118.71	121.75	127.60	126.91	131.29
55	99	57		73	74	75	92	77	78	62		81	82	83	84	85	98
Cs	Ba	*La	Ht	Ta	*	Re	Os	Ir	Pt	Au	Hg	I	Pb	Bi	P_0	At	Rn
132.91	137.33		178.49	_	183.85	186.21	190.2	192.2	195.08	196.97	200.59	204.38	207.2	208.98	(209)	(210)	(222)
87			104	105	106	107	108	109	110	1111							
Fr	Ra	$^{\dagger}\mathrm{Ac}$	Rf	Db	Sg	Bh	Hs	Mt	Ds	Rg							
(223)	226.02	227.03	(261)	(262)	(366)	(264)	(277)	(268)	(271)	(272)							

	58	59	09	61	62	63	64	65	99	29	89	69	70	71
thanide Series	Ce	Pr	Nd	Pm		Eu	Сd	$\mathbf{T}\mathbf{p}$	Dy	H_0	Er		Χb	Lu
	140.12	140.91	144.24	(145)		151.97	157.25	158.93	162.50	164.93	167.26		173.04	174.97
	06	91	92	93		95	96	26	86	66	100		102	103
ctinide Series	Th	Pa	n	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No	L
	232.04 2.	231.04	238.03	(237)		(243)	(247)	(247)	(251)	(252)	(257)		(259)	(262)

STANDARD REDUCTION POTENTIALS IN AQUEOUS SOLUTION AT 25°C

$\begin{array}{cccccccccccccccccccccccccccccccccccc$		Half-re	action		E°(V)
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	F_2	$(g) + 2e^{-}$	\rightarrow	2F ⁻	2.87
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Со	$^{3+} + e^-$	\rightarrow	Co^{2+}	1.82
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Au	$3^{3+} + 3e^{-}$	\rightarrow	Au(s)	1.50
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cl ₂	$(g) + 2e^{-}$	\rightarrow	2Cl ⁻	1.36
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	O_2	$(g) + 4H^+ + 4e^-$	\rightarrow	$2{\rm H}_2{\rm O}(l)$	1.23
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Br	$(2(l) + 2e^{-}$	\rightarrow	2Br ⁻	1.07
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2 H	$1g^{2+} + 2e^{-}$	\rightarrow	Hg_2^{2+}	0.92
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Нд	$e^{2+} + 2e^{-}$	\rightarrow	Hg(l)	0.85
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Ag	$e^{+} + e^{-}$	\rightarrow	Ag(s)	0.80
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Нg	$e_2^{2+} + 2e^-$	\rightarrow	$2 \operatorname{Hg}(l)$	0.79
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Fe ²	$^{3+} + e^{-}$	\rightarrow	Fe ²⁺	0.77
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	I ₂ ($(s)+2e^{-}$	\rightarrow	2I ⁻	0.53
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cu	$^{+} + e^{-}$	\rightarrow	Cu(s)	0.52
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cu	$^{2+} + 2e^-$	\rightarrow	Cu(s)	0.34
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cu	$^{2+} + e^{-}$	\rightarrow	Cu ⁺	0.15
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Sn	$^{4+} + 2e^{-}$	\rightarrow	Sn^{2+}	0.15
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	S(s	$(x) + 2H^+ + 2e^-$	\rightarrow	$H_2S(g)$	0.14
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	2 H	$I^+ + 2e^-$	\rightarrow	$H_2(g)$	0.00
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Pb	$^{2+} + 2e^{-}$	\rightarrow	Pb(s)	-0.13
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Sn	$^{2+} + 2e^{-}$	\rightarrow	Sn(s)	-0.14
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Ni	$^{2+} + 2e^{-}$	\rightarrow	Ni(s)	-0.25
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Со	$^{2+} + 2e^-$	\rightarrow	Co(s)	-0.28
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Cd	$2^{2+} + 2e^{-}$	\rightarrow	Cd(s)	-0.40
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cr	$^{3+} + e^{-}$	\rightarrow	Cr ²⁺	-0.41
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Fe	$^{2+} + 2e^{-}$	\rightarrow	Fe(s)	-0.44
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Cr	$^{3+} + 3e^{-}$	\rightarrow	Cr(s)	-0.74
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Zn	$^{2+} + 2e^-$	\rightarrow	Zn(s)	-0.76
$Al^{3+} + 3e^{-} \longrightarrow Al(s) \qquad -1.66$ $Be^{2+} + 2e^{-} \longrightarrow Be(s) \qquad -1.70$ $Mg^{2+} + 2e^{-} \longrightarrow Mg(s) \qquad -2.37$ $Na^{+} + e^{-} \longrightarrow Na(s) \qquad -2.71$ $Ca^{2+} + 2e^{-} \longrightarrow Ca(s) \qquad -2.87$ $Sr^{2+} + 2e^{-} \longrightarrow Sr(s) \qquad -2.89$ $Ba^{2+} + 2e^{-} \longrightarrow Ba(s) \qquad -2.90$ $Rb^{+} + e^{-} \longrightarrow Rb(s) \qquad -2.92$ $K^{+} + e^{-} \longrightarrow K(s) \qquad -2.92$ $Cs^{+} + e^{-} \longrightarrow Cs(s) \qquad -2.92$	2 H	$I_2O(l) + 2e^-$	\rightarrow	$H_2(g) + 2OH^-$	-0.83
$Be^{2^{+}} + 2e^{-} \longrightarrow Be(s) \qquad -1.70$ $Mg^{2^{+}} + 2e^{-} \longrightarrow Mg(s) \qquad -2.37$ $Na^{+} + e^{-} \longrightarrow Na(s) \qquad -2.71$ $Ca^{2^{+}} + 2e^{-} \longrightarrow Ca(s) \qquad -2.87$ $Sr^{2^{+}} + 2e^{-} \longrightarrow Sr(s) \qquad -2.89$ $Ba^{2^{+}} + 2e^{-} \longrightarrow Ba(s) \qquad -2.90$ $Rb^{+} + e^{-} \longrightarrow Rb(s) \qquad -2.92$ $K^{+} + e^{-} \longrightarrow K(s) \qquad -2.92$ $Cs^{+} + e^{-} \longrightarrow Cs(s) \qquad -2.92$	Mı	$n^{2+} + 2e^-$	\rightarrow	Mn(s)	-1.18
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Al	$^{3+} + 3e^{-}$	\rightarrow	Al(s)	-1.66
	Ве	$^{2+} + 2e^-$	\rightarrow	Be(s)	-1.70
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Mg	$g^{2+} + 2e^{-}$	\rightarrow	Mg(s)	-2.37
$Sr^{2+} + 2e^{-}$ \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	Na	$e^{+} + e^{-}$	\rightarrow	Na(s)	-2.71
$Sr^{2+} + 2e^{-}$ \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	Ca	$^{2+} + 2e^-$	\rightarrow	Ca(s)	-2.87
$Rb^{+} + e^{-} \longrightarrow Rb(s) \qquad -2.92$ $K^{+} + e^{-} \longrightarrow K(s) \qquad -2.92$ $Cs^{+} + e^{-} \longrightarrow Cs(s) \qquad -2.92$	Sr ²	$^{2+} + 2e^{-}$	\rightarrow		-2.89
$K^{+} + e^{-} \qquad \rightarrow \qquad K(s) \qquad -2.92$ $Cs^{+} + e^{-} \qquad \rightarrow \qquad Cs(s) \qquad -2.92$	Ва	$^{2+} + 2e^-$	\rightarrow	Ba(s)	-2.90
$Cs^+ + e^- \rightarrow Cs(s)$ -2.92	Rb	$^{+} + e^{-}$	\rightarrow	Rb(s)	-2.92
$Cs^+ + e^- \rightarrow Cs(s)$ -2.92	K ⁺	$+e^{-}$	\rightarrow	K(s)	-2.92
	Cs	$^{+} + e^{-}$	\rightarrow		-2.92
D = D = D = D = D = D = D = D = D = D =	Li⁴	$+e^{-}$	\rightarrow	Li(s)	-3.05

ADVANCED PLACEMENT CHEMISTRY EQUATIONS AND CONSTANTS

ATOMIC STRUCTURE

$$E = hv c = \lambda v$$

$$\lambda = \frac{h}{mv} p = mv$$

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

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 Δn = moles product gas – moles reactant gas

THERMOCHEMISTRY/KINETICS

$$\Delta S^{\circ} = \sum S^{\circ}$$
 products $-\sum S^{\circ}$ reactants
 $\Delta H^{\circ} = \sum \Delta H_{f}^{\circ}$ products $-\sum \Delta H_{f}^{\circ}$ reactants
 $\Delta G^{\circ} = \sum \Delta G_{f}^{\circ}$ products $-\sum \Delta G_{f}^{\circ}$ reactants

$$\Delta G^{\circ} = \Delta H^{\circ} - T \Delta S^{\circ}$$
$$= -RT \ln K = -2.303 RT \log K$$
$$= -n \mathcal{F} E^{\circ}$$

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

$$q = mc\Delta T$$

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

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

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

$$E = \text{energy}$$
 $v = \text{velocity}$
 $v = \text{frequency}$ $n = \text{principal quantum number}$
 $\lambda = \text{wavelength}$ $m = \text{mass}$
 $p = \text{momentum}$

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

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

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

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

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

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

Equilibrium Consta	nts
K_a (weak acid)	
K_b (weak base)	
K_w (water)	

 K_p (gas pressure) K_c (molar concentrations)

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

 H° = standard enthalpy

 G° = standard free energy

 E° = 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,
$$\mathcal{F} = 96,500$$
 coulombs per mole of electrons

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

= 0.0821 L atm mol⁻¹ K⁻¹
= 62.4 L torr mol⁻¹ K⁻¹
= 8.31 volt coulomb mol⁻¹ K⁻¹

GASES, LIQUIDS, AND SOLUTIONS

$$PV = nRT$$

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

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

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

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

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

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

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

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

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

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

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

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

$$\text{molarity, } M = \text{ moles solute per kilogram solvent}$$

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

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

$$\pi = iMRT$$

$$A = abc$$

OXIDATION-REDUCTION; ELECTROCHEMISTRY

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

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

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

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

P = pressureV = volumeT = temperaturen = number of moles D = densitym = massv = velocity u_{rms} = root-mean-square speed KE = kinetic energyr =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 = absorbancea = molar absorptivityb = path lengthc = concentrationQ = reaction quotientI = current (amperes)q = charge (coulombs)t = time (seconds)

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

 $= 0.0821 \,\mathrm{L}\,\mathrm{atm}\,\mathrm{mol}^{-1}\,\mathrm{K}^{-1}$
 $= 62.4 \,\mathrm{L}\,\mathrm{torr}\,\mathrm{mol}^{-1}\,\mathrm{K}^{-1}$
 $= 8.31 \,\mathrm{volt}\,\mathrm{coulomb}\,\mathrm{mol}^{-1}\,\mathrm{K}^{-1}$
Boltzmann's constant, $k = 1.38 \times 10^{-23} \,\mathrm{J}\,\mathrm{K}^{-1}$
 $K_f \,\mathrm{for}\,\mathrm{H}_2\mathrm{O} = 1.86 \,\mathrm{K}\,\mathrm{kg}\,\mathrm{mol}^{-1}$
 $K_b \,\mathrm{for}\,\mathrm{H}_2\mathrm{O} = 0.512 \,\mathrm{K}\,\mathrm{kg}\,\mathrm{mol}^{-1}$
 $1 \,\mathrm{atm} = 760 \,\mathrm{mm}\,\mathrm{Hg}$
 $= 760 \,\mathrm{torr}$
STP = $0.00 \,\mathrm{^{\circ}C}$ and $1.0 \,\mathrm{atm}$
Faraday's constant, $\mathcal{F} = 96,500 \,\mathrm{coulombs}\,\mathrm{per}\,\mathrm{mole}$

 E° = standard reduction potential K = equilibrium constant

of electrons

CHEMISTRY

Section II

(Total time—95 minutes)

Part A

Time—55 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.

1. Answer the following questions about the solubility of the salts Li₃PO₄ and PbCl₂. Assume that hydrolysis effects are negligible.

The equation for the dissolution of $Li_3PO_4(s)$ is shown below.

$$\text{Li}_{3}\text{PO}_{4}(s) \rightleftharpoons 3 \text{Li}^{+}(aq) + \text{PO}_{4}^{3-}(aq)$$
 $K_{sp} = 3.2 \times 10^{-9} \text{ at } 25^{\circ}\text{C}$

- (a) Write the equilibrium-constant expression for the dissolution of $\text{Li}_3\text{PO}_4(s)$.
- (b) Assuming that volume changes are negligible, calculate the maximum number of moles of $\text{Li}_3\text{PO}_4(s)$ that can dissolve in
 - (i) 0.50 L of water at 25°C
 - (ii) 0.50 L of 0.20 M LiNO₃ at 25°C

The equation for the dissolution of PbCl₂ is shown below.

$$PbCl_2(s) \rightleftharpoons Pb^{2+}(aq) + 2 Cl^{-}(aq)$$
 $K_{sp} = 1.6 \times 10^{-5} \text{ at } 25^{\circ}C$

- (c) Calculate the concentration of Cl⁻(aq) in a saturated solution of PbCl₂ at 25°C.
- (d) An open container holds 1.000 L of 0.00400 M PbCl₂, which is unsaturated at 25°C. Calculate the minimum volume of water, in mL, that must evaporate from the container before solid PbCl₂ can precipitate.

- 2. Answer the following using chemical concepts and principles of the behavior of gases.
 - (a) A metal cylinder with a volume of 5.25 L contains 3.22 g of He(g) and 11.56 g of $N_2(g)$ at 15.0°C.
 - (i) Calculate the total pressure in the cylinder.
 - (ii) Calculate the partial pressure of $N_2(g)$ in the cylinder.
 - (b) A 1.50 L container holds a 9.62 g sample of an unknown gaseous saturated hydrocarbon at 30°C and 3.62 atm.
 - (i) Calculate the density of the gas.
 - (ii) Calculate the molar mass of the gas.
 - (iii) Write the formula of the hydrocarbon.
 - (iv) Calculate the root-mean-square speed of the gas molecules in the container at 30° C. (Note: $1 \text{ J} = 1 \text{ kg m}^2 \text{ s}^{-2}$)
- 3. A student performs a titration in which a 10.00 mL sample of 0.0571 *M* HCl is titrated with a solution of NaOH of unknown concentration.
 - (a) Describe the steps that the student should take to prepare and fill the burst for the titration given a wet 50.00 mL burst and the materials listed below.

0.0571 *M* HCl solution Indicator solution
NaOH(*aq*) (unknown concentration) Distilled water
10.5 *M* NaOH solution 100 mL beaker

- (b) Calculate the pH of the 0.0571 M HCl.
- (c) A volume of 7.62 mL of the NaOH solution was needed to reach the end point of the titration. Calculate the molarity of the NaOH solution used in the titration.

In a different titration using a different NaOH solution, the concentration of NaOH was determined by the student to be 0.0614 *M*.

- (d) Given that the actual concentration of the NaOH solution was 0.0627 *M*, calculate the percent error of the student's result.
- (e) Calculate the volume of 10.5 M NaOH that is needed to prepare 250.0 mL of 0.0627 M NaOH.

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—40 minutes NO CALCULATORS MAY BE USED FOR PART B.

Answer Question 4 below. The Section II score weighting for this question is 10 percent.

EXAMPLE:

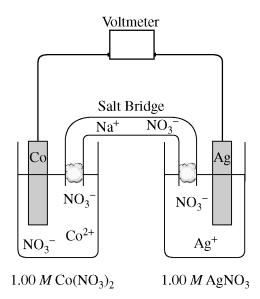
4. For each of the following three reactions, in part (i) write a balanced equation and in part (ii) answer the question about the reaction. In part (i), coefficients should be in terms of lowest whole numbers. Assume that solutions are aqueous unless otherwise indicated. Represent substances in solutions 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 graded.

Mg + 2 Ag ⁺ \longrightarrow Mg ²⁺ + 2 Ag
hich substance is oxidized in the reaction?
Mg is oxidized.
volumes of 0.1 <i>M</i> hydrofluoric acid and 0.1 <i>M</i> potassium hydroxide are combined.
Balanced equation:

(b)	Solic	d calcium metal burns in air.
	(i)	Balanced equation:
	(ii)	Predict the algebraic sign of ΔH° for the reaction. Explain your prediction.
(c)	Samj	ples of nitrogen monoxide gas and oxygen gas are combined.
	(i)	Balanced equation:
	(ii)	If the reaction is second order with respect to nitrogen monoxide and first order with respect to oxygen, what is the rate law for the reaction?

Answer Question 5 and Question 6. The Section II score weighting for these questions is 15 percent each.

Your responses to these questions 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.



- 5. Answer the following questions relating to the galvanic cell shown in the diagram above.
 - (a) Write the balanced equation for the overall cell reaction.
 - (b) Calculate the value of E° for the cell.
 - (c) Is the value of ΔG° for the overall cell reaction positive, negative, or 0 ? Justify your answer.
 - (d) Consider the cell as it is operating.
 - (i) Does E_{cell} increase, decrease, or remain the same? Explain.
 - (ii) Does ΔG of the overall cell reaction increase, decrease, or remain the same? Explain.
 - (iii) What would happen if the NaNO₃ solution in the salt bridge was replaced with distilled water? Explain.
 - (e) After a certain amount of time, the mass of the Ag electrode changes by x grams. Given that the molar mass of Ag is 108 g mol^{-1} and the molar mass of Co is 59 g mol^{-1} , write the expression for the change in the mass of the Co electrode in terms of x.

- 6. Answer each of the following using principles of atomic or molecular structure, and/or intermolecular or intramolecular forces.
 - (a) Explain why the H–O–H bond angle in H₂O is less than the H–N–H bond angles in NH₃, as shown in the table below.

H–O–H	H–N–H
Bond Angle in H ₂ O	Bond Angles in NH ₃
104.5°	107°

(b) Explain why the radius of the Br atom is less than the radius of the Br ion, as shown in the table below.

Radius of Br	Radius of Br ⁻
0.111 nm	0.196 nm

(c) Explain why the dipole moment of HI is less than the dipole moment of HCl, as shown in the table below.

Dipole Moment of HI	Dipole Moment of HCl
0.42 debye	1.08 debyes

(d) Explain why the normal boiling point of Ne is less than the normal boiling point of Kr, as shown in the table below.

Normal Boiling Point of Ne	Normal Boiling Point of Kr
27 K	121 K

STOP

END OF EXAM

IF YOU FINISH PART B OF SECTION II BEFORE TIME IS CALLED, YOU MAY RETURN TO PART A OF SECTION II IF YOU WISH, BUT YOU MAY NOT USE A CALCULATOR.

Name: _____

AP® Chemistry Student Answer Sheet for Multiple-Choice Section

No.	Answer
2	
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No.	Answer
31	
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No.	Answer
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AP® Chemistry Multiple-Choice Answer Key

3. T	Correct
No.	Answer
1	D
2	В
3	A
1 2 3 4 5 6 7	E A
5	Α
6	С
7	D
8	В
9	С
10	A
11	Е
12	D
13	В
14	С
15	С
16	Е
17	С
18	Е
19	D
20	В
21	Е
22	D
23	С
24	С
25	С
26	A
27	D
28	Е
28 29	В
30	D

	Correct
No.	Answer
31	A
32	A C
33	В
34	В
35	В
36	С
37	Е
38	D
39	С
40	Е
41	С
42	B A C
43	A
44	С
45	D
46	C
47	Е
48	D
49	A
50	С
51	В
52	Е
53	В
54	D
55	D
56	С
57	Е
58	D
56 57 58 59	В
60	С

	Correct
No.	Answer
61	D
62	В
63	В
64	D
65	Е
66	D
67	E
68	С
69	A
70	Е
71	С
72	D
73	С
74	В
75	С

General Scoring Principles

In regard to mathematical errors, a 1-point deduction is made; this deduction may be applied only once per question. In regard to errors in reporting significant figures, a 1-point deduction is made; this deduction may be applied only once per question. A leeway of plus or minus one significant figure different from the correct number of significant figures is allowed before a 1-point deduction is made.

For questions including parts that refer back to previous parts of the same question, a wrong answer that derives from the correct use of a previously calculated incorrect answer should not be counted as wrong. The emphasis of scoring is on *process*, and an error made early on in a multipart calculation should not jeopardize the earning of full credit for the correct use of that incorrect value later in the same question.

Question 1

Answer the following questions about the solubility of the salts Li₃PO₄ and PbCl₂. Assume that hydrolysis effects are negligible.

The equation for the dissolution of $Li_3PO_4(s)$ is shown below.

$$\text{Li}_{3}\text{PO}_{4}(s) \rightleftharpoons 3 \text{Li}^{+}(aq) + \text{PO}_{4}^{3-}(aq)$$
 $K_{sp} = 3.2 \times 10^{-9} \text{ at } 25^{\circ}\text{C}$

(a) Write the equilibrium-constant expression for the dissolution of $\text{Li}_3\text{PO}_4(s)$.

- (b) Assuming that volume changes are negligible, calculate the maximum number of moles of $\text{Li}_3\text{PO}_4(s)$ that can dissolve in
 - (i) 0.50 L of water at 25°C

Let x represent the molar concentration of Li_3PO_4 present in a saturated solution of Li_3PO_4 . Then $K_{sp}=3.2\times 10^{-9}=(3x)^3(x)=27x^4$, thus $x=\sqrt[4]{\frac{3.2\times 10^{-9}}{27}}=3.3\times 10^{-3}M$. Therefore the number of moles of $\text{Li}_3\text{PO}_4(s)$ that can dissolve in 0.50 L of water is

 $0.50 L \times \frac{3.3 \times 10^{-3} \text{ mol Li}_3 PO_4}{1.0 L} = 1.7 \times 10^{-3} \text{ mol}$

One point is earned for a correct setup using the K_{sp} expression from part (a).

One point is earned for a correct calculation of moles of Li₃PO₄.

(ii) 0.50 L of 0.20 M LiNO₃ at 25°C

In 0.20 M LiNO₃, [Li⁺] = 0.20 M. Assume that the amount of Li⁺(aq) contributed to the solution by the Li₃PO₄(s) that dissolves is negligibly small compared to 0.20 M. Then let y represent the molar concentration of PO₄³⁻(aq) present in the solution due to the Li₃PO₄(s) that dissolves in the 0.20 M LiNO₃. Then

$$K_{sp} = 3.2 \times 10^{-9} = (0.20)^3 (y) \implies y = \frac{3.2 \times 10^{-9}}{(0.20)^3} = 4.0 \times 10^{-7} M.$$

So
$$[PO_4^{3-}(aq)] = [Li_3PO_4(aq)] = 4.0 \times 10^{-7} M \text{ in } 0.20 M \text{ LiNO}_3(aq).$$

Therefore, the number of moles of $Li_3PO_4(s)$ that can dissolve in

0.50 L of water is
$$0.50 L \times \frac{4.0 \times 10^{-7} \text{ mol Li}_3 PO_4}{1.0 L} = 2.0 \times 10^{-7} \text{ mol}$$

One point is earned for recognizing that [Li⁺] is approximately equal to 0.20 M in the solution to which the $\text{Li}_3\text{PO}_4(s)$ was added.

One point is earned for a correct calculation of moles of Li_3PO_4 .

Question 1 (continued)

The equation for the dissolution of PbCl₂ is shown below.

$$PbCl_2(s) \rightleftharpoons Pb^{2+}(aq) + 2 Cl^{-}(aq)$$
 $K_{sp} = 1.6 \times 10^{-5} \text{ at } 25^{\circ}C$

(c) Calculate the concentration of Cl⁻(aq) in a saturated solution of PbCl₂ at 25°C.

Let $z = [Pb^{2+}]$ in a saturated solution of $PbCl_2$.

Then $K_{sp} = [Pb^{2+}][Cl^{-}]^{2} = (z)(2z)^{2} = 4z^{3}$.

Thus $1.6 \times 10^{-5} = 4z^3 \implies z = \sqrt[3]{\frac{1.6 \times 10^{-5}}{4}}$

 $\Rightarrow z = [Pb^{2+}] = 1.6 \times 10^{-2} M$

 \Rightarrow [Cl⁻] = 2[Pb²⁺] = 2(1.6 × 10⁻² M) = **3.2** × **10⁻²** M

One point is earned for a correct setup.

One point is earned for a correct calculation of the value of [Cl⁻].

(d) An open container holds 1.000 L of 0.00400 M PbCl₂, which is unsaturated at 25°C. Calculate the minimum volume of water, in mL, that must evaporate from the container before solid PbCl₂ can precipitate.

 $1.000 \text{ L of } 0.00400 \text{ M PbCl}_2(aq) \text{ contains } 0.00400 \text{ mol of PbCl}_2(aq),$ thus it contains $0.00400 \text{ mol Pb}^{2+}(aq)$ and $0.00800 \text{ mol Cl}^{-}(aq)$.

Let V = volume of the solution at saturation, then

 $K_{sp} = 1.6 \times 10^{-5} = \left(\frac{0.00400 \text{ mol}}{V}\right) \left(\frac{0.00800}{V}\right)^2 = \frac{2.6 \times 10^{-7}}{V^3}$

 $\Rightarrow V = \sqrt[3]{\frac{2.6 \times 10^{-7}}{1.6 \times 10^{-5}}} = 0.25 \text{ L} = 250 \text{ mL at saturation}$

Thus the volume of water that must evaporate = 1,000. - 250 = 750 mL

OR

From part (c), the amount of PbCl₂ dissolved in 1 L of saturated PbCl₂(aq)

is $1.6 \times 10^{-2} M$. Let V = volume of the solution at saturation, then

 $\frac{0.016 \text{ mol PbCl}_2}{1.000 \text{ mL}} = \frac{0.00400 \text{ mol PbCl}_2}{V} \implies V = 250 \text{ mL}$

Thus the volume of water that must evaporate = 1,000. - 250 = 750 mL

One point is earned for the calculation of the saturation volume.

One point is earned for subtracting the saturation volume from 1,000 mL.

Question 2

Answer the following using chemical concepts and principles of the behavior of gases.

- (a) A metal cylinder with a volume of 5.25 L contains 3.22 g of He(g) and 11.56 g of $N_2(g)$ at 15.0°C.
 - (i) Calculate the total pressure in the cylinder.

$$3.22 \text{ g He} \times \frac{1.00 \text{ mol He}}{4.00 \text{ g He}} = 0.805 \text{ mol He}$$

$$11.56 \text{ g N}_2 \times \frac{1.00 \text{ mol N}_2}{28.02 \text{ g N}_2} = 0.4126 \text{ mol N}_2$$

total moles of gas = (0.805 + 0.4126) = 1.218 mol

$$P = \frac{nRT}{V} = \frac{(1.218 \text{ mol})(0.0821 \text{ L atm mol}^{-1} \text{ K}^{-1})(15 + 273) \text{ K}}{5.25 \text{ L}}$$

P = 5.49 atm

One point is earned for the calculations of moles and adding them together.

One point is earned for the correct substitution into the gas law equation.

One point is earned for the correct answer.

(ii) Calculate the partial pressure of $N_2(g)$ in the cylinder.

pressure N₂ =
$$\frac{\text{moles N}_2 \text{ gas}}{\text{total moles of gas}} \times \text{total pressure in flask}$$

= $\frac{0.4126 \text{ mol}}{1.218 \text{ mol}} \times 5.49 \text{ atm} = 1.86 \text{ atm}$

One point is earned for the calculation of the mole fraction of $\,N_2\,$.

One point is earned for the correct answer.

- (b) A 1.50 L container holds a 9.62 g sample of an unknown gaseous saturated hydrocarbon at 30°C and 3.62 atm.
 - (i) Calculate the density of the gas.

$$D = \frac{m}{V} = \frac{9.62 \,\mathrm{g}}{1.50 \,\mathrm{L}} = 6.41 \,\mathrm{g} \,\mathrm{L}^{-1}$$

One point is earned for the correct answer.

Question 2 (continued)

(ii) Calculate the molar mass of the gas.

Let $M = \text{molar mass, then } PV = nRT = \left(\frac{m}{M}\right)RT \implies M = \frac{mRT}{PV} = \frac{(9.62 \text{ g})(0.0821 \text{ Latm K}^{-1}\text{mol}^{-1})(303 \text{ K})}{(3.62 \text{ atm})(1.50 \text{ L})} = 44.1 \text{ g mol}^{-1}$

One point is earned for the correct answer.

(iii) Write the formula of the hydrocarbon.

Saturated hydrocarbons have the generic formula C_nH_{2n+2} ,

therefore let 44.1 g = $12(n) + 1(2n + 2) = 14n + 2 \implies$

 $42.1 = 14n \implies 3 = n \implies \mathbf{C_3H_8}$

One point is earned for the correct answer.

(iv) Calculate the root-mean-square speed of the gas molecules in the container at 30°C. (Note: $1 \text{ J} = 1 \text{ kg m}^2 \text{ s}^{-2}$)

 $v_{\text{rms}} = \sqrt{\frac{3RT}{M}} = \sqrt{\frac{3(8.31 \text{ kg m}^2 \text{s}^{-2} \text{ mol}^{-1} \text{ K}^{-1})(303 \text{ K})}{0.0441 \text{ kg mol}^{-1}}}$

 $= 414 \text{ m s}^{-1}$

One point is earned for the correct setup using the molar mass in kilograms.

One point is earned for the correct answer.

Question 3

A student performs a titration in which a 10.00 mL sample of 0.0571 M HCl is titrated with a solution of NaOH of unknown concentration.

(a) Describe the steps that the student should take to prepare and fill the buret for the titration given a wet 50.00 mL buret and the materials listed below.

 $0.0571\,M$ HCl solution Indicator solution NaOH(aq) (unknown concentration) Distilled water $10.5\,M$ NaOH solution 100 mL beaker

Rinse the buret with some distilled water and then pour some of the NaOH solution of unknown concentration into the beaker and use it to rinse the buret two times.

Use the beaker to carefully fill the buret with the NaOH solution of unknown concentration.

Put the beaker under the buret and drain enough solution to remove any air bubbles in the neck and tip of the buret. One point is earned for rinsing the buret with the titrant solution.

One point is earned for removing air bubbles from the neck and tip of the buret.

(b) Calculate the pH of the $0.0571\,M$ HCl.

HCl is a strong acid \Rightarrow [H ⁺] in 0.0571 M HCl = 0.0571 M
$pH = -\log [H^+] = -\log (0.0571) = 1.243$

One point is earned for the correct answer.

(c) A volume of 7.62 mL of the NaOH solution was needed to reach the end point of the titration. Calculate the molarity of the NaOH solution used in the titration.

mol HCl titrated =
$$10.00 \text{ mL} \times \frac{0.0571 \text{ mol HCl}}{1,000 \text{ mL}} = 0.000571 \text{ mol};$$

ratio HCl:NaOH in neutralization is 1:1, so 0.000571 mol NaOH reacted;

$$\frac{0.000571 \text{ mol NaOH}}{0.00762 \text{ L}} = 0.0749 M \text{ NaOH}$$

One point is earned for calculating the moles of HCl in the sample that was titrated.

One point is earned for calculating the molarity of the NaOH solution.

Question 3 (continued)

In a different titration using a different NaOH solution, the concentration of NaOH was determined by the student to be $0.0614\,M$.

(d) Given that the actual concentration of the NaOH solution was 0.0627 *M*, calculate the percent error of the student's result.

percent error = $\frac{ 0.0614 - 0.0627 }{0.0627} \times 100 = 2.1 \%$	One point is earned for the correct answer.
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(e) Calculate the volume of 10.5 M NaOH that is needed to prepare 250.0 mL of 0.0627 M NaOH.

$250.0 \text{ mL} \times \frac{0.0627 \text{ mol}}{1,000. \text{ mL}} = 0.0157 \text{ mol NaOH needed}$	One point is earned for calculating the moles of NaOH needed.
$0.0157 \text{ mol NaOH} \times \frac{1,000. \text{ mL}}{10.5 \text{ mol NaOH}} = 1.49 \text{ mL}$	One point is earned for the correct answer.

Question 4

For each of the following three reactions, in part (i) write a balanced equation and in part (ii) answer the question about the reaction. 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 graded.

EXAMPLE:

A strip of magnesium metal is added to a solution of silver(I) nitrate.

(i) Balanced equation:

(ii) Which substance is oxidized in the reaction?

- (a) Equal volumes of 0.1 M hydrofluoric acid and 0.1 M potassium hydroxide are combined.
 - (i) Balanced equation:

$$HF + OH^- \rightarrow F^- + H_2O$$

(ii) Draw the complete Lewis electron-dot diagram for the reactant that is the Brønsted-Lowry base in the forward reaction.

- (b) Solid calcium metal burns in air.
 - (i) Balanced equation:

$$2 \text{ Ca} + \text{O}_2 \rightarrow 2 \text{ CaO}$$

(ii) Predict the algebraic sign of ΔH° for the reaction. Explain your prediction.

The sign of ΔH° will be negative because ΔG° is negative (the reaction occurs) and ΔS° is negative (a solid and a gas react to form a solid). According to the Gibbs-Helmholtz equation, $\Delta H^{\circ} = \Delta G^{\circ} + T\Delta S^{\circ}$. Therefore ΔH° is the sum of two negative quantities and as such must be negative.

Question 4 (continued)

- (c) Samples of nitrogen monoxide gas and oxygen gas are combined.
 - (i) Balanced equation:

$$2 \text{ NO} + \text{O}_2 \rightarrow 2 \text{ NO}_2$$

(ii) If the reaction is second order with respect to nitrogen monoxide and first order with respect to oxygen, what is the rate law for the reaction?

$$rate = k[NO]^2[O_2]$$

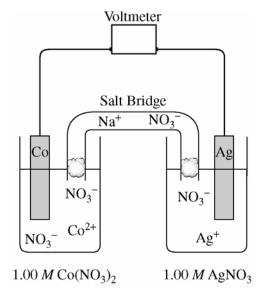
General Scoring Notes for Question 4

Five points are earned for each of parts (a), (b), and (c), distributed as follows.

Four points are earned for part (i): one point for the correct reactants, two points for the correct product(s), and one point for the correct coefficients in the balanced equation.

One point is earned for the correct answer in part (ii).

Question 5



Answer the following questions relating to the galvanic cell shown in the diagram above.

(a) Write the balanced equation for the overall cell reaction.

$2 \operatorname{Ag^{+}}(aq) + \operatorname{Co}(s) \rightarrow 2 \operatorname{Ag}(s) + \operatorname{Co^{2+}}(aq)$	One point is earned for the correct equation.
--	---

(b) Calculate the value of E° for the cell.

E° 0.00 (0.20) 1.00 V	One point is earned for the correct value of
$E_{cell}^{\circ} = 0.80 - (-0.28) = 1.08 \text{ V}$	E_{cell}° .

(c) Is the value of ΔG° for the overall cell reaction positive, negative, or 0 ? Justify your answer.

The value of ΔG° for the overall reaction must be negative because the cell reaction occurs (is spontaneous) as the cell operates.

OR

Since E_{cell}° is positive and $\Delta G^{\circ} = -nFE^{\circ}$, the value of ΔG° must be negative.

One point is earned for the correct answer, including a valid justification.

Question 5 (continued)

- (d) Consider the cell as it is operating.
 - (i) Does ${\cal E}_{cell}$ increase, decrease, or remain the same? Explain.

As the cell operates, the concentration of Ag^+ decreases and the concentration of Co^{2+} increases \Rightarrow the ratio $Q = \frac{[Co^{2+}]}{[Ag^+]^2}$ increases \Rightarrow $\ln Q$ increases \Rightarrow

 $E_{cell} = E_{cell}^{\circ} - \frac{RT}{nF} \ln Q$ becomes smaller (decreases).

One point is earned for the correct answer, including a valid justification.

(ii) Does ΔG of the overall cell reaction increase, decrease, or remain the same? Explain.

The value of ΔG for the system increases (becomes less negative) as the cell operates and the system approaches equilibrium (when $\Delta G = 0$).

One point is earned for the correct answer, including a valid justification.

(iii) What would happen if the NaNO₃ solution in the salt bridge was replaced with distilled water? Explain.

The cell would not operate. The voltage of the cell is too small to overcome the electrical resistance of distilled water, which is a poor conductor due to its very low concentration of ions (about $10^{-7} M \text{ H}^+(aq)$ and $10^{-7} M \text{ OH}^-(aq)$) that could "carry" the current from one cell to the other.

One point is earned for the correct answer, including a valid justification.

(e) After a certain amount of time, the mass of the Ag electrode changes by x grams. Given that the molar mass of Ag is 108 g mol^{-1} and the molar mass of Co is 59 g mol^{-1} , write the expression for the change in the mass of the Co electrode in terms of x.

-44-

 $\Delta \mod Ag = \Delta \max Ag \times \frac{1 \mod Ag}{108 \text{ g Ag}} = x \times \frac{1}{108} = \frac{x}{108}$

 $\Delta \text{ mol Co} = -\Delta \text{ mol Ag} \times \frac{1 \text{ mol Co}}{2 \text{ mol Ag}} = -\frac{x}{108} \times \frac{1}{2} = -\frac{x}{216}$

 Δ mass Co = Δ mol Co $\times \frac{59 \text{ g Co}}{1 \text{ mol Co}} = -\frac{x}{216} \times 59 = -\frac{59}{216} x$

One point is earned for using the correct mole ratio of Co to Ag.

One point is earned for the correct answer (negative sign is not required).

Question 6

Answer each of the following using principles of atomic or molecular structure and/or intermolecular or intramolecular forces.

(a) Explain why the H–O–H bond angle in H₂O is less than the H–N–H bond angles in NH₃, as shown in the table below.

H-O-H	H–N–H
Bond Angle in H ₂ O	Bond Angles in NH ₃
104.5°	107°

Both molecules have tetrahedral electron-domain geometries and might be expected to have bond angles of 109.5°. However, electron domains for nonbonding pairs of electrons exert a greater repulsion on adjacent pairs of electrons than do electron domains for bonding pairs. Thus, in the $\rm H_2O$ molecule with its $\rm \underline{two}$ nonbonding pairs of electrons, the electron domains of bonding pairs are compressed to a greater extent than they are in the $\rm NH_3$ molecule, which has only $\rm \underline{one}$ nonbonding pair of electrons.

One point is earned for citing the difference in number of nonbonding pairs of electrons.

One point is earned for citing the greater repulsion from nonbonding pairs as compared with bonding pairs.

(b) Explain why the radius of the Br atom is less than the radius of the Br ion, as shown in the table below.

Radius of Br	Radius of Br ⁻
0.111 nm	0.196 nm

The nuclear charge (+35) is the same for both the Br and Br⁻ species, but the "extra" electron in Br⁻ causes the electron cloud to expand due to an increase in mutual repulsions among the electrons that make up the cloud.

One point is earned for recognition that Br and Br⁻ have the same nuclear charge.

One point is earned for citing increased repulsion among electrons.

Question 6 (continued)

(c) Explain why the dipole moment of HI is less than the dipole moment of HCl, as shown in the table below.

Dipole Moment of HI	Dipole Moment of HCl
0.42 debye	1.08 debyes

Iodine, having a lower electronegativity than chlorine has, forms a bond with hydrogen that is less polar than the bond between chlorine and hydrogen in HCl. The lower polarity of the H–I bond means that the dipole moment of the bond is smaller than that of the H–Cl bond.

One point is earned for citing the difference in electronegativity between I and Cl.

One point is earned for the comparison of the polarity of the two bonds.

(d) Explain why the normal boiling point of Ne is less than the normal boiling point of Kr, as shown in the table below.

Normal Boiling Point of Ne	Normal Boiling Point of Kr
27 K	121 K

The intermolecular forces among atoms in liquid Ne are the same type of forces as those among atoms in liquid Kr, namely London (dispersion) forces. However, the magnitudes of these forces are smaller in Ne because the electron clouds of Ne atoms are smaller and less polarizable than the electron clouds of Kr atoms.

<u>Note</u>: An explanation that cites only periodic trends or only the relative masses of Ne and Kr does not earn credit.

One point is earned for mentioning that intermolecular forces involved are London (dispersion) forces.

One point is earned for mentioning the relative polarizability of the electron clouds of the atoms.