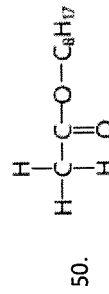
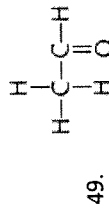
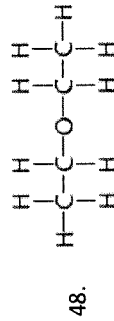
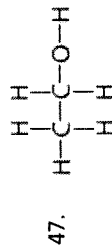
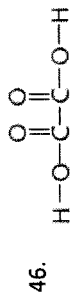


## Day 10: Continue

Questions 46 through 50 refer to types of organic compound below.

- (A) alcohol
- (B) aldehyde
- (C) carboxylic acid
- (D) ester
- (E) ether



Day 10

STOP. Correct your answers and note how many correct points

## Day 10: Answers and Explanations

### Answer: Quick Check

1. A 2. A 3. B 4. A 5. C 6. C 7. C 8. A 9. E 10. D  
 11. A 12. A 13. E 14. A 15. D 16. C 17. B 18. B 19. A 20. E  
 21. B 22. B 23. B 24. A 25. E 26. A 27. C 28. E 29. B 30. C  
 31. B 32. E 33. E 34. C 35. B 36. C 37. B 38. D 39. D 40. D  
 41. C 42. D 43. B 44. D 45. B 46. C 47. A 48. E 49. B 50. D

### Answers and Explanations

#### Questions 1 through 5: Le Chatelier's Principle, equilibrium reactions

1. A **Note:** N<sub>2</sub>O in the equation is a reactant.

**Note:** To reestablish equilibrium from adding more reactant, the added reactant must be consumed.

**Relate:** Speeding up in the forward direction (**shift to the right**) allow the reaction to consume the added reactant.

2. A **Note:** O<sub>2</sub> in the equation is a product

**Note:** To reestablish equilibrium from removal of a product, the reaction will make more of that product.

**Relate:** Speeding up in the forward direction (**shift to the right**) allows the reaction to produce more of the removed product.

3. B **Note:** A decrease in volume means pressure is increased on the reaction.

**Recall:** To reestablish equilibrium from a pressure increase, more of the substances on the side that has fewer total moles must be produced.

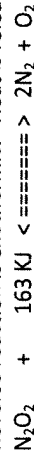
**Note:** In the equation, the reactant side has fewer total moles (2)

**Relate:** A decrease in volume speeds up the reverse reaction (**shift to the left**) in order to produce more N<sub>2</sub>O

4. A **Note:** The reaction has + ΔH value. This means :

Forward reaction is endothermic. Heat is absorbed as a reactant.

Reverse reaction is exothermic. Heat is released as a product



**Recall:** To reestablish equilibrium from increased temperature, endothermic (forward) reaction is favored (**Shift to the right**) so the added heat can be consumed.

5. C **Recall:** Addition of a catalyst lowers the activation energies for both the forward and reverse reactions. As a result, both the forward and reverse reactions are increased equally, and there is **no shift in either direction**.

## Day 10: Answers and Explanations

### Questions 6 through 8: Electron configuration, atomic structure

**6. C** *Recall:* The electron configuration of a paramagnetic element must have a sublevel with at least one unpaired e-

*Note:* The configuration  $1s^2 2s^2 2p^6 3s^2 3p^4$   
 $\uparrow\downarrow \uparrow\downarrow \uparrow\downarrow \uparrow\downarrow \uparrow\downarrow \uparrow\downarrow \uparrow\downarrow \uparrow$

has **two unpaired** electrons in the 3p sublevel.

**7. D** *Note:* Both potassium ion (K<sup>+</sup>) and chlorine ion (Cl<sup>-</sup>) have 18 e<sup>-</sup>. Both have the same configuration  $1s^2 2s^2 2p^6 3s^2 3p^6$ , which is the same as that of the noble gas element closest to them (Ar)

**8. A** *Note:* The atom of this element (Ne) has only 2 electron shells. All of the others have 3 or more shells.

### **9. E** Solution – Beer's Law relationship

*Recall:* According to Beer's law, absorbency is directly proportional to the concentration of a solution.

*Note:* To measure absorbance, a spectrometer is needed. An appropriate reagent must be a colored solution

*Determine:* Ni(NO<sub>3</sub>)<sub>2</sub>, which contains a transition metal, is the only colored solution listed.

### **10. D** Solubility Guideline, precipitate

*Note:* A precipitate is formed when an insoluble compound is formed from mixing two solutions.

*Note:* Mixing NaCl with Pb(NO<sub>3</sub>)<sub>2</sub> produces PbCl<sub>2</sub>(s).

*Recall solubility guidelines:* Chloride ion is mostly soluble, EXCEPT when it combines with **Pb<sup>2+</sup>**, **Ag<sup>+</sup>**, or **Hg<sup>2+</sup>**

## Day 10: Answers and Explanations

### 11. A Boiling point–concentration relationship, van't Hoff factor

*Recall:* The degree of boiling point elevation is related to the number of moles of ions in the solution. Solution with the greatest number of moles of ions will have the highest boiling point.

*Note:* Based on information given, the number of moles of ions in each solution can be calculated using the equation:

Moles of ion = Molarity x Volume x i (van't Hoff factor)

**(A)** Moles in KNO<sub>3</sub> = **0.50** x **.250** x **2 ions** = **.250 moles**

**(B)** Moles in Al(NO<sub>3</sub>)<sub>3</sub> = 0.10 x .400 x 4 ions = .160 moles

**(C)** Moles in NH<sub>4</sub>NO<sub>3</sub> = 0.20 x .500 x 2 ions = .200 moles

**(D)** Moles AgNO<sub>3</sub> = 0.30 x .300 x 2 ions = .180 moles

**(E)** Moles in Ni(NO<sub>3</sub>)<sub>2</sub> = 0.10 x .200 x 3 ions = .060 moles

*Note:* Choice A solution has the highest total moles of ions, therefore, highest boiling point

### 12. A Ion concentration in solution

*Note:* Highest NO<sub>3</sub><sup>-</sup> concentration can be calculated for each solution using the equation:

[NO<sub>3</sub><sup>-</sup>] = M x moles of NO<sub>3</sub><sup>-</sup> ions

**(A)** [NO<sub>3</sub><sup>-</sup>] in KNO<sub>3</sub> = **0.50** x **1** = **.50 M**

**(B)** [NO<sub>3</sub><sup>-</sup>] in Al(NO<sub>3</sub>)<sub>3</sub> = 0.10 x 3 = .30 M

**(C)** [NO<sub>3</sub><sup>-</sup>] in NH<sub>4</sub>NO<sub>3</sub> = 0.20 x 1 = .20 M

**(D)** [NO<sub>3</sub><sup>-</sup>] in AgNO<sub>3</sub> = 0.30 x 1 = .30 M

**(E)** [NO<sub>3</sub><sup>-</sup>] in Ni(NO<sub>3</sub>)<sub>2</sub> = 0.10 x 2 = .20 M

*Note:* Choice A has highest [NO<sub>3</sub><sup>-</sup>]

## Day 10: Answers and Explanations

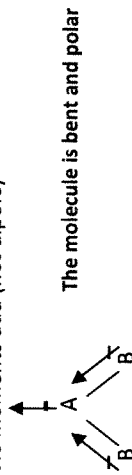
### 13. E Factors affecting molecular shape

**Note:** Shape of  $AB_2$  molecule can be determined if dipole moments of the molecule are known.

If dipole moments cancel out:



If dipole moments add (net dipole)



**Note:** Non of the other properties (alone) will be enough to determine the shape of the molecule

### 14. A Nuclear decay, emission particles

**Note:** The difference between  ${}^{19}_8\text{O}$  and  ${}^{19}_9\text{F}$  is  ${}^0_{-1}\text{e}$  (beta)

**Note:** Since this is a decay, the answer is beta emission, NOT, electron capture

### 15. D Interpreting potential energy diagram

**Note:** To determine the incorrect statement, consider each choice and eliminate those that are true of the energy diagram.

**Note:** **Eliminate Choice A.** True b/c product energy is higher than reactant's

**Eliminate Choice B.** True b/c Points 1 and 3 represent intermediates

**Eliminate Choice C.** True b/c the transitional point is measured at the activated complex

**Eliminate Choice E.** True b/c increasing energy at point 2 will result in a longer reaction pathway, hence, slower rate.

**Note:** **Choice D is false:** Equilibrium concentration of products depends on the initial concentration of reactants and final concentration of products only. Therefore, raising energy of intermediate 3 will not affect the equilibrium concentration of the product

## Day 10: Answers and Explanations

### 16. C Energy change and pressure calculation, thermodynamic

**Recall:** Recognize that based on information given, change in energy,  $\Delta E$ , can be calculated using the First Law of thermodynamics equation below.

$$\Delta E = q + w \quad \text{Note: } w = -(P_{\text{ext}} \times \Delta V)$$

Rewrite equation, substitute factors into equation, and solve

$$\Delta E = q - (P_{\text{ext}} \times \Delta V) \quad 101.32 \text{ J} \cdot \text{L}^{-1} \cdot \text{atm}^{-1}$$

$$\Delta E = 505.64 \text{ J} - (2.0 \text{ atm} \times 1 \text{ L}) \quad 101.32 \text{ J} \cdot \text{L}^{-1} \cdot \text{atm}^{-1}$$

$$\Delta E = 505.64 \text{ J} - 202.64 \text{ J} = \mathbf{303.00 \text{ J}}$$

### 17. C Interpreting reaction equation

**Note:** In the reaction,  $\text{NH}_4^+$  has 1 more  $\text{H}^+$  (proton) than  $\text{NH}_3$

**Recall:** A species in a reaction with one more  $\text{H}^+$  than another is the conjugate acid

**Note:** Non of the other choices is true or can be infer about the reaction based on just the equation given

### 18. B Molecular shapes-hybridization relationship

**Recall** the relationship between electron pairs, electron geometry and hybridization:

Electron pairs	electron geometry	hybridization
2	linear	$sp$
3	trigonal planer	$sp^2$
4	tetrahedral	$sp^3$
5	trigonal bipyramidal	$sp^3 d$
6	octahedral	$sp^3 d^2$

### 19. A Interpreting Gibb's free energy

**Recall:** Free energy  $\Delta G = \Delta H - T\Delta S$

**Note:** Since  $\Delta H$  is + and  $\Delta S$  is - as stated in the question,  $\Delta G = +\Delta H + T\Delta S = +\Delta G$  (nonspontaneous) regardless of temperature (T)

## Day 10: Answers and Explanations

### 20. E Common-ion effect, solubility

**Note:** Silver hydroxide (AgOH) is a base.

**Relate:** Its solubility will be hindered the most when placed in a solution with the highest OH<sup>-</sup> concentration (highest pH)

### 21. B Quantum numbers interpretation

**Note:** The set of numbers listed in the choices represents

$n$ ,  $l$ ,  $m_l$ , and  $m_s$ , respectively

**Recall** what each quantum number represents:

$n$  = shells       $n$  value range: 1 to 7  
 $l$  = subshells     $l$  value range: 0 to  $(n - 1)$   
 $m_l$  = orbitals     $m_l$  value range:  $-l$  to  $+l$   
 $m_s$  = spin         $m_s$  values:  $-\frac{1}{2}$  or  $+\frac{1}{2}$

**Note:** Based on information above,  $m_l$  value should never be greater than  $l$

**Note:** In Choice B, the  $m_l$  value (1) is greater than the  $l$  value (0). Therefore, B has the impossible set of quantum numbers.

### 22. B Percent yield calculation, equation writing

**Step 1:** Write a balanced equation for the reaction described.



**Step 2:** Determine theoretical yield of 1-fluoroethane from 96-g  $\text{CH}_4$ .

$$96 \text{ g CH}_4 \times \frac{1 \text{ mol CH}_4}{16 \text{ g CH}_4} \times \frac{2 \text{ mol CH}_3\text{CH}_2\text{F}}{4 \text{ mol CH}_4} \times \frac{48 \text{ g CH}_3\text{CH}_2\text{F}}{1 \text{ mol CH}_3\text{CH}_2\text{F}} = 144 \text{ g CH}_3\text{CH}_2\text{F}$$

**Step 3:** Calculate percent yield

$$\% \text{ yield} = \frac{\text{actual yield}}{\text{theoretical yield}} \times 100$$

$$\% \text{ yield} = \frac{96}{144} \times 100 = 66\%$$

## Day 10: Answers and Explanations

### 23. B Total pressure–mole fraction relationship

**Recognize** that based on information given, total pressure ( $P_{\text{total}}$ ) can be calculated using mole fraction equation

$$P_{\text{total}} = X_A P_A + X_B P_B \quad X = \text{mole fraction} \quad P = \text{partial pressure}$$

**Rewrite** equation, substitute factors from question, and solve.

$$P_{\text{total}} = \left( \frac{\text{mole of A}}{\text{total moles}} \right) P_A + \left( \frac{\text{mole of B}}{\text{total moles}} \right) P_B$$

$$P_{\text{total}} = \left( \frac{1.50}{7.0} \right) 75 \text{ mm Hg} + \left( \frac{5.50}{7.0} \right) 25 \text{ mm Hg}$$

$$P_{\text{total}} = 35.7 \text{ mm Hg}$$

### 24. A Solution properties

**Note:** When solute particles (in this question, potassium and chloride ions) are present in water, they interact and interfere with certain physical and chemical properties of water.

**Recall:** The present of solute in water allows the solution to have: **Higher boiling point, lower freezing point, and lower vapor pressure**

### 25. E Percent composition

**Step 1:** Write the correct formula for calcium hydroxide:  $\text{Ca}(\text{OH})_2$

**Step 2:** Determine molar mass of  $\text{Ca}(\text{OH})_2$

$$\text{Molar mass} = 1 \text{ Ca} + 2 \text{ O} + 2 \text{ H}$$

$$\text{Molar mass} = 1(40) + 2(16) + 2(1) = 74 \text{ g/mol Ca}(\text{OH})_2$$

**Step 3:** Determine mass of Oxygen in 148 g  $\text{Ca}(\text{OH})_2$

$$\text{Mass of O} = 148 \text{ g Ca}(\text{OH})_2 \times \frac{32 \text{ g O}}{74 \text{ g Ca}(\text{OH})_2} = 64 \text{ g O}$$

## Day 10: Answers and Explanations

### 26. A Equilibrium expression

$$\text{Recall: } K_{eq} = \frac{[\text{Products}]}{[\text{Reactants}]} = \frac{[\text{CO}_2]^2}{[\text{CO}]^2}$$

**Note:** Solids are not included in the equilibrium expression because they have constant concentrations

The exponents are the coefficients of the substances in the balanced equation.

### 27. C Understanding Titration process

**Note:** **Choice C** is the only correct statement. This statement will be true for all acid-base titration graphs.

**Note:** **Choice A** is false because the equivalent point of weak acid – strong base titration occurs at pH above 7.

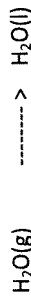
**Choice B** is false because pH does not depend on indicator

**Choice D** is false because a basic salt will be produced, resulting in a sharp increase in pH

**Choice E** is false because  $[H^+]$  is equal to the ionization constant,  $K_a$ , at the half-equivalence point; NOT at the equivalence point.

### 28. E Phase change – Energy – Entropy – Free energy relationship

**Note** the followings about the phase change of water given.



Gas to liquid = **Exothermic** (heat is released)      $\Delta H = -$

Gas to liquid = **Entropy decreases** (less disorder)      $\Delta S = -$

Gas to liquid = **Spontaneously** (occurs by itself at room temp)      $\Delta G = -$

## Day 10: Answers and Explanations

### 29. B Half-reaction interpretation

**Note** the followings about the half-reaction equation:



It represents reduction as written ( $e^-$  is a reactant).

Gold changes from +3 to 0 (a reduction of oxidation state)

**Recall:** Reduction occurs at cathode **Choice B**

**Note:** Chloride ion does not change its oxidation state. (Eliminate D & E)

### 30. C Coulombs law, factor-labeling

**Recognize** that based on information given, grams of gold produced can be calculated by setting up factor-labeling. When all units are crossed out, what's left is the correct setup.

$$\begin{array}{l} 4.0 \text{ C} \quad 30 \text{ min} \quad 60 \text{ s} \quad 1 \text{ mol } e^- \quad 1 \text{ mol Au} \quad 197 \text{ g Au} \\ \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \\ \text{s} \quad 1 \quad 1 \text{ min} \quad 96,500 \text{ C} \quad 3 \text{ mol } e^- \quad 1 \text{ mol Au} \\ \\ 4.0 \text{ C} \quad 30 \text{ min} \quad 60 \text{ s} \quad 1 \text{ mol } e^- \quad 1 \text{ mol Au} \quad 197 \text{ g Au} \\ \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \quad \times \text{-----} \\ \text{s} \quad 1 \quad 1 \text{ min} \quad 96,500 \text{ C} \quad 3 \text{ mol } e^- \quad 1 \text{ mol Au} \end{array}$$

### 31. C Atomic radius – nuclear charge relationship, ionic size comparison

**Recall:** The size (radius) of atoms depends on factors such as:

Number of electron shells

Number of electrons

Number of protons (nuclear charge)

**Note:** All of the ions listed are isoelectronic, meaning they have the same number of electrons (10), therefore, same number of electron shells.

**Relate:** The difference in size of these ions depends mostly on their nuclear charges. The greater the nuclear charge, the smaller the radius. Therefore:

Order of increasing size = Order of decreasing nuclear charge

Smallest radius :  $Mg^{2+}$  (+12 nuclear charge)

$F^-$  (+9 nuclear charge)

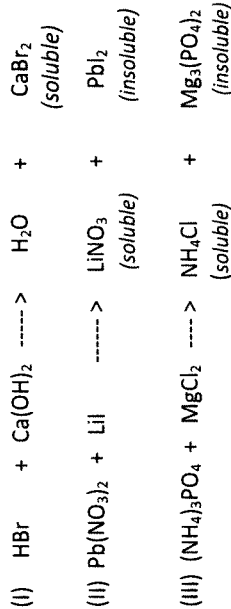
Largest radius :  $O^{2-}$  (+8 nuclear charge)

## Day 10: Answers and Explanations

### 32. E Completing equation, Solubility rules, precipitate

**Recall:** Precipitate is formed when an insoluble compound is formed from mixing two solutions.

Determine product for each reaction, then use solubility rules to determine which product is insoluble



**Note:** Both II and III produce insoluble (precipitate) products

### 33. E Le Chatelier's Principle

**Note:** All five reactions involve gaseous reactants and products.

**Recall:** Equilibrium concentration of substances in a gaseous reaction WILL NOT change when the total moles of reactants and products are equal.

**Note:** Reaction for Choice E is the only one listed with total moles of reactants and products being equal (3 moles on each side).

### 34. C Equation writing, mass – mole calculation in equation

**Step 1:** Write a correct balanced equation for the reaction described.



**Step 2:** Use factor-labeling to solve by utilizing mole ratio in equation

$$\text{mole of O}_2 = 25.5 \text{ g Al}_2\text{O}_3 \times \frac{1 \text{ mol Al}_2\text{O}_3}{102 \text{ g Al}_2\text{O}_3} \times \frac{3 \text{ mol O}_2}{2 \text{ mol Al}_2\text{O}_3}$$

$$\text{mole of O}_2 = \mathbf{0.375 \text{ mol}}$$

## Day 10: Answers and Explanations

### 35. B Oxidation of acids, acid formulas, acid strength

**Note:** When an oxyacid (acid with oxygen) is oxidized, the number of oxygen of the acid increases. The new acid formed is stronger.

**Note:** When an oxyacid has the maximum number of oxygen that it could have, that acid can't be oxidized any further.  
Ex: H<sub>2</sub>SO<sub>3</sub> can be oxidized to form H<sub>2</sub>SO<sub>4</sub>

But

H<sub>2</sub>SO<sub>4</sub> can't be oxidized any further.

**Note:** Of all the acids listed, only HNO<sub>2</sub> (a weaker acid) can be oxidized further (to HNO<sub>3</sub>, a stronger acid)

### -36. C Enthalpy change/Energy change in equations, thermodynamics

**Recognize** that based on information given, the equation

$$\Delta H = \Delta E + P\Delta V \text{ (law of thermodynamic)}$$

must be considered in order to correctly determine the reaction in which  $\Delta H$  and  $\Delta E$  are nearly equal.

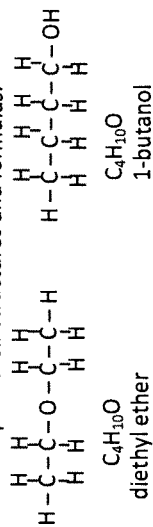
**Note:** Based on the equation,  $\Delta H$  and  $\Delta E$  will be equal only when  $P\Delta V$  is equal to Zero. This will be the case if  $\Delta V$  (change in volume) of the system is zero.

**Relate:** A reaction will have  $\Delta V$  equal to zero if there are no gaseous substances involve in the reaction.

**Note:** Choice C is the only reaction of those listed in which there no gaseous substances are involved.

### 37. B Comparing properties of organic compounds, isomers

**Recognize** that one way to choose the correct statement about the boiling points of these two substances is to correctly draw and compare their structures and formulas.



**Note:** Based on the structures and formulas, these two substances are isomers. (Eliminate Choice E)

**Note:** Based on the structures, diethyl ether is nonpolar (symmetrical) while 1-butanol is polar (asymmetrical).

**Recall:** Polar substances have higher boiling point than nonpolar.

## Day 10: Answers and Explanations

### 38. D Molar mass - rate relationship, comparing rate of gases

**Recall:** Rate of effusion of gases is related to their molar masses (M) according to the equation below:

$$\frac{V_1}{V_2} = \sqrt{\frac{M_2}{M_1}}$$

**Note:** The mass of  $H_2 = 2 \text{ g}$

**Relate:** A gas that travels at  $\frac{1}{3}$  the speed of  $H_2$  must have a molar mass that is 9 times as great as the molar mass of  $H_2$

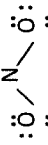
**Note:**  $H_2O$  has a molar mass of 18 g (9 times as great as 2 g)

### 39. C Bond angle-Molecular shape relationship

**Note** that drawing the Lewis diagram for the nitrate ion is one way of determining its shape, as well as the N - O bond angle.

Determine number of electrons available for bonding in  $NO_3^-$ , and then draw the Lewis structure for  $NO_3^-$

$$\begin{array}{l} N = 1(5) = 5 \text{ e-} \\ O = 3(6) = 18 \text{ e-} \\ -1 = 1 \text{ e-} \\ \hline \text{Total e-} = 24 \text{ e-} \end{array}$$



**Note:** The shape of  $NO_3^-$  is trigonal planer. Bond angles of this shape is always  $120^\circ$

### 40. D Titration data interpretation, sources of errors

**Note:** Only the NaOH volume for the first sample is, in terms of titration, significantly different from the rest.

**Note:** The only reasonable explanations is that the pipet that was used to transfer the acetic acid in the first sample was not rinsed with the acid.

**Note:** All other errors listed will not affect the volume of the titrant (NaOH)

## Day 10: Answers and Explanations

### 41. C Entropy Change calculation

**Step 1:** Recall entropy change,  $\Delta S^\circ$ , equation

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

**Step 2:** Substitute numbers from table into equation. Be sure to take number of moles of substances in the balanced equation into account.

$$3 \text{ Mg} + \text{Al}_2\text{O}_3 \quad 2 \text{ Al}(s) + 3 \text{ MgO}$$

$$\Delta S^\circ = [3(33.0) + 51.0] - [2(28.0) + 3(27.0)]$$

$$\Delta S^\circ = 150 - 137 = 13.0 \text{ J/(mole}\cdot\text{K)}$$

### 42. D Flame test

**Recall** your flame test colors: Lithium always red

### 43. B Percent abundance of isotopes, Atomic mass calculation

**Note:** Since the atomic mass (63.55) is closer to the mass of  $^{63}\text{Cu}$ , the percent of  $^{63}\text{Cu}$  will be greater than that of  $^{65}\text{Cu}$

*Eliminate choice C and D (these choices reflect greater % of  $^{65}\text{Cu}$ )*

**Note:** One way to quickly solve this problem is to set up the problem as follows:

Let  $x$  = decimal fraction of percent of  $^{63}\text{Cu}$   
then  $1 - x$  = decimal fraction of percent of  $^{65}\text{Cu}$

**Recall:** Atomic mass is the sum of the products of decimal fraction x mass number of the isotopes

$$63(x) + 65(1 - x) = 63.55$$

$$63x + 65 - 65x = 63.55$$

$$2x = 1.45$$

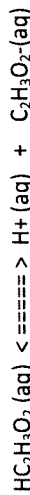
$$x = .73 \approx 70\% \text{ of } ^{63}\text{Cu}$$

## Day 10: Answers and Explanations

### 44. D Equilibrium expression calculation, dissociation

Recognize that  $K_a$  can be determined from equilibrium expression.

Step 1: Write equation for the dissociation of  $\text{HC}_2\text{H}_3\text{O}_2(\text{aq})$



Step 2: Write equilibrium expression for above equation

$$K_a = \frac{[\text{H}^+][\text{C}_2\text{H}_3\text{O}_2^-]}{[\text{HC}_2\text{H}_3\text{O}_2]}$$

Step 3: Determine concentrations:

$$[\text{H}^+] = -\log \text{pH} = -\log(2) = .01 \text{ M}$$

$$[\text{C}_2\text{H}_3\text{O}_2^-] = [\text{H}^+] = .01 \text{ M} \quad (\text{based on mole ratio of } 1\text{H}^+ : 1\text{C}_2\text{H}_3\text{O}_2^-)$$

$$[\text{HC}_2\text{H}_3\text{O}_2] = 1.0 \text{ M} \quad (\text{concentration of a weak acid like acetic stays nearly unchanged b/c weak acids dissociate very slightly})$$

Step 4: Substitute [ ] into equation and solve for  $K_a$

$$K_a = \frac{(0.01)(0.01)}{1.0} = .0001 = 1.0 \times 10^{-4}$$

### 45. B Percent calculation

$$\% \text{ dissociation} = \frac{[\text{Part}]}{[\text{Whole}]} \times 100$$

$$\% \text{ dissociation} = \frac{.01 \text{ M}}{1.0 \text{ M}} \times 100 = 1.0 \%$$

## Day 10: Answers and Explanations

### Questions 46 through 50: Organic functional groups

Recall the functional group associated with each class of organic compound listed as choices

class functional group

(A) alcohol -OH

(B) aldehyde  $\begin{array}{c} \text{O} \\ \parallel \\ \text{C}-\text{H} \end{array}$

(C) carboxylic acid  $\begin{array}{c} \text{O} \\ \parallel \\ \text{C}-\text{OH} \end{array}$

(D) ester  $\begin{array}{c} \text{O} \\ \parallel \\ -\text{C}-\text{O}- \end{array}$

(E) ether -O-

46. C Note: Structure given contains a *carboxylic acid* functional group

47. A Note: Structure given contains an *alcohol* functional group

48. E Note: Structure given contains an *ether* functional group

49. B Note: Structure given contains an *aldehyde* functional group

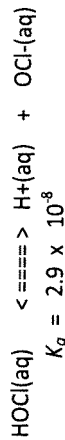
50. D Note: Structure given contains an *ester* functional group



## Day 11: Answers and Scoring Guidelines

(see important scoring guideline information on pg i)

1. Hypochlorous acid, HOCl, is a weak acid that ionizes in water, as shown in the equation below



(a) Calculate the  $[\text{H}^+]$  in a HOCl solution that has a pH of 5.24.

Recall:  $\text{pH} = -\log [\text{H}^+]$

$$[\text{H}^+] = 10^{-\text{pH}}$$

$$[\text{H}^+] = 10^{-5.24} = 5.75 \times 10^{-6}$$

1 point is earned for correctly calculating the  $[\text{H}^+]$

(b) Using information provided above:

(i) Write the equilibrium expression ( $K_a$ ) for the ionization of HOCl in water.

$$\text{Recall: } K_a = \frac{[\text{Products}]}{[\text{Reactants}]}$$

$$K_a = \frac{[\text{H}^+][\text{OCl}^-]}{[\text{HOCl}]}$$

1 point is earned for the correct equilibrium expression

(ii) Calculate the concentration of  $\text{OCl}^-\text{(aq)}$  in a HOCl solution that has  $[\text{H}^+]$  equal to  $2.4 \times 10^{-5} \text{ M}$ .

Note:  $[\text{OCl}^-] = [\text{H}^+]$  because they are at 1 : 1 ratio in the equation.

$$[\text{HOCl}] = \frac{[\text{H}^+][\text{OCl}^-]}{K_a}$$

$$[\text{HOCl}] = \frac{(2.4 \times 10^{-5} \text{ M})(2.4 \times 10^{-5})}{2.9 \times 10^{-8}}$$

$$[\text{HOCl}] = 2.0 \times 10^{-2} \text{ M}$$

1 point is earned for the correct setup to calculate  $[\text{HOCl}]$

1 point is earned for  $[\text{HOCl}]$  that is consistent with the setup

### Day 11: Answers and Scoring Guidelines

<p>(c) A solution of <math>\text{Ba}(\text{OH})_2</math> is titrated into a solution of <math>\text{HOCl}</math>.</p> <p>(i) Calculate the volume of <math>0.200 \text{ M Ba}(\text{OH})_2(\text{aq})</math> needed to reach the equivalence point when titrated into a <math>75.0 \text{ mL}</math> sample of <math>0.150 \text{ M HOCl}(\text{aq})</math>.</p> <p><i>Recall:</i> (moles of <math>\text{OH}^-</math>)(<math>M_b</math>)(<math>V_b</math>) = (moles of <math>\text{H}^+</math>)(<math>M_a</math>)(<math>V_a</math>)</p> <p><i>Note:</i> <math>\text{Ba}(\text{OH})_2</math> contains 2 moles of <math>\text{OH}^-</math>  <math>\text{HOCl}</math> contains 1 mole of <math>\text{H}^+</math></p> <p><i>Substitute values from question into equation and solve for volume of <math>\text{Ba}(\text{OH})_2</math> (<math>V_b</math>)</i></p> <p>(moles of <math>\text{OH}^-</math>)(<math>M_b</math>)(<math>V_b</math>) = (moles of <math>\text{H}^+</math>)(<math>M_a</math>)(<math>V_a</math>)</p> <p>(2)(<math>0.200 \text{ M}</math>)(<math>V_b</math>) = (1)(<math>0.150 \text{ M}</math>)(<math>0.075 \text{ L}</math>)</p> <p><math>V_b = 0.0281 \text{ L Ba}(\text{OH})_2</math></p>		<p><b>1 point</b> is earned for the correct volume</p>
<p>(ii) Write the equilibrium expression, <math>K_b</math>, for the titration reaction that occurs.</p> <p><i>Note:</i> <math>\text{OCl}^- + \text{H}_2\text{O} \rightleftharpoons \text{HOCl} + \text{OH}^-</math></p> <p><math>K_b = \frac{[\text{HOCl}][\text{OH}^-]}{[\text{OCl}^-]}</math></p>		<p><b>1 point</b> is earned for the correct expression</p>
<p>(iii) Calculate <math>k_b</math> of <math>\text{OCl}^-</math></p> <p><i>Recall:</i></p> <p><math>K_w = K_b \times k_a</math></p> <p><math>k_b = \frac{K_w}{K_a} = \frac{1.0 \times 10^{-14}}{2.9 \times 10^{-8}} = 3.4 \times 10^{-7}</math></p>		<p><b>1 point</b> is earned for the correct <math>k_b</math></p>

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<p>(iv) Calculate the pH at the equivalence point.</p> <p><i>Note:</i> Once <math>[\text{OH}^-]</math> is known, pH can be calculated.</p> <p><i>Note:</i> <math>[\text{OH}^-]</math> can be calculated by substituting values from the question and calculations (i – iii) into the <math>k_b</math> equation</p> <p><i>Calculate <math>[\text{OCl}^-]</math></i></p> <p><math>[\text{OCl}^-] = \frac{\text{moles of OCl}^-}{\text{Total volume}} = \frac{\text{Molarity} \times \text{Volume}}{(V_a + V_b)}</math></p> <p><math>[\text{OCl}^-] = \frac{(0.150 \text{ mol.L}^{-1})(0.075 \text{ L})}{(0.075 \text{ L} + 0.0281 \text{ L})} = 0.109 \text{ M}</math></p> <p><i>Determine concentrations</i></p> <p><math>[\text{OH}^-] = X</math></p> <p><math>[\text{HOCl}] = X</math> because they are at a 1 : 1 ratio</p> <p><i>Substitute [ ] into <math>k_b</math> expression and solve for X</i></p> <p><math>K_b = \frac{[\text{HOCl}][\text{OH}^-]}{[\text{OCl}^-]}</math></p> <p><math>3.4 \times 10^{-7} = \frac{X^2}{0.100}</math></p> <p><math>1.93 \times 10^{-4} = X = [\text{OH}^-]</math></p> <p><i>Calculate pOH</i></p> <p><math>\text{pOH} = -\log [\text{OH}^-] = -\log (1.93 \times 10^{-4}) = 3.72</math></p> <p><i>Calculate pH</i></p> <p><math>\text{pH} = 14 - \text{pOH} = 14 - 3.71 = 10.28</math></p>		<p><b>1 point</b> is earned for setup that leads to calculation of <math>[\text{OH}^-]</math> or <math>[\text{H}^+]</math></p>
<p><b>1 point</b> is earned for correctly calculating the pH</p>		

## Day 11: Answers and Scoring Guidelines

(d)  $\text{HClO}_3$  is a stronger acid than  $\text{HOCl}$ . Account for this fact in terms of molecular structure.

Note: In both  $\text{HOCl}$  and  $\text{HClO}_3$ ,  $\text{H}-\text{O}$  bond must be broken for the acids to ionize.

The weaker the  $\text{H}-\text{O}$  bond, the easier it is for the acid to ionize, and the stronger the acid.

$\text{HClO}_3$  is a stronger acid than  $\text{HOCl}$  because the  $\text{H}-\text{O}$  bond in  $\text{HClO}_3$  is weaker than the  $\text{H}-\text{O}$  bond in  $\text{HOCl}$ . This is due to the fact that the additional  $\text{O}$  atoms that are bonded to the central  $\text{Cl}$  atom in  $\text{HClO}_3$  weakens its  $\text{H}-\text{O}$  bond.

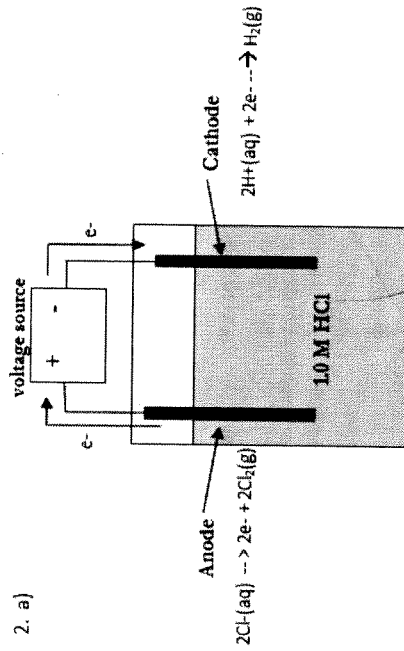
**1 point** is earned for the correct explanation.

## Day 11: Answers and Scoring Guidelines

2. Electrical current is passed through a  $1.0\text{M}$  solution of  $\text{HCl}(\text{aq})$  by means of two nonreactive electrodes immersed into the solution, with the electrodes connected to opposing terminals of a voltage source.

(8 points)

(a) (i) Sketch and label the diagram of the electrolytic cell, (ii) labels for anode and cathode, (iii) write half-reaction occurring at each electrode, and (iv) direction of electron flow.



Your diagram will vary greatly from the one above. Points are based on the followings:

(i) A correctly sketched diagram must have: **one cell (beaker)**, a **voltage source**, **two bars connected to the positive and negative ends of the voltage source and submerged in HCl solution.**

(ii) Metal bar connected to + is labeled **"Anode"** and Metal bar connected to - is labeled **"Cathode"**

(iii) Half-reaction at + electrode:  **$2\text{Cl}^{-}(\text{aq}) \rightarrow 2\text{e}^{-} + 2\text{Cl}_2(\text{g})$**  and Half-reaction at - electrode:  **$2\text{H}^{+}(\text{aq}) + 2\text{e}^{-} \rightarrow \text{H}_2(\text{g})$**

(iv) **Direction of  $\text{e}^{-}$**  is shown from bar at + to bar at -

**1 point** for a correct diagram

**1 point** for correctly labeled anode and cathode

**1 point** for correct and balanced half-reactions at + and - electrodes

**1 point** for correct direction of  $\text{e}^{-}$

### Day 11: Answers and Scoring Guidelines

(b) 4.5 amperes of current is applied to the reaction for 20 minutes.

(i) How many Coulombs pass through the cell during that time?

Note: 4.5 amp = 4.5 coulomb/sec (1 amp = 1 coulomb/sec)  
 20 minutes =  $1.2 \times 10^3$  sec

Note:  $I = \frac{q}{t}$  (see Reference Materials pg 340)

Charge (q) =	time (t)	x	current (I)	<b>1 point</b> is earned for the correct charge
Charge (q) =	$1.2 \times 10^3$ sec	x	4.5 coulombs sec	
Charge (q) =	<b><math>5.4 \times 10^3</math> coulombs</b>			

(ii) How many moles of electrons pass through the cell during that time?

Note: 1 mole of e<sup>-</sup> = 96500 coulombs (Faraday's constant)

moles of e <sup>-</sup> =	charge	x	Faraday's constant	<b>1 point</b> is earned for correct moles of e <sup>-</sup>
moles of e <sup>-</sup> =	$5.4 \times 10^3$ coulombs	x	$\frac{1 \text{ mole e}^-}{96500 \text{ coulombs}}$	
moles of e <sup>-</sup> =	<b><math>5.6 \times 10^{-2}</math> mol</b>			

iii) If the system is at STP, how much volume of hydrogen gas is produced during that time?

Recall: 1 mole of gas = 22.4 L at STP

Determine moles of H <sub>2</sub> using mole proportion in the half-reaction equation:	$2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2$	<b>1 point</b> is earned for calculating moles of H <sub>2</sub>
$5.6 \times 10^{-2}$ mol e <sup>-</sup> x	$\frac{1 \text{ mole H}_2}{2 \text{ moles e}^-} = 0.028 \text{ moles H}_2$	
Calculate volume of H <sub>2</sub>		<b>1 point</b> is earned for correct volume of H <sub>2</sub>
Volume of H <sub>2</sub> =	moles of H <sub>2</sub> x 22.4 L/mol	
Volume of H <sub>2</sub> =	<b><math>0.028 \text{ mol} \times 22.4 \text{ L/mol} = 0.627 \text{ L}</math></b>	

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(c) What happens to the pH of the HCl solution as current passes through it within the cell? Justify your response with explanation.

Recall: pH is related to H<sup>+</sup> concentration of a solution.

Note: As current passes through the solution, H<sup>+</sup> ion of the HCl solution is converted to H<sub>2</sub>. As a result, [H<sup>+</sup>] decreases.

<b>pH increases</b>	<b>1 point</b> is earned for indicating that pH increases
<b>pH increases because the H<sup>+</sup> concentration decreases as the electrolytic processes is taking place. A decrease in H<sup>+</sup> resulted in an increase in phi</b>	<b>1 point</b> is earned for explanation that is consistent with change in pH indicated.

## Day 12: Answers and Scoring Guidelines

(see important-scoring guideline information on pg 1)

1.	<p>(a) Dinitrogen oxide is mixed with water?</p> <p style="text-align: center;"><b>(15 points)</b></p> <p><i>Recall:</i> Nonmetal oxide combines with water to produce an acid.  <i>Note:</i> The acid (HNO<sub>3</sub>) formed is left in an ionized form because nitric acid is a strong acid. Strong acids ionize completely.</p>	<p>(i) Balanced equation</p> $\text{N}_2\text{O}_5 + \text{H}_2\text{O} \rightarrow 2\text{H}^+ + 2\text{NO}_3^-$ <p style="text-align: center;"><i>nonmetal oxide</i>      <i>water</i>      <i>acid</i></p> <p><b>1 point</b> is earned for correct reactants  <b>2 points</b> are earned for correct products  <b>1 point</b> is earned for correctly balancing the equation</p>
	<p>(ii) Would the pH of the solution that is produced less than 7, equal to 7, or greater than 7. Explain your answer.</p>	<p><b>1 point</b> is earned for the correct pH with explanation</p>
	<p>(b) Carbon dioxide gas is heated in the presence of solid magnesium oxide.</p> <p><i>Note:</i> This is a combination reaction in which a salt is formed from a basic and an acid anhydrides.</p>	<p>(i) Balanced equation</p> $\text{CO}_2 + \text{MgO} \rightarrow \text{MgCO}_3$ <p style="text-align: center;"><i>Basic anhydride</i>      <i>Acid anhydride</i>      <i>Salt</i></p> <p><b>1 point</b> is earned for correct reactants  <b>2 points</b> are earned for correct products  <b>1 point</b> is earned for correctly balancing the equation</p>

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(ii)	<p>How many grams of magnesium oxide must completely react with 11 grams of the carbon dioxide?</p>	<p><i>Determine moles of CO<sub>2</sub> in 11 g.</i></p> $11 \text{ g CO}_2 \times \frac{1 \text{ mole CO}_2}{44 \text{ g CO}_2} = 0.25 \text{ mol CO}_2$ <p><i>Determine moles of MgO based on proportion</i></p> $0.25 \text{ mol CO}_2 \times \frac{1 \text{ mol MgO}}{1 \text{ mol CO}_2} = 0.25 \text{ mol MgO}$ <p><i>Calculate grams of MgO</i></p> $0.25 \text{ mol MgO} \times \frac{40 \text{ g MgO}}{1 \text{ mol MgO}} = 10 \text{ g MgO}$	<p><b>1 point</b> is earned for correctly calculating the grams of magnesium oxide</p>
	<p>(c) Small pieces of aluminum are added to a solution of copper(II) sulfate.</p> <p><i>Note:</i> This is a single replacement redox reaction  <i>Note:</i> Sulfate ions is unchanged in the reaction. Therefore, it is not included in the equation.</p>	<p>(i) Balanced equation</p> $2 \text{ Al} + 3 \text{ Cu}^{2+} \rightarrow 2 \text{ Al}^{3+} + 3 \text{ Cu}$	<p><b>1 point</b> is earned for correct reactants  <b>2 points</b> are earned for correct products  <b>1 point</b> is earned for correctly balancing the equation</p>
	<p>(ii) Write the correct oxidation and reduction half equations for the reaction that occurs.</p>	<p><i>Oxidation-half:</i> <math>\text{Al} \rightarrow \text{Al}^{3+} + 3\text{e}^-</math>  <i>Reduction-half:</i> <math>\text{Cu}^{2+} + 2\text{e}^- \rightarrow \text{Cu}</math></p>	<p><b>1 point</b> is earned for the correct oxidation and reduction half equations</p>

## Day 12: Answers and Scoring Guidelines

<p>2. Use your knowledge of redox and electrochemistry to answer the following questions. <b>(8 points)</b></p> <p>(a) What is the function of a salt bridge in a galvanic cell, and why is it necessary in order for the cell work?</p> <p><i>Recall:</i> A salt bridge connects the two compartments of a galvanic cell</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; padding: 5px;"> <p>A salt bridge allows anions ( - ) and cations ( + ) to flow between the two compartments.</p> <p>A salt bridge is required for a galvanic cell to work because <b>the flow of the ions maintains cell neutrality</b> .</p> </td> <td style="width: 50%; padding: 5px;"> <p><b>1 point</b> is earned for correct function</p> <p><b>1 point</b> is earned for correct explanation.</p> </td> </tr> </table>	<p>A salt bridge allows anions ( - ) and cations ( + ) to flow between the two compartments.</p> <p>A salt bridge is required for a galvanic cell to work because <b>the flow of the ions maintains cell neutrality</b> .</p>	<p><b>1 point</b> is earned for correct function</p> <p><b>1 point</b> is earned for correct explanation.</p>	<p>(b) Explain why, when a piece of nickel is placed in hydrochloric acid, a reaction occurs, but when a piece of copper wire is placed in hydrochloric acid, no reaction occurs.</p> <p><i>Recall:</i> Spontaneous redox reaction occurs when potential (E) is +.</p> <p><b>A reaction occurs between the nickel (Ni) and HCl solution because the reaction has a positive potential (+E).</b></p> <p>According to the Table of Reduction Potentials:</p> $E_{\text{oxi}} \quad \text{Ni} \text{ ----} > \text{Ni}^{2+} + 2e^- = +0.25 \text{ V}$ $E_{\text{red}} \quad 2 \text{H}^+ + 2e^- \text{ ----} > \text{H}_2 = 0.00 \text{ V}$ <p>E for reaction = <b>+ 0.25 V</b></p> <p><b>A reaction did not occur between the copper wire (Cu) and the HCl solution because the reaction has a negative potential (-E).</b></p> $E_{\text{oxi}} \quad \text{Cu} \text{ ----} > \text{Cu}^{2+} + 2e^- = -0.34 \text{ V}$ $E_{\text{red}} \quad 2 \text{H}^+ + 2e^- \text{ ----} > \text{H}_2 = 0.00 \text{ V}$ <p>E for reaction = <b>- 0.34 V</b></p>
<p>A salt bridge allows anions ( - ) and cations ( + ) to flow between the two compartments.</p> <p>A salt bridge is required for a galvanic cell to work because <b>the flow of the ions maintains cell neutrality</b> .</p>	<p><b>1 point</b> is earned for correct function</p> <p><b>1 point</b> is earned for correct explanation.</p>		

## Day 12: Answers and Scoring Guidelines

<p>(c) <math>\text{Cu}^{2+} + \text{Pb}(s) \text{ ----} &gt; \text{Cu}(s) + \text{Pb}^{2+}</math></p> <p>(i) Give the standard cell potential for the reaction above.</p> <p><i>Recall:</i> <math>E_{\text{cell}} = E_{\text{oxi}} + E_{\text{red}}</math></p> <p><i>Note:</i> Use Table of Reduction Potential on pg 338 to get E for each half-reaction. Be sure to reverse the sign for oxidation half.</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; padding: 5px;"> <math display="block">E_{\text{oxi}} \quad \text{Pb} \text{ ----} &gt; \text{Pb}^{2+} + 2e^- = +0.13 \text{ V}</math> </td> <td style="width: 50%; padding: 5px;"> <p><b>1 point</b> is earned for the correct cell potential</p> </td> </tr> <tr> <td style="padding: 5px;"> <math display="block">E_{\text{red}} \quad \text{Cu}^{2+} + 2e^- \text{ ----} &gt; \text{Cu} = +0.34 \text{ V}</math> </td> <td></td> </tr> <tr> <td style="padding: 5px;"> <math display="block">E_{\text{cell}} \text{ for reaction} = \mathbf{+0.47 \text{ V}}</math> </td> <td></td> </tr> </table>	$E_{\text{oxi}} \quad \text{Pb} \text{ ----} > \text{Pb}^{2+} + 2e^- = +0.13 \text{ V}$	<p><b>1 point</b> is earned for the correct cell potential</p>	$E_{\text{red}} \quad \text{Cu}^{2+} + 2e^- \text{ ----} > \text{Cu} = +0.34 \text{ V}$		$E_{\text{cell}} \text{ for reaction} = \mathbf{+0.47 \text{ V}}$		<p>(ii) What happens to the cell potential in (i) when <math>[\text{Pb}^{2+}]</math> is increased? Explain.</p> <p><b>The <math>E_{\text{cell}}</math> will decrease</b> because <math>[\text{Pb}^{2+}]</math> is a product, and increasing [ ] of a product forces the reaction to shift left (favors the reverse reaction). This decreases the voltage that is produced by the cell.</p> <p><b>1 point</b> is earned for the correct change in <math>E_{\text{cell}}</math> with explanation</p>
$E_{\text{oxi}} \quad \text{Pb} \text{ ----} > \text{Pb}^{2+} + 2e^- = +0.13 \text{ V}$	<p><b>1 point</b> is earned for the correct cell potential</p>						
$E_{\text{red}} \quad \text{Cu}^{2+} + 2e^- \text{ ----} > \text{Cu} = +0.34 \text{ V}$							
$E_{\text{cell}} \text{ for reaction} = \mathbf{+0.47 \text{ V}}$							

<p>(iii) What is the value of <math>\Delta G^\circ</math> for the cell?</p> <p><i>Note:</i> <math>\Delta G^\circ = -nFE</math> (See Reference Materials on pg 339)</p> <p><i>Note:</i> n = moles of electrons = 2 (see equations in (i) )</p> <p style="text-align: center;">F = Faraday's constant = 96500 C/mol (see Reference materials)</p>	<p><math>\Delta G^\circ = -nFE</math></p> <p><math>\Delta G^\circ = -(2)(96500 \text{ C/mol})(+0.47 \text{ V})</math></p> <p><math>\Delta G^\circ = -90710 \text{ C}\cdot\text{V/mol}</math></p> <p style="text-align: center;"><b>- 90710 J/mol or 90.7 KJ/mol</b></p> <p><b>1 point</b> is earned for correct setup</p> <p><b>1 point</b> is earned for correctly calculating <math>\Delta G^\circ</math></p>
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