



CHEMISTRY

BOOKS - CENGAGE CHEMISTRY (HINGLISH)

ELECTROCHEMISTRY

Illustration

1. Consider the following cell reaction

$$Zn + 2Ag^{\oplus} \rightarrow Zn^{2+} + 2Ag.$$

Given that

$$E^{c} \cdot Zn^{2+}(aq)Zn(s) = -0.76V$$

- $E^{c} \cdot Ag \oplus (aq) Ag(s) = 0.80V$
- a. Calculate the standard EMF fo the cell.
- b. Which ion is more powerful oxidizing agent ?
- c. Which metal is more powerful reducing agent ?



3. Construct a cell using given electrodes at 298K and also calculate its standart *EMF*. Given : $E^{c-} \cdot Zn \mid Zn^{2+} = 0.76V$ $E^{c-} \cdot Cu^{2+} \mid Cu = 0.34V$

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4. If $E^{c} \cdot (Ag | Ag^{\oplus}) = -0.8V$ and $E^{c} \cdot (H_2 | 2H^{\oplus}) = 0V$, in a cell arrangement using these two electrodes, find $E^{c} \cdot cell$ and find out which electrode acts as anode and which acts as cathode.

5. Calculate E^{C^-} for each cell and write the equation for each cell precess. Explaing the significance of any negative. E^{c^-} value.

a.
$$Cd(s) |Cd^{2+}(1.0M)| |AgNO_{3}(1.0m)|Ag(s)$$

b. $Fe(s) |FeSO_{4}(1.0M)| |ZnSO_{4}(1.0m)|Zn(s)$
c. $Pt, Cl_{2}(g)(1atm)|NaCl(1.0M)|Hg_{2}Cl_{2}(s)| Hg(s)$
Given $E^{C^{-}} \cdot Cd = -0.40V, E^{C^{-}} \cdot (Fe) = -0.41V, E^{C^{-}} \cdot (Zn) = -0.76V$
 $E^{C^{-}} \cdot (Ag) = +0.80V, E^{C^{-}} \cdot (2Cl^{C^{-}}|Cl) = -1.36V.$
 $E^{C^{-}} \cdot (Hg|Hg_{2}Cl_{2}) = -0.27V.$

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6. For the reduction of $NO_3^{c^-}$ ion in an aqueous solution, E^{c^-} is +0.96V, the values of E^{c^-} for some metal ions are given below :

i.
$$V^{2^+}(aq) + 2e^- \rightarrow V$$
, $E^{c^-} = -1.19V$
ii. $Fe^{3^+}(aq) + 3e^- \rightarrow Fe$, $E^{c^-} = -0.04V$
iii. $Au^{3^+}(aq) + 3e^- \rightarrow Au$, $E^{c^-} = +140V$
iv. $Hg^{2^+}(aq) + 2e^- \rightarrow Hg$, $E^{c^-} = +0.86V$

The pair(s) of metals that is / are oxidized by NO_3^{c-} in aqueous solution is

/ are



7. The following electrochemical cell is represented as :

$$Pt(1) | Hg^{2+}, Hg_2^{2+}(a=1) | | Ce^{4+}, Ce^{3+}(a=1) | Pt(2)$$

a. If an ammeter is connected between two platinum electrodes, predict the direction of flow of current.

b. Will the current decrease or increase with time?

Given :
$$E^{c} \cdot {}_{2Ha^{2^+} | Ha_2^{2^+}} = 0.92V, E^{c} \cdot {}_{Ce^{4^+}/Ce^{3^+}} = 1.61V$$

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8. The following reaction: occurs in the dry cell, called flash light battery, which is used to power radios, clocks, and flashlights : Φ

$$2MnO_2(s) + 8NH_4 + Zn(s) \rightarrow 2Mn^{3+} + 8NH_3 + An^{2+} + 4H_2O$$

a. Write the anode and cathode reactions.

b. Calculate $E^{c} \cdot_{cell}$ of the dry cell if the electrode potential of cathode $\left(E^{c} \cdot_{reduction}\right)$ varies between 0.49V and 0.74V and that of anode $E^{c} \cdot_{reduction}$ is -0.76V.

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9. Represent the cell in which following reaction takes place :

 $Mg(s) + 2Ag^{\oplus}(0.0001M) \rightarrow Mg^{2+}(0.130M) + 2Ag(s)$ calculate its E_{cell} if $E^{c-} \cdot_{cell} = 3.17V.$

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10. Calculate the reduction potential of the following electrodes :

- a. $Pt, H_2(4atm) \mid H_2SO_4(0.01M)$
- b. $Pt, H_2(1atm) \mid HCl(0.2M)$
- *c*. Calculate the potential of hydrogen electrode in contact with a solution whose

ipH = 5 *ii.pOH* = 4



11. Calculate the equilibrium constant of the reaction :

 $Cu(s) + 2Ag(aq) \Leftrightarrow Cu^{2+}(aq) + 2Ag(s)$

 E^{c} . cell = 0.46V

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12. The standard electrode potential for Deniell cell is 1.1V. Calculate the

standart Gibbs energy for the reaction.

 $Zn(s) + Cu^{2+}(aq) \Leftrightarrow Zn^{2+}(aq) + Cu(s)$

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13. For the cell :

$$Zn(s)\left|Zn^{2+}(aM)\right|\left|Ag^{\oplus}(bM)\right|Ag(s).$$

a. Write Nernst equation to show how $E_{\mbox{\it cell}}$ vary with concentration of

Zn^{2+}	and	Ag^{\oplus}	ions.	Given
		0		

$$E^{c^{-}} \cdot (Zn^{2^{+}} | Zn) = 0.76V, E^{-} \cdot (Ag^{\oplus} | Ag) = 0.80V.$$

b. Find E_{cell} for $[Zn^{2^{+}}] = 0.01M$ and $[Ag^{c^{-}}] = 0.02M.$

c. For what values of Q will the cell EMF be

i. 0.0V ii. 0.97V

14. Calculate the potential of the following half - cells | cells:
a.
$$Cr |Cr^{3+}(0.1M)||Fe^{2+}(0.01M)|Fe$$

Given : $E^{c-} \cdot Cr^{3+} |Cr| = -0.74VE^{c-} \cdot Fe^{2+} |Fe| = -0.44V$
b. $6e^{c-} + BrO_3^{c-}(aq) + 3H_2O \rightarrow Br^{c-}(aq) + 6OH(aq)$
Given : $E^{c-} \cdot (BrO_3^{c-} |Br^{c-}) = 0.61V$,
 $[BrO_3^{c-}] = 2.5 \times 10^{-3}M, [Br^{c-}] = 5.0 \times 10^{-3}M, pH = 9.0$
 $c. Ag |Ag^{\oplus}(0.1M)||Cl^{c-}(0.02M)|Cl_2(g)(0.5atm)| Pt$
Given $E (Ag^{\oplus} |Ag) = 0.80V, E^{c-} ((Cl_2 | 2Cl^{c-}) = 1.36V)$
 $d. NO_3^{c-}(aq) + 2H^{\oplus}(aq) + e^{-} \rightarrow NO_2 + H_2O$
Given : $E^{c-} \cdot NO_3^{c-} |NO_2 = 0.78V$

What will be the reductino potential of the half cell in neutral solution ? Assuming all the other species to be at unit concentration.

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15. The solution of $CuSO_4$ in which Cu ord is dipped is diluted to 10 times,

the reduction electrode potential will :

- *a*. Decrease by 0.03V *b*. Decrease by 0.059V
- *c*. Increase by 0.03V *d*. Increase by -.059V

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$$TI \left| TI \stackrel{\oplus}{=} (10^{-3}M) \right| \left| Cu^{2+} (10^{-1}M) \right| Cu$$

 E_{cell} can be increased by

16. For the cell:

- a. Decreasing $\left[Cu^{2^+}\right]$. b. Decreasing $\left[TI^{\oplus}\right]$
- c. Increasing $\begin{bmatrix} Cu^{2+} \end{bmatrix}$ d. Increasingby[TI^(o+)]`

17. A zinc electrode is placed in 0.1M solution of $ZnSO_4$ at $25 \degree C$. Assuming salt is dissociated to the extent of 20 % at this dilution. The potential of this electrode at this temperature is :

$$\left(E_{Zn^{2+}\mid Zn}^{\circ}=-0.76V\right)$$

 $a.\,0.79V$. b. - 0.79V c. - 0.81V. $d.\,0.81V$

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18. If excess of Zn is added to 1.0M solution of $CuSO_4$, find the concentration of Cu^{2+} ions at equilibrium.

Given:
$$E^{c^{-}} \cdot (Zn^{2^{+}} | Zn) = -0.76V$$

 $E^{c^{-}} \cdot Cu^{2^{+}} = (E^{c^{-}} \cdot (Cu^{2^{+}} | Cu)) = 0.34V$

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19. Find the reduction potential of $AsO_4^{3-} | AsO_2^{c-}$ in a solution when 18mL of 0.1N solution of *NaI* is added to 20mL of $0.1NNa_3AsO_4$ solution

at pH = 5. The standard reduction potential of $AsO_4^{3-} | AsO_2^{c-} = -0.70V$.



20. Calculate the potential of the following cell :

$$Pt(s) \begin{vmatrix} Ce^{3+} & (2M) \\ Ce^{4+} & (1M) \end{vmatrix} \begin{vmatrix} Cr^{3+}(2M) \\ Cr_2O_7^{2-}(1M) & H^{\oplus}(1M) \end{vmatrix} Pt(s)$$

Given : $E^{c^-} \cdot Ce^{3^+} | Ce^{4^+} = -1.7, E^{c^-} \cdot Cr_2O_7^{2^-} | Cr^{3^+} = 1.3V$

(Take0.059 ≈ 0.06)

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21. Explain the construction and working of Westron standard cell.



22. The *EMF* of Westron standard cell is 1.0153 at 20 $^{\circ}C$ and 1.01807 at

25 ° C. Calculate ΔG , ΔH , and ΔS for the cell reaction at 25 ° C.

23. The *EMF* of the cell :

 $Cd \mid CdCl_2($ solution $)(1atm)\mid AgCl(s)\mid Ag$ is 0.675 at 25 °C. The temperature coefficient of the cell is -6.5×10^{-4} degree⁻¹. Find the change in heat content and entropy for the electrochemical reaction that occurs when 1F of electricity is drawn for it.

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24. At 25 ° *C* , the free energy of formation of $H_2O(l)$ is -56, 700*Calmol*⁻¹. The free energy of ionization of water to H^{\oplus} and OH is 19050*calmol*⁻¹. What is the reversible *EMF* of the following cell at 25 ° *C*: $H_2(g)(1atm) | H^{\oplus} | | OH, O_2(g), 1atm$

25. Find K_c for the complex:

$$\begin{bmatrix} Ag(NH_3)_2 \end{bmatrix}^{\oplus} \Leftrightarrow Ag^{\oplus} + 2NH_3$$
$$E^{C^-} \cdot \begin{pmatrix} Ag^{\oplus}/Ag \end{pmatrix} = 0.8V \text{ and } E^{C^-} \cdot \begin{bmatrix} Ag(NH_3)_2 \end{bmatrix}^{\oplus} |Ag|NH_3 = 0.37V$$

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26. The standard free energy of formation of AgCl(s) at $25 \degree C$ is $-109.7kJmol^{-1}$ and $\left[H^{\oplus} + Cl^{c^{-}}\right](aq)$ is $-131.2kJmol^{-1}$. Find $E^{c^{-}}$ of a cell up cells, with standard hydrogen electrode, and $Cl^{c^{-}}|Ag|AgCl(s)$.

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27. The *EMF* of the following cellis 1.05V at $25 \degree C$:

Pt, *H*₂(*g*)(1.0*atm*)|*NaOH*(0.1*m*), *NaCl*(0.1*M*)|*AgCl*(*s*), *Ag*(*s*)

a. Write the cell reaction,

b. Calculate pK_w of water.

28. If the oxidation of oxalic acid by acidic MnO_4^{c-} solution is carried out in a reversible cell, then what is the electrode reaction and equilibirum constant of the cell reaction.

Given :

$$E^{C^{-}} \cdot \left(MnO_{4}^{C^{-}} \mid Mn^{2^{+}} \right) = 1.51V$$
$$E^{C^{-}} \cdot \left(CO_{2} \mid C_{2}O_{4}^{2^{-}} \right) = 0.49V$$

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29. Find the standard electrode potential of $MnO_4^{c-} | MnO_2$. The standard electrode potential of $MnO_4^{c-} | Mn^{2+} = 1.51V$ and $MnO_2 | MnO_2 | Mn^{2+} = 1.23V$.

30. Find the *EMF* of the concentration cell represented as given below:





31. Calculate the *EMF* of the following concentration cells a $30 \degree C$ and predict whether the cells are exergonic or endergonic. [Assume Kw does not chage at 30 $^{\circ}C$ a. $Pt \left| H_2(g)(1atm) \right| H^{\oplus} \left(10^{-6}M \right) \right| \left| H^{\oplus} \left(10^{-4}M \right) \right| H_2(g)(1atm) \right| Pt$ b. $Pt || Hg_2(1atm) |NaOH(10^{-4}M)|| H^{\oplus}(10^{-5}M) |H_2(g), (1atm)| Pt$ c. $Pt|H_2(g)(1atm)|H_2SO_4(0.05M)||KOH(10^{-3}M)|H_2(g)(1atm)|Pt$ $d. Pt \left| H_2(g)(1atm) \right| CH_2COOH\left(10^{-2}M\right) \right| \left| CsOH\left(10^{-3}M\right) \right| H_2(g)(1atm) \left| Pt\left(pK_a of Charge (10^{-3}M) \right) \right| CH_2COOH\left(10^{-2}M\right) \right| CSOH\left(10^{-3}M\right) \left| H_2(g)(1atm) \right| CH_2COOH\left(10^{-2}M\right) \right| CSOH\left(10^{-3}M\right) \left| H_2(g)(1atm) \right| CH_2COOH\left(10^{-2}M\right) \left| H_2(g)(1atm) \right| CH_2COOH\left(10^{-2}M\right) \right| CSOH\left(10^{-3}M\right) \left| H_2(g)(1atm) \right| CH_2COOH\left(10^{-2}M\right) | CH_2COOH$



32. Calculate the *EMF* of the following concentration cells at 30 $^{\circ}C$ and

predict whether the cells are exergonic or endergonic.

a. Pt $H_2(g)$	CH ₃ COONa	NH	4NO3(0	.2 M)	$H_2(g)$	Pt
(1 atm)	$(10^{-2}M)$	p <i>K</i>	, of NH ₄	ОН	(1 atm)	
	pK _a of	= 4	.74			
	CH ₃ COOH					
	= 4.74					
b. Pt $H_2(g)$	CH ₃ COONH	[4	CH ₃ CC)ONa	H ₂ (g)	Pt
(<u>1</u> _atm)	pK _a CH ₃ COC)H =	$(10^{-3}M)$)	(1 atm)	
	pK _b of NH ₄ O	H				
No.	= 4.74					
c. Pt $H_2(g)$	CH ₃ COOH	NH ₄	OH(10 ⁻²	² M)	$H_2(g),$	Pt
(1 atm)	(0.1 M)	р <i>К</i> _b =	= 4.74		(1 atm)	
	$\mathbf{p}K_{\mathbf{a}} = 4.74$					
d. Ag (s) $ Ag^{\oplus}(0.1 \text{ M}) Ag^{\oplus}(1 \text{ M}) Ag (s)$						
e. Pt $ Cl_2(g) $	$Cl^{\oplus}(10^{-3}M)$	Cl®	$(10^{-2} \mathrm{M})$		(g) Pt	
(1 atm)				(1 a	tm)	
f. Pt $ Cl_2(g) C$	$l^{\oplus}(10^{-3}M) \parallel 0$	Cl [⊜] (1	$(0^{-2} M)$	Cl ₂ (g) Pt	
2 atm				(1 at	m)	
					•	

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33. The *EMF* of the following cell is 0.180V at 30 ° *C*.

Find EMF cell when

a. 40mLof0.2MNaOH is added to the negative terminal of the battery .

b. 50mL of 0.2MNaOH is added to the negative terminal of the battery .

c. 50mL OF 0.2MNaOH is added to 100mL of H_2SO_4 at the positive terminal of the battery.

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34. A hydrogen electrode placed in a solution containing CH_3COOK and CH_3COOH in the ration a:b and b:a has electrode potential values of -1.59 and +1.0V, respectively. Calculate pK_a of CH_3COOH .

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35. Two buffer solutions A and B each made with benzoic acid and sodium benzoate differ in their pH by two units. A has salt: acida: b. B has salt: acid = b:a. If a > b, then the value of a:b is a. 3.17 b.10.0 c.3.0 d. 6.0 **36.** Calculate the charge on mercurous ion and its magnetic moment. *EMF* of the cell given below is 0.0295V at $25 \degree C$.

	0.01Mmercurous	0.1Mmercurous	
Pt, Hg(l)	nitratesolution	nitratesolutionin	Hg(l), Pt
	in0.1MHNO ₃	0.1 <i>MHNO</i> 3	5.77

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37. The quinhydrone electrode $(Q, H^{\oplus} | H^2Q)$ is used in conjunction with a saturated calomel electrode, as represented below:

 $EMF_{cell} = 0.264V$ at 30 ° C. Calculate the pH of unknown solution at this temperature.

Given :
$$E_{calomel} = 0.24V$$
 and $E^{c} \cdot {}_{2H}^{\oplus}, Q \mid H_2Q = 0.7V$
Hg(l), Hg₂Cl₂(s) $\begin{vmatrix} KCl \text{ (saturated } \\ solution \end{pmatrix} \mid \begin{vmatrix} H^{\oplus} \\ (Unknown) \end{vmatrix} Q, H_2Q, Pt$
Calculate the pH of unknown

Glass electrodeBuffer solutionStandard
$$pH = ?$$
calomel electrode

38.

If the *EMF* of the above cell is 0.03V, $E_{SCE} = 0.24V$, E^{c} . $_{glass} = 0.51V$, then calculate the *pH* of buffer solution at 30 ° C.

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39. For galvanic cell,

$$Ag|AgCl(s), CsCl(0.1M)||CsBr(10^{-3}M), AgBr(s)|Ag.$$

Calculate the *EMF* generated and assign correct polarity to each electrode for spontaneous or exergonic process at $25 \degree C$.

Given
$$\therefore K_{sp}$$
 of $AgCl = 3.0 \times 10^{-10}$, K_{sp} of $AgBr = 4.0 \times 10^{-13}$.

40. EMF of the cell

|Ag|AgNO₃(0.1M) | |KBr(1N), AgBr(s)|Agis - 0.6V at 298K

AgNO₃ is 80 % and KBr is 60 % dissociated.

Calculate a. Solubility and

 $b. K_{sp}$ of AgBr at 298K.

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41. Estimate the $E^{c^{-}}$ reduction for $Cu \mid CuS$ electrode.

Given :
$$K_{sp}$$
 of $CuS = 8.0 \times 10^{-36}$, $E^{c} \cdot (Cu \mid Cu^{2+}) = -0.34V$

A. -0.71V

B. 0.71V

C. -1.42V

D. 1.42V

Answer: A

42. 1F of electricity is passed through 10L of a solution of aqueous solution of NaCl. Calculate the pH of the solution.



43. A current strength of 96.5A is passed for 10s through 1L of a solution

of 0.1M aqueous $CuSO_4$. Calculate the pH of the solution.

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44. A solution of $CuSO_4$ is electrolyzed for 10 min with a current of 1.5*A*. What is the mass of *Cu* deposited at the cathode? [Atomic mass of Cu = 63q]

45. In the electrolysis of 7.2*L* aqueous solution of $CuSO_4$, a current of 9.65*A* passed for 2 hours.

a. Calculate the weight of *Cu* deposited at cathode.

b. Calculate the volume of O_2 produced at anode at 27 °C and 1*atm* pressure.

c. Calculate the *pH* of the solution after electrolysis.

[Atomic mass of Cu = 63g]

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46. How many grams of $Cl_2(g)$ can be produced by the electrolysis molten

NaCl with a current of 5.5*A* for 25*min*? [Atomic weight of *Cl* = 35.5*amu*]

A. 3.017g

B. 3.017g

C.

D.

47. What is the volume of O_{20} liberated at anode at *STP* in the electrolysis

of CdSO₄ solution when a current of 2A is passed for 8min?

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48. What weight of Ni is plated out in an electrolysis of aqueous $NiSO_4$ solution that it takes place to deposit 2G OF Ag in a silver coulometer that is arranged in series with $NiSO_4$ electrolytic cell. [Atomic weight of Ag = 107.8amu, atomic weight of Ni = 58.7amu]

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49. What is the amount of Al deposited on the electrolysis of molten Al_2O_3 when a current of 9.65A is passed for 10.0s .

50. Calculate the number of Faradays required to electrolyze 6.35g of $Cu^{\oplus}(aq)$ ions from an aqueous solution.

51. 100mL of $0.3MFe^{3+}(aq)$ ions were electrolyzed by a charge of 0.072F. In electrolysis, metal was deposited and $O_2(g)$ was evolved. At the end of electrolysis, it is desired to oxidize the un - electrolyzed metal ion. $Fe^{3+} + e^- \rightarrow Fe^{2+}$ $Fe^{2+} + 2e^- \rightarrow Fe$ The moles of Fe^{2+} ions left un - electrolyzed in the solution is

a. 0.009 b.0.021 c.0.072 d.0.042

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52. Marshall's acid $(H_2S_2O_8)$ or peroxodisulphuric acid is prepared by the

electrolytic oxidation of $mmolH_2SO_4$ as :

 $2H_2SO_4 \rightarrow H_2S_2O_8 + 2H^{\oplus} + 2e^-$

 $O_2(g)$ and $H_2(g)$ are obtained as byproducts. In such electrolysis 4.48L of $H_2(g)$ and 1.12L or $O_2(g)$ were produced at *STP*. The weight of $H_2S_2O_8$ formed is

a.9.7g b.19.4g c.14.5g d.29.1g

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53. What volume of $0.2MFeSO_4$ can be oxidized by a current of 0.965 ampere hour?

A. 0.07 L

B. 0.08 L

C. 0.09L

D. 0.1 L

Answer: C

54. $100mLof \ 1M$ solution of $CuBr_2$ was electrolyzed with a current of 0.965 ampere hour. What is the normality of the remaining $CuBr_2$ solution ?

a. 1.64 b.3.28 c.0.82 d.4.92

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55. In a 500mL of $0.5MCuSO_4$ solution, during electrolysis 1.5×10^{23} electron were passed using copper electrodes. Assume the volume of solution remains unchanged during electrolysis. Which of the following statements is/ are correect?

- a. At the end of electrolysis, the concentration of the solution is 0.5M.
- b. 7.9g of Cu is deposited on the cathode.
- c. 4g of Cu is dissolved from the anode.
- d. 7.9g of Cu ions are discharged.



56. A Zn rod weighing 1.0g is taken in 100mL of $1MCuSO_4$ solution . After some time, $\begin{bmatrix} Cu^{2+} \end{bmatrix}$ in solution = 0.9M(atomic weight of Zn = 65.5g). Which of the following statements is / are correct ?

a. 0.655g of Zn was lost during the reaction.

b. 0.327gof Zn was lost during the reaction .

- c. There is no change in the molarity of SO_4^{2-} ion.
- d. There is a change in the molarity of SO_4^{2-} ion.

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57. A sodium salt of ternary acid of molybdenum (atomic mass = 96) has the formula Na_2MoO_n . When an acidified solution of Na_2MoO_n is electrolyzed, O_2 gas is liberated corresponding to a volume of 0.112L at *STP* and 0.32*q* of *Mo* is deposited. Find the formula of salt.

58. Calculate the quantity of electricity required to reduct 24.6g of nitrobenzene to aniline if the current efficienty is 75 %. If the potential drop across the cell is 4.0V, how much energy is consumed (*Mw* of $C_6H_5NO_2 = 123gmol^{-1}$)

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59. Find the volume of gases evolved by passing 0.1A of current for 965s,

through an aqueous solution of sodium succinate at 27 \degree C and 1*atm*.



60. A current strength of 1.0*A* is passed for 96.5*s* through 200*mL* of a solution of 0.05*MKCl*. Find

c -

- a. The amoudn of gases produced
- b. The concentration of final solution w. r. t. OH ions

c. pH of the solution.



61. $100mLCuSO_4(aq)$ was electrolyzed using inert electrodes by passing 0.965Atill the *pH* of the resulting solution was 1. The solution after electrollysis was neutralized, treated with excess *KI* and titrated with $0.04MNa_2S_2O_3$. Volume of $Na_2S_2O_3$ required was 35mL. Assuming no volume change during electrolysis, calculate:

(a) duration of electrolysis if current efficiency is 80 %

(b) initial concentration (M) of $CuSO_4$.

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62. A current of 5.0*A* is passed through a 100*L* aqueous solution of sodium soccinate for 4.0*h*. The gases produced at anode are collected separately at 25 ° *C* and 1.0*atm* pressure. Find the volume of gases. Also find the *pH* of solution at the end of electrylysis . (Assume that at the start of electrolysis, *pH* of solution is 7.0 at 25 ° *C*)

63. A lead storage cell is discharged which causes H_2SO_4 electrolyte to change from a concentration of 40 % by weight (density $1.25gmL^{-1}C$ to one of 30 % by weight. The original volume of electrolyte is 1L. How many Faradays have left the anode of battery. Overall reaction of lead storage cell is : ltbr. $Pb(s) + PbO_2 + 2H_2SO_4(l) \rightarrow 2PbSO_4(s) + 2H_2O$

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64. The density of copper is $8.95gmL^{-1}$. Find out the number of coulombs needed to plate an area of $100cm^2$ to a thickness of $10^{-2}cm$ using $CuSO_4$ solution as electrolyte. (Atomic weight of Cu = 63.5g)



65. What is the (*i*) volume of gases evolved at STP and (*ii*)pH of solution on the electrolysis of 10L of the following solutions when 1F of electricity

is passed :

- a. Aqueous solution of Na_2SO_4
- b. Aqueous solution of CH₃COONa
- c. Aqueous solution of HCOOK



66. 18.97*g* fused $SnCl_{20}$ was electrolyzed using inert electrodes. 1.187*gSn* was deposited at cathode. If nothing is obtained during electrolysis, calculate the ration of weight of $SnCl_2$ and $SnCl_4$ in fused state after electrolysis.

Given:

Atomic weight of Sn = 118.7, $Mwof SnCl_2 = 189.7$, $Mw of SnCl_4 = 260.7$

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67. The resistance of a conductivity cell filled with 0.1MKCl solution is 100Ω . If R of the same cell when filled with 0.02MKCl solution is 520Ω , calculate the conductivity and molar conductivity of 0.02MKCl solution. The conductivity of 0.1MKClsolution is $1.29Sm^{-1}$.

68. The electrical resistance of a column of 0.05MNaOH solution of diameter 1cm and length 50cm is $5.55 \times 10^{3}ohm$. Calculate its resistivity, conductivity, and molar conductivity.

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69. The resistance of 0.01N solution of an electrolysis is 210Ω at 298K with a cell constant of $0.88cm^{-1}$. Calculate the conductivity and equivalent conducitivity of the solution.

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70. The resistance and conductivity of 0.02MKCl solution are 82.4ohm and $0.002768Sch^{-1}$ respectively . When filled with $0.005NK_2SO_4$, the solution had a resistance of 324ohm. Calculate :

a. Cell constant

- b. Conductance of K_2SO_4 solution
- c. Conductivity of K_2SO_4 solution
- d. Equivalent conductivity of K_2SO_4 solution
- e. Molar conductivity of K_{20SO_4} solution.



71. The resistance of decinormal solution of a salt occupying a volume between two platinum electrodes 1.80cm apart and $5.4cm^2$ area was found to be 32ohm. Calculate k and Λ_{eq} .



72. Λ_{eq} of 0.10N solution of CaI_2 is $100.0Scm^2eq^{-1}$ at 298K. G^* of the cell = $0.25cm^{-1}$. How much current will flow potential difference between the electrode is 5V?

A. 0.4

B. 0.6

C. 0.2

D. 0.8

Answer: B

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73. The resistance of a solution A is 50*ohm* and that of solution B is 100*ohm*, both solutions are taken in the same conductivity cell. If equal volumes of solutionA and B are mixed, what is the resistance of the mixture using the same cell ? (Assume there is no change or increase in the \propto of A and B on mixing).

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74. Arrange the following compounds in the order of increasing conductance : *HCl*, *LiCl*, *NaCl*, *KCl*.

75. Arrange the following compounds in the order of decreasing molar conductivity in aqueous solution.

$$i. a. K \Big[Co \Big(NH_3 \Big)_2 \Big(NO_2 \Big)_4 \Big]$$

$$b. \Big[Cr \Big(NH_3 \Big)_3 \Big(NO_2 \Big)_3 \Big]$$

$$c. \Big[Cr \Big(NH_3 \Big)_5 \Big(NO_2 \Big) \Big]_3 \Big[Co \Big(NO_2 \Big)_6 \Big]_2$$

$$d. \Big[Mg \Big(Cr \Big(NH_3 \Big)_5 \Big) \Big]$$

$$ii. a. Li \Big[Co \Big(NH_3 \Big)_2 \Big(NO_2 \Big)_4 \Big] \quad b. \quad Na \Big[Co \Big(NH_3 \Big)_2 \Big(NO_2 \Big)_4 \Big]$$

$$ii. a. Li \Big[Co \Big(NH_3 \Big)_2 \Big(NO_2 \Big)_4 \Big] \quad d. \quad Cs \Big[Co \Big(NH_3 \Big)_2 \Big(NO_2 \Big)_4 \Big]$$

$$iii. a. Pt \Big(NH_3 \Big)_6 Cl_4 b. \quad Cr \Big(NH_3 \Big)_6 Cl_3$$

$$c. \quad Co \Big(NH_3 \Big)_4 Cl_3 \quad d. K_2 PtCl_6$$

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76. The molar conductivity of *KCl* solution at different concentrations at 298*K* is given below :

 $c \text{ or } M(molL^{-1}) \qquad \wedge_m(Scm^2mol^{-1})$ $0.000198 \qquad 148.61$ $0.000309 \qquad 148.29$ $0.000521 \qquad 147.81$ $0.000989 \qquad 147.09$ Show that a plot between \wedge_m and \sqrt{c} is a straight line. Determine the value of \wedge_m° and A for KCl.

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77. The conductivity (k) of a saturated solution of AgBr at 298K is $8.5 \times 10^{-7}Scm^{-1}$. If $\lambda^{\circ} \cdot_{Ag^{\oplus}}$ and $\lambda^{\circ} \cdot_{Br^{c-}}$ are 62 and $78Scm^2mol^{-1}$, respectively, then calculate the solubility and K_{sp} of AgBr.

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78. $\Lambda^{\circ} \cdot_{m}$ for $CaCl_{2}$ and $MgSO_{4}$ from the given data. $\lambda^{\circ}_{Ca^{2+}} = 119.0Scm^{2}mol^{-1}$ ltbr. $\lambda^{\circ}_{Cl^{c-}} = 76.3Scm^{2}mol^{-1}$ $\lambda^{\circ}_{Mg^{2+}} = 106.0Scm^{2}mol^{-1}$ $\lambda^{\circ}_{SO_{4}^{2-}} = 160.0cm^{2}mol^{-1}$
79. Λ_m° for *NaCl*, *HCl*, and *NaAc* are126.4, 425.9, and 91.0Scm²mol⁻¹, respectively. Calculate Λ° for *Hac*.

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80. The conductivity of 0.001028M acetic acid is $4.95 \times 10^{-5}Scm^{-1}$. Calculate dissociation constant if Λ_m° for acetic acid is $390.5Scm^2mol^{-1}$.

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81. The molar conductivity of acetic at infinite dilution is 390.7 and for 0.01M acetic acid is $3.9.7Scm^2mol^{-1}$. Calculate (*a*) α and (*b*)*pH* of solution.

82. The ionic equivalent conductivities of $C_2 O_4^{2-}$, K^{\oplus} , and Na^{\oplus} ions are *x*, *y*, and $zScm^2Eq^{-1}$ respectively. Calculate Λ_{ea}° of (*NaOOC* - *COOK*).



83. The ionic molar conductivities of $C_2 O_4^{2^-}$, K^{\oplus} , and Na^{\oplus} ions are x', y', and $z'Scm^2mol^{-1}$, respectively. Calculate Λ_m° and Λ_{eq}° of (*NaOOC* - *COO*)

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84. The ionic equivalent conductivities of K^{\oplus} , Al^{3+} , and SO_4^{2-} ions are x, y, and $zScm^2Eq^{-1}$, respectively. Calculate Λ_m° and Λ_{eq}° for $\left(K_2SO_4.Al_2\left(SO_4\right)_3.24H_2O\right)$ (Potash alum).

85. From the following molar conductivities at infinite dilution :

$$\Lambda_{m}^{\circ} \text{ for } Ba(OH)_{2} = 457.6\Omega^{1}cm^{2}mol^{-1}$$
$$\Lambda_{m}^{\circ} \text{ for } BaCl_{2} = 240.6\Omega^{-1}cm^{2}mol^{-1}$$
$$\Lambda_{m}^{\circ} \text{ for } NH_{4}Cl = 129.8\Omega^{-1}cm^{2}mol^{-1}$$
Calculate Λ_{m}° for $NH_{4}OH$.

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Solved Examples (Electrochemical Cell)

1. Calculate EMF of the following half cells :

a. Pt, $H_2(2atm) \mid HCl(0.02M) \quad E^{c-} = 0V$

b. Pt, $Cl_2(10atm) \mid HCl(0.1M) \quad E^{-c} = 1.36V$

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2. Calculate *pH* of the half cell :

 $Pt, H_2(1atm) \mid H_2SO_4 \quad E^0 = -0.3V$

A.	2
В.	5

C. 10

D. 1

Answer: B

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3. A graph is plotted between E_{cell} and $\log \cdot \frac{\left[Zn^{2^+}\right]}{\left[Cu^{2^+}\right]}$. The curve is linear

with intercept on E_{cell} axis equals to 1.10V. Calculate E_{cell} for the cell.

$$Zn(s) || Zn^{2+}(0.1M) || Cu^{2+}(0.01M) | Cu^{2+}(0.01$$

4. Given :

 $NO_3^{c^-} \rightarrow NO_2$ (acidic medium), $E^{c^-} = 0.8V$ $NO_3^{c^-} \rightarrow NO_3OH$ (acidic medium) $E^{c^-} = 0.74V$ At what *pH* the above two half reactions will have same *EMF* values ? Assume the concentration of all the species to be unity. (Take 0.059 \approx 0.06)

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5. The standard potential of a cell using the reaction $2Ni(s) + O_2 + 2H_2O \implies 2Ni(OH)_2$ is 1.12. The

heat of the reaction is $-504.2kJmol^{-1}$ at $25 \degree C$. Calculate the entropy change.

6. The standard potential of a cell using the reaction $2MnO_4^{\oplus}(aq) + 3Hg(l) + H_2O \implies 2MnO_2(s)$

 $+3HgO(s) + 2\begin{pmatrix} c-\\OH \end{pmatrix}(aq)$ is 0.489V at 25 °C. What is the equilibrium

constant of the reaction ?

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7. The EMF of the cell :

 $Ag|AgCl, 0.1MKCl| | 0.1MAgNO_3 | Ag$ is 0.45V. 0.1MKClis 85% dissociated and 0.1MAgNO_3 is 82% dissociated. Calculate the solubility product of AgCl at 25 ° C.

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8. Calculate the potential corresponding to the following cell. Given

$$Pt \left| Co^{2+}(2.0M), Co^{3+}(0.01M) \right| \left| Cr^{3+}(0.5M), Cr_2O_7^{2-}(4.0M), H^{\oplus}(1.5M) \right| Pt$$

$$E^{c} \cdot c_{o^{2^{+}} \mid Co^{3^{+}}} = -1.82V$$

$$E^{c} \cdot c_{r_2} O_7^{2^{-}} \mid Cr^{3^{+}} = +1.33V$$
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9. Estimate the standard reduction potential for the copper / copper

sulphide electrode. For *CuS*, $K_{sp} = 8.5 \times 10^{-36}$, $E^{c-} \cdot_{Cu \mid Cu^{2+}} = -0.34V$.

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10. Knowing that K_{sp} for AgCl is 1.0×10^{-10} , calculate E for a silver / silver chloride electrode immersed in 1.00MKCl at $25 \degree C$. $E^{c} \cdot Ag \circledast | Ag = 0.799V$.

11. Consider the following half reactions :

$$PbO_{2}(s) + 4H^{\oplus}(aq) + SO_{4}^{2-} + 2e^{-} \rightarrow PbSO_{4}9s + 2H_{2}O \quad E^{c-} = +1.70V$$
$$PbSO_{4}(s) + 2e^{-} \rightarrow Pb(s) + SO_{4}^{2-}(aq) \quad E^{c-} = -0.31V$$

a. Calculate the value of E^{c-} for the cell.

b. Calculate the voltage generated by the cell if $\left[H^{\oplus}\right] = 0.10M$ and $\left[SO_4^{2-}\right] = 2.0M$

c. What voltage is generated by the cell when it is at chemical equilibrium ?

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12. The *EMF* of the cell

 $Ag \left| 0.1(N)AgNO_3 \right| |1(N)KBr, AgBr(s)|Ag$ was found to be -0.64V at 298K. $0.1NAgNO_3$ is 81.3 % dissociated and 1NKBr is 75.5 % dissociated. Calculate :

- a. Solubility
- b. Solubility product of AgBr at 298K

13. For the cell $Zn \left| ZnCl_2(m) \right| AgCl$, E is 1.24V at 25 ° C and 1.260V at 35 ° C of $m = 10^{-3}$. Write down the cell reaction and calculate ΔG , ΔH , and ΔS at 25 ° C.



14. A saturated calomel electrode is coupled through a salt bridge with a quinhydrone electrode dipping in $0.1MNH_4Cl$. The observed *EMF* at 25 °*C* is 0.152*V*. Find the dissociated constant of NH_4OH . The oxidation potential of saturated calomel electrode = -0.699V at 25 °*C*.

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15. Two weak acid solutions HA_1 and HA_2 with the same concentration and having pK_a values 3 and 5 are placed in contact with hydrogen electrode (1*atm* and 25 ° *C*) and are interconnected through a salt bridge. Find the *EMF* of the cell. **16.** Find the solubility of AgCl in $0.1MCaCl_2$. $E^{C^-} \cdot_{Ag} \oplus |_{Ag} = 0.799V$ and that of AgCl | Ag = 0.222V.



17. The *EMF* of the cell :

 $Ag \left| Ag_2 CrO_4(s), K_2 CrO_4(0.1M) \right| \left| AgNO_3(0.1M) \right| | Ag \text{ is } 206.5mV. \text{ Calculate}$

the solubility of Ag_2CrO_4 in $1MNa_2CrO_4$ soluion.

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18. The *EMF* of a galvanic cell $Pt | H_2(1atm) | HCl(1M) | Cl_2(g) | Pt$ is 1.29V.

Calculate the partial pressure of $Cl_2(g)$. $E^{c^-} \cdot Cl_2 | Cl^{c^-} = 1.36V$.

19. Calculate the potential of silver electrode in a saturated solution of $AgBr(K_{sp} = 6 \times 10^{-13})$ containing 0.1*MKBr*. E^{c} . $Ag^{\oplus}|_{Ag} = 0.80V$. Watch Video Solution **20.** A solution of Fe^{2+} is titrated potentiaometrically using Ce^{4+} solution . Calculate the *EMF* of the redox electrode thus formed when a. 50 % of Fe^{2+} is titrated b. 90 % of Fe^{2+} is titrated c. 110 % titration is done Given : $E^{C^-} \cdot_{Fe^{2^+} | Fe^{3^+}} = -0.77V$ and $Fe^{2^+}Ce^{4^+} \rightarrow Fe^{3^+} + Ce^{3^+}, K = 10^{14}$ View Text Solution 25 ° C. Find the *EMF* of the cell at 21.

DecinormalBufferQuinhydronecalomel electrodepH = 3.5electrode



 E^{c} ·*red*(calomel electrode) = +0.268V·



22. The *EMF* of the following cell is found to be -0.46V:

Pt, H₂
$$\begin{vmatrix} NaHSO_4 \\ (0.4 \text{ M}) \end{vmatrix} \begin{vmatrix} Na_2SO_4 \\ (6.44 \times 10^{-3} \text{ M}) \end{vmatrix} \begin{vmatrix} Zn^{+2} \\ (0.3 \text{ M}) \end{vmatrix}$$
 Zn

If the standard electrode potential of Zince Is -0.763V, find the value of K_2

for
$$H_2SO_4$$
, where $K_2 = \frac{\left[H^{\oplus}\right]\left[SO_4^{2^-}\right]}{\left[HSO_4^{c^-}\right]}$

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23. The *EMF* of the following cell is observed to be 0.118V at $25 \degree C$:

Pt,
$$H_2$$
 HA
 H ^{\oplus}
 H_2 , Pt

 (1 atm)
 (100 mL of 0.1 M)
 (0.1 M)
 (1 atm)

If 30mL of 0.2MNaOH is added to the negative terminal of the battery, find the *EMF* of the cell.

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24. Find the standard electrode potential of $I_2 \mid 2I^{c-}$ if the equilibrium constant for the reaction $I_2 + I^{c-} \rightarrow \frac{c^-}{3}$ is 703. The standard electrode potential of $I_3^{c-} \mid 3I^{c-}$ is 0.5355V. Also give the electrode reaction.

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25. A silver electrode dipping in $AgNO_3$ solution (0.1*M*) is combined salt bridge with a hydrogen electrode dipping in a solution of $pH = 3(at25 \,^{\circ}C)$. If the standard reduction potential of the silver electrode is 0.799V, what is the *EMF* of the cell ?

26. The *EMF* of a galvanic cell composed of two hydrogen electrodes is 177mV. If the solution at one of the electrode has $\left[H^{\oplus}\right] = 10^{-3}$, find the $\left[H^{\oplus}\right]$ at the other electrode.



27. The *EMF* of the cell :

 $Pt \left| Ce^{4+} (90\%), Ce^{3+} (10\%) \right|$ Normal calomel electrode is 1.464V at 25°C

. Find the value of equilibrium constant of the reaction :

 $2Ce^{3+} + 2H^{\oplus} \rightarrow 2Ce^{4+} + H_2$

The electrode potential of the normal calomel electrode is +0.28V.

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28. A hydrogen electrode placed in a buffer solution of sodium cyanide and HCN in the ration of x: y and Y: x has electrode potential value a and *b* volts, respectively, at 25 ° *C*. If the difference a - b = 35.52mV, what is the ratio of *y*:*x*.

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29. Two electrochemical cells are assembled in which the following reactions occur : $V^{2} = VO^{2+} + 2H^{\oplus} \rightarrow 2V^{3+} + H_{2}O$ $V^{3+} + Ag^{\oplus} + H_{2}O \rightarrow VO^{2+} + 2H^{\oplus} + Ag(s)$ Calculate E^{c-} for half reaction $V^{3+} + e^{-} \rightarrow V^{2+}$ Given : $E^{c-} \cdot (Ag^{\oplus} | Ag) = 0.799$ $E^{c-} = E^{c-} \cdot V^{4+} | V^{3+} - E^{c-} \cdot V^{3+} | V^{2+} = 0.616V$ $E^{c-} = E^{c-} \cdot Ag^{\oplus} | Ag - E^{c-} \cdot V^{4+} | V_{3+} = 0.439V$

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30. Predict whether or not Cl_2 would disproportionate in cold alkaline medium. The standard reduction potentials of $Cl_2 \mid Cl^{c-}$ and $ClO^{c-} \mid Cl_2$ are 1.36V and 0.40V, respectively.

31. What would be the electrode potential of a silver electrode dipped in a saturated solution of AgCl in contact with 0.1MKCl solution at 25 ° C?

$$E^{c}$$
-._{Ag} \oplus | Ag = 0.799V

 K_{sp} of $AgCl = 1 \times 10^{-10}$

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32. $E^{c^{-}}$ of some elements are given as : $I_{2} + 2e^{-} \rightarrow 2I^{c^{-}}$ $E^{c^{-}} = 0.54V$ $MnO_{4}^{c^{-}} + 8H^{\oplus} + 5e^{-} \rightarrow Mn^{2^{+}} + 4H_{2}O$ $E^{c^{-}} = 1.52V$ $Fe^{3^{+}} + e^{-} \rightarrow Fe^{2^{+}}$ $E^{c^{-}} = 0.77V$ $Sn^{4^{+}} + 2e^{-} \rightarrow Sn^{2^{+}}$ $E^{c^{-}} = 0.1V$

a. Select the stronges reductant and weakes oxidant among these elements.

b. Select the weakest reductant and strongest oxidant among these elements.

33. Select the spontaneous rreactions from the changes given below

a. $Sn^{4+} + 2Fe^{2+} \rightarrow Sn^{2+} + 2Fe^{3+}$ b. $2Fe^{2+} = I_2 \rightarrow 2Fe^{3+} + 2I^{c-}$ c. $Sn^{4+} + 2I^{c-} \rightarrow Sn^{2+} + I_2$ d. $Sn^{2+} + I_2 \rightarrow Sn^{4+} + 2I^{c-}$

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34. Two metals M_1 and M_2 have $E^{c-} \cdot_{red} = -0.76V$ and 0.80V, respectively. Which will liberate $H_2(g)$ from H_2SO_4 ?

Given :

 $M_1^{n^+} + ne^- \rightarrow M_1 \qquad E^{c^-} \cdot_{red} = -0.76V$ $M_2^{n^+} + ne^- \rightarrow M_2 \qquad E^{c^-} \cdot_{red} = 0.80V$ $2H^{\oplus} + 2e^- \rightarrow H_2(g) \qquad E^{c^-} \cdot_{red} = 0.0V$

35. Determine whether $O_2(g)$ can oxidize sulphate (SO_4^{2-}) ion to peroxodisulphate $(S_2O_8^{2-})$ ion or not in an acidic solution with $O_2(g)$ being reduced to water.

Given :
$$E^{C^-} \cdot O_2 | H_2O = 1.20V$$
 and $E^{C^-} \cdot S_2O_8^{2^-} | 2SO_4^{2^-} = 2.0V$

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36. Quinones are good electron acceptors, party because reduction restores aromaticity.



 $Q + 2e^- + 2H^{\oplus} \rightarrow H_2Q$

Give the decreasing order of E^{c-} . *reduction* of the following quinones :



Solved Examples(Electrolysis And Electrolytic Cells)

1. Express each of the following combinations of electrical units as a single unit:

a. Volt - ampere



2. A resistance heater was wound around a 5.0g metallic cylinder. A current of 0.84A was passed through the heater for 20s while the drop in voltage across the heater was 50V. The temperature change of the cylinder was from 25 °C before the heating period and 35 °C at the end. If the heat loss is neglected, what is the specific heat of the cylinder metal in $calg^{-1}K^{-1}$.

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3. In a zinc manganese dioxide dry cell, the anode is madeup of Zn1 and cathode of carbon rod surrounded by a mixture of MnO_2 , carbon, NH_4Cl , and $ZnCl_2$ in aqueous base. The cathodic reaction is :

 $ZnMnO_2(s) + Zn^{2+} + 2e^- \rightarrow ZnMn_2O_4(s)$

8.7*g* of MnO_2 is taken in the cathodic compartment. How many days will the dry cell continue to give a current of $9.65 \times 10^{-3}A$? (Atomic weight of Mn = 55)(Mw of $MnO_2 = 87gmol^{-1}$)

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4. An aqueous solution of *NaCl* on electrolysis gives $H_2(g)$, $Cl_2(g)$, and *NaOH* accroding to the reaction :

$$2Cl^{c}(aq) + 2H_2O \rightarrow 2OH(aq) + H_2(g) + Cl_2(g)$$

A direct current of 25*A* with a current efficiency of 62 % is passed through 20*L* of *NaCl* solution (20 % by weight). Write down the reactions taking place at the anode and cathode. How long will it take to produce 1kg of Cl_2 ? What will be the molarity of the solution with respect to hydroxide ion ? (Assume no loss due to evaporation .)



5. A constant current was flowen for 1min through a solution of Kl. At the end of experiment, liberated I_2 consumed 150mL of 0.01M solution of $Na_2S_2O_3$ following the reaction :

 $I_2 + 2S_2O_3^{2-} \rightarrow 2I^{c-} + S_4O_6^{2-}$

What was the average rate of current flow in ampere ?

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6. Aqueous solution of m - dinitroso benzene was electrolyzed for 2 hors passing current of 2A with efficiency of 90 %. Calculate the amound of 3 - aminoaniline.

$$(Mw = 108gmol^{-1})$$

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7. A 35 % solution of *LiCl* was electrolyzed by using a 2.5*A* current for 0.8*h*. Assuming the current efficiency of 90 % , find the mass of *LiOH* produced at the end of electrolysis. (Atomic mass of Li = 7)

8. A current of 10A is employed to plate nickel in $NiSO_4$ bath. The current efficiency with respect to Ni plating is 60 % .

a. How many grams of Ni is plated on the cathode per hour ?

b. What is the thickness of the plating if the cathode consists of a sheet of metal $4.0cm^2$ which is coated on both faces ?

c. What is the volume of $H_2(STP)$ evolved during above electrolysis ?

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9. A certain amount of charge is passed through acidulated water. A total of 504mL of hydrogen and oxygen were collected at STP. Find the magnitude of charge that is passed during electrolysis in coulombs.

10. During the electrolysis of water, a total volume of 33.6*mL* of hydrogen and oxygen gas was collected at *STP*. Find the amound of electricity that passed during electrolysis.

11. During an electrolysis of conc H_2SO_4 , perdisulphuric acid $(H_2S_2 - (8))$ and O_2 are formed in equimolar amount. The moles of H_2 that will be formed simultaneously will be

A. Thrice that of O_2

B. Twice that of O₂

C. Equal to that of O_2

D. Half of that of O₂

Answer: A

12. A current of 1.0A is passed for 96.5s trhough a 200mL solution of

0.05MLiCl solution. Find

a. The volume of gases produced at STP

b. The pH of solution at the end of electrolysis

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Ex 3.1 (Objective)

1. If the temperature coefficient of *EMF* if $-0.125VK^{-1}$, ΔS for the given cell at 25 ° *C* is :

$$Fe\left|Fe^{2+}(aq)\right|\left|Cd^{2+}(aq)\right|Cd$$

A. - 26.125*kJK*⁻¹

B. - 24.125*kJK*⁻¹

C. - 22.125*kJK*⁻¹

D. - 20.125*kJK*⁻¹

Answer: b

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- 2. Which of the following is (are) function (s) of salt bridge?
 - A. It completes the electrical circuit with electrons flowing from one

electrode to other through wires and flow of ions between the two

compartments through salt bridge.

B. It prevents the accumulation of the ions.

C. Both (a) and (b)

D. None of the above.

Answer: C

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3. $Cu^{2+} + 2e^- \rightarrow Cu$. On increasing $\begin{bmatrix} Cu^{2+} \end{bmatrix}$, electrode potential

A. Increases

B. Decreases

C. No change

D. First increases, then decreases

Answer: a

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4. Consider the following E° values $E_{Fe^{3+}/Fe^{2+}} \circ = +0.77V^{\circ}$

 $E_{Sn^{2+}/Sn}^{\circ} = -15$. V The E_{cell}° for the reaction ,

`Sn (s) + 2Fe_(aq)^(3+) rarr 2 Fe_(aq.)^(2+) + Sn_(aq.)^(2+) is :

A. -0.58V

B.-0.30V

C.+0.30V

D. +0.58V

Answer: c

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5. $E^{c^{-}}$ of some elements are given as : $I_{2} + 2e^{-} \rightarrow 2I^{c^{-}}$ $E^{c^{-}} = 0.54V$ $MnO_{4}^{c^{-}} + 8H^{\oplus} + 5e^{-} \rightarrow Mn^{2^{+}} + 4H_{2}O$ $E^{c^{-}} = 1.52V$ $Fe^{3^{+}} + e^{-} \rightarrow Fe^{2^{+}}$ $E^{c^{-}} = 0.77V$ $Sn^{4^{+}} + 2e^{-} \rightarrow Sn^{2^{+}}$ $E^{c^{-}} = 0.1V$

a. Select the stronges reductant and weakes oxidant among these elements.

b. Select the weakest reductant and strongest oxidant among these elements.

A. Zn

B. *Cr*

 $C.H_2$

D. Fe

Answer: a



6. For hydrogen oxygen fuel cell with reaction

 $2H_2(g) + O_2(g) \rightarrow 2H_2O(l)$ $\Delta G_f^{c-}(H_2O) = -237.2kJmol^{-1}$. Hence, *EMF* of the fuel cell is A. +2.46V B. -2.46V C. +1.23V

D. - 1.23V

Answer: c

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7. A metal - insoluble salt electrode consists of

A. A piece of metal placed in a solution containing a sparingly soluble

salt.

B. Crystals of an insoluble salt coated with a metal.

C. A piece of metal coated with one of its insoluble salts.

D. A metal fixed with an insoluble salt at high temperature.

Answer: c

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8. Which of the following is the most powerful reducing agent ?

A. *F*^{*c*} -

B. *Cl^c*-

C. *Br^{c-}*

D. *I^c* -

Answer: d

9. If all species are in their standard states, which of the following is the strongest oxidizing agent ?

A. *Br^c* -

B. Zn^{2+}

C. *CO*³⁺

D. Fe^{2+}

Answer: c

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10. The standard *EMF* of a galvanic cell involving cell reaction with n = 2 is found to be 0.295*V* at 25 ° *C* . The equilibrium constant of the reaction would be

A. 4.0×10^{12}

B. 1.0×10^{2}

 $C. 1.0 \times 10^{10}$

D. 2.0×10^{11}

Answer: c

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11. The correct order of reactivity of K, Mg, Zn and Cu with water according to the electrochemical series is

A. K > Mg > Zn > Cu

B. Mg > Zn > Cu > K

C.K > Zn > Mg > Cu

D. Cu > Zn > Mg > K

Answer: a

12. For $Pt(H_2) \mid H_2O$, reduction potential at 298K and 1*atm* is :

A. -0.23V

B.-0.41V

C. 0.41V

 $D.\,0.00V$

Answer: b

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13. Consider the cell:

$$Pt\Big|H_2\Big(p_1atm\Big)\Big|H^{\oplus}\Big(x_1M\Big) | \Big|H^{\oplus}\Big(x_2M\Big)\Big|H_2\Big(p_2atm\Big)Pt.$$

The cell reaction be spontaneous if

A. $p_1 = p_2$

B. $p_1 > p_2$

 $C. p_2 > p_1$

D. $p_1 = 1$ *atm*

Answer: b



14. If
$$E^{c} \cdot Fe^{3+} | Fe$$
 and $E^{c} \cdot Fe^{2+} | Fe$ are = -0.36V and -0.439V,
respectively, then the value of $E^{c} \cdot Fe^{3+} | Fe^{2+}$

A. $3x_2 - 2x_1$

- **B.** *x*₂ *x*₁
- $C. x_2 + x_1$

D. $2x_2 + 3x_2$

Answer: a

15. $Pt(Cl_2)(p_1)|HCl(0.1M)|(Cl_2)(p_2), Pt$ cell reaction will be endergonic if

A. $p_1 = p_2$ B. $p_1 > p_2$ C. $p_2 > p_1$

D. $p_1 = 1$ *atm*

Answer: c

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16. Consider the following cell with hydrogen electrodes at difference pressure p_1 and p_2 .

Pt, H2 (
$$p_1$$
)H ^{\oplus} (aq)H2 (p_2), Pt1 M1 M

The EMF of the cell is given by

A.
$$\frac{RT}{F}$$
ln. $\frac{p_1}{p_2}$
B. $\frac{RT}{2F}$ ln. $\frac{p_1}{p_2}$
C. $\frac{RT}{F}$ ln. $\frac{p_2}{p_1}$
D. $\frac{RT}{2F}$ ln. $\frac{p_2}{p_1}$

Answer: b

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17. Consider the following cell reaction

 $Zn + 2Ag^{\oplus} \rightarrow Zn^{2+} + 2Ag.$

Given that

 $E^{C^{-}} \cdot _{Zn^{2^{+}}(aq)Zn(s)} = -0.76V$
$E^{c} \cdot Ag \oplus (aq) Ag(s) = 0.80V$

- *a*. Calculate the standard *EMF* fo the cell.
- b. Which ion is more powerful oxidizing agent ?
- c. Which metal is more powerful reducing agent ?

A. x + 2y

B. 2x + y

C. y - x

D. y - 2x

Answer: c

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18. The standard reduction potentials of three metals A, B, and C are +0.5V, -3.0V, and -1.2V, respectively. The order of reducing power of these metals is

A. B > C > A

 $\mathsf{B}.A > B > C$

 $\mathsf{C}.\, C > B > A$

D.A > C > B

Answer: a

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19. Calculate the maximum work that can be obtained from the decimolar

Daniell cell at 25 ° C.

Given
$$E^{C^{-}} \cdot (Zn^{2^{+}} | Zn) = -0.76V$$
 and $E^{C^{-}} \cdot (Cu^{2^{+}} | Cu) = 0.34V$

A. 193.0kJ

B. 212.3*kJ*

C. 81.06kJ

D. 40.53kJ

Answer: B



20. Stronger the oxidizing agent, greater is the

A. Standard reduction potential

B. Standard oxidation potential

C. lonic nature

D. None

Answer: a

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21. Consider the cell $Ag(s) |AgBr(s)Br^{c-}(aq)| |AgCl(s), Cl^{c-}(aq)| Ag(s)$ at 298K. The K_{sp} of AgBr and AgCl, respectively are 5×10^{-13} and 1×10^{-10} . At what ratio of $[Br^{c-}]$ and $[Cl^{c-}]$ ions, EMF_{cell} would be zero ?

A. 200:1

B.1:200

C. 1:100

D.1:500

Answer: a

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22. The *pH* of *LHE* in the following cell is :

 $Pt, H_{2}(1atm) \left| H^{\oplus}(xM) \right| \left| H^{\oplus}(0.1M) \right| H_{2}(0.1atm) Pt \text{ E}_(\text{cell})=0.295 \text{V}.$

A. 6.5

B. 6.0

C. 5.5

D. 4.0

Answer: c

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23. At what concentration of $\begin{bmatrix} c - \\ OH \end{bmatrix}$ does the following half reaction has a

potential of 0V when other species are at 1M?

 $NO_3^{c-} + H_2O + 2e^- \rightarrow NO_2^{c-} + 2OH, E^{c-} \cdot_{cell} = -0.06V$

A. 2.0M

B. 1.0M

C. 0.1M

D. 0.01M

Answer: c



24. If hydrogen electrodes dipped in two solutions of pH = 3 and pH = 6 are connected by a salt bridge, the EMF_{cell} is

A. 0.052V

B.0.104V

C. 0.177*V*

D. 0.3V

Answer: C

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25. Consider the cell reaction :

$$Mg(s) + Cu^{2+}(aq) \rightarrow Cu(s) + Mg^{2+}(aq)$$

If $E^{c-} \cdot Mg^{2+} | Mg(s)$ and $E^{c-} \cdot Cu^{2+} | Cu(s)$ are -2.37 and 0.34V, respectively.
 $E^{c-} \cdot Cu^{2+} | S$

A. 2.03V

B. - 2.03V

C. - 2.17V

D. 2.71V

Answer: d



$$E^{c} \cdot Cu^{2+} | Cu = 0.34V, E^{c} \cdot Zn^{2+} | Zn = -0.76V.$$
$$E^{c} \cdot Ag \oplus | Ag = 0.80V, E^{c} \cdot Mg^{2+} | Mg = -2.37V.$$

In which cell ΔG^{c-} is most negative ?

A.
$$Zn |Zn^{2+}(1M)| |Mg^{2+}(1M)| Mg$$

B. $Zn |Zn^{2+}(1M)| |Ag^{\oplus}(1M)| Ag$
C. $Cu |Cu^{2+}(1M)| |Ag^{\oplus}(1M)| Ag$
D. $Ag |Ag^{\oplus}(1M)| |Mg^{2+}(1M)| Mg$

Answer: b

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27. For the reaction :

$$A + 2B^{3+} \rightarrow 2B^{2+} + A^{2+}$$

$$E^{0} \text{ of the given redox reaction is :}$$

$$A^{2+} + 2e^{-} \rightarrow A E^{0} = xV$$

$$B^{3+} + e^{-} \rightarrow B^{2+} E^{0} = yV$$

$$A \cdot x - 2y$$

$$B \cdot x + y/2$$

$$C \cdot x - y$$

$$D \cdot y - x$$

Answer: D



28. Excess of solid AgCl is added to a 0.1M solution of Br^{c-} ions. E^{c-} for

half cell is :

 $AgBr + e^- \rightarrow Ag + Br^{c-}, E^{c-} = 0.095V$

 $AgCl + e^- \rightarrow Ag + Cl^{c-}, E^{c-} = 0.222V$

The value of $\left[Br^{c}\right]$ ion at equilibrium is :

[Given : Antilog(2.152) = 142]

A. 0.0317M

B. 0.013M

C. 0.99M

D. 0.099M

Answer: d

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Ex 3.2 (Objective)

1. During the electrolysis of acidified water, O_2 gas is formed at the anode. To produce O_2 gas at the anode at the rate of 0.224Ml per second at *STP*, the current passed is A. 0.224A

B. 2.24A

C. 9.64A

D. 3.86A

Answer: d

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2. Number of electrons lost dring electrolysis of 0.355g of Cl^{c-} is $(N_A =$

Avogadro's number)

A. 0.01

B. 0.01N_A

C. 0.02N_A

D.
$$\frac{0.01}{2N_A}$$

Answer: b

3. How many faradays are required to reduce 1 mol of MnO_4^- to Mn^{2+} ?

A. 1 B. 5 C. 3 D. 2

Answer: B

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4. Copper containing zinc as impurity is refined by electrolysis. The cathode and anode used are

A.	Cathode	Anode
	Purecopper	Purezinc
B.	Cathode	Anode
	Purezinc	Purecopper

C.	Cathode	Anode
	Purecopper	Impurecopper
D.	Cathode	Anode
	Purezinc	Impurezinc

Answer: c

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5. Which of the following reactions is / are possible at the anode ?

A.
$$F_2 + 2e^- \rightarrow 2F^{c^-}$$

B. $2H^{\oplus} + \frac{1}{2}O(2) + 2e^- \rightarrow H_2O$
C. .2 Cr 3 + + 7 H 2 O \rightarrow Cr 2 O 2 - 7 + 14 H \oplus + 6 e -
D. $Fe^{2^+} \rightarrow Fe^{3^+} + e^-$

Answer: c,d

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6. The number of moles of Zn^{2+} ions deposited when a current of 1.5A is passed for 4 hours through a molten solution of a zince salt. (Assume current efficiency to be 90 %)

A. 6.35

B. 0.1

C. 0.4

D. None of these

Answer: b

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7. Molten NaCl is electrolyzed in a cell called

A. Downs cell

B. Castner cell

C. Kellner cell

D. Hall cell

Answer: a., b,c., d.



8. A dilute aqueous solution of sodium fluoride is electrolyzed, the products at the and cathode are

A. O_2 and K

B. O_2 and F_2

 $\operatorname{C.}H_2$ and F_2

D. O_2 and H_2

Answer: D

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9. If 0.224*L* of $H_2(g)$ is formed at the cathode of one cell at *STP*, how much of *Mg* is formed at the cathode of the other electrolytic cell arranged in series ?

A. 0.24g

B. 2.4g

C. 0.48g

D. 4.8g

Answer: a

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10. A certain amount of charge is passed through acidulated water. A total of 504mL of hydrogen and oxygen were collected at STP. Find the magnitude of charge that is passed during electrolysis in coulombs.

B.965C

C. 482.5*C*

D. 241.2*C*

Answer: b

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11. 1*L* of 1*MCuSO*₄ solution is electrolyzed using *Pt* cathode and *Cu* anode. After passing 2*F* of electricity, the $[Cu^{2+}]$ will be

A. 0

 $\mathsf{B}.M$

C. M/2

D.*M*/4

Answer: b

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12. In a *Ni* - *Cd* battery (more than one correct)

A. All the reactants and products in the overall reaction are in the

solid state.

B. The voltage of the cell changes rapidly.

C. The elctrolyte used is an alkali solution.

D. All of the above are true.

Answer: a,c

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13. Rusting of iron is

A. A decomposition process

B. A photochemical process

C. An electrochemical

D. A reduction process

Answer: c



14. In H_2 - O_2 fuel cell, the reaction occurring at cathode is

A. $2H_2O(l) \rightarrow 2H_2(g) + O_2(g)$

 $\mathsf{B.} 2H_2(g) + O_2(g) \rightarrow 2H_2O(l)$

 $\mathsf{C.} CH_4(g) + O_2(g) \rightarrow CO_2(g) + H_2O(l)$

 $\mathsf{D}.\ C(s) + O_2(g) \rightarrow CO_2(g)$

Answer: b

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15. The cathode reaction during the charging of a lead - acid battery leads to the

A. Formation of PbSO₄

- B. Reduction of Pb^{2+} to Pb
- C. Formation of PbO₂

D. Deposition of *Pb* at the anode.

Answer: b

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16. Which of the following cells is rechargeable ?

A. Lead storage cell

B. Ni - Cd cell

C. Edison cell (Iron - nickel cell)

D. All of these

Answer: d



D. All of the above are true.

Answer: d



18. Explain how rusting of iron is envisaged as setting up of an electroCHMemical cell.

A. Fe^{2+} | Fe B. O_2 | H_2O C. Fe^{3+} | F^{2+} D. Fe | Fe^{3+}

Answer: b

Watch Video Solution

19. How many Faradays are required to reduce $1 molof BrO_3^{c^-}$ to Br^{c^-} in basic medium ?

A. 6

B. 5

C. 4

D. 3

Answer: a

20. Which of the following aqueous solutions remains neutral after electrolysis ?

A. K_2SO_4

B. NaCl

 $C.AgNO_3$

D. CuSO₄

Answer: a

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21. Same quantity of current is passed through molten NaCl and molten cryolite containing Al_2O_3 . If 4.6g of Na was liberated in one cell, the mass of Al liberated in the other cell is

A. 0.9g

B. 1.8g

C. 2.7g

D. 3.6g

Answer: b



22. In the electrolysis of a $40LCuSO_4$ solution, there are two possible reactions at anode :

$$i. Cu \rightarrow Cu^{2+} + 2e^{-}$$

 $ii. 2H_2O \rightarrow 4H^{\oplus} + 4e^- + O_2$

A current of 1.07A is passed for 2 hours. The loss in the mass of Cu at anode was 1.27g.

(Atomic weight of $Cu = 63.5 gmol^1$).

Which of the following statement(s) is / are correct ?

A. 0.08mol of electrons are passed through the solution during entire

electrolysis

B. 224mL of $O_2(g)$ is liberated at STP at anode.

C. Fraction of current in the production of Cu^{2+} ions = 0.5.

D. pH drops to 3

Answer: a,b,c,d

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23. Two platinum electrodes were immersed in a solution of $CuSO_4$ and electric current was passed through the solution. After some time, it was fournd that the colour of $CuSO_4$ disappeared with the evolution of $O_2(g)$ at the electrode.

Which statement is true regarding the resultant solution having pH = X?

A.
$$\left[Cu^{2^+}\right] > \left[SO_4^{2^-}\right]$$

B. $\left[Cu^{\oplus}\right] > \left[Cu^{2^+}\right]$

$$\mathsf{C}. \begin{bmatrix} c^{-} \\ OH \end{bmatrix} > \begin{bmatrix} H^{\oplus} \end{bmatrix}$$
$$\mathsf{D}. 7 > X$$

Answer: d

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24. Aluminium oxide may be electorlysed at 1000 ° C to furnish aluminim metal (Atomic Mass = 27 amu, 1F = 96, 500C). The cathode reaction is $Al^3 + 3d^- \rightarrow Al$ °

To prepare 5.12kg of aluminimu metal by this method woold require .

A. 5.49 × $10^{10}C$

B. $1.83 \times 10^7 C$

C. 5.49 × $10^4 C$

D. 5.49 × $10^{7}C$

Answer: d

25. The density of copper is $8.95gmL^{-1}$. Find out the number of coulombs needed to plate an area of $100cm^2$ to a thickness of $10^{-2}cm$ using $CuSO_4$ solution as electrolyte. (Atomic weight of Cu = 63.5g)

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26. *a*. In the following reactions, what weight of substance would be liberated if 1F of electricity were passed through the cell :

$$i. Cu^{2+} + 2e^- \rightarrow Cu \ ii. Al^{3+} + 3e^- \rightarrow Al$$

iii. $2Cl^{c^-} \rightarrow Cl_2 + 2e^-$ iv. $Ag^{\oplus} + e^- \rightarrow Ag$

b. In the series of cathodes / anodes given above, how many coulombs are needed to produce 1g of each of

A. Cu

B.Al

C. Cl₂

Answer: a. i. 31.75g ii. 9g iii. 35.5g iv. 108g b. i. 3039C ii. 10722.2C iii. 2718.3C iv. 893.5C



1. Arrange the following compounds in the order of increasing conductance : *HCl*, *LiCl*, *NaCl*, *KCl*.

A. LiCl > NaCl > KCl

B. KCl > NaCl > LiCl

C. NaCl > KCl > LiCl

D.LiCl > KCl > NaCl

Answer: B

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2. Which of the following alkali metal ions has the lowest ionic mobility in aqueous solutions?

A. *Li* [⊕] B. *Na* [⊕] C. *K* [⊕]

D. Rb ⊕

Answer: A

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3. For a $0.01MCH_2COOH$ solution, $\Lambda_m = 7.8\Omega^{-1}cm^2mol^{-1}$ if $\Lambda_m^{\circ} = 390\Omega^{-1}cm^2mol^{-1}$. What is the degree of the dissociation (α) of acetic acid ?

A. 0.20

B. 0.48

C. 0.02

D. 0.05

Answer: C

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4. The resistance of 1N solution of acetic acid is 250 *ohm*, when measured in a cell of cell constant 1.15 cm⁻¹. The equivalent conductance (in $ohm^{-1}cm^2eq^{-1}$) of 1N acetic acid is

A. 46.0

B. 9.2

C. 18.4

D. 2.3

Answer: A



5. The equivalent conductance of Ba^{2+} and Cl^{c-} are $63.5ohm^{-1}cm^2eq^{-1}$ and $76ohm^{-1}cm^2eq^{-1}$, respectively, at infinite dilution . The equivalent conductance (in $oh^{-1}cm^2$) of $BaCl_2$ at infinite dilution will be

A. 139.5

B. 203.0

C. 279.0

D. 101.15

Answer: A



6. The quivalnet conductance of any electrolyte MA at infinite dilution

 Λ° . (MA) is equal (more than one correct answer)

A.
$$\wedge \circ \cdot_{(MA)} = \wedge^{c} \cdot_{(MCl)} + \wedge \circ \cdot_{(NaA)} + \wedge \circ \cdot_{(NaCl)}$$

B. $\wedge \circ \cdot_{(MA)} = \wedge^{c} \cdot_{(MCl)} + \wedge \circ \cdot_{(NaA)} - \wedge \circ \cdot_{(NaCl)}$
C. $\wedge \circ \cdot_{(MA)} = \lambda \circ \cdot_{(M^{\oplus})} + \lambda \circ \cdot_{(A^{c} -)}$
D. $\wedge \circ \cdot_{(MA)} = \wedge^{c} \cdot_{(MCl)} + \wedge \circ \cdot_{(NaA)} - \wedge \circ \cdot_{(NaCl)}$

Answer: B,C



7. Which of the following alkali metal ions has the lowest ionic mobility in aqueous solutions?

A. Na ⊕

B.*F^c*-

 $C.SO_4^2$

с-D. ОН

Answer: D

8. $\wedge \circ ._{aq}$ of $BaCl_2, H_2SO_4$, and HCl(aq) solutions are x_1, x_2 , and x_3 , respectviley. $\wedge \circ ._m(BaSO_4)$ is :

A. $x_1 + x_2 - x_3$ B. $x_1 - x_2 - x_3$ C. $x_1 + x_2 - 2x_3$ D. $x_1 - 2x_2 + x_3$

Answer: A

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9. $\Lambda \circ ._m$ of $BaCl_2, H_2SO_4$, and HCl(aq) solutions are x_1, x_2 , and x_3 respectively. $\Lambda_m(BaSO_4)$ is :

A. $x_1 + x_2 - x_3$

B. $x_1 - x_2 - x_3$ C. $x_1 + x_2 - 2x_3$ D. $x_1 - 2x_2 + x_3$

Answer: C

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10. Which of the following statemetn is / are correct ?

I. Van't Hoff factor for $10^{-3}MCH_{3}COOH$ is $39/35(\wedge^{\circ}._{m} = 350Scm^{2}mol^{-1})$ and $k = 4 \times 10^{-5}Scm^{-1})$. *II*. If $\wedge^{\circ}._{m}$ of *KCl*, *LiCl*, and *KNO*₃ are 150, 115, and $145Scm^{2}mol^{-1}$,

then that of $LiNO_3$ will be $110Scm^2mo^{-1}$.

III. In general, with increase in concentration, the specific conductance increases and reaches a maximum and then decreases with further increase in concentration.

A. I, II, and III

B. I and II

C. Only I

D. Only II

Answer: A

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Exercise(Linked Comprehension)

1. An aqueous solution containing $0.1MFe^{3+}$ and $0.01MFe^{2+}$ was titrated with a concentrated solution of *NaOH* at 30 °*C*, so that changes in volumes were negligible. Assuming that the new species formed during titration are *Fe*(*OH*)₃ and *Fe*(*OH*)₂ only.

Given $E^{c} \cdot Fe^{3+} | Fe^{2+} = 0.80V$, $K_{spFe(OH)_3} = 10^{-37}$, and $K_{spFe(OH)_2} = 10^{-19}$ The redox potential of $Fe^{3+} | Fe^{2+}$ electrode at pH = 6 is B. 0.5V

C. 0.2V

 $D.\,0.1V$

Answer: a

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2. An aqueous solution containing $0.1MFe^{3+}$ and $0.01MFe^{2+}$ was titrated with a concentrated solution of *NaOH* at 30 °*C*, so that changes in volumes were negligible. Assuming that the new species formed during titration are *Fe*(*OH*)₃ and *Fe*(*OH*)₂ only.

Given $E^{c} \cdot Fe^{3+} | Fe^{2+} = 0.80V$,

$$K_{spFe(OH)_3} = 10^{-37}$$
, and $K_{spFe(OH)_2} = 10^{-19}$

The redox potential of Fe^{3+} | Fe^{2+} electrode at pH = 6 is

A. 0.8V

 $B.\,0.5V$

C. 0.2V

 $D.\,0.1V$

Answer: C

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3. An aqueous solution containing $0.1MFe^{3+}$ and $0.01MFe^{2+}$ was titrated with a concentrated solution of *NaOH* at 30 °*C*, so that changes in volumes were negligible. Assuming that the new species formed during titration are *Fe*(*OH*)₃ and *Fe*(*OH*)₂ only.

Given E^{c} . $_{Fe^{3+} | Fe^{2+}} = 0.80V$,

 $K_{spFe(OH)_3} = 10^{-37}$, and $K_{spFe(OH)_2} = 10^{-19}$

The redox potential of Fe^{3+} | Fe^{2+} electrode at pH = 6 is

A. 0.8V

 $B.\,0.5V$

C. 0.2V
D.0.1V

Answer: c



4. Calculate
$$\Delta_r G^{c^-}$$
 of the reaction :
 $Ag^{\oplus}(aq) + Cl^{c^-}(aq) \rightarrow AgCl(s)$
Given : $\Delta_f G^{c^-} \cdot AgCl} = -109kJmol^{-1}$
 $\Delta_f G^{c^-} - ((Cl^{c^-})) = -129kJmol^{-1}$
 $\Delta_f G^{c^-} \cdot (Ag^{\oplus}) = -77kJmol^{-1}$
A. $-97kJmol^{-1}$
B. $-57kJmol^{-1}$
C. $57kJmol^{-1}$

D. 97kJmol⁻¹

Answer: b

5. E^{c-} . _{cell} of the reaction above in Question is

A. 0.59V

B.-0.59V

C. 0.295V

D.-0.295V

Answer: a

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6. K_{sp} of AgCl is

A. 10⁻¹³

B. 10⁻¹²

C. 10⁻¹¹

D. 10⁻¹⁰

Answer: d

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7. $6.537 \times 10^{-2} f$ of metallic Zn was added to 100 mL of saturated solution

of
$$AgCl$$
. Calculate log. $\frac{\left[Zn^{2+}\right]}{\left[Ag^{\oplus}\right]}$.
Given $: E^{c^{-}} \cdot_{Ag^{\oplus} \mid Ag} = 0.80V, E^{c^{-}} \cdot_{Zn^{2+} \mid Zn} = -0.763V$.
 K_{sp} of $AgCl \approx 10^{-10}$, atomic weight of $Zn = 65.37$

A. 26.5

B. 13.24

C. 53

D. 106

Answer: c



8. Calculate the number of moless of Ag formed above in Question.

Answer: c

9. The magnitude (but not the sign) of the standard reduction potentials of two metals X and Y are :

$$Y^{2} + 2e^{-} \rightarrow Y \left| E_{1}^{c^{-}} \right| = 0.34V$$
$$X^{2} + 2e^{-} \rightarrow X \left| E_{2}^{c^{-}} \right| = 0.25V$$

When the two half cells of X and Y are connected to construct a cell,

eletrons flow from X to Y. When X is connected to a standard hydrogen electrode (*SHE*),electrons flow from X to *SHE*.

If a half call $X \mid X^2(0.1M)$ is connected to another half cell $Y \mid Y^{2+}(1.0M)$ by means of a salt bridge and an external circuit at 25 ° *C*, the cell voltage would be

A. 0.06V

B.0.12V

C. 0.62V

D. 0.72V

Answer: c

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10. The magnitude (but not the sign) of the standard reduction potentials of two metals X and Y are :

$$Y^{2} + 2e^{-} \rightarrow Y \left| E_{1}^{c^{-}} \right| = 0.34V$$
$$X^{2} + 2e^{-} \rightarrow X \left| E_{2}^{c^{-}} \right| = 0.25V$$

When the two half cells of X and Y are connected to construct a cell, eletrons flow from X to Y. When X is connected to a standard hydrogen electrode (*SHE*),electrons flow from X to *SHE*.

If standard emf (E^{c^-}) of a half cell $Y^2 \mid Y^{\oplus}$ is 0.15V, the standard emf of the half cell $Y^{\oplus} \mid Y$ will be

A. 0.19V

 $B.\,0.53V$

C. 0.49V

D. 0.64V

Answer: b

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11. The magnitude (but not the sign) of the standard reduction potentials of two metals X and Y are :

$$Y^2 + 2e^- \rightarrow Y \left| E_1^{c^-} \right| = 0.34V$$

$$X^2 + 2e^- \rightarrow X \left| E_2^{c^-} \right| = 0.25V$$

When the two half cells of X and Y are connected to construct a cell, eletrons flow from X to Y. When X is connected to a standard hydrogen electrode (*SHE*),electrons flow from X to *SHE*.

Given the following half cell : $YI + e^- \rightarrow Y - I^{c^-}$: $E^{c^-} = -0.27V$

Solubility product of the iodide salt YI is

A. 2×10^{-3}

B. 2×10^{-12}

 $C.2 \times 10^{-14}$

D. 6.8×10^{-16}

Answer: c

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12. A volataic cell consists of an electode of solide silver immerse in a $0.10MAgNO_3$ solution and an electrode of unknown metal 'X' immersed in a 0.10M solution $X(NO_3)_2$. A porous barrier separates the two half of

the cell. Also given :

$$E^{C^{-}} \cdot \left(Ag^{\oplus} \mid Ag\right) = 0.80V \text{ and } E^{C^{-}} \cdot_{cell} = 1.05V \text{ at } 25 \circ C$$
$$X(s) \left| X^{2^{+}}(1.0M) \right| \left| Ag^{\oplus}(0.1M) \right| Ag(s)$$

Which of the following statements regarding the cell and X is incorrect?

A. Standard
$$EMF(E^{c-})$$
 of $X^{2+} \mid X$ is -0.25V at 25 °C.

- B. X is the stronger reducing agent than $H_2(g)$.
- C. As the cell operates , the concentration of both X^2 and Ag^{\oplus} increase in their respective half cells.
- D. As the cell operates, the concentration of X^{2+} increases in the anode chamber while the concentration of Ag^{\oplus} decreases in the cathode chamber.

Answer: c



13. A volataic cell consists of an electode of solide silver immerse in a $0.10MAgNO_3$ solution and an electrode of unknown metal 'X' immersed in a 0.10M solution $X(NO_3)_2$. A porous barrier separates the two half of the cell. Also given :

$$E^{c^{-}} \cdot \left(Ag^{\oplus} \mid Ag\right) = 0.80V \text{ and } E^{c^{-}} \cdot_{cell} = 1.05V \text{ at } 25 \circ C$$
$$X(s) \left| X^{2^{+}} (1.0M) \right| \left| Ag^{\oplus} (0.1M) \right| Ag(s)$$

If $Ag \oplus |Ag$ half cell in the above voltaic cell is replaced by $Zn^{2+} | Zn$ half cell $\left(E^{c-} \cdot Zn^{2+} | Zn = -0.76V\right)$

A. The direction of current flow will remain same.

B. Polarity of the electrodes will be reversed.

C. Cell will stop working.

D. EMF of the cell will increase.

Answer: b

14. Fuel cells : Fuel cells are galvanic cells in which the chemical energy of fuel cell is directly converted into electrical energy. A type of fuel cell is a hydrogen - oxygen fuel cell. It consists of two electrodes made up of two porous graphite impregnated with a catalyst (platinum, silver, or metal oxide). The electrodes are placed in aqueous solution of *NaOH*. Oxygen and hydrogen are continuously fed into the cell. Hydrogen gets oxidized to H^{\oplus} which is neutralized by *OH*, *i. e.*, anodic reaction.

 $H_{2} \rightleftharpoons 2H^{\oplus} + 2e^{-}$ $2H^{\oplus} + 2\overset{\odot}{O}H \rightleftharpoons 2H_{2}O$ $H_{2} + 2\overset{\odot}{O}H \rightleftharpoons 2H_{2}O + 2e^{-}$ At cathode, O₂ gets reduced to $\overset{\odot}{O}H$ i.e., O₂ + 2H₂O + 4e⁻ \implies 4\overset{\odot}{O}H Hence, the net reaction is $2H_{2} + O_{2} \rightleftharpoons 2H_{2}O$

At cathode, O_2 gets reduced to OH

 $O_2 + 2H_2O + 4e^- \implies 4OH \text{ has } E^{\odot} = 0.40 \text{ V},$ then E^{\odot} for $2H_2O + 2e^- \implies H_2 + 2OH \text{ will be}$ **a.** 0.41 V **b.** 0.83 V **c.** -0.41 V **d.** -0.83 V

с-

Hence, the net reaction is

The overall reaction has

 $\Delta H = -285.6 k Jmol^{-1}$ and $\Delta G = -237.4 k Jmol^{-1}$ at 25 ° C

If the cell voltage is 1.23V for the H_2 - O_2 fuel cell and for the half cell :

A. 0.41V

B. 0.83V

C. -0.41V

D. 0.83V

Answer: d

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15. Fuel cells : Fuel cells are galvanic cells in which the chemical energy of fuel cell is directly converted into electrical energy. A type of fuel cell is a hydrogen - oxygen fuel cell. It consists of two electrodes made up of two porous graphite impregnated with a catalyst (platinum, silver, or metal oxide). The electrodes are placed in aqueous solution of *NaOH*. Oxygen and hydrogen are continuously fed into the cell. Hydrogen gets oxidized to H^{\oplus} which is neutralized by *OH*, *i. e.*, anodic reaction.

H₂ \implies 2H[⊕] + 2e⁻ 2H[⊕] + 2OH \implies 2H₂O H₂ + 2OH \implies 2H₂O + 2e⁻ At cathode, O₂ gets reduced to OH i.e., O₂ + 2H₂O + 4e⁻ \implies 4OH Hence, the net reaction is 2H₂ + O₂ \implies 2H₂O At cathode, O₂ gets reduced to OH Hence, the net reaction is The overall reaction has $\Delta H = -285.6 k Jmol^{-1}$ and $\Delta G = -237.4 k Jmol^{-1}$ at 25 °C

What is the value of ΔS^{c-} for the fuel cell at 25 ° C?

A. - 1600*JK*⁻¹

B. - 160*JK*⁻¹

C. 160*JK*⁻¹

D. 1600*JK*⁻¹

Answer: b

16. Fuel cells : Fuel cells are galvanic cells in which the chemical energy of fuel cell is directly converted into electrical energy. A type of fuel cell is a hydrogen - oxygen fuel cell. It consists of two electrodes made up of two porous graphite impregnated with a catalyst (platinum, silver, or metal oxide). The electrodes are placed in aqueous solution of *NaOH*. Oxygen and hydrogen are continuously fed into the cell. Hydrogen gets oxidized to H^{\oplus} which is neutralized by *OH*, *i. e.*, anodic reaction.

 $H_{2} \rightleftharpoons 2H^{\oplus} + 2e^{-}$ $2H^{\oplus} + 2\overset{\odot}{O}H \rightleftharpoons 2H_{2}O$ $H_{2} + 2\overset{\odot}{O}H \rightleftharpoons 2H_{2}O + 2e^{-}$ At cathode, O₂ gets reduced to $\overset{\odot}{O}H$ i.e., O₂ + 2H₂O + 4e⁻ \implies 4\overset{\odot}{O}H Hence, the net reaction is $2H_{2} + O_{2} \rightleftharpoons 2H_{2}O$

с-

At cathode, O_2 gets reduced to OH

Hence, the net reaction is

The overall reaction has

 $\Delta H = -285.6 k Jmol^{-1}$ and $\Delta G = -237.4 k Jmol^{-1}$ at 25 ° C

Suppose the concentration of hydroxide ioin in the cell is doubled, then

the cell voltage will be

A. Reduced by half

B. Increased by a factor of 2

C. Increased by a factor of 4

D. Unchanged

Answer: d



17. Fuel cells : Fuel cells are galvanic cells in which the chemical energy of fuel cell is directly converted into electrical energy. A type of fuel cell is a hydrogen - oxygen fuel cell. It consists of two electrodes made up of two porous graphite impregnated with a catalyst (platinum, silver, or metal oxide). The electrodes are placed in aqueous solution of *NaOH*. Oxygen and hydrogen are continuously fed into the cell. Hydrogen gets oxidized to H^{\oplus} which is neutralized by *OH*, *i. e.*, anodic reaction.

 $H_{2} \rightleftharpoons 2H^{\oplus} + 2e^{-}$ $2H^{\oplus} + 2OH \rightleftharpoons 2H_{2}O$ $H_{2} + 2OH \rightleftharpoons 2H_{2}O + 2e^{-}$ At cathode, O_{2} gets reduced to OHi.e., $O_{2} + 2H_{2}O + 4e^{-} \rightleftharpoons 4OH$ Hence, the net reaction is $2H_{2} + O_{2} \rightleftharpoons 2H_{2}O$ At cathode, O_{2} gets reduced to OHHence, the net reaction is The overall reaction has $\Delta H = -285.6kJmol^{-1}$ and $\Delta G = -237.4kJmol^{-1}$ at $25 \circ C$

A fuel cell is

I. A voltaic cell in which continuous supply of fuels are sent at anode to perform oxidation.

II. A voltaci cell in which fuels such as $: CH_4, H_2$, and CO are used up at anode.

III. One which involves the reaction of H_2 - O_2 fuel cell such as :

Anode : $2H_2O + 4OH \rightarrow 4H_2O(l) + 4e^{c}$

Cathode : $O_2 + 2H_2O(l) = 4e^- \rightarrow 4OH$

IV. The efficiency of H_2 - O_{20} fuel cell is 70 to 75 %

A. I, III

B. I, III, IV

C. I, II, III, IV

D. I, II, III

Answer: c



18. A conductivity cell is used to measure the conductance of electrolyte .It makes use of conductivity of water which does not contain any ions.The cell constant of conductivity cell is determined.

If the cell constant is 0.40cm⁻¹, the conductivity of 0.051MNaCl solution

having R = 1850ohm is equal to

```
A. 1.08 \times 10^{-4} Scm^{-1}
```

B. $4.32 \times 10^{-4} Scm^{-1}$

C. 2.16 × 10^{-4} Scm⁻¹

D. $5.04 \times 10^{-5} Scm^{-1}$

Answer: c

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19. A conductivity cell is used to measure the conductance of electrolyte . It makes use of conductivity of water which does not contain any ions. The cell constant of conductivity cell is determined. Calculate α of CH_3COOH if $\Delta^{\infty} \cdot_m$ for $HCl, NaCl, CH_3COOHNa$ are

426, 126, $91Scm^2mol^{-1}$, respectively, and $\Lambda_m = 14.4Scm^2mol^{-1}$ at 0.015M concentration.

A. 0.037

B. 0.018

C. 0.37

D. 0.18

Answer: a

 $3CH_3CH_2OH(aq)Ehanol + 2Cr_2O_7^{2-}(aq) + 16H^{\oplus}(aq) \rightarrow 3CH_3COH(aq)Ethanoicacian$ The breathalyzer measures the colour change and produces a metrereading calibrated in the terms of blood alcohol content.

If $E^{c^-} \cdot CH_3COOH \mid C_2H_5OH = 0.06V$ and $E^{c^-} \cdot Cr_2O_7^{2^-} \mid Cr^{3^+} = 1.33V$,

then E^{c-}._{cell} of the reaction taking place in alcohol metre is

A. 1.39V

 $B.\,1.27V$

C. - 1.39V

D. - 1.51V

Answer: b

 $3CH_3CH_2OH(aq)Ehanol + 2Cr_2O_7^{2-}(aq) + 16H^{\oplus}(aq) \rightarrow 3CH_3COH(aq)Ethanoicacide$ The breathalyzer measures the colour change and produces a metrereading calibrated in the terms of blood alcohol content.

Colour of the testing solution changes from

A. Orange to green

B. Colourless to green

C. Orange to green

D. Yellow to blue

Answer: c

 $3CH_3CH_2OH(aq)Ehanol + 2Cr_2O_7^{2-}(aq) + 16H^{\oplus}(aq) \rightarrow 3CH_3COH(aq)Ethanoicacide$ The breathalyzer measures the colour change and produces a metrereading calibrated in the terms of blood alcohol content.

The *EMF* of the reaction when the concentration of all the species are 1.0M and *pH* is 4.0 is

A. 1.64

B. 0.31

C. - 1.01V

D. 0.95V

Answer: d

 $3CH_3CH_2OH(aq)Ehanol + 2Cr_2O_7^{2-}(aq) + 16H^{\oplus}(aq) \rightarrow 3CH_3COH(aq)Ethanoicacide$ The breathalyzer measures the colour change and produces a metrereading calibrated in the terms of blood alcohol content.

What is the ethanol ethanoic acid ratio if the breathalyzer records 1.33Vand other species are at 1M?

A. 10⁴

B. 10⁻⁴

C. 10¹²

D. 10⁻¹²

Answer: a

24. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



A. I

B. *I*, *II*

C. *I*, *II*, *III*

D. II, III, IV

Answer: b

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25. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



Given : $Co^{2^+} + 2e^- \rightarrow Co \quad E^{c^-} = -0.28V$ Ag^(o+)+e^(-) rarr Ag" "E^(c-)=0.80V2Ag^(o+)+Corarr Co^(2+)+2Ag" "E^(c-). (cell)=1.08VF $\in ally, thell`$ compartment contains

A. I

B. *I*, *II*,

C. *I*, *II*, *III*

D. II, III, IV

Answer: c

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26. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



A. I

B. *I*, *II*,

C. I, II, III

D. II, III, IV

Answer: d

27. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



A. 0.105, 0.005, 0.0M

B. 0.005, 0.105, 0.0M

C. 0.105, 0.0, 0.005M

D. 0.0,0.005,0.105M`

Answer: a

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28. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



Given : $Co^{2^+} + 2e^- \rightarrow Co \quad E^{c^-} = -0.28V$

Ag^(o+)+e^(-) rarr Ag" "E^(c-)=0.80V2Ag^(o+)+Corarr Co^(2+)+2Ag"

"E^(c-)._(cell)=1.08VThef \in alconcentrationofNO_(3)^(c-) \in I,II, and III`

compartment is

A. 0.100, 0. .210, 0.0900M

B. 0.210, 0.100, 0.0900M

C. 0.0900, 0.210, 0.100M

D. 0.900, 0.100, 0.210M

Answer: b

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29. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by

passign 0.1F of electricity.



A. 0.090, 0.0, 0.0100M

B. 0.0, 0.0100, 0.090M

C. 0.0, 0.090, 0.0100M

D. 0.100, 0.90, 0.0M

Answer: c

30. A cell, as shown below, consists of there compartments separated by porous pots. The first contains a cobalt eletrode in 5.0*L* of $0.10MCo(NO_3)$, the second contains 5.0*L* of $0.10MKNO_3$, the third contains an *Ag* electrode in 5.0*L* of $0.10MAgNO_3$. Assuming that current with in the cell iis carried equally by the negative and positive ions by passign 0.1F of electricity.



A. 0.0, 0.0, 0.08M

B. 0.0, 0.08, 0.0M

C. 0.08, 0.0, 0.0M

D. 0.0M in all compartments

Answer: a

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Exercisemultiple Correct Ansers

1. Consider the following concentration cell:

$$Zn(s) \left| Zn^{2+}(0.024M) \right| \left| Zn^{2+}(0.480M) \right| Zn(s)$$

which of the following statements is / are correct?

A. The *EMF* of the cell at 25 $^{\circ}C$ is nearly +0.039V.

B. The *EMF* of the cell at $25 \degree C$ is nearly -0.039V.

C. If water is added in *LHE*, so that the $\left[Zn^{2+}\right]$ is reduced to 0.012*M*,

the cell voltage increases.

D. If water is added in LH, so that the $\left[Zn^{2+}\right]$ is reduced to 0.012M,

the cell voltage decreases.

Answer: a,c

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2. Given :

Oxidation $H_2O_2 \rightarrow O_2 + 2H^{\oplus} + 2e^- E^{c^-} = -0.69V$, $2F^{c^-} \rightarrow F_2 + 2e^- E^{c^-} = -287V$, Reduction $:H_2O_2 + 2H^{\oplus} + 2e^- \rightarrow 2H_2O E^{c^-} = 1.77V$, $2I^(c_-)rarr I_(2)+2e^(-)" "E^(c_-)=-0.54V$, Which of the follow $\in gstatements is //^{\sim}$ are correct ?

A. H_2o_2 behaves as an oxidant for I^{c-1}

B. H_2O_2 behaves as a reductant for $1I_2$

C. H_2O_2 behaves as an oxidant for F^{C-}

D. H_2O_2 behaves as a reductant for F_2 .

Answer: a,d

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3. Peroxodisulphate salts, $(e. g., Na_2S_2O_8)$ are strong oxidizing agents used be bleaching agents for fats, oils, ets. Given : $O_2(g) + 4H^{\oplus}(aq) + 4e^- \rightarrow 2H_2O(l) \quad E^{c-} = 1.23V$

 $S_2O_8^{2^-}(aq) + 2e^- \rightarrow 2SO_4^{2^-}(aq) \quad E^{C^-} = 2.01V$

Which of the following statement is (are) correct ?

A. Oxygen gas can oxidize sulphate ion to per - oxo disulphate ion

$$\left(S_2 O_8^{2^-}\right)$$
 in acidic solution.

- B. $O_2(g)$ is reduced to water.
- C. Water is oxidized to O_2
- D. $S_2O_8^{2-}$ ions are reduced to SO_4^{2-} ions.

Answer: c,d

4. A current is passed through 500mL of an aqueous solution of CaI_2 . After sometime, it is observed that 50 millimoles of I_2 have been formed. Which of the following statements is (are) correct ?

A. The number of faradays of charge passed through the solution is

0.10*F*.

- B. The volume of dry H_2 at *STP* that has been formed during electrolysis is 1120mL.
- C. The pH of the solution is nearly 0.7
- D. The mass of calcium produced is 2.0g.

Answer: a,b

5. Which of the following statements is / are correct ?

A. The cell constant of an electrolytic cell is measured as the product

: *kR* rather using *l*/*a*.

B. As an electrolytic solution is diluted, its conductance, equivalent

conductance, and molar conductance increase.

C. Kohlraushc's law may be applied to calculate molar conductance at

infinite dilution for both weak and strong electrolytes.

D. Kohlrausch's law may also be applied at any concentrated of the

electrolyte.

Answer: a,b,c



6. Predict which of the following reactions would proceed spontaneously

at 298K?

A.
$$Co(s) + Fe^{2+}(aq) \rightarrow Co^{2+}(aq) + Fe(s)$$

 $E^{c-} \cdot Co^{2+}/Co} = -0.28V$
B. $Cd^{2+}(aq) + Fe(s) \rightarrow Cd(s) + Fe^{2+}(aq)$
 $E^{c-} \cdot Cd^{2+}/Cd} = -0.4V$
C. $Cd(s) + Co^{2+}(aq) \rightarrow Cd^{2+}(aq) + Co(s)$
 $E^{c-} \cdot Fe^{2+}/Fe} = -0.44V$
D. $Zn^{2+}(aq) + H_2 \rightarrow Zn(s) + 2H^{\oplus}$
 $E^{c-} \cdot Zn^{2+}/Zn} = -0.76V$

Answer: b,c

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7. The gas X at 1*atm* is bubbled through a solution containing a mixture of $1MY^{c-}$ and $1MZ^{c-}$ at 25 °C. If the order of reduction potential is Z > Y > X, then

A. Y will oxidize X only
B. *Y* will oxidize *Z* only

- C. Z will oxidize X and Y
- D. Z will reduce both X and Y

Answer: a,c

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8. During the electrolysis of aqueous zinc nitrate.

A. Zince plates out at the cathode

B. Zince plates out at the anode

C. Hydrogen gen H_2 is evolved at the anode.

D. Oxygen gas O_2 is evolved at anode

Answer: a,d

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9. Which of the following changes will increase the *EMF* of the cell : $Co(s) \left| CoCl_2(M_2) \right| \left| HCl(M_2) \right| + (H_2, g) Pt$

A. Increase the volume of $CoCl_2$ from 100mL to 200mL

B. Increase M_2 from 0.1M to 0.50M.

C. Increase the pressure of the $H_2(g)$ from 1.0 to 2.0*atm*.

D. Increase M_1 from 0.01M to 0.50M.

Answer: a,b

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10. Given :

 $E^{c} \cdot A_{g} \oplus A_{g} = 0.80V, E^{c} \cdot A_{g}^{2+} A_{g} = -2.37V,$

 $E^{c} \cdot Cu^{2+} \mid Cu = 0.34V, E^{c} \cdot Hg^{2+} \mid Hg = 0.79V$

Which of the following statements is / are incorrect ?

A. $AgNO_3$ can be stored in copper vessel.

B. $Cu(NO_3)_2$ can be stored in copper vessel.

C. $CuCl_2$ can be stored in silver vessel.

D. $HgCl_2$ can be stored in copper vessel.

Answer: a,b,d

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11. Iron can be prevented from rusting by

A. Connecting iron to more electropositive metal - a case of cathodic

protection

B. Connecting iron to more electropositive metal - a case of anodic

protection.

C. Connecting iron to less electropositive metal - a case of anodic

protection

D. Connecting iron to less electropositive metal - a case of cathodic

protection.

Answer: a,c

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12. 100mL of buffer of $1MNH_3(aq)$ and $1MNH_4^{\oplus}(aq)$ are placed in two compartments of a voltaic cell separately. A current of 1.5A is passed through both cells for 20min. If only electrolysis of water takes place, then

A. pH of LHE half cell will increase

B. pH of RHE half cell will increase

C. pH of both half cell will increase

D. *pH* of both half cell will decrease

Answer: B

13. In the following electrochemical cell :

$$Zn \left| Zn^{2+} \right| \left| H^{\oplus} \right| \left(H_2 \right) Pt$$

$$E_{cell} = E^{c-} \cdot_{cell}. \text{ This will be when}$$

$$A. \left[Zn^{2+} \right] = \left[H^{\oplus} \right] = 1M \text{ and } p_{H_2} = 1atm$$

$$B. \left[Zn^{2+} \right] = 0.01M, \left[H^{\oplus} \right] = 0.1M, \text{ and } p_{H_2} = 1atm$$

$$C. \left[Zn^{2+} \right] = 1M, \left[H^{\oplus} \right] = 0.1M, \text{ and } p_{H_2} = 1atm$$

D. None of the above.

Answer: a,b

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14. For the electrochemical cell,
$$(M | M^{\oplus}) | | (X^{c-} | X),$$

 $E^{c-} \cdot (M^{\oplus} | M) = 0.44V$, and $E^{c-} \cdot (X | X^{c-}) = -0.33V^{\bullet}$
From this data, one can conclude that

A. $M + X \rightarrow M^{\oplus} + X^{c-}$ is a spontaneous reaction

B. $M^{\oplus} + X^{c-} \rightarrow M + x$ is the spontaneous reaction

$$C. E^{c-}._{cell} = 0.77V$$

D.
$$E^{c-}$$
. cell = -0.77V

Answer: b,c

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15. For a strong electrolyte, equivalent conductance increases slowly with dilution and can be expressed by the relationship: $\Lambda_m = \Lambda^{\circ} \cdot_m - A\sqrt{c}$ Which electrolyte (s) have same value of *A* ?

A. NaCl

B. $CaCl_2$

C. ZnCl₂

D. $MgSO_4$

Answer: b,c



16. During electrolysis, $O_2(g)$ is evolved at anode in

A. Dilute H_2SO_4 with Pt electrode

B. Aqueous AgNO₃ with Pt electrode

C. Dilute H_2SO_4 with Cu electrode

D. Fused NaOH with an Fe cathode and Ni anode

Answer: a,b

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17. During electrolysis of aqueous CuBr₂ using Pt electrode,

- A. $Br_2(g)$ is evolved at anode
- B. Cu(s) is deposited at cathdoe

C. $Br_2(g)$ is evolved at anode and $H_2(g)$ at cathode

D. $H_2(g)$ is evolved at anode

Answer: a,b



18. A current of 2.68A is passed for 1.0 hour through an aqueous solution

of CuSO₄ using copper electrodes.

Which of the following statements is / are correct ?

A. Increase in the mass of cathode = 3.174g

B. Decrease in the mass of anode = 3.174g

C. No change in the mass of electrodes

D. The ration between the change in the mass of cathode to anode is

Answer: a,b

^{1:2}

19. The *EMF* of the following cell :

 $Cd(s) \left| CdCl_2(0.10M) \right| AgCl(s) \mid Ag(s)$ is 0.6915V at 0 °C and 0.6753V at 25 °C. The ΔH of reaction in kJ at 25 °C is

A. - 176

B. - 234.7

C. 123.5

D. - 167.6

Answer: d

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20. When 4.0A of current is passed through a 1.0L, $0.10MFe^{3+}(aq)$ solution for 1.0 hour, it is partly reduced to Fe(s) and partly of $Fe^{2+}(aq)$. The correct statements (s) is (are):

A. 0.10*mol* of electrons are required to convert all Fe^{3+} to Fe^{2+}

B. 0.025mol of Fe(s) will be deposited.

C. 0.075*mol* of iron remains as Fe^{2+} .

D. 0.050*mol* of iron remains as Fe^{2+}

Answer: a,b,c

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21. In which of the following cells, *EMF* is greater than E^{c-} .

A.
$$Pt, H_2(g) | H^{\oplus}(pH = 5) | | H^{\oplus}(pH = 3) | | H_2(g), Pt$$

B. $Zn(s) | Zn^{2+}(0.2M) | | Cu^{2+}(0.1M) | Cu(s)$
C. $Cr(s) | Cr^{3+}(0.1M) | | Cu^{2+}(0.2M) | Cu(s)$
D. $Pt, H_2(g) | H^{\oplus}(pH = 4) | | H^{\oplus}(pH = 6) | H_2(g) | Pt$

Answer: a

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22. In the atmosphere of industrial smog, copper corrodes to form

A. Basic copper carbonate

B. Copper sulphide

C. Basic copper sulphate

D. Copper oxide

Answer: a,c

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23. The tarnishing of silver ornaments in atmosphere is due to

A. Ag_2O

 $B.Ag_2S$

 $C.Ag_2CO_3$

 $D.Ag_2SO_4$

Answer: a,b



24. If
$$A + B \rightleftharpoons C + D$$
; $K_C = K_1$ and $E^{\textcircled{o}} = a \vee A$
24. If $2A + 2B \rightleftharpoons 2C + 2D$; $K_C = K_2$ and $E^{\textcircled{o}} = b \vee A$
then
A. $a = b$
B. $K_2 = K_1^2$
C. $a = 2b$

Answer: a,b

D. $b = a^2$

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25. Rusting of iron is catalyzed by

A. H ⊕

- B. Dissolved CO_2 in water
- C. O₂
- D. Impurities present in Fe

Answer: a,b,c,d

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26. Select the wrong relation (s).

A.
$$\Delta S = \left(\frac{\partial E}{\partial T}\right)_P \times nF$$

B. $-\Delta S = \left(\frac{\partial E}{\partial T}\right)_P \times nF$
C. $\left(\frac{\partial E}{\partial T}\right)_P = \left(\frac{\partial \Delta S}{\partial T}\right)$
D. $\left(\frac{\partial E}{\partial T}\right)_P = \frac{\Delta H + nEF}{T}$

Answer: a,d



27. Select the correct statements (*s*) about *NHE*.

A. E^{c-} of *SHE* is arbitrarily assumed to be zero.

B. E^{c} of SHE is equal to zero.

C. *SHE* refers as *Pt*, $H_2(g)$ _{1bar} | $H^{\oplus}(aq)a = 1$ at 25 °*C*.

D. SHE is very susceptible to dissolved O_2 , H_{20S} and all other reducing

agents.

Answer: a,c,d

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28. In which of the following salt bridge is not needed ?

A.
$$Pb \left| PbSO_4(s) \right| H_2SO_4 \left| PbO_2(s) \right| Pb$$

B. $Cd|CdO(s)|KOH(aq)|NiO_2(s)|Ni$

C. $Fe(s)|FeO(s)|KOH(aq), Ni_2O_3(s) | Ni$

D. Zn
$$ZnSO_4$$
 CuSO₄ | Cu

Answer: a,b,c

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29. Select the correct statements if 9.65A current is passed for 1 hour through the cell :

 $\begin{array}{l} Ag \left| Ag^{\oplus}(1M) \right| \left| Cu^{2+}(1M) \right| Cu. \\ \\ A. Ag \text{ will oxidize to } Ag^{\oplus} \text{ and new } \left[Ag^{\oplus} \right] = 1.36M. \\ \\ B. Ag^{\oplus} \text{ will reduce to } Ag \text{ and new } \left[Ag^{\oplus} \right] = 0.64M \\ \\ C. Cu^{2+} \text{ will reduce to } Cu \text{ and new } \left[Cu^{2+} \right] = 0.82M. \\ \\ D. Cu \text{ will oxidize to } Cu^{2+} \text{ and new } \left[Cu^{2+} \right] = 0.82M. \end{array}$

Answer: a,c

30. The temperature coefficient of the cell is $\left(\frac{\partial E}{\partial T}\right)_P$. Choose the correct

statements (s).

A. When
$$\left(\frac{\partial E}{\partial T}\right)_{P} = 0$$
, then $\Delta H = -nFE$
B. When $\left(\frac{\partial E}{\partial T}\right)_{P} < 0$, then $|nFE| > |\Delta H|$
C. When $\left(\frac{\partial E}{\partial T}\right)_{P} > 0$, then $|nFE| < |\Delta H|$ Exothermic reaction
D. When $\left(\frac{\partial E}{\partial T}\right)_{P} = 0$, then $|\Delta H| > |nFE|$ Endothermic reaction.

Answer: a,b,c

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31. During discharging of alead storage battery

A. The reaction at anode is $Pb \rightarrow Pb^{2+} + 2e^{-}$

B. The reaction taking place at cathode is

 $Pb^{2^+} + 2e^- \rightarrow Pb$

C. The overall reaction is

 $Pb + PbO_2 + 4H^{\oplus} + 2SO_4^{2-} \rightarrow 2PbSO_4 + 2H_2O$

D. The reaction taking place at cathode is

 $PbO_2 + 4H^{\oplus} + 2e^- \rightarrow Pb^{2+} + 2H_2O$

Answer: a,c,d

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32. Which of the following statements is / are correct ?

A. F_2 is the strongest oxidizing agent. F

B. Li is the strongest reducing agent.

C. Li^{\oplus} is the weakest oxidizing agent.

D. F_2 has a highest reduction potential.



33. Identify the correct statements (*s*):

- A. Λ_m increases with increase in temperature.
- B. Λ_m decreases with increase in concentration.
- C. Specific conductance increase with increase in concentration.
- D. Specific conductance decreases with increse in temperature.

Answer: a,b,c



34. Which of the following cells is / are rechargeable or secondary cell (s)

A. Ni - Cd cell

B. Mercury cell

C. Lead storage cell

D. Lithium battery

Answer: a,c,d

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35. For $I_2 + 2e^- \rightarrow 2I^{c^-}$, standard reduction potential = + 0.54V. For $2Br^{c^-} \rightarrow Br_2 + 2e^-$, standard oxidation potential = - 1.09V. For $Fe \rightarrow Fe^{2^+} + 2e^-$, standard oxidation potential = + 0.44V. Which of the following reactions is (are) spontaneous ?

A. $Br_2 + 2I^{c-} \rightarrow 2Br^{c-} + I_2$

 $B. Fe + Br_2 \rightarrow Fe^{2+} + 2Br^{C-}$

 $\mathsf{C}. Fe + I_2 \rightarrow Fe^{2+} + 2I^{C-}$

 $\mathsf{D}.\,I_2 + 2Br^{c-} \rightarrow 2I^{c-} + Br_2$

Answer: a,b,c



36. Consider the cell :

$$Pt\Big|H_2\Big(p_1atm\Big)\Big|H^{\oplus}\Big(x_1M\Big)\Big|\Big|H^{\oplus}\Big(x_2M\Big)\Big|H_2\Big(p_2atm\Big)Pt.$$

The cell reaction be spontaneous if

A.
$$p_1 = p_2$$
 and $x_1 < x_2$

B.
$$p_1 = p_2$$
 and $x_1 < x_2$

C.
$$x_1 = x_2$$
 and $p_1 > p_2$

D.
$$x_1 = x_2$$
 and $p_1 < p_2$

Answer: b,c

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37. Which of the following changes will cause the free energy of a cell reaction to decrease ?

$$Zn \left| ZnSO_4(aq) \left(x_1 M \right) \right| \left| HCl(aq) \left(x_2 M \right) \right| H_2(g), Pt.$$

A. Increase in the volume of *HCl* solution from 100*mL* to200*mL*

B. Increase in the pressure of hydrogen from 1atm to 2atm

C. Increase in molarity x_2 from 0.1 to 1M

D. Increase in molarity x_1 from 1*M* to 0.1*M*.

Answer: c,d

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38. During the working of a galvanic cell and with the passage of time.

A. Spontaneity of the cell reaction decreases, E_{cell} decreases

B. Reaction quotient Q decreases, E_{cell} increases

C. Reaction quotient Q increases, E_{cell} decreases

D. At equilibrium, $Q = K_{eq}$, $E_{cell} = 0$

Answer: a,c,d



39. In the following electrochemical cell :

$$Zn \left| Zn^{2+} \right| \left| H^{\oplus} \right| (H_2) Pt$$

$$E_{cell} = E^{c-} \cdot_{cell}.$$
 This will be when
$$A. \left[Zn^{2+} \right] = \left[H^{\oplus} \right] = 1M \text{ and } p_{H_2} 1atm$$

$$B. \left[Zn^{2+} \right] = 0.01M, \left[H^{\oplus} \right] = 0.1M, \text{ and } p_{H_2} = 1atm$$

$$C. \left[Zn^{2+} \right] = 1m, \left[H^{\oplus} \right] = 0.1M, \text{ and } p_{H_2} = 0.01atm$$

$$D. \left[Zn^{2+} \right] = \left[H^{\oplus} \right] = 0.1M \text{ and } p_{H_2} = 0.1atm$$

Answer: a,b,c,d

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40. Consider the cell :

$$Cd(s) \left| Cd^{2+}(1.0M) \right| \left| Cu^{2+}(1.0M) \right| Cu(s)$$

If we wish to make a cell with a more positive voltage using the same substances, we should

A. Increase both
$$[Cd^{2+}]$$
 and $[Cu^{2+}]$ to 2.0*M*
B. Decrease the $[Cd^{2+}$ to 1.0*m*
C. Increase the $[Cu^{2+}]$ to 2.0*M*
D. Decrease both the $[Cd^{2+}]$ and $[Cu^{2+}]$ to 0.01*M*

Answer: b,c

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41. Electrolysis of aqueous solutions of which of the following substances

results in only the decomposition of water ?

A. Potassium chloride

B. Zinc sulphate

C. Potassium hydroxide

D. Sodium phosphate

Answer: c,d

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42. When an aqueous solution of $CaCl_2$ is electrolyzed using inert electrods, which of the following is (are) true ?

A. Calcium deposites on cathode.

B. Calcium deposits an anode

C. Chloride is liberated on anode

D. Calcium hydroxide precipitates near cathode on prolonged

Answer: c,d

hydrolysis

43. On passing $0.5m \infty l$ of electrons through $CuSO_4$ and $Hg_2(NO_3)_2$ solutions in series using inert electrodes

A. 0.5mol of Cu is deposited

B. 0.5mol of Hg is deposited

C. 0.125*mol* of O_2 is produced

D. 0.5mol of O_2 is produced

Answer: b,c

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44. Which of the following statements is / are correct ?

A. The electrolysis of concentrated H_2SO_4 at $0-5^{\circ}C$ using a Pt

electrode produces $h_2S_2O_8$.

B. The electrolysis of a brine solution produces *NaClO*₃ and *NaClO*.

C. The electrolysis of $CuSO_4$ solution using Pt electrodes causes the

liberation of O_2 at anode and the deposition of copper at cathode.

D. All electrolytic reactions are redox reactions.

Answer: a,c,d

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45. If same quantity of electricity is passed through three electrolytic cells containing $FeSO_4$, $Fe_2(SO_4)_3$, and $Fe(NO_3)_3$, then

A. The amount of iron deposited in $FeSO_4$ and $Fe_2(SO_4)_3$ are equal.

B. The amount of iron deposited in $FeSO_4$ is 1.5 times of the amount

of iron deposited in $Fe(NO_3)_3$

C. The amount of iron deposited in $Fe_2(SO_4 - (3) \text{ and } Fe(NO_3)_2$ are equal.

D. The same amount of gas is evolved in all three cases of the anode.

Answer: b,c,d



46. Which of the following aqueous solutions remain alkaline after electrolysis ?

A. CH₃COONa

B. KNO₃

C. NaCl

D. LiF

Answer: a,c

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47. A solution containing Na^{\oplus} , $NO_3^{c^-}$, Cl^{c^-} , and $SO_4^{2^-}$ ions, all at unit concentrations, is electrolyzed between nickel anode and platinum

cathode. As the current is passed through the cell

A. Ph of the cathode increases

B. Oxygen is the major product at anode

C. Nickel is deposited at cathode.

D. Chlorine is the major product at anode.

Answer: a,d



48. To obeserve the effect of concentration on the conductivity, electrolytes of different natures are taken in two vessels A and B, A contains weak electrolyte, $e.g., NH_4OH$ and B contains strong electrolyte, e.g., NaCl. In both containers, the concentration of respective electrolyte is increased and the conductivity observed :

A. In A conductivity increases, in B conductivity decrease

B. In A conductivity decreases while, in B conductivity decrease

C. In both A and B conductivity increases

D. In both A and B conductivity decreases

Answer: c

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49. Consider the following concentration cell:

$$Zn(s) |Zn^{2+}(0.024M)| |Zn^{2+}(0.480M)| Zn(s)$$

which of the following statements is / are correct?

- A. The *EMF* of the cell at $25 \degree C$ is nearly 0.038V.
- B. The *EMF* of the cell at $25 \degree C$ is nearly 0.038V.
- C. If water is added in *LHE*, so that the $\left[Zn^{2+}\right]$ is reduced to 0.012*M*,

the cell voltage increases.

D. If water is added in *LHE*, so that the $\left[Zn^{2+}\right]$ is reduced to 0.12*M* ,

the cell voltage remains same.

Answer: a,c



50. Given :

Oxidation $H_2O_2 \rightarrow O_2 + 2H^{\oplus} + 2e^- E^{c^-} = -0.69V$, $2F^{c^-} \rightarrow F_2 + 2e^- E^{c^-} = -287V$, Reduction $:H_2O_2 + 2H^{\oplus} + 2e^- \rightarrow 2H_2O E^{c^-} = 1.77V$, $2I^(c)$ rarr $I_(2)+2e^(-)$ " "E^(c-)=-0.54V,Whichofthefollow \in gstatementsis//^ are correct ?

A. H_2O_2 behaves as an oxidant for I_2/I^{c-1}

B. H_2O_2 behaves as an reductant for I_2/I^{c-1}

C. I^{c-}/I_2 behaves as an reductant for H_2O_2

D. None of these is correct

Answer: a,c

51. Which of the following statements regarding rusting of iron is / are correct ?

A. It takes place is moist air.

B. It is stopped in CO_2 atmosphere

C. If produces *Fe*(*III*) oxide.

D. It is an electrochemical process.

Answer: a,c,d

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52. Given :

 $A^{2^+} + 2e^- \rightarrow A(s) \quad E^{c^-} = 0.08V$

 $B^{\oplus} + e^- \rightarrow B(s) \quad E^{c-} = -0.64V$

 $X_2(g) + 2e^- \rightarrow 2X^{c-} E^{c-} = 1.03V$

Which of the following statements is / are correct ?

- A. $X_2(g)$ will oxidize both (A) and (B).
- B. A^{2+} will oxidize both (A) and (B)

C. The reaction

 $2X^{c}(1.0M) + A^{2+}(1.0M) \rightarrow X_{2}(1atm) + A(s)$

will be spontaneous.

D. The oxidizing power of A^{2+}, B^{\oplus} , and $X_2(g)$ is in the order

 $X_2 > A^{2+} > B^{\oplus}$

Answer: a,b,d

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53. Which of the following statements is / are correct?

A. Rust is Fe_2O_3

B. Zn - Cu cell is called Daniell cell

C. Saline water slows down rusting.

D. Pure metals undergo corrosino faster then impure metals.

Answer: a,c,d



Exercises Ingle Correct

1. Which of the following solutions can be safely stored in a copper vessel

?

A. ZnSO₄

 $B.AgNO_3$

C. AuCl₃

D. All of them.

Answer: a

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2. If X is the specific resistance of the solution and N is the normality of the solution, the equivalent conductivity of the solution is given by

A.
$$\frac{1000x}{N}$$

B. $\frac{1000}{Nx}$
C. $\frac{1000N}{x}$
D. $\frac{Nx}{1000}$

Answer: b



3. By virtue of Faraday's second law of electrolysis, the electrochemical equivalent of the two metals liberated at the electrodes has the same ratio as that of their

A. Atomic masses

B. Molecular masses

C. Equivalent masses

D. Any of three

Answer: c

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4. The units of conductivity of the solution are

A. ohm ⁻¹

B. ohms

C. *ohm*⁻¹*cm*⁻¹

D. *ohm*⁻¹*eq*⁻¹

Answer: c

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5. According of Kohlrausch law, the limiting value of molar conductivity of an electrolyte A_2B is

A.
$$\lambda^{\infty} \cdot_{A^{+}} + \lambda^{\infty} \cdot (B^{-})$$

B. $\lambda^{\infty} \cdot_{A^{+}} - \lambda^{\infty} \cdot_{B^{-}}$
C. $2\lambda^{\infty} \cdot_{A^{+}} + \frac{1}{2}\lambda^{\infty} \cdot (B^{-})$
D. $2\lambda^{\infty} \cdot_{A^{+}} + \lambda^{\infty} \cdot_{B^{-}}$

Answer: d

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6. The values of Λ_m^{∞} for NH_4Cl , NaOH, and NaCl are, respectively, 149.74, 248.1, and 126.4 ohm⁻¹cm²eq⁻¹. The value of $\Lambda_{eq}^{\infty}NH_4OH$ is

A. 371.44

B. 271.44

C. 71.44
D. It cannot be calculated from the data given.

Answer: b

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7. 0.5F of electricity is passed through 500m of copper sulphate solution. The amount of copper which can be deposited will be

A. 63.5g

B. 31.75g

C. 15.8g

D. Unpredictable

Answer: c

8. On carrying out the electrolysis of acidified water, the volume of hydrogen liberated at *STP* condition is 22.4*L*. The volume of oxygen liberated is

A. 22.4L

B. 44.8L

C. 11.2*L*

D. 2.24L

Answer: c

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9. During the electrolysis of the aqueous solution of copper sulphate using Pt electrode, the reaction taking place at anode electrode is

A.
$$Cu^{2+} + 2e^- \rightarrow Cu$$

 $B. Cu \rightarrow Cu^{2+} + 2e^{-}$

$$C. 2H_2O \rightarrow 4H^{\oplus} + O_2 + 4e^-$$

$$D. H_2O + e^- \rightarrow OH + 1/2H_2$$

Answer: c

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10. In passing 3F of electricity through three electrolytic cells connect in series containing Ag^{\oplus} , Ca^{2+} , and Al^{3+} ions, respectively. The molar ratio in which the three metal ions are liberated at the electrodes is

A. 1:2:3 B. 2:3:1 C. 6:3:2

D.3:4:2

Answer: c

11. Given that $I_2 + 2e^- \rightarrow 2I^{c^-}$, $E^{c^-} = 0.54V$

 $Br_2 + 2e^- \rightarrow 2Br^-, E^{c-} = 1.69V$

Predict which of the following is true.

A. *I^{c-}* ions will be able to reduce bromine.

B. Br^{c-} ions will be able to reduce iodine.

C. Iodine will be able to reduce broming.

D. Bromine will be able to reduce iodide ions.

Answer: a

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12. The increase in the molar conductivity of HCl with dilution is due to

A. Increase in the self ionization of water

B. Hydrolysis of HCl

C. Decrease \in theselfionizationofwater

D. Decrease in the interionic forces.

Answer: D

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13. An electrochemical cell stops working after some time because

A. Electrode potential of both the electrodes becomes zero.

B. Electrode potential both the electrons becomes equal.

C. One of the electrode is eaten away.

D. The reaction starts proceeding in opposite direction.

Answer: b

14. Which of the following statements is correct for a galvanic cell?

A. Reduction occurs at cathode

B. Oxidation occurs at anode

C. Electrons flow from anode to cathode

D. All the statements are correct.

Answer: d

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15.

Given

 $E^{c} \cdot Ag \oplus |Ag = +0.80V, E^{c} \cdot Co^{2}|Co = -0.28V, E^{c} \cdot Cu^{2+}|Cu = +0.34V, E^{c} \cdot Cn^{2}$

Which metal will corrode fastest ?

A. Ag will oxidize to Ag
$$\oplus$$
 and new $\left[Ag \oplus \right] = 1.36M$.

В. Си

С. Со

Answer: d



16. Red hot carbon will remove oxygen from the oxides XO and Yo but not from ZO. Y will remove oxygen from XO. Use this evidence to deduce the order of activity of the three metals X, Y, and Z, putting the most reactive first.

A. *X*, *Y*, *Z*

B. Z, Y, X

C. *Y*, *X*, *Z*

D. *Z*, *X*, *Y*

Answer: b

17. Among *Na*, *Hg*, *S*, *Pt* and graphite which can be used as electodes in electrolytic cell having aqueous solutions?

A. Na and S

B. Hg, pT and S

C. Na, Hg, and S

D. Hg, Pt, and graphite

Answer: d

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18. In an electrolytic cell current flows

A. From cathode to anode in outer circuit

B. From anode to cathode outside the cell

C. From cathode to anode inside the cell

D. None of the above.

Answer: a



19. The reaction
$$Cu^{2+}(aq) + 2Cl^{c-}(aq) \rightarrow Cu(s) + Cl_2(g)$$
 has

A. Can be made to produce electricity in voltaic cell

B. Can be made to occur in an electrolytic cell

C. Can occur in acidic medium only

D. Can occur in basic medium only.

Answer: b



20. Which statements is true about a spontaneous cell reaction in

galvanic cell?

A.
$$E_{cell}^{\Theta} > 0, \Delta G^{\Theta} < 0, Q < K_c$$

B. $E_{cell}^{\Theta} < 0, \Delta G^{\Theta} > 0, Q < K_c$
C. $E_{cell}^{\Theta} > 0, \Delta G^{\Theta} > 0, Q > K_c$
D. $E_{cell}^{\Theta} > 0, \Delta G^{\Theta} > 0, Q < K_c$

Answer: A

_



21. *Zn* acts as sacrifical or cathodic protect iont to prevent rusting of iron because

A.
$$E^{c^{-}}$$
. $_{OP}$ of $Zn < E^{c^{-}}$. $_{OP}$ of Fe

B. E^{c^-} . OP of $Zn > E^{c^-}$. OP of Fe

C. E^{c^-} . _{OP} of $Zn = E^{c^-}$. _{OP} of Fe

D. Zn is cheaper than iron

Answer: b

22. The oxidation potential of a hydrogne electrode at pH = 10 and

 $p_{H_2} = 1atm$ is

A.-0.59V

 $\mathsf{B.}\,0.00V$

C. + 0.59V

D. 0.059V

Answer: c

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23. E^{c-} of $Mg^{2+} | Mg, Zn^{2+} | Zn$, and $Fe^{2+} | Fe$ are -2.37V, -0.76V, and

-0.44V, respectively. Which of the following is correct ?

A. Mg oxidize Fe

B. Zn oxidizes Fe

C. Zn reduces Mg^{2+}

D. Zn reduces Fe^{2+}

Answer: d

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24. If the solution of the $CuSO_4$ in which copper rod is immersed is diluted to 10 times, the electrode potential :

A. Increases by 30mV

B. Decreases by 30mV

C. Increases by 59mV

D. Decrease by 59mV

Answer: b

25. Deduce from the following E^{c-} values of half cells, what combination

of two half cells would results in a cell with the largest potential?

Ι.	$A + e(-) \rightarrow A^{c-}$	$E^{c-} = -0.24V$
II.	$B^{c-} + e^- \rightarrow B^{2-}$	$E^{c-} = + 1.25V$
III.	$C^{c-} + 2e^- \rightarrow C^{3-}$	$E^{c-} = -1.25V$
IV.	$D + 2e^{c} \rightarrow D^2$	$E^{c-} = +0.68V$

A. II, IV

B. II, III

C. III, IV

D. *I*, *II*

Answer: b

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26. Calculate the solubility product of $Co_2[Fe(CN)_6]$ in water at 25 ° C. Given, conductivity of saturated solutions of $Co_2[Fe(CN)_6]$ is $2.06 \times 10^{-6} \Omega^{-1} cm^{-1}$ and that of water used is $4.1 \times 10^{-7} \Omega^{-1} cm^{-1}$. The ionic molar conductivities of Co^{2+} and $\left[Fe(CN)_6\right]^4$ are $86.0\Omega cm^2 mol^{-1}$ and $444.0\Omega^{-1} cm^2 mol^{-1}$, respectivly.

A. 7.87×10^{-7} B. 7.87×10^{-6} C. 7.87×10^{-8}

D. 7.87×10^{-9}

Answer: a

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27. Given $E^{c} \cdot Fe^{2+} | Fe$ and $E^{c} \cdot Fe^{3+} | Fe^{2+}$ are -0.44 and 0.77V respectively. If Fe^{2+}, Fe^{3+} and Fe blocks are kept together, then $[Fe^{3+}]$and $[Fe^{2+}]$

B. Fe^{3+} decreases

C. Fe^{2+} , Fe^{3+} remain unchanged

D. Fe^{2+} decreases

Answer: b

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28. The dissociation of a wask electrolyte obeys the law of mass action. It

was found by

A. Ostwald

B. Arrhenius

C. Berzelius

D. None of these

Answer: a

29. During electrolysis of fused calcium hydride, the hydrogen is produced

at

A. Cathode

B. Anode

C. Hydrogen is not liberated at all

D. H_2 produced reacts with oxygen to form water.

Answer: b

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30. The increase in the value of molar conducitivity of acetic acid with dilution is due to

A. Decrease in interionic forces and increases in α

B. Increase in the degree of ionization and interionic forces.

C. Increase in self ionization of water.

D. None of these

Answer: a



31. Which of the following is anodic reaction.

A.
$$SO_4^{2-} + H_2O \rightarrow H_2SO_4 + 1/2O_2 + 2e^{-}$$

$$B.H^{\oplus} + e^- \rightarrow 1/2OH_2$$

$$\mathsf{C}.Ag \oplus + e^- \to Ag$$

D. None of these

Answer: a



32. The value of molar conductivity of HCl is greater than that of NaCl at

a particular temperature because

A. Molecular mass of *HCl* is less thatn that of *NaCl*

B. Velocity of H^{\oplus} ions is more than that of Na^{\oplus} ions.

C. HCl is stongly acidic.

D. Ionization of *HCl* is larger than that of *NaCl*.

Answer: b

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33. A galvanic cell is set up from a zinc bar weighing 100g and 1.0L of $1.0MCuSO_4$ solution. How long would the cell run if it is assumed to deliver a steady current of 1.0A. (Atomic mass of Zn = 65).

A. 1.1 hours

B. 46 hours

C. 53.6 hourse

D. 24 hours.

Answer: c

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34. The charge required for the reduction of $1 \mod Cr_2 O_7^{2-}$ ions to Cr^{3+} is

A. 96500C

B. 2 × 96500*C*

C. 3 × 96500*C*

 $\mathsf{D.6} \times 96500C$

Answer: d

35. In order to completely oxidize 0.1mol of MnO_4^{2-} to permanganate ion.

The quantity of electricity required is

A. 96500C

B. 2 × 96500*C*

C. 9650C

D. 96.50C

Answer: c

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36. Which of the following reactions occurs at the anode during the recharging of lead storage battery ?

A.
$$Pb + SO_4^2 \rightarrow PbSO_4 + 2e^2$$

 $\mathsf{B}.\mathit{Pb} + \mathit{PbO}_2 + \mathit{H}_2\mathsf{SO}_4 \rightarrow \mathit{2PbSO}_4 + \mathit{2H}_2\mathsf{O}$

 $\mathsf{C.} PbSO_4 + 2e^- \rightarrow Pb + SO_4^{2-}$

$$D. PbSO_4 + 2H_2O \rightarrow Pb + PbO_2 + 2H_2SO_4 + 2e^{-1}$$

Answer: c



37. In H_2 - O_2 fuel cell, the reaction occurring at cathode is

A.
$$2H_2 + O_2 \rightarrow 2H_2O(l)$$

B. $H^{\oplus} + OH \rightarrow H_2O$
C. $O_2 + 2H_2O + 4e^- \rightarrow 4OH$
D. $H^{c^-} + e^- \rightarrow \frac{1}{2}H_2$

Answer: c

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38. The thermodynamic efficiency of cell is given by

A. $\Delta H/\Delta G$

B. $\frac{Nfe}{\Delta G}$ C. $\frac{-nFE}{\Delta H}$

D. nFE^{c-}

Answer: c



39. 10800*C* of electricity passed through the electrolyte deposited 2.977*g* of metal with atomic mass $106.4 gmol^{-1}$. The charge on the metal cation is

A. +4

B. + 3

C. + 2

D. +1

Answer: a

40. In which of the following will the corrosion of iron be most rapid?

A. In pure water

B. In pure oxygen

C. In air and moisture

D. In air and saline water

Answer: d

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41. The two *Pt* electrodes fitted in a conductance cell are 1.5*cm* apart while the cross - sectional area of each electrode is 0.75*cm*. What is the cell constant?

B. 0.5cm

C. 2.0*cm*⁻¹

D. 0.2*cm*⁻¹

Answer: c

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42. For a reaction $A(s) + 2B^{\oplus} \rightarrow A^{2+} + 2B$

 K_c has been found to be 10^{12} . The E^{c} . cell is

A. 0.354V

 $\mathsf{B.}\,0.708V$

 $\mathsf{C.}~0.0098V$

D. 1.36V

Answer: a

43. Rust is a mixture of

A. FeO and $Fe(OH)_2$

B. FeO and $Fe(OH)_3$

C. Fe_2O_3 and $Fe(OH)_3$

D. Fe_3O_4 and $Fe(OH)_3$

Answer: c

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44. Galvanized iron sheets are coated with

A. Copper

B. Nickel

C. Zinc

D. Carbon

Answer: c



46. For an electrolyte solution of $0.05molL^{-1}$, the conductivity has been

found to be 0.0110Scm⁻¹. The molar conductivity is

A. 0.05Scm²mol⁻¹

- B. 550*Scm*²*mol*⁻¹
- C. 0.22Scm²mol⁻¹
- D. 220Scm²mol⁻¹

Answer: d

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47. How much will the reduction potential of a hydrogen electrode change when its solution initially at pH = 0 is neutralized to pH = 7?

A. Increase by 0.059V

B. Decrease by 0.059V

C. Increase by 0.41V

D. Decrease by 0.41V

Answer: d

48. If $E^{c} \cdot Fe^{3+} | Fe$ and $E^{c} \cdot Fe^{2+} | Fe$ are = -0.36V and -0.439V, respectively, then the value of $E^{c} \cdot Fe^{3+} | Fe^{2+}$

A. (-36-0.439)V

B. [3(- 0.36 - 0.436)]V

C.(-0.36 = 0.436)V

D. [3(-0.36) - 2(-0.439)]V

Answer: d

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49. The standard electrode of a metal ion $(Ag \mid Ag^{\oplus})$ and metal - insoluble salt anion $(Ag|AgCl|Cl^{c})$ are related as

A. $E^{c} \cdot Ag \oplus Ag = E^{c} \cdot Cl^{c} \cdot Agcl Ag + \frac{RT}{F} \ln K_{sp}$

$$\mathsf{B}. E^{c} \cdot ._{Cl^{c}} |_{AgCl} |_{Ag} = E^{c} \cdot ._{Ag} \oplus_{Ag} + \frac{RT}{F} \ln K_{sp}$$

$$C. E^{c} \cdot_{Ag} \oplus |_{Ag} = E^{c} \cdot_{Cl^{c}} |_{AgCl|Ag} + \frac{RT}{F} \ln \cdot \frac{\left[Cl^{c}\right]}{K_{sp}}$$
$$D. E^{c} \cdot_{Cl^{c}} |_{AgCl|Ag} = E^{c} \cdot_{Ag} \oplus |_{Ag} + \frac{RT}{F} \ln \cdot \frac{K_{sp}}{\left[Cl^{c}\right]}$$

Answer: b

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50. The standard reduction potential at 25 ° *C* for the reaction, $2H_2O + 2e^- \implies H_2 + 2OH$ is -0.8277 volt. The

equilibrium constant for the reaction :

$$2H_2O \Longrightarrow H_3O^{\oplus} + OH \text{ at } 25^{\circ}C \text{ is}$$

A. 10⁻¹²

B. 10⁻¹⁴

C. 10⁻¹¹

D. 10⁻¹¹

Answer: b



51. What would be the magnitude of *EMF* of the following cell:



at 25 ° C ? The ionization constant of acetic acid, $K_a \sim 10^{-5}$, while that of

```
formic acid, K_a \sim 10^{-4}
```

A. 0.0295V

B. 0.059V

C. -0.059V

D. -0.0295V

Answer: b

52. The rusting of iron takes place as follows :

$$2H^{\oplus} + 2e^{-} + \frac{1}{2}O_2 \rightarrow H_2O(l), \quad E^{c-} = +1.23V$$

 $Fe^{2^+} + 2e^- \rightarrow Fe(s), \quad E^{c-} = -0.44V$

Calculae ΔG^{c^-} for the net process.

A. - 322kJmol⁻¹⁰

B. - 161*kJmol*⁻¹

C. - 152kJmol⁻¹

D. - 76*kJmol*⁻¹

Answer: a



53. For the electrolytic production of $NaClO_4$ from $NaClO_3$ according to

the reaction $NaClO_3 + H_2O \rightarrow NaClO_4 + H_2$. How many faradays of

electricity would be required to produce 0.5mole of NaClO₄?

B. 2 C. 3 D. 1.5

A. 1

Answer: a

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54. If the specific conductane of $1MH_2SO_4$ solution is $26 \times 10^2 Scm^2$, then

the equivalent conductivity would be

A. $1.3 \times 10^2 Scm^{-1}$

B. $1.6 \times 10^2 Scm^{-2}$

C. 13*Scm*²*mol*⁻¹

D. $1.3 \times 10^{3} Scm^{2} mol^{-1}$

Answer: a



55. The highest electrical conductivity of the following aqueous solutions

is of

A. 0.1M acetic acid

B. 0.1M chloro acetic acid

C. 0.1M fluoro acetic acid

D. 0.1M difluoro acetic acid

Answer: d



56. Which of the following statements is wrong ?

- A. The conductance of $1cm^3$ of a material is called specific conductance.
- B. Specific conductance increases while equilvalent conductance decreases on progressive dilution.
- C. The limiting equivalent conductances of weak electrolytes cannot

be determined by the extrapolation of the plot of Λ against concentration.

D. The conductivity of metals is due to the movement of electrons.

Answer: b

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57. Which of the following statement is correct?

A. Specific conductance of a solution decreases with dilution, whereas

molar conductance increases with dilution.

B. Specific conductance of a solution increases with dilution, whereas

molar conductance decreases with dilution

- C. Both specific conductance and molar conductance decrease with dilution.
- D. Both specific conductance and molar conductance increase with dilution.

Answer: a

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58. For a dilute solution of a strong electrolyte, the variation of molar conductivity with concentration is given by

A.
$$\Lambda_m = \Lambda^{\circ} \cdot_m + Ac$$

B. $\Lambda_m = \Lambda^\circ \cdot_m - Ac$

D.
$$\Lambda_m = \Lambda^{\circ} \cdot_m - A\sqrt{c}$$

Answer: D



59. How many coulombs are required for the oxidation of 1mol of H_2O to

*O*₂?

A. 93000C

B. $1.93 \times 10^5 C$

 $C.9.65 \times 10^{40C}$

D. $19.3 \times 10^{2}C$

Answer: b
60. On electrolysis of a solution of dilute H_2SO_4 between platinum electrodes, the gas evolved at the anoe is

A. *SO*₂

 $B.SO_3$

C. O₂

 $D.H_2$

Answer: c

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61. In electrolysis of very dilute of NaOH using platinum electrodes

A. H_2 is evolved at cathode and O_2 at anode

- B. NH₃ is produced at anode
- C. Cl_2 is obtained at cathode
- D. O_2 is produced at cathode and H_2 at anode.

Answer: a



62. During the electrolysis of fused NaCl, which reaction occurs at anode

?

A. Chloride ions are oxidized

B. Chloride ions ar reduced

C. Sodium ions ar oxidized

D. Sodium ions are reduced

Answer: a



63. Two platinum electrodes were immersed in a solution of $CuSO_4$ and

electric current was passed through the solution. After some time, it was

found that colour of $CuSO_4$ disappeared with evolution of gas at the electrode. The colourless solution contains.

A. Platinum sulphate

B. Copper hydroxide

C. Copper sulphate

D. Sulphuric acid

Answer: d

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64. In an experiment setup for the measurement of *EMF* of a half cell using a reference electrode and a salt bridge, when the salt bridge is removes, the voltage

A. Does not change

B. Increase to maximum

C. Decreases to half the value

D. Drops to zero

Answer: d



65. The reference calomel electrode is made from which of the following ?

A. $ZnCl_2$

B. $CuSO_4$

 $C.Hg_2Cl_2$

D. HgCl₂

Answer: c



66. When electricity is passed through a solution of $AlCl_3$ and 13.5g of Al is deposited, the number of *Faradayofe* \leq *ctricitypassedµstbe*......F`.

A. 0.5

B. 1.0

C. 1.5

D. 2.0

Answer: c

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67. What weight of copper will be deposited by passing 2 faradays of electricity through a cupric salt (atomic weight of Cu = 63.5)?

A. 2.0

B. 3.175

C. 63.5

D. 127.0

Answer: c



68. A cell constant is generally found by measuring the conductivity of aqueous solution of

A. BaCl₂

B. KCl

C. NaCl

D. $MgCl_2$

Answer: b

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69. A solution of sodium sulphate was electrolyzed using some inert electrode. The product at the electrodes are

A. O₂, H₂
B. O₂, Na
C. O₂, SO₂

D. $O_2, S_2 O_8^2$

Answer: a

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70. A solution of 9.65A flowing for 10min deposits 3.0g of the metal which

is monovalent. The atomic mass of the metal is

A. 10

B. 50

C. 30

D. 96.5

Answer: b



71. A certain current liberates 0.5*g* of hydrogen in 2 hours. How many grams of copper can be liberated by the same current flowing for the same time in a copper sulphate solution ?

A. 12.7g

B. 15.9g

C. 31.8g

D. 63.5g

Answer: b

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72. The mass of copper that will be deposited at cathode in electrolysis of 0.2*M* solution of copper sulphate when a quantity of electricity equal to that required to liberate 2.24*L* of hydrogen from 0.1*M* aqueous H_2SO_4 is passed (atomic mass of Cu = 63.5) will be

A. 1.59g

B. 3.18g

C. 6.35g

D. 12.70g

Answer: c

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73. Which of the following is a strong electrolyte ?

A.
$$Ca(NO_3)_2$$

B. HCN

 $C.H_2SO_3$

 $D. NH_4OH$

Answer: a

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74. The standard reduction potential of $Li \oplus |Li, Ba^{2+}| Ba, Na \oplus |Na|$ and $Mg^{2+}|Mg|$ are -3.05, -2.71, -2.71, and -2.37 volts, respectively. Which one of the following is the strongest oxidizing agent ?

A. Na [⊕]

B. *Li* ⊕

C. *Ba*²⁺

D. Mg^{2+}

Answer: d

75. The resistance of 1*N* solution of acetic acid is 250*ohm*, when measured in a cell of cell constant $1.15cm^{-1}$. The equivalent conductance (in $ohm^{-1}cm^2eq^{-1}$) of 1*N* acetic acid is

A. 4.6

B. 9.2

C. 18.4

D. 0.023

Answer: a

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76. What volume of $0.1NFeSO_4$ can be oxidized by a current of 2 ampere

hours ?

A. 0.746L

B. 7.46L

C. 1.482L

D. 0.373L

Answer: a

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77. 500mL of 1N solution of $CuCl_2$ was electrolyszed with a current fo 2 amperes for 1 hour. What is the normality of the remaining $CuCl_2$ solution ?

A. 0.85

B. 0.15

C. 0.30

D. 1.0

Answer: a



78. Ionic strength of $0.4MCaCl_2$ is

A. 1.2

B. 1.0

C. 0.9

D. 0.8

Answer: a

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79. Ionic strength of $0.4MCaCl_2$ is

A. 2.8

B. 1.2

C. 1.0

D. 1.8

Answer: b

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80. Ionic strength of a solution made by mixing equal volumes of 0.01*MNaCl* and 0.02*MAlCl*₃

A. 0.065

B. 0.13

C. 0.0325

D. 0.0216

Answer: a

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81. Marshell's acid is prepared by the electrolytic oxidation of H_2SO_4 as

 $2H_2SO_4 \rightarrow H_2S_2O_8 + 2H^{\oplus} + 2e^{-1}$

Oxygen and hydrogen are byproducts. In such electrolysis 2.24L of H_2 and 0.56L of O_2 were product at *STP*. The weight of $H_2S_2O_8$ fromed is

A. 9.7g

B. 19.4g

C. 14.55g

D. 29.1g

Answer: a

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82. The volume of gases evolved at *STP* by passing 0.1*A* of current for 965*g*, through an aqueous solution of potassium acetate

A. 22.4mL

B. 11.2mL

C. 89.6mL

D. 44.8*mL*

Answer: d

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83. The mass of gases evolved in Question is

A. 0.06g

B. 0.6g

C. 6.0g

D. 60g

Answer: a

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84. The volume of gases evolved at STP by passing 0.2A of current for

965s through an aqueous solution of sodium furmarate is

A. 22.4mL

B. 11.2*mL*

C. 89.6mL

D. 44.8mL

Answer: c

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85. The products obtained at cathode and anode on electrolysis of aqueous sodium succinate are

A. Anode : Ehene+ CO_2 , Cathode : H_2

B. Anode : Ethyne+ CO_2 ,Cathode : H_2

C. Anode: Ethene+ H_2 , Cathode : Na

D. Anode: Ethyne $+H_2$, Cathode: Na

Answer: a



86. What is the electrode potential of a gasous hydrogen electrode dipped in a solution pH = 5.0 relative to the calomel electrode with an electrode potential of +0.28V?

A. 0.0125V

B. + 0.575V

C. +0.015V

D. -0.575V

Answer: d

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87. The *EMF* of concentration cell consisting of two zinc electrodes, one dipping into M/4 solution of $ZnSO_4$ and the other into M/16 solution of the same salt at 25 ° C is

A. 0.0125V

B. 0.0250V

 $C.\,0.0178V$

D. 0.0356V

Answer: c

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88. A certain electrode has standard (reduction potential) of 0.384V. The potential when measured against a normal calomel electrode (with electrode potential = 0.28V) is

A. 0.104

B. 0.664

C. 0.3322

D. 0.218

Answer: a

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89. The potential of a hydrogen electrode in a solution with pOH = 4 at

25 ° C is

A. +0.59

B. - 0.59

C.-0.295

D. 0.295

Answer: c

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90. How much will the reduction potential of a hydrogen electrode change when its solution initially at pH = 0 is neutralized to pH = 7?

A.-0.059

B. 0.059

C.-0.59

D. 0.59

Answer: c

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91. Consider the electrode $Ag \mid AgCl(s), Cl^{c-}(0.1M), i. e.$, silver electrode in contact with 0.1MKCl solution saturated with AgCl. If it is combined with the electrode $Ag \mid Ag^{\oplus}(0.1M)$ to form a complete cell, the *EMF* would be $\left(K_{sp}ofAgCl = 10^{10} \text{ at } 25 \degree C\right)$

A. 0.799

B.-.6363

C. 0.59

D. 0.472

Answer: d

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92. A current strength of 1.0A is passed for 96.5s through 100mL of a

solution of 0.05*MKCl*. The concentration of the final solution with respect *c*to *OH* ions is

A. 0.005M

B. 0.05M

C. 0.01*M*

D. 0.001M

Answer: c

93. In Question *pH* of the final solution will be

A. 12

B. 2

C. 11.7

D. 3

Answer: a

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94. The standard *EMF* of decinormal calomel electrode is 0.268V. The

EMF is

A. -0.298V

B. 0.327V

C.-0.327V

D. 0.298V

Answer: b

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95. The satandard *EMF* of quinhydrone is 0.699V. The *EMF* of the quinhydrone electrode dipped in a solution with pH = 10 is

A. 0.109V

B.-0.109V

C. 1.289V

D. - 1.289V

Answer: a

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96. A hydrogen electrode placed in a solution containing sodium acetate and acetic acid in the ratio of x:y and y:x has an electrode potential value E_1 and E_2 volts, respectively, at 25 °C. The pK_a value of acetic acid is

A.
$$\frac{-(E_1 + E_2)}{2 \times 0.059}$$

B.
$$\frac{E_1 + E_2}{2 \times 0.059}$$

C.
$$\frac{E_2 - E_1}{2 \times 0.059}$$

D.
$$\frac{-(E_1 + E_1)}{0.059}$$

Answer: a

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97. Given standard E^{c-} :

 $Fe^{3+} + 3e^- \rightarrow Fe, \quad E^{c-} = -0.036$

Fe⁽²⁺⁾+2e⁽⁻⁾rarr Fe," "E^(c-)=-0.440V*The*E^(c-)*of*Fe⁽³⁺⁾+e⁽⁻⁾ rarr Fe⁽²⁺⁾

A. -0.476V

B.-0.404V

C.0.404V

D. 0.772V

Answer: d

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98. Given the ionic equivalent conductivities for the following ions :

$$\lambda^{\circ} \cdot_{eq} K^{\oplus} = 73.5 cm^2 ohm^{-1} eq^{-1}$$
$$\lambda^{\circ} \cdot_{eq} A l^{3+} = 149 cm^2 ohm^{-1} eq^{-1}$$

 $\lambda \circ ._{eq} SO_4^{2-} = 85.8 cm^2 ohm^{-1} eq^{-1}$

The Λ_{eq}° for potash alum $\left(K_2SO_4, Al_2(SO_4)_3, 24H_2O\right)$ is :

A. 215.92

B. 348.3

C. 368.2

Answer: a

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99. For the cell $Zn(s) |Zn^{2+}| |Cu^{2+}| Cu(s)$, the standard cell voltage, $E^{c-} \cdot_{cell}$ is 1.10V. When a cell using these reagents was prepared in the lab, the measured cell voltage was 0.98V. One possible explanatino for the observed voltage is

A. There were 2.00*mol* of Zn^{2+} but only 1.00*mol* of Cu^{2+}

B. The Zn electrode had twice the surface of the Cu electrode.

- C. The $\left[Zn^{2+}\right]$ was larger than the $\left[Cu^{2+}\right]$.
- D. The volume of the Zn^{2+} solution was larger than the volume of the

 Cu^{2+} solution.

Answer: c



100.
$$\Delta G = \Delta H - T\Delta S$$
 and ItBRgt $\Delta G = \Delta H + T \left[\frac{d(\Delta G)}{dT} \right]_p$, then $\left(\frac{dE_{cell}}{dT} \right)$ is



D. nFE_{cell}

Answer: a

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101.
$$Zn + Cu^{2+}(aq) \Leftrightarrow Cu + Zn^{2+}(aq)$$
.

Reaction quotient is $Q = \frac{\left[Zn^{2+}\right]}{\left[Cu^{2+}\right]}$. Variation of E_{cell} with log Q is of the



A.
$$[Cu^{2+}]/[Zn^{2+}] = 0.01$$

B. $[Zn^{2+}]/[Cu^{2+}] = 0.01$
C. $[Zn^{2+}]/[Cu^{2+}] = 0.1$
D. $[Zn^{2+}]/[Cu^{2+}] = 1$

Answer: b

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102. The standard reduction potentials of $Cu^{2+} | Cu$ and $Cu^{2+} | Cu^{\oplus}$ are 0.337V and 0.153V, respectively. The standard electrode potential fo $Cu^{\oplus} | Cu$ half cell is

A. $x_2 - 2x_2$ B. $x_1 + 2x_2$ C. $x_1 - x_2$ D. $2x_2 - x_1$

-

Answer: d

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103. $Zn \mid Zn^{2+} \left(C_1 \mid Zn^{2+} \left(C_2 \mid Zn. \text{ For this cell } \Delta G \text{ is negative if:} \right) \right)$

A. $c_1 = c_2$

B. $c_1 > c_2$

 $C. c_2 > c_1$

D. None

Answer: c

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104. $Pt(H_2)(p_1)|H^{\oplus}(1M)|(H_2)(p_2)$, Pt cell reaction will be exergonic if

A. $p_1 = p_2$

- **B**. $p_1 > p_2$
- $C. p_2 > p_1$

D. $p_1 = 1$ *atm*

Answer: b

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105. $Pt(Cl_2)(p_1)|HCl(0.1M)|(Cl_2)(p_2), Pt$ cell reaction will be endergonic if

A. $p_1 = p_2$

B. $p_1 > p_2$

 $C. p_2 > p_1$

D. $p_1 = p_2 = 1$ atm

Answer: c

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106.
$$Pt(H_2)(1atm) \mid H_2O$$
, electrode potential at 298K is

A.-0.2364V

B.-0.4137V

C. 0.4137V

 $\mathsf{D}.\,0.00V$

Answer: c



107. For the half cell



At pH = 2, the electrode potential is

A. 1.36V

 $B.\,1.30V$

C. 1.42V

D. 1.20V

Answer: c



108. Which of the following changes will increase the EMF of the cell :

$$Co(s) \left| CoCl_2(M_2) \right| \left| HCl(M_2) \right| + (H_2, g) Pt$$

A. Increae in the volume of $CoCl_2$ solution from 100mL to 200mL.

B. Increase M_2 from 0.1M to 0.50M.

C. Increase the pressure of the $H_2(g)$ from 1.0 to 2.0*atm*.

D. Increase M_1 from 0.01M to 0.50M.

Answer: b



109. $Ag \mid Ag^{\oplus}(1M)$ 1Lsolution $\mid Ag^{\oplus} \mid Ag$ 1Lsolution

0.5F electricity in the LHS(anode) and 1F of electricity in the RHS(

cathode) is first passed making them independent electrolytic cells at 298*K*. *EMF* of the cell after electrolysis will be

A. Increased

B. Decreased

C. No change

D. Time is also required

Answer: c

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110. The cell EMF is independent of the concentration of the species of

the cell in

A. Fe⁻ FeO(s)|KOH(aq)|NiO(s) | Ni₂O₃(s)
$$\stackrel{\oplus}{|}$$
 Ni
B. Pt $|(H_2)|HCl | (Cl_2)Pt$
C. Zn $|Zn^{2+}||Cu^{2+}|Cu$

D. Hg, $Hg_2Cl_2|KCl||AgNO_3|Ag$

Answer: a

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111. Assume that during the electrolysis of AgNO + (3), only H_2O is electrolyzed and O_2 is formed as $2H_2O \rightarrow 4H^{\oplus} + O_2 + 4e(-)$

 O_2 formed at *NTP* due to passage of 2 amperes of current for 96 second is

A. 0.112L

B. 0.224L

C. 11.2*L*

D. 22.4L

Answer: a
112. During electrolysis of acidified water, O_2 gas is formed at the anode . To produce O_2 gas at the anode at the rate of 0.224mL per second at *STP*, current passed is

A. 0.224A

B. 2.24*A*

C. 9.65A

D. 3.86A

Answer: d

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113. The gas evolved at the anode when $K_2SO_4(aq)$ is electrolyzed between *Pt* electrode is

 $\mathbf{B}.H_2$

C. *SO*₂

D. *SO*₃

Answer: a

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114. A quantity of electrical charge that brigns about the depositiion of 4.5*gAl* from Al^{3+} at the cathode will also produce the following volume (*STP*) of $H_2(g)$ from H^{\oplus} at the cathode.

A. 44.8L

B. 22.4L

C. 11.2*L*

D. 5.6L

Answer: d



115. 100mL of buffer of $1MNH_3(aq)$ and $1MNH_4^{\oplus}(aq)$ are placed in two compartments of a voltaic cell separately. A current of 1.5A is passed through both cells for 20min. If only electrolysis of water takes place, then

A. *pH* of *LHS* half cell will increase.

B. pH of RHS half cell will increase

C. pH of both half cell will increase

D. Ph of both half cell will decrese

Answer: b

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116. Given the ionic conductance of $COO^{c-} | coo^{c-}, K^{\oplus}$, and Na^{\oplus} are 74, 50, and $73cm^2ohm^{-1}eq^{-1}$, respectviley. The equivalent conductance at infinite dilution of the salt COONa | cook is

A. $197 cm^2 ohm^{-1} eq^{-1}$

- B. $172 cm^2 ohm^{-1} eq^{-1}$
- C. 135.5*cm*²*ohm*⁻¹*eq*⁻¹
- D. 160.5*cm*²*ohm*⁻¹*eq*⁻¹

Answer: c



117. The variation equivalent conductance of stronge electrolyte with

 $\sqrt{Concentration}$ is correctly shown in the figure.





Answer: b

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118. Given the standard potential of the following at $25 \degree C$.

 $MnO_2 \rightarrow Mn^{3+}, \quad lE^{c-} = 0.95V$

 $Mn^{3+} \rightarrow Mn^{2+}, E^{c-} = 1.51V$

The standard potential of $MnO_2 \rightarrow Mn^{2+}$ is

A. -0.56V

B.-2.46V

C. - 1.23*V*

D. 1.23V

Answer: d

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119. The potential of the following cell at 25 $^{\circ}C$ is

A. -0.059V

 $\mathsf{B.}\,0.059V$

C. -0.59V

D. 0.5V

Answer: c

120. Given the following cell at 25 $^{\circ}C$

What will be the potential of the cell ?

Given pK_a of $CH_3COOH = 4.74$

A. -0.42V

B. 0.42V

C. -0.19V

D. 0.19V

Answer: a

121. What is the potential of the cell containing two hydrogen electrode

as represented below ?

$$Pt, \frac{1}{2}H_2(g) |H_2O| |H^{\oplus}(0.01M)| 1/2H_2(g)Pt$$

A. -0.236V

B.-0.0591V

C. 0.236V

D. 0.0591V

Answer: c

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122. Given electrode potentials asre $Fe^{3^+} + e^- \rightarrow Fe^{2^+}, \quad E^{c^-} = 0.771V$ I_(2)+2e^(-) rarr 2l^(c-)," "E^(c-)=0.536VE^(c-)._(cell)f or thecellreaction, Fe^(3+)+2l^(c-) rarr Fe^(2+)+I_(2)` is A. $(2 \times 0.771 - 0.536) = 1.006V$

- B. $(0.771 0.5 \times 0.536) = 0.503V$
- C.(0.771 0.536) = 0.235V

D.(0.536 - 0.771) = 0.236V

Answer: c

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123. The following facts are availabel :

- $2A^{c-} + B_2 \rightarrow 2B^- + A_2,$
- $2C^{c-} + B_2 \rightarrow Noreaction,$

 $2D^{c-} + A_2 \rightarrow 2A^{c-} + D_2$

Which of the following statement is correct ?

$$\begin{aligned} \mathbf{A} \cdot E^{c^{-}} \cdot C^{c^{-}} \mid C_{2} &\geq E^{c^{-}} \cdot B^{c^{-}} \mid B_{2} &\geq E^{c^{-}} \cdot A^{c^{-}} \mid A_{2} &\geq E^{c^{-}} \cdot D^{c^{-}} \mid D_{2} \\ \mathbf{B} \cdot E^{c^{-}} \cdot C^{c^{-}} \mid C_{2} &\leq E^{c^{-}} \cdot B^{c^{-}} \mid B_{2} &\leq E^{c^{-}} \cdot A^{c^{-}} \mid A_{2} &\leq E^{c^{-}} \cdot D^{c^{-}} \mid D_{2} \\ \mathbf{C} \cdot E^{c^{-}} \cdot C^{c^{-}} \mid C_{2} &\leq E^{c^{-}} \cdot B^{c^{-}} \mid B_{2} &\geq E^{c^{-}} \cdot A^{c^{-}} \mid A_{2} &\geq E^{c^{-}} \cdot D^{c^{-}} \mid D_{2} \end{aligned}$$

$$\mathsf{D}. E^{c^{-}} \cdot C^{c^{-}} | C_2 > E^{c^{-}} \cdot B^{c^{-}} | B_2 < E^{c^{-}} \cdot A^{c^{-}} | A_2 < E^{c^{-}} \cdot D^{c^{-}} | D_2$$

Answer: b



124. The potential the cell at 25
$$^{\circ}C$$
 is

$$Pt \begin{vmatrix} H_2 \\ (1 \text{ atm}) \end{vmatrix} \begin{vmatrix} NH_4 OH \\ (10^3 M) \end{vmatrix} \begin{vmatrix} NaOH \\ (10^{-3} M) \end{vmatrix} \begin{vmatrix} H_2 \\ (1 \text{ atm}) \end{vmatrix} Pt$$

Given pK_b of $NH_4OH = 4.74$

A. 0.05V

B.-0.05V

C. -0.28V

D. 0.28V

Answer: b

125. The potential of the cell at 25 $^{\circ}C$ is

Given pK_a of CH_3COOH and pK_b of $NH_4OH = 4.74$

A. -0.04V

 $\mathsf{B.}\,0.04V$

C. -0.189V

D.0.189V

Answer: c

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126. Which metal can deposit copper from copper sulphate solution ?

A. Mercury

B. Iron

C. Gold

D. Platinum

Answer: b

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127. On the basis of position in the electrolchemical series, the metal which does not displace H_2 from water and acid is :

A. Hg

B.Al

C. Pb

D. Ba

Answer: a

128. A dilute aqueous solution of sodium fluoride is electrolyzed, the products at the and cathode are

A. O₂, H₂

B. *F*₂, *Na*

C. O₂, Na

 $D.F_2, H_2$

Answer: a

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129. Copper can be deposited from acidified copper sulphate and alkaline

cuprous cyanide. If the same current is passed for a definite time :

A. The amount of copper deposited from acidic copper sulphate will

be higher.

B. The amount of copper deposited from alkaline cuprous cyanide will

be higher

C. The same amount of copper will be deposited.

D. None of these

Answer: b

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130. Silver is removed electrolytically from 200mL of a 0.1N solution of $AgNO_3$ by a current of 0.1A. How long will it take to remove half of the silver from the solution ?

A. 0.1s

B. 100s

C. 965s

Answer: d

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131. Chromium plating can involve the electrolysis of an electrolyte of an acidified mixture of chromic acid and chromium sulphate. If during electrolysis the article being plated increases in mass by 2.6g and $0.6dm^3$ of oxygen are evolved at an inert anode, the oxidation state of chromium ions being discharged must be : (assuming atomic weight of Cr = 52 and 1mole of gas at room temperature and pressure occupies a volume at $24dm^3$)

A. - 1

B. Zero

C. +1

D. + 2

Answer: c



132. Which of the following does not evolve oxygen at anode when the electrolysis is carried out of

A. Dilute H_2SO_4 with Pt electrode

B. Fused sodium hydroxide with Pt electrodes

C. Acidic water with Pt electrodes

D. Dilute sulphuric acid using Cu electrodes

Answer: d



133. Calculate the potential of the following cell :

$$Pt \left| Co^{2^{+}}, Co^{3^{+}} (2.0M) (1.0M) \right| Cr^{3^{+}}, Cr_{2}O_{7}^{2^{-}}, H^{\oplus} (1.0M) (4.0M) (1.0M) \mid Pt$$

$$E^{c^{-}} \cdot Co^{2^{+}} \mid Co^{3^{+}} = -2V, E^{c^{-}} \cdot Cr_{2}O_{7}^{2^{-}} \mid Cr^{3^{+}} = +1.0V$$

A. 1.024V

B.-1.024V

C. 0.976V

D.-0.976V

Answer: d

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134. The electricity conductivity of a solution serves as a means of determing the end point in a chemical reaction, involved in the titration of acids, bases, or precipitation. Which of the following conductometric titrations represent the curve of $HClvsNH_4OH$

```
x - axis \Rightarrow Volume of alkali added
```





Answer: b

135. A constant current was passed through a solution of $AuCl_4^{c^-}$ ion between gold electrodes. After a period of 10.0*min*, the increase in the weight of cathode was 1.314*g*. The total charge passed through solution is (atomic weight of $AuCl_4^{c^-} = 339$)

A. $1.16 \times 10^{-2}F$

B. $3.5 \times 10^{-2}F$

 $C.2 \times 10^{-2}F$

D. 4 × 10⁻³*F*

Answer: c

$$K_{\infty} \text{ for } Fe^{-3} + Y^{4} \iff Fe Y^{\textcircled{o}} = \frac{[FeY^{\textcircled{o}}]}{[Fe^{+3}][Y^{4-}]} = 1.3 \times 10^{25}$$

$$K_{N} \text{ for } Fe^{-2} + Y^{4-} \iff Fe Y^{2-} = \frac{[FeY^{2-}]}{[Fe^{+2}][Y^{4-}]} = 2.1 \times 10^{14},$$

$$Cell \text{ reaction is:}$$

$$Fe^{-3} + X^{4-} \iff Fe Y^{\textcircled{o}}$$

$$Fe Y^{2-} \iff Fe^{+2} + X^{4-}$$

$$Fe^{+3} + Fe Y^{2-} \longrightarrow Fe^{+2} + Fe Y^{\textcircled{o}}$$

$$K_{eq} = \frac{K_{eq_{1}}}{K_{eq_{2}}}$$

$$= \frac{1.3 \times 10^{25}}{2.1 \times 10^{14}}$$

136.

 E^{c-} for $FeY^{c-} + e^{c-} \rightarrow FeY^{2-}$

A. 0.13V

B.-0.636V

C. + 0.636V

D. 1.41V

Answer: a

137. Calculate E^{c-} for the reactions :

 $ZnY^{2^-} \Leftrightarrow Zn(s) + Y^{4^-}$ where Y^{4^-} is the completely deprotonated anion of EDTA. The formation constant for ZnY^{2^-} is 3.2×10^{16} and E^{c^-} for $Zn \rightarrow Zn^{2^+} + 2e^-$ is 0.76V.

A. - 1.25V

B.0.48V

C. + 0.68V

D. -0.27V

Answer: a

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138. The solubility product of $Pb_3(AsO_4)_2$ is 4.1×10^{-36} . E^{c^-} for the reaction :

$$Pb_{3}(AsO_{4})_{2}(s) + 6e^{-} \Leftrightarrow 3Pb(s) + 2AsO_{4}^{2}$$

$$E_{(Pb)2^{+}|Pb}^{\Theta} = -0.13V$$

A. +0.478V
B. -0.13V
C. -0.478V
D. +0.13V

Answer: c

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139. A cell is to be constructed to show a redox change :

 $Cr + 2Cr^{3+} \Leftrightarrow 3Cr^{2+}$. The number of cells with different E^{c-} an n but same value of ΔG^{c-} can be made (Given $E^{c-} \cdot Cr^{3+} | Cr^{2+} = -0.40V, E^{c-} \cdot Cr^{3+} | Cr = -0.74V,$ adn $E^{c-} \cdot Cr^{2+} | Cr = -0.91V$)

140. E^{c^-} for $Cr^{3^+} + 3e^- \rightarrow Cr$ and $Cr^{3^+} + e^- \rightarrow Cr^{2^+}$ are -0.74V and -0.40V, respectively, E^{c^-} for the reaction is $Cr^{+2} + 2e^- \rightarrow Cr$

A. -0.91V

B.+0.91V

C. - 1.14V

D.+0.34V

Answer: a

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141. The efficiency of a fuel cell si 80 % and the standard heat of reaction is -300kJ. The reaction involves two electrons in redox change. E^{c-} for the cell is

A. 1.24V

B.2.48V

C.0V

D. 0.62V

Answer: a

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142. The E_{cell} for a given cell is 1.2346 and 1.2340V at 300K and 310K, respectively. Calculate the change in entropy during the cell reaction if the redox change involves three electrons.

A. - 17.37*JK*⁻¹

B. +17.37*JK*⁻¹

C. 173.7*JK*⁻¹

D. 5.79*JK*⁻¹

Answer: a

143. A curent of 3A was passed for 1 hour through an electrolyte solution of $A_x B_y$ in water. If 2.977g of A(atomic weight 106.4) was deposited at cathode and B was a monovalent ion, the formula of electrolyte was

A. AB_2

B.AB

 $C.AB_3$

 $D.AB_4$

Answer: d

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144. The E^{c^-} for Cu^{2^+}/Cu^{\oplus} , Cu^{\oplus}/Cu , Cu^{2^+}/Cu , are 0.15V, 0.50V and 0.325V, respectively. The redox cell showing redox reaction $2Cu^{\oplus} \rightarrow Cu^{2^+} + Cu$ is made. E^{c^-} of this cell reaction and ΔG^{c^-} may be

A. $E^{c-} = 0.175V$ or $E^{c-} = 0.350V$

B. n = 1 or 2

 $C. \Delta G^{c-} = -33.775 kJ$

D. All of these

Answer: d

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145. Total charge required to convert three moles of Mn_2O_4 to $MnO_4^{C^-}$ in

present of alkaline medium

A. 10*F*

B. 20*F*

C. 30*F*

D. 40F`

Answer: c

146. A current of 965A is passed for 1s through 1L solution of $0.02NNiSO_4$

using Ni electrodes. What is the new concentration of NiSO₄?

A. 0.01N

B. 0.01M

C. 0.003M

D. 0.02M

Answer: b

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147. For the given cell $Pt_{D_2 | D^{\oplus}} | | H^{\oplus} | Pt_{H_2}$, if $E^{c-} \cdot_{D_2 | D^{\oplus}} = 0.003V$, , what will be the ratio of D^{\oplus} and H^{\oplus} at 25 °C when the reaction $D_2 + 2H^{\oplus} \rightarrow 2D^{\oplus} + H_2$ attains equilibrium

A. 1.34

B. 1.24

C. 1.124

D. 1.45

Answer: c

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148. What is $E^c \cdot_{red}$ for the reaction: $Cu^{2^+} + 2e^- \rightarrow Cu$ in the half cell

 $Pt_{S^2|CuS|Cu}$ if $E^{c} \cdot Cu^{2+}|Cu}$ is 0.34V and K_{sp} of $CuS = 10^{-35}$?

A. 0.34V

B.-0.6925V

C. + 0.6925V

D. -0.66V

Answer: b

149. The combustion of butane in O_2 at 1 bar and 298K shows a decrease in free energy equal to $2.95 \times 10^3 k Jmol^{-1}$ in a fuel cell. K and E^{c-} of the fuel cell are

A. 9.55×10^{482} , 1.096V

B. 9.55, 1.096V

C. 1.023×10^{966} , 2.85V

D. 5.5×10^{484} , 0.55V

Answer: a

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150. A half cell reaction $Ag_2S(s) + 2e^- \rightarrow 3Ag(s) + S^{2-}$ is carried out in a

half cell $Pt_{Ag_2S \mid Ag, H_2S(0.1M)}$, at $\left[H^{\oplus}\right] = 10^{-3}$, The *EMF* of the half cell is $\left[E^{c-} \cdot_{Ag^{\oplus} \mid Ag} = 0.80V, K_a(H_2S) = 10^{-21}$, and K_{sp} of $Ag_2S = 10^{-49}$]

A. -0.1735V

B.-0.19V

C.+0.1735V

D. + 0.19V

Answer: a

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151. Which one is wrong if electrolysis of $CH_3COONa(aq)$ is made using Pt

electrodes ?

A. *pH* of solution increases.

B. Molar ratio of gases at anode and cathode is 3:1

C. $\begin{bmatrix} CH_3 COO^{c-} \end{bmatrix}$ in solution decreases.

D. The molar ration of gases at anode and cathode is 2:1

Answer: d

152. The calomel and quinhydrone electrodes are reversible with respect to which ions, respectively ?

A. Cl^{c-} , H^{\oplus} B. H^{\oplus} , Cl^{c-} C. Hg_2^{2+} , OHD. Hg_2^{2+} , H^{\oplus}

Answer: a

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153. The EMF of Ni - Cad battery is dependent of :

A. Cd(OH)₂

B. $Ni(OH)_2$

с-С. ОН

D. None of these

Answer: d

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154. The electrode with reaction
:
$$Cr_2O_7^{2-}(aq) + 14H^{\oplus}(aq) + 6e^- \rightarrow 2Cr^{3+}(aq) + 7H_2O$$
 can be represented

as

A.
$$Pt \mid H^{\oplus}(aq), Cr_2O_7^{2-}(aq)$$

B. $Pt \mid H^{\oplus}(aq), Cr_2O_7^{2-}(aq), Cr^{3+}(aq)$
C. $Pt_{H2} \mid H^{\oplus}(aq), Cr_2O_7^{2-}$
D. $Pt_{H2} \mid H^{\oplus}(aq), Cr_2O_7^{2-}(aq), Cr^{3+}(aq)$

Answer: d

155. For a given reaction $:M^{(x+n)} + ne^- \to M^{c+}, E^{c-}.$ (*red*) is known along with M^{x+n} and M^{x+} ion concentrations. Then

A. n can be evaluated

B. x can be evaluated

C. (x + n) can be evaluated

D. n, x, (x + n) can be evaluated

Answer: a

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156. Select the wrong statement.

A. The electrolysis of molten CaH_2 liberates H_2 at cathode

B. During the discharge of lead storage battery, sulphuric acid is

consumed.

C. Sulphur acts as a polymerizing agent in the vulcanization of rubber.

D. Galvanization of iron denotes coating with Zn.

Answer: a

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157. Select the correct statement.

A. Faraday represents 96500 coulombs per second.

B. Coulomb represents one ampere for 1/2 second.

C. Coulomb represents 1/2 ampere for 1 second

D. Coulomb represnts charge of one mole electrone.

Answer: d

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158. From the following information, calculate the solubility product of

AgBr. AgBr(s) + $e^- \rightarrow Ag(s) + Br^{c-}(aq), \quad E^{c-} = 0.07V$ Ag $^{\oplus}(aq) + e^- \rightarrow Ag(s), \quad E^{c-} = 0.080V$ A. 4×10^{-13} B. 4×10^{-10} C. 4×10^{-17} D. 4×10^{-7}

Answer: a

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159. The strongest oxidizing agent among the following is

A. $K_2Cr_2O_7$

B. MnO_2

C. $KMnO_4$

 $D.Cl_2$

Answer: c

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160. The weakest oxidizing agent among the following is

A. O₃

 $\mathbf{B}.F_2$

C. $KMnO_4$

D. *Cl*₂

Answer: d
161. Suppose that gold is being plated onto another metal in a electrolytic cell. The half - cell reaction producing the Au(s) is $AuCl_4^{C^-} \rightarrow Au(s) + 4Cl^{C^-} + 3e^-$

If a 0.30 - A current runs for 1.50min , what mass of Au(s) will be plated, assuming all the electrons are used in the reduction of $AuCl_4$?

A. 0.184g

B. 0.551g

C. 1.84g

D. 0.613g

Answer: a

:



162. Chromium plating is applied by electrolysis to objects suspended in a dichlromate solution , according to following (unbalanced) hald reaction

$$Cr_2O_7^{2^-}(aq) + e^- + H^{\oplus}(aq) \rightarrow Cr(s) + H_2O(l)$$

How many hours would it take to apply a chromium plating of thickness $2.0 \times 10^{-2} mm$ to a car bumper of suface area $0.25m^2$ in an electrolysis cell carrying a current of 75.0A?

[Density of chromium is $7.19gcm^{-3}$]

A. 2.2h

B. 1.5h

C. 3.0h

D. 0.25h

Answer: b

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163. 20Ml of *KOH* solution was titrated with $0.20MH_2SO_4$ solution in a conductivity cell. The data obtained were plotted to give the graphe shown below.

The concentration of the KOH solution was



A. 0.30*molL*⁻¹

- B. 0.15*molL*⁻¹
- C. 0.12*molL*⁻¹
- D. 0.075molL⁻¹

Answer: a

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164. What is the cell entropy change (in JK^{-1}) of the following cell :

$$Pt(s) \left| H_2(g)p = 1 a t m \right| CH_2 COOH, HCl0.1M \left| KCl(aq)0.1M \right| Hg_2 Cl_2(s) \mid Hg$$

The *EMF* of the cell is found to be 0.045V at 298K and temperature coefficient if $3.4 \times x10^{-4}VK^{-1}$

(Given :
$$K_a(CH_3COOH) = 10^{-5}M$$
)

A. 60

B. 65.2

C. 69.2

D. 63.5

Answer: b

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165. $k = 4.95 \times 10^{-5} Scm^{-1}$ for a 0.001*M* solution. The reciprocal of the degree of dissociation of acetic acid, if Λ_m° for acetic acid is

 $400Scm^{-2}mol^{-1}$ is :

A. 7

B. 8

C. 9

D. 10

Answer: b

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166. What is the value of $pK_b(CH_3COO^{c-})$ if $\land \circ \cdot_m = 390Scm^{-1}mol^{-1}$ and $\land_m = 7.8Scm^2mol^{-1}$ for 0.04*M* of CH_3COOH at 25 ° *C*?

A. 9.3

B. 9.2

C. 4.7

D. 4.8

Answer: b



167. In Hall's process, in the production of *Al*, carbon is used as the anode material. The reactions are

A. Anode
$$:Al^{3+} + 3e^- \rightarrow Al$$

Cathode : $C(s) + 2O^2 \rightarrow CO_2(s) + 4e^-$

B. Anode:
$$C(s) + 2O^{2^-} \rightarrow CO_2(g) + 4e^-$$

Cathode :
$$Al^{3+} + 3e^- \rightarrow Al$$

C. Anode
$$:Al \rightarrow Ae^{3+} + e^{-}$$

Cathode : $CO_2 + 4e^- \rightarrow C(s) + 2O^{2-}$

D. None of these

Answer: b

168. ΔG^{c-} or the reaction is ,

$$4Al + 3O_{2} + 6H_{2}O + 4OH \rightarrow 4Al(OH)_{4}^{c}$$

$$E^{c} \cdot_{cell} = 2.73V$$

$$\Delta_{f}G^{c} \cdot \begin{pmatrix} c \\ OH \end{pmatrix} = -157kJmol^{-1}$$

$$\Delta_{f}G^{c} \cdot \begin{pmatrix} c \\ OH \end{pmatrix} = -237kJmol^{-1}$$
A. -3.16 × 10³kJmol^{-1}
B. -0.79 × 10³kJmol^{-1}
C. -0.263 × 10³kJmol^{-1}
D. +0.263 × 10³kJmol^{-1}

Answer: a



169. $Cu^{2^+} + 2e^- \rightarrow Cu$. For this, graph between E_{red} versus $\ln \left[Cu^{2^+} \right]$ is a

straight line of intercept 0.34V, then the electrode oxidation potential of

the half cell $Cu \mid Cu^{2+}(0.1M)$ will be

A.
$$0.34 + \frac{0.0591}{2}$$

B. $-0.34 - \frac{0.0591}{2}$
C. 0.34
D. $-0.34 + \frac{0.0591}{2}$

Answer: d

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170. A cell $Cu | Cu^{2+} | | Ag^{\oplus} | Ag$ inintially contains $2MAg^{\oplus}$ and $2MCu^{2+}$ ion in 1*L* solution each. The change in cell potential after it has supplied 1A current for 96500s is

A. -0.003V

B.-0.02V

C. -0.04V

D. None of these

Answer: b



171. The value of reaction quotient Q for the cell

$$Zn(s) \left| Zn^{2+}(0.01M) \right| \left| Ag^{\oplus}(1.25M) \right| Ag(s) is$$

A. 156

B. 125

C. 1.25xx10^(-2)

D. 64xx10^(-3)

Answer: d

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172. In acid medium, MnO_4^{c-} is an oxidizing agent.

 $MnO_4^{c^-} + 8H^{\oplus} + 5e^- \rightarrow Mn^{2+} + 4H_2O$

If H^{\oplus} ion concentration is doubled, electrode potential of the half cell MnO_4^{c-} , $Mn^{2+} \mid Pt$ will

A. Increase by 28.36mV

B. Decrease by 28.36mV

C. Increase by 14.23mV

D. Decrease by 142.30mV

Answer: a

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173. During the electrolysis of $AgNO_3$, the volume of O_2 formed at *STP* due to passage of 2A of current for 965s is

A. 0.112L

B. 0.224L

C. 11.2*L*

D. 22.4L

Answer: a

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174. CH_3COOH is titrated with NaOH solution. Which of the following statements is true ?

A. Conductance decreases upto equivalence point, after which it increases.

- B. Conductance increases upto equivalence point,after which it decreases.
- C. Conductance first decreases (but not rapidly) and then increases

upto equibalence point and then increases rapidley after

equivalence point.

D. None of these

Answer: c

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175. The rusting of iron takes place as follows :

 $2H^{\oplus} + 2e^{-} + \frac{1}{2}O_2 \rightarrow H_2O(l), \quad E^{c^-} = +1.23V$ $Fe^{2^+} + 2e^- \rightarrow Fe(s), \quad E^{c^-} = -0.44V$

Calculae ΔG^{c-} for the net process.

A. - 322kJmol⁻¹

B. - 152*kJmol*⁻¹

C. - 76kJmol⁻¹

D. - 161*kJmol*⁻¹

Answer: a



176. The number of atoms of Ca that will be deposited from a solution of

 $CaCl_2$ by a current of 25mA for 60s will be

A. 4.68 \times 10^{18}

B. 4.68×10^{15}

 $C. 4.68 \times 10^{10}$

 $D. 2.34 \times 10^{15}$

Answer: a

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177. A reaction :

$$\frac{1}{2}H_2(g) + AgCl(s) \Leftrightarrow H^{\oplus}(aq) + Cl^{\Theta}(aq) + Ag(s)$$

occurs in a galvanic cell. The structure of the cell will be

A. Ag, AgCl(s)|KCl(sol)|AgNO₃(sol), Ag

B. Pt, $H_2(g)|HCL(sol)|AgNO_3(sol)$, Ag

C. Pt, $H_2(g)|HCl(sol)|AgCl(s)$, Ag

D. Pt, $H_2(g)|KCl(sol)|AgCl(s), Ag$

Answer: c



178. During the electrolysis of aqueous solution of *HCOOK*, the number of gases obtained at cathode, anode, and total number of gases are

A. 1, 2, 3

B. 1, 2, 2

C. 2, 1, 3

D. 2, 1, 2

Answer: b

Exerciseassertion -Reasoning

1. Assertion(A): Whne acidified $ZnSO_4$ solution is electrolyzed between Zn electrodes, it is Zn that is deposited at the cathode and $H_2(g)$ is not evolved.

Reason (R): The electrode potential of Zn is more negative than hydrogen as the overpotential for hydrogen evolution in Zn is quite large.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

B. If both (A) and (R) are correct, but (R) is not the correct explanation of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: C

⁽A).

2. Assertion (*A*): A saturated solution of *KCl* is used in making salt bridge.

Reason (*R*): Ionic mobilities of K^+ and Cl^- are comparable.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: A

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3. Assertion (*A*): When an aqueous solution of KNO_3 is electrolyzed, potassium is liberated at the cathode.

Reason (R): K^+ ions are discharged at cathode and NO_3^- at anode.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: D

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4. Assertion (A): For four half - cell reactions involving different number

of electrons,

 $E_4 = E_1 + E_2 + E_3$

Reaction (R): $\Delta G_4 = \Delta G_1 + \Delta G_2 + \Delta G_3$

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: D

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5. Assertion (A): 1 Faraday of electricity deposits 1g equivalent of Ag, Cu

or Al.

Reason (R): $1mo \le of$ electrons are required to reduce $1moleofAg^{\oplus}$ or $\frac{1}{2}moleofCu^{2+}$ or $\frac{1}{3}$ mole of Al^{3+} ions.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: B

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6. Assertion (A): The mobility of Na^{\oplus} is lower than that of K^{\oplus} ion.

Reason (R): The ionic mobility depends upon the effective radius of the ion.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: C

Watch Video Solution

7. Assertion (*A*): Equivalent conductance increase with dilution for an electrolyte solution.

Reason (R): The number of ions per litre of electrolyte increases with dilution.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: A

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8. Assertion (A):
$$\Lambda_m(H^{\oplus})$$
 and $\Lambda_m(C^{\circ}_{OH})$ ions are very much higher than

those of other ions.

Reason (R): It is due to proton jump from one water molecule to another resulting in a more rapid transfer of positive charge form one region to another.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

- B. If both (A) and (R) are correct, but (R) is not the correct explanation
 - of (*A*).
- C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: A



9. Assertion (*A*): Ionic conductivities increase with increase of temperature and pressure.

Reason (R): Viscosity of water decreases with increasew of temperature and increases with the increases of pressure.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

- B. If both (A) and (R) are correct, but (R) is not the correct explanation of (A).
- C. If (A) is correct, but (R) is incorrect.
- D. If (A) is incorrect, (R) is correct.

⁽A).

Answer: D

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10. Assertion (A: Cell constant is the *EMF* of a cell.

Reason (R): Cell constant is determined by using saturated KCl solution.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: D

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11. Assertion (A): At the end of electrolysis using Pt electrodes, an aqueous solution of $CuSO_4$ turns colourless.

Reason (R): $CuSO_4$ chnages to $Cu(OH)_2$ during electrolysis.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: c

Watch Video Solution

12. Assertion (A): Cu liberates $H_2(g)$ from a dilute solution of HCl.

Reason (R): Hydrogen is below Cu in the electrochemical series.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If both (A) and (R) are incorrect.

Answer: e

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13. Assertion(A): Na^{\oplus} ions are discharged in preference to H^{\oplus} ions at Hg cathode.

Reason (R): The nature of the cathode can affect the order of discharge of ions.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: a

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14. Assertion (*A*): *Cu* gets readily corroded in acidic aqueous solution.

Reason (R): Free energy of the process is positive.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If both (A) and (R) are incorrect.

Answer: e



15. Assertion (*A*): Galvanized iron does not rust.

Reason (R): Zn has a more negative electrode potential than Fe.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: a

16. Assertion (A): For a Daniell cell :

 $Zn \left| Zn^{2+} \right| \left| Cu^{2+} \right| Cu$ with $E_{cell} = 1.1V$, the application of opposite potential greater than 1.1V results into the flow of electron from cathod to anode.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: b



17. Assertion (A): The electrolysis of NaCl solution gives $H_2(g)$ at cathode

and $Cl_2(g)$ at anode.

Reason (R): Cl_2 has higher oxidation potential than H_2O

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: c

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18. Assertion (A): Fe is protected from corrosing by connecting Mg metal

with it.

Reason (R): Fe acts as cathode and Mg as anode which gradully

disappears.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: a

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19. Assertion (A): The Daniell cell becomes dead after sometimes.

Reason (R): The oxidation protential of Zn anode decreases and that of Cu increases.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: a

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20. Assertion (A): $\Lambda \circ \cdot_{eq}(CH_3COOH)$ cannote be determines experimentally.

Reason (R): CH_3COOH is a weak acid and DebyeHuckel Onsager equation cannot be used. Extrapolation method cannot be employed.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: a

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21. Assertion (A): The presence of CO_2 in the air accelerates corrosion.

Reason (R): CO_2 is a poisonous gas.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: c

22. Assertion (A): $(H_2 + O_2)$ fuel cell gives a constant voltages throughout its life.

Reason (*R*): In this fuel cell, H_2 reacts with OhH ions, yet the over all $\begin{bmatrix} c - \\ OH \end{bmatrix}$ does not change.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (A).

- C. If (A) is correct, but (R) is incorrect.
- D. If (A) is incorrect, (R) is correct.

Answer: a

23. Assertion (*A*): In a Daniell cell, if the concentration of Cu^{2+} and Zn^{2+} ions are doubled, the *EMF* of the cell will be doubled.

Reason (R): If the concentration of ions in contact with metals is doubled, the electrode potential is doubled.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: d



24. Assertion (*A*): The ration of specific conductivity to the observed conductance does not depend upon the concentration of the solution taken in the conductivity cell.

Reason (R): Specific conductivity decreases with dilution whereas observed conductance increases with the dilution.

A. If both (A) and (R) are correct, and (R) is the correct explanation of

(A).

B. If both (A) and (R) are correct, but (R) is not the correct explanation of (A).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: b

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25. Assertion(*A*): The rusting on the surface of iron involves following reaction :

 $a. Fe(s) \rightarrow Fe^{2+}(aq) + 2e^{-}$ (at anodic site) $b. O_2(g) + 4H^{\oplus}(aq) + 4e^{-} \rightarrow 2H_2O(l)$ (at cathodic site) $c. 4Fe^{2+}(aq) + O_2(g) + 4H_2O(l) \rightarrow 2Fe_2O_3(s) + 8H^{\oplus}$ $d. Fe_2O_3(aq) + xH_2O(l) \rightarrow Fe_2O_3. xH_2O$

Reason (R): Rusting is accelerated in the presence of NaCl and CO_2

A. If both (A) and (R) are correct, and (R) is the correct explanation of

B. If both (A) and (R) are correct, but (R) is not the correct explanation

of (*A*).

C. If (A) is correct, but (R) is incorrect.

D. If (A) is incorrect, (R) is correct.

Answer: b


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2. How many Faradays are required to reduce $1 molof BrO_3^{c-}$ to Br^{c-} in

basic medium ?

- **3.** The total number of Faradays required to oxidize the following separately:
- a. $1 molof S_2 O_3^{2-}$ in acid medium
- b. 1 Equivalent of $S_2 O_3^{2^-}$ in acid medium
- c. $1 molof S_2 O_3^{2-}$ in basic medium.

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4. For the oxidation of ferric oxalate to CO_2 , 18F of electricity is required.

How mnay moles of ferric oxalate is oxidized ?

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5. During the discharge of a lead storage battery, the density of $40 \% H_2SO_4$ by weight fell from 1.225 to 0.98(which is 20 % by weight). What is the change in molarities of H_2SO_4 ?

6. In Question above, the number of ampere hours for which the battery is used containing 1L of the acid is 16.08x ampere hour. Calculate the value of x.

7.
$$\Delta G$$
 for the reaction :

$$\frac{4}{3}Al + O_2 \rightarrow \frac{2}{3}Al_2O_3$$

is -772kJmol⁻¹ of O_2 .

-

Calculate the minimum EMF in volts required to carry out an electrolysis

of Al_2O_3

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8. When electrolysis of KCl is doen in alkaline medium, 10g of $KClO_3$ is produced as follows :

$$Cl^{c-} + 6OH \rightarrow ClO_3 + 3H_2O + 6e^{-}$$





1. When an aqueous solution of LiCl is electrolyzed using graphite electrodes, as the current flows, the pH of the solution around cathode

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2. Given $E^{c^-} \cdot Fe^{2^+} | Fe$ and $E^{c^-} \cdot Fe^{3^+} | Fe^{2^+}$ are -0.44 and 0.77V respectively. If Fe^{2^+}, Fe^{3^+} and Fe blocks are kept together, then $\left[Fe^{3^+}\right]$ and $\left[Fe^{2^+}\right]$

vvalui	VIUCU	Solution	

3. A certain current liberates 0.504g of $H_2(g)$ in 2 hours. The weight of Cu

deposited by same current flowing for the same time in $CuSO_4$ solution is

......g.

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4. Number of Faradays required to reduce 3mol of MnO_4^{c-} to MnO^{2+} is

.....F

Watch Video Solution

5. The gas *X* at 1*atm* is bubbled through a solution containing a mixture of $1MY^{c-}$ and $1MZ^{c-}$ at 25 °*C*. If the order of reduction potential is Z > Y > X, then

6. The *EMF* of the cell :

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7. The electrochemical equivalent for zinc (atomic weight = 6.4) is

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8. Given :

 $Fe^{2+}(aq) + 2e^- \rightarrow Fe(s), \quad E^{c-} = -0.44V$

 $Al^3 + 3e^- \rightarrow Al(s), \quad E^{c-} = -1.66V$

 $Br^{2+} + 2e^- \rightarrow 2Br^{c-}(aq) \quad E^{c-} = -1.08V$

The decreasing order of reducing power is



> Watch Video Solution

11. 13g of metal M is deposited at cathode by passing 0.4F of electricity .

The cathodic reaction is :

 $M^{n^+} + ne^- \rightarrow M$

The formula of metal chloride (Aw = 65) is



12. If the temperature coefficient $\left(\frac{\partial E}{\partial T}\right)$ is zero for a cell reaction then

out of ΔS , ΔH , and ΔG , the is zero.

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13. The ionization constant (K_a) of a weak electrolyte is 2.5×10^{-7} , while

 $\Lambda_{\it eq}$ of its 0.01M solution is 19.65Scm^2eq^{-1} \wedge °._{\it eq} is

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14. When dilute H_2SO_4 is electrolyzed between Pt electrodes, the gas

liberated at the anode will be.....





21. Cl_{20} is stronger agent then Br_2 and I_2 as its
reduction potential is than that of Br_2 and I_2
Vatch Video Solution
22. The Nernst equation gives <i>EMF</i> of the cell and in a Deniell
cell current flows from to to
Watch Video Solution
23. Dry cell is a cell and lead storage cell is a cell.
Watch Video Solution
24. The density of H_2SO_4 in lead storage cell during discharging and calomel electrode is a electrode.



3. Sodium cannot be obtained by the electrolysis of aqueous solution of

NaCl using *Pt* electrodes.



4. The standard potential of $Cl^{c-}|AgCl|Ag$ half cell is related to that of

 $Ag \oplus |Ag$ though the expression.

 $E^{c} \cdot Ag \oplus |Ag = E^{c} \cdot Cl^{c} |AgCl|A + \frac{RT}{F} \ln K_{sp}(AgCl)$

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5. In a Daniell cell, electrons flow from zinc electrode to copper electrode outside the cell.



6. Compounds of active metals (Zn, Na, Mg) are reducible by H_2 whereas

those of noble metals (Cu, Ag, Au) are not reducible.



discharged at the electrodes.

10. Cations having more negative potential than -0.828V are reduced in

preference to water.

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11. At the anode, the species having minimum reduction potential is formed form the oxidation of corresponding oxidizable species.

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12. In a galvanic cell, the half cell with higher potential provides a reducing agent.



13. In highly alkaline medium, the anodic process during the electrolytic

process is

с- $4OH \rightarrow O_2 + 2H_2O + 4^-$

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14. Metallic anodes smore reactive than platinum tend to pass into the solution instead of O_2 being produced.

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15. The mass of a substance deposited on the cathode or anode during electrolysis is given as m = QM/Fz.

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16. The cell potential becomes half if the cell reaction is divided by 2 througout.

17. Rusting of Fe is quicker in saline water than in ordinary water.

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18. F_2 is prepared by the electrolysis of molten KHF_2 and anhydrous HF.
Watch Video Solution

19. In Hall's process for the preparation of *Al*, graphite anode has a long life.

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20. In Down's process for the manufacture of sodium, CaCl₂, is added to

increase its melting point.

21. In Castner - Kellner cell for the manufacture of NaOH(caustic soda

), *Hg* acts as cathode as well as anode.



22. The amount of charge carried by an electrone is the same as carried by a K^{\oplus} ion.

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23. The same quantity of electricity is passed through $Al_2(SO_4)_3$ and $AgNO_3$ solution with platinum electrodes. If *n* number of *Al* atoms are deposited on the cathode, 3n number of *Ag* atoms will be deposited on the cathode.







28. Lead storage battery has anode and cathode made up of Pb.

29. Electrode potential for the electrode $M^{n+} \mid M$ with concerntration is

given by the expression under STP conditions :

$$E = E^{c-} + \frac{0.059}{n} \log \left[M^{n+} \right]$$

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30. In the electroplating of silver, $AgNO_3$ solution is usually used as an electrolyte.

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31. The conductance of electrolyte solution increases wih temperature.

32. Resistivity is reciprocal of molar conductivity of electrolyte.

Watch Video Solution
33. Cell constant has unit m^{-1} .
Watch Video Solution
34. The conductivity of molten <i>KCL</i> is due to the movement of K^{\oplus} and
Cl^{c-} ions.
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35. Solid *KCl* is a good conductor of electricity.

36. Molten Na_2SO_4 is a good conductor because of mobile electrons.

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37. Cathode is negative terminal both in electrochemical and electrolytic

cells.

Watch Video Solution

38. Reduction occurs at cathdooe both in galvanic as well as in electrolytic cell.

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39. The chemical change in an electrolytic cell is non - spontaneous.

40. The cell voltage is independent of the size of the cell or electrodes.

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Archieves (Linked Comprehension)

1. Tollen reagent is used for the detection of aldehydes. When a solution of $AgNO_3$ is added to glucose with NH_4OH , then gluconic acid is formed. $Ag^{\oplus} + e^- \rightarrow Ag, \quad E^{c^-} \cdot_{red} = 0.8V$ $C_6H_{12}O_6 \rightarrow C_6H_{12}O_7 + Gluconicacid2H^{\oplus} + 2e^-, \quad E^{c^-} \cdot_{oxid} = -0.05V$ $\left[Ag(NH_3)_2\right]^{\oplus} + e^- \rightarrow Ag(s) + 2NH_3, \quad E^{c^-} \cdot_{red} = 0.337V$ $\left[Use2.303 \times \frac{RT}{F} = 0.0592 \text{ and } \frac{F}{RT} = 38.92at298K\right]$ $2Ag^{\oplus} + C_6H^{12}O_6 + H_2O \rightarrow 2Ag^s + C_6H_{12}O_7 + 2H^{\oplus}$ Find $\ln K$ of this reaction.

A. 66.13

B. 58.38

C. 28.30

D. 46.29

Answer: b

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2. Tollen reagent is used for the detection of aldehydes. When a solution of $AgNO_3$ is added to glucose with NH_4OH , then gluconic acid is formed. $Ag^{\oplus} + e^- \rightarrow Ag$, $E^{c^-} \cdot_{red} = 0.8V$ $C_6H_{12}O_6 \rightarrow C_6H_{12}O_7 + Gluconicacid2H^{\oplus} + 2e^-$, $E^{c^-} \cdot_{oxid} = -0.05V$ $\left[Ag(NH_3)_2\right]^{\oplus} + e^- \rightarrow Ag(s) + 2NH_3$, $E^{c^-} \cdot_{red} = 0.337V$ $\left[Use2.303 \times \frac{RT}{F} = 0.0592$ and $\frac{F}{RT} = 38.92at298K\right]$ When ammonia is added to the solution, *pH* is raised to 11. Which half

cell reaction is affected by pH and by how much ?

A. E_{oxid} will increase by a factor of 0.65 from E^{c-} . $_{oxid}$.

B. E_{oxid} will decrease by a factor of 0.65 from E^{c} . $_{oxid}$.

C. E_{red} will increase by a factor of 0.65 from E^{c-} .

D. E_{red} will decrease by a factor of 0.65 from E^{c-} .

Answer: c

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3. Tollen reagent is used for the detection of aldehydes. When a solution of $AgNO_3$ is added to glucose with NH_4OH , then gluconic acid is formed. $Ag^{\oplus} + e^- \rightarrow Ag, \quad E^{c^-} \cdot_{red} = 0.8V$ $C_6H_{12}O_6 \rightarrow C_6H_{12}O_7 + Gluconicacid2H^{\oplus} + 2e^-, \quad E^{c^-} \cdot_{oxid} = -0.05V$ $\left[Ag\left(NH_3\right)_2\right]^{\oplus} + e^- \rightarrow Ag(s) + 2NH_3, \quad E^{c^-} \cdot_{red} = 0.337V$ $\left[Use2.303 \times \frac{RT}{F} = 0.0592 \text{ and } \frac{F}{RT} = 38.92at298K\right]$

Ammonia is always added in this reaction. Which of the followijng must be wrong ?

A. NH_3 combines with Ag^{\oplus} to form a complex.

B. $\left[Ag(NH_3)_2\right]^{\oplus}$ is a stronger oxidizing reagent than Ag^{\oplus}

C. In the absence of NH_3 , a silver salt of gluconic acid is formed.

D. NH_3 has affected the standard reduction potential of glucose /

gluconic acid electrode.

Answer: d

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4. Chemical reactions involve interation of atoms and molecules. A large number of atoms / molecules (approximately 6.023×10^{23}) are present in a few grams of any chemical compound varying with their atomic / molecular masses. To handle such large numbers conveniently, the mole concept was introduced. This concept has implications in diverse areas such as analytical chemistry, biochemistry, electrochemistry, and radiochemistry. The following example illustrates a typical case, involving chemical / electrochemical reaction, which requires a clear understanding of the mole concept.

A 4.0M aqueous solution of NaCl is prepared and 500mL of this solution

is electrolyzed. This leads to the evolution of chlorine gas at one of the electrodes (atomic mass of Na is 23 and Hg is 200)(1F = 96500C). The total number of moles of chlorine gas evolved is

A.	0	•	5	

- B. 1.0
- **C**. 2.0
- D. 3.0

Answer: b

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5. Chemical reactions involve interation of atoms and molecules. A large number of atoms / molecules (approximately 6.023×10^{23}) are present in a few grams of any chemical compound varying with their atomic / molecular masses. To handle such large numbers conveniently, the mole concept was introduced. This concept has implications in diverse areas such as analytical chemistry, biochemistry, electrochemistry, and

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A 4.0*M* aqueous solution of *NaCl* is prepared and 500*mL* of this solution is electrolyzed. This leads to the evolution of chlorine gas at one of the electrodes (atomic mass of *Na* is 23 and *Hg* is 200)(1*F* = 96500*C*). If the cathode is an *Hg* electrode, the maximum weight ($\in g$) of

amalgam formed from this solution is

A. 200

B. 225

C. 400

D. 446

Answer: d

6. Chemical reactions involve interation of atoms and molecules. A large number of atoms / molecules (approximately 6.023×10^{23}) are present in a few grams of any chemical compound varying with their atomic / molecular masses. To handle such large numbers conveniently, the mole concept was introduced. This concept has implications in diverse areas such as analytical chemistry, biochemistry, electrochemistry, and radiochemistry. The following example illustrates a typical case, involving chemical / electrochemical reaction, which requires a clear understanding of the mole concept.

A 4.0*M* aqueous solution of *NaCl* is prepared and 500*mL* of this solution is electrolyzed. This leads to the evolution of chlorine gas at one of the electrodes (atomic mass of *Na* is 23 and *Hg* is 200)(1*F* = 96500*C*). The total charge (coulomb) required for complete electrolysis is

A. 24125

B. 48250

C. 96500

D. 193000

Answer: d

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7. Redox reactions play a pivotal role in chemistry and biology. The values standard redox potential (E^{c-}) of two half cell reactions decided which way the reaction is expected to preceed. A simple example is a Daniell cell in which zinc goes into solution and copper sets deposited. Given below are a set of half cell reactions (acidic medium) along with their $E^{c-}(V$ with respect to normal hydrogen electrode) values. Using this data, obtain correct explanations for Question.

$$\begin{split} &I_2 + 2e^- \rightarrow 2I^{c^-}, \quad E^{c^-} = 0.54 \\ &Cl_2 + 2e^- \rightarrow 2Cl^{c^-}, \quad E^{c^-} = 1.36 \\ &Mn^{3^+} + e^- \rightarrow Mn^{2^+}, \quad E^{c^-} = 1.50 \\ &Fe^{3^+} + e^- \rightarrow Fe^{2^+}, \quad E^{c^-} = 0.77 \\ &O_2 + 4H^{\oplus} + 4e^- \rightarrow 2H_2O, \quad E^{c^-} = 1.23 \end{split}$$

Among the following, identify the correct statement.

A. Chloride ion is oxidized by O_2 .

B. Fe^{2+} is oxidized by iodine.

C. lodide ion is oxidized by chlorine

D. Mn^{2+} is oxidized by chlorine.

Answer: c

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8. Redox reactions play a pivotal role in chemistry and biology. The values standard redox potential (E^{c-}) of two half cell reactions decided which way the reaction is expected to preceed. A simple example is a Daniell cell in which zinc goes into solution and copper sets deposited. Given below are a set of half cell reactions (acidic medium) along with their $E^{c-}(V$ with respect to normal hydrogen electrode) values. Using this data, obtain correct explanations for Question.

 $I_2 + 2e^- \rightarrow 2I^{c^-}, \quad E^{c^-} = 0.54$ $Cl_2 + 2e^- \rightarrow 2Cl^{c^-}, \quad E^{c^-} = 1.36$ $Mn^{3^+} + e^- \rightarrow Mn^{2^+}, \quad E^{c^-} = 1.50$ $Fe^{3^{+}} + e^{-} \rightarrow Fe^{2^{+}}, \quad E^{c^{-}} = 0.77$ $O_{2} + 4H^{\oplus} + 4e^{-} \rightarrow 2H_{2}O, \quad E^{c^{-}} = 1.23$ While $Fe^{3^{+}}$ is stable, $Mn^{3^{+}}$ is not stable in acid solution because A. O_{2} oxidizes $Mn^{2^{+}}$ to $Mn^{3^{+}}$ B. O_{2} oxidizes both $Mn^{2^{+}}$ to $Mn^{3^{+}}$ and $Fe^{2^{+}}$ to $Fe^{3^{+}}$ C. $Fe^{3^{+}}$ oxidizes $H_{2}O$ to O_{2} D. $Mn^{3^{+}}$ oxidized $H_{2}O$ to O_{2}

Answer: d

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9. Redox reactions play a pivotal role in chemistry and biology. The values standard redox potential (E^{c-}) of two half cell reactions decided which way the reaction is expected to preceed. A simple example is a Daniell cell in which zinc goes into solution and copper sets deposited. Given below are a set of half cell reactions (acidic medium) along with their $E^{c-}(V$ with respect to normal hydrogen electrode) values. Using this data,

obtain correct explanations for Question.

$$\begin{split} I_{2} + 2e^{-} &\rightarrow 2I^{c^{-}}, \quad E^{c^{-}} = 0.54 \\ Cl_{2} + 2e^{-} &\rightarrow 2Cl^{c^{-}}, \quad E^{c^{-}} = 1.36 \\ Mn^{3^{+}} + e^{-} &\rightarrow Mn^{2^{+}}, \quad E^{c^{-}} = 1.50 \\ Fe^{3^{+}} + e^{-} &\rightarrow Fe^{2^{+}}, \quad E^{c^{-}} = 0.77 \\ O_{2} + 4H^{\oplus} + 4e^{-} &\rightarrow 2H_{2}O, \quad E^{c^{-}} = 1.23 \end{split}$$

Sodium fusion extract obtained from aniline on treatment with iron (II) sulphate and H_2SO_4 in the presence of air gives a Prussion blue precipitate. The blue colour is due to the formation of

A.
$$Fe_4 [Fe(CN)_6]_3$$

B. $Fe_3 [Fe(CN)_6]_2$
C. $Fe_4 [Fe(CN)_6]_2$
D. $Fe_3 [Fe(CN)_6]_3$

Answer: a

10. The concentration of potassium ions inside a biological cell is at least 20 times higher than outside. The resulting potential difference across the cell is important in several processes such as transmission of nerve impulses and maintaining the ion balance. A simplel model for a concentration cell involving a metal *M* is

$$M(s) \mid M^{\oplus}(aq, 0.05 \text{ molar}) \mid M^{\oplus}(aq, 1 \text{ molar}) \mid M(s)$$

For the abov electrolytic cell, the magnitude of the cell potential is $\left|E_{cell}\right| = 70 mV.$

For the above cell

A.
$$E_{cell} < 0, \Delta G > 0$$

B. $E_{cell} > 0, \Delta G < 0$
C. $E_{cell} < 0, \Delta G^{c-} > 0$
D. $E_{cell} > 0, \Delta G^{c-} < 0$

Answer: b

11. The concentration of potassium ions inside a biological cell is at least 20 times higher than outside. The resulting potential difference across the cell is important in several processes such as transmission of nerve impulses and maintaining the ion balance. A simplel model for a concentration cell involving a metal *M* is

 $M(s) \mid M^{\oplus}(aq, 0.05 \text{ molar}) \mid M^{\oplus}(aq, 1 \text{ molar}) \mid M(s)$

For the abov electrolytic cell, the magnitude of the cell potential is $\left|E_{cell}\right| = 70 mV.$

If the 0.05 moolar solution of M^{\oplus} is replaced by a 0.0025 molar M^{\oplus} solution, then the magnitude of the cell potential would be

A. 35mV

B. 70mV

C. 140mV

D. 700mV

Answer: c

12. The electrochemical cell shown below is a concentration cell. $M \mid M^{2+}($ saturated solution of sparingly soluble salt, $MX_2 \mid \left| M^{2+} \left(0.001 moldm^{-3} \right) \right| M$

The *emf* of the cell depends on the difference in the concentration of M^{2+} ions at the two electrodes. The *emf* of the cell at 298 is 0.059V.

The solubility product $(K_{sp}, mol^3 dm^{-9})$ of MX_2 at 298 based on the information available the given concentration cell is (Take $2.303 \times R \times 298/F = 0.059V$)

A. 1×10^{-15}

B. 4×10^{-15}

 $C.1 \times 10^{-12}$

D. 4×10^{-12}

Answer: b
13. Consider $Zn + Cu^{2+} \rightarrow Zn^{2+} + Cu$

If the standard emf is $E_{cell} = 2.0V\&F = 96500C$

Find $\Delta G^{\circ}(KJmol)$

A. - 5.7

B. 5.7

C. 11.4

D. - 11.4

Answer: d

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Archieves Multiple Correct Ansers

1. For the reduction of $NO_3^{c^-}$ ion in an aqueous solution, E^{c^-} is +0.96V, the values of E^{c^-} for some metal ions are given below :

 $i. V^{2^+}(aq) + 2e^- \rightarrow V, \quad E^{c^-} = -1.19V$

ii. $Fe^{3+}(aq) + 3e^- \rightarrow Fe$, $E^{c-} = -0.04V$ iii. $Au^{3+}(aq) + 3e^- \rightarrow Au$, $E^{c-} = +140V$ iv. $Hg^{2+}(aq) + 2e^- \rightarrow Hg$, $E^{c-} = +0.86V$

The pair(s) of metals that is / are oxidized by NO_3^{c-1} in aqueous solution is

/ are

A. Fe and Au

B. Hg and Fe

C. V and Hg

D. Fe and V

Answer: b,c,d

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2. In a galvanic cell, the salt bridge.

A. Does not participate chemically in the cell reaction

B. Stops the diffusion of ions from one electrode to another

C. Is necessary for the occurrence of the cell reaction

D. Ensures mixing of the two electrolytic solutions

Answer: a,b,c

D Watch Video Solution

Archieves Single Correct

1. The standard reduction potentials at 298K for the following half reactions are given against each

Which is the strongest reducting agent?

A. Zn(s)

B. *Cr*(*s*)

 $C. H_2(s)$

D. $Fe^{2+}(aq)$

Answer: a



Answer: c



3. A solution containing 1*mol* per litre of each $Cu(NO_3)_2$, $AgNO_3$, and $Hg_2(NO_3)_2$ is being electrolyzed by using inert electrodes. The values of standard electrode potentials in volts (reduction potential) are

$$Ag | Ag^{\oplus} = +0.80, 2Hg | Hg_2^{2+} = +0.79$$
$$Cu | Cu^{2+} = +0.34, Mg | Mg^{2+} = -2.37.$$

With increasing voltage, the sequence of deposition of metals at the cathode will be

A. Ag, Hg, Cu, Mg

B. Mg, Cu, Hg, Ag

C. Ag, Hg, Cu, Mg

D. Cu, Hg, Ag, Mg

Answer: a

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4. The electrice charge for eeletrode deposition of 1g equivalent of a substance is

A. 1 ampere per second

B. 96500 Coulombs per second

- C. 1 ampere for one hour
- D. The charge on one mole of electrons

Answer: d

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5. The reaction

 $1/2Hg(g) + AgCl(s) \rightarrow H^{\oplus}(aq) + Cl^{c^-}(aq) + Ag(s)$ occurs in the galvanic cell.

A.
$$Ag|AgCl(s)|KCl(sol)|AgNO_3|Ag$$

B. $Pt|H_2(g)|HCl(sol)|AgNO_3(SOL)|Ag$
C. $Pt|H_2(g)|HCl(sol)|AgCl(s)|Ag$
D. $Pt|H_2(g)|KCl(sol)|AgCl(s)|Ag$

Answer: c

6. When a lead storage battery is discharged.

A. SO_2 is evolved

B. Lead is formed

C. Lead sulphate is consumed

D. Sulphuric acid is consumed

Answer: d

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7. The standard reductino potentials E^{c-} for the half reactinos are as

follows:

 $Zn \rightarrow Zn^{2+} + 2e^{-} E^{c-} = +0.76V$

 $Fe \rightarrow Fe^{2+} + 2e^- \quad E^{c-} = 0.41V$

The EMF for the cell reaction

 $Fe^{2+} + Zn \rightarrow Zn^{2+} + Fe$ is A. -0.35V B. +0.35V

C. +1.17V

D. - 1.17V

Answer: b

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8. The standard reduction potentials of $Cu^{2+} | Cu$ and $Cu^{2+} | Cu^{\oplus}$ are 0.337V and 0.153V, respectively. The standard electrode potential fo $Cu^{\oplus} | Cu$ half cell is

A. 0.184V

B.0.827V

C. 0.521V

D. 0.490V

Answer: c



9. The standard reduction potential values of three metallic cations, X, Y, and Z are 0.52, - 3.03, and -0.18V, respectively. The order of reducing power of the corresponding metal is

A. Y > Z > X

 $\mathsf{B}.\, X > Y > Z$

 $\mathsf{C}.\, Z > Y > X$

D. Z > X > Y

Answer: a

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10. The gas*X* at 1atm is bubbled through a solution containing a mixture of $1MY^{c-}$ and $1MZ^{c-}$ at $25 \degree C$. If the order of reduction potential is Z > Y > X, then

A. Y will oxidize X and not Z

B. Y will oxidize Z and not X.

C. *Y* will oxidize both *X* and *Z*.

D. Y will reduct both X and Z.

Answer: b

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11. For the electrochemical cell,

$$(M | M^{\oplus}) | | (X^{c-} | X), E^{c-} \cdot (M^{\oplus} | M) = 0.44V$$
 and
 $E^{c-} \cdot (X | X^{c-}) = 0.334V$
A. $M + X \rightarrow M^{\oplus} + X^{c-}$ is a spontaneous reaction.

B. $M^{\oplus} + X^{c-} \rightarrow M + X$ is the spontaneous reaction

 $C. E_{cell} = 0.77V$

D. $E_{cell} = -0.77$

Answer: b

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12. Arrange the following compounds in the order of increasing conductance : *HCl*, *LiCl*, *NaCl*, *KCl*.

A. LiCl > NaCl > KCl

B. KCl > NaCl > LiCl

C. NaCl > KCl > LiCl

D. LiCl > KCl > NaCl

Answer: b

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13. A standard solution of KNO_3 is used to make salt bridge, because

A. The velocity of K^{\oplus} is greater than that of NO_3^{\oplus} .

B. The velocity of NO_3^{\oplus} is greater than that of K^{\oplus}

C. The volocities of K^{\oplus} and NO_3^{\oplus} same

D. KNO₃ is highly solubel in water.

Answer: c



14. Standard electrode potential data are useful for understanding the suitability of an oxidant in a redox titration. Some half cell reaction and their standard potentials are given below:

$$MnO_{4}^{-}(aq) + 8H^{+}(aq) + 5e^{-} \rightarrow Mn^{2+}(aq) + 4H_{2}O(l)E^{\circ} = 1.51V$$

$$Cr_{2}O_{7}^{2-}(aq) + 14H^{+}(aq) + 6e^{-} \rightarrow 2Cr^{3+}(aq) + 7H_{2}O(l), E^{\circ} = 1.38V$$

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

$$CI_{2}(g) + 2e^{-} \rightarrow 2CI^{-}(aq), E^{\circ} = 1.40V$$

Identify the only correct statement regarding quantitative estimation of aqueous $Fe(NO_3)_2$

A. MnO_4^{c-} can be used in aqueous HCl

B. CrO_4^{2-} can be used in aqueous HCl

C. MnO_4^{c-} can be used in aqueous H_2SO_4

D. $Cr_2O_7^{2-}$ can be used in aqueous H_2SO_4

Answer: a

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15. In an electrolytic cell, the flow of electrons is form

A. Cathode to anode in solution

B. Cathode to anode through external supply

C. Cathode to anode through internal supply

D. Anode to cathode through internal supply.

Answer: c



16.
$$Zn \left| Zn^{2+} (a = 0.1M) \right| \mid Fe^{2+} (a = 0.01M)Fe.$$

The EMF of the above cell is 0.2905. The equilibrium constant for the cell reaction is

A. $10^{0.32/0.0591}$

B. 10^{0.32/0.0295}

C. $10^{0.26/0.0295}$

D. $e^{0.32/0.2995}$

Answer: b



17. The rusting of iron takes place as follows :

$$2H^{\oplus} + 2e^{-} + \frac{1}{2}O_2 \rightarrow H_2O(l), \quad E^{c^-} = +1.23V$$

 $Fe^{2^+} + 2e^- \rightarrow Fe(s), \quad E^{c^-} = -0.44V$

Calculae ΔG^{c-} for the net process.

A. - 322*kJmol*¹

B. - 161*kJmol*⁻¹

C. - 152kJmol⁻¹

D. - 76*kJmol*⁻¹

Answer: a

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18. Electrolysis of dilute aqueous *NaCl* solution was carried out by passing 10mA current. The time required to liberate 0.01mol of H_2 gas at the cathode is $(1F = 96500Cmol^{-1})$

A. $9.65 \times 10^4 s$ B. $19.3 \times 10^4 s$ C. $28.95 \times 10^4 s$ D. $38.6 \times 10^4 s$

Answer: b

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19. Consider the following cell reation :

$$2Fe(s) + O_2(g) + 4H^{\oplus}(aq) \rightarrow 2Fe^{2+}(aq) + 2H_2O(l) \quad E^{c-} = 1.67V$$
$$At\left[Fe^{2+}\right] = 10^{-3}M, p\left(O_2\right) = 0.1atm \text{ and } pH = 3.$$

The cell potential at 25 $^{\circ}C$ is

A. 1.47V

 $B.\,1.77V$

C. 1.87V

D. 1.57V

Answer: d



20. $AgNO_2(aq)$ was added to an aqueous *KCl* solution gradually and the conductivity of the solution was measured. The plot conductivity of the solution was measured. The plot of conductance (Λ) versus the volume of $AgNO_3$ is



A. P	
В. <i>Q</i>	
C. <i>R</i>	

D. S

Answer: d

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21. given $E_{S_2O_8^{2^-}/SO_4^{2^-} \land (\circ) = 2.05V} E_{Br_2/Br^-}^{\circ} = 1.40V E_{Au^{3^+}/Au}^{\circ} = 1.10V$, brgt $E_{O_2/H_2O}^{\circ} = 1.20V$ Which of the following is the strongest oxidizing agent ?

A. *Cr*³⁺

B. Mn^{2+}

 $C.MnO_4^{c}$

D. *Cl^c*-

Answer: c Watch Video Solution 22. The metal that cannot obtained by electrolysis of an aqueous solution of its salts is : A. Cu B. Cr C.AqD. Ca

Answer: d



23. The resistance of a conductivity cell filled with 0.1MKCl solution is 100Ω . If *R* of the same cell when filled with 0.02MKCl solution is 520Ω ,

calculate the conductivity and molar conductivity of 0.02MKCl solution. The conductivity of 0.1MKCl solution is $1.29Sm^{-1}$.

A. 5×10^{3} B. 5×10^{2} C. 5×10^{-4} D. 5×10^{-3}

Answer: c

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24. The equivalent conductance of *NaCl* at concentration of *C* and at infinite dilution are λ_C and λ_{∞} respectively. The correct relationship between λ_C and λ_{∞} is given as :

(where the constant B is positive)

A.
$$\lambda_C = \lambda_{\infty} - (B)\sqrt{C}$$

B. $\lambda_C = \lambda_{\infty} + (B)\sqrt{C}$

 $C. \lambda_C = \lambda_\infty + (B)C$

D. .λC=λ∞−(B)C

Answer: a

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25. Find the standard electrode potential of $MnO_4^{c-} \mid MnO_2$. The standard electrode potential of $MnO_4^{c-} \mid Mn^{2+} = 1.51V$ and $MnO_2 \mid MnO_2 \mid Mn^{2+} = 1.23V$.

A. -0.33V, the reaction will not occur

B. -0.33V, the reaction will occur

C. -2.69V, the reaction will not occur

D. - 2.69V, the reaction will occur

Answer: c

Archieves Fill in The Rian	
	5

1. Galvanization of iron denotes coating with

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2. The more the standard reduction potential, the

..... its ability to displace hydrogen from acids.

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3. The electrical conductivity of a solution of acetic acid will be if a solution of sodium hydroxide is added.

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Archieves Subjective

1. A current of 3.7 A is passed for 6hrs. Between Ni electrodes in 0.5 L of 2 M solution of $Ni(NO_3)_2$. What will be the molarity of solutionn at the end of electrolysis?

2. Consider the cell:

 $Zn \left| Zn^{2+}(aq)(1.0M) \right| \left| Cu^{2+}(aq)(1.0M) \right| | Cu$

Thee standard reduction potentials are 0.350V for

 $Cu^{2+}(aq) + 2e^{-} \rightarrow Cu$ and -0.763V for

$$Zn^{2^+}(aq) + 2e^- \rightarrow Zn$$

- a. Write the cell reaction.
- b. Calculate the EMF of the cell.
- c. Is the reaction spontaneous or not?



3. In an electrolysis experiment, current was passed for 5h through two cells connected in series. The first cell contains a solution of gold and second contains copper sulphate solution. In the first cell, 9.85*g* of gold was deposited. If the oxidation number of gold is +3, find the amount of copper deposited at the cathode of the second cell. Also calculate the

magnitude of the current in ampere, (Atomic weight of Au is 1197 and
atomic weight of <i>Cu</i> is 63.5).
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4. How long a current of 3A has to be passed through a solution of silver nitrate to coat a metal surface of $80cm^2$ with a 0.005 - mm - thick layer ? The density of silver is $10.5qcm^{-3}$.

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5. Give reasons in one or two sentences : anhydrous *HCl* is a bad conductor of electricity but aqueous *HCl* is a good conductor.

0	Watch Video Solution	ו

6. The *EMF* of the following cellis 1.05V at $25 \degree C$:

 $Pt, H_2(g)(1.0atm)|NaOH(0.1m), NaCl(0.1M)|AgCl(s), Ag(s)$

- a. Write the cell reaction,
- b. Calculate pK_w of water.

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7. During the discharge of a lead storage battery, the density of sulphuric acid fell from $1.294gmL^{-1}$ to $1.139gmL^{-}$. Sulphuric acid of density $1.294gmL^{-1}$ is 39 % by weight and that of density $1.139gmL^{-1}$ is 20 % by weight. The battery hold 3.5 litre of acied and discharge. Calculate the no. of ampere hour for which the battery must have been used. The charging and discharging reactions are:

$$Pb + SO_4^2 \rightarrow PbSO_4 + 2e$$
 (charging)

 $PbO_2 + 4H^+ + SO_4^{2-} + 2e \rightarrow PbSO_4 + 2H_2O$ (discharging)

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8. A 100 - W, 100 - V incardescent lamp is connected in series with an electrolyte cell containing cadmium sulphate solution. How much cadmium will be deposited by the current flowing for 10h?

9. A cell contains two hydrogen electrodes. The negative electrode is in contact with a solution of $10^{-6}M$ hydrogen ions. The *EMF* of the cell is 0.118V at 25 °C. Calculate the concentration of hydrogen ions at the positive electrode.

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10. In a fuel cell, hydrogen and oxygen react to produce electricity. In process, hydrogen gas is oxidized at the anode and oxygen at the cathode. If 67.2L of H_2 at *STP* reacts in 15min, what is the average current produced ? If the entire current is used for electro - deposition of copper from copper (*II*) solution, how many grams of copper will be deposited ?

Anode reaction : $H_2 = 2OH \rightarrow 2H_2O + 2e^-$

Cathode reaction : $O_2 + 2H_2O + 2e^- \rightarrow 4OH$

11. An acidic solution of Cu^{2+0} salt containing 0.4g of Cu^{2+} is electrolyzed until all the copper is deposited. The electrolysis is continued for seven more minutes with the volume of solution kept at 100mL and the current at 1.2A. Calculate the volume of gases evolved at *STP* during the entire electrolysis.

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12. The standard reduction potential at 25 ° C of the reaction

 Θ 2 $H_2O + 2e^- \Leftrightarrow H_2 + 2OH$ is -0.8277V. Calculate the equilibrium constant

for the reaction.

 $\begin{array}{c} \Theta \\ 2H_2O \Leftrightarrow H_3O^{\oplus} + OH \text{ at } 25 °C \,. \end{array}$

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13. The standard reduction potential of $Cu^{2+} | Cu$ and $Ag^{\oplus} | Ag$ electrodes are 0.337 and 0.799V, respectively. Construct a galvanic cell using these electrodes so that its standard *EMF* is positive. For what concentration of Ag^{\oplus} will the *EMF* of the cell , at 25 ° C, be zero if the concentration fo Cu^{2+} is 0.01*M*?

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14. Calculate the quantity of electricity that would be required to reduce 12.3g of nitrobenzene to aniline, if the current efficiency for the process is 50%. If the potential drop across the cell is 3.0V, how much energy will be consumed?

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15. Zinc granules are added in excess to 500 mL of 1M $Ni(NO_3)_2$ solution of 25 ° C untill the equilibrium is reached. If $E_{Zn^{2+}/Zn}^{\circ}$ and $E_{Ni^{2+}/Ni}^{\circ}$ are -0.75V and -0.24V respectively, find out the $[Ni^{2+}]$ at equilibrium. **16.** A current of 1.70*A* is passed trhough 300.0*mL* of 0.160*M* solution of $ZnSO_4$ for 230*s* with a current efficiency of 90 %. Find out the molarity of Zn^{2+} after the deposition of *Zn*. Assume the volume of the solution to remain constant during the electrolysis.

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17. For the galvanic cell: Ag|AgCI(s)|KCI(0.2M)||KBr(0.001M)|AgBr(s)|Ag, calculate the EMF generated and assign correct polarity to each electrode for a spontaneous process after taking into account the cell reaction at 25 ° *C*.

$$\left[K_{sp(AgCI)} = 2.8 \times 10^{-10}, K_{sp(AgBr)} = 3.3 \times 10^{-13}\right]$$

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18. An aqueous solution of *NaCl* on electrolysis gives $H_2(g)$, $Cl_2(g)$, and *NaOH* accroding to the reaction :

$$c^{-}$$

2 $Cl^{c^{-}}(aq) + 2H_2O \rightarrow 2OH(aq) + H_2(g) + Cl_2(g)$

A direct current of 25*A* with a current efficiency of 62 % is passed through 20*L* of *NaCl* solution (20 % by weight). Write down the reactions taking place at the anode and cathode. How long will it take to produce 1kg of Cl_2 ? What will be the molarity of the solution with respect to hydroxide ion ? (Assume no loss due to evaporation .)

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19. The standard reduction potential for the half cell :

 $NO_3(aq) + 2H^+ + e^- \rightarrow NO_2(g) + H_2O$ is 0.78V.

a. Calculate the reduction potential in $8MH^{\oplus}$.

b. What will be the reduction potential of the half cell in a neutral solution ? Assume all the other species to be at unit concentration.

20. Chromium metal can be plated out from an acidic solution containing

 CrO_3 according to the following equation :

 $CrO_3(aq) + 6H^{\oplus} + 6H^{\oplus}(aq) + 6e^- \rightarrow Cr(s) + 3H_2O$

a. How many grams of chromium will be plated out by 24000C?

b. How long will take to plate out 1.5*g* of chromium by using 12.5*A* current ?



21. The standard reduction potential of the $Ag \oplus |Ag|$ electrode at 298K is 0.799V. Given that for AgI, $K_{sp} = 8.7 \times 10^{-17}$, evaluate the potential of the $Ag \oplus |Ag|$ electrode in a saturated solution of AgI. Also calculate the standard reduction potential of the $I^{c-}|AgI|Ag$ electrode.

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22. The Edison storage cell is represented as :

 $Fe(s)|FeO(s)|KOH(aq)|Ni_2O_3(s)|Ni(s)$

The half - cell reactions are :

$$Ni_{2}O_{3}(s) + H_{2}O(1) + 2e^{-} \xrightarrow{\sim} 2NiO(s) + 2OH;$$

$$E^{\odot} = +0.40 V$$

$$FeO(s) + H_{2}O(1) + 2e^{-} \xrightarrow{\sim} Fe(s) + 2OH; E^{\odot} = -0.87 V$$

a. What is the cell reaction ?

b. What is the cell *EMF* ? How does it depend on the concentration of *KOH* ?

c. What is maximum amount of electrical energy that can be obtained from 1mol of Ni_2O_3 ?

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23. An excess of liquid mercury is added to an acidicfied solution of $1.0 \times 10^{-3} MFe^{3+}$. It is found that 5 % of Fe^{3+} remains at equilibrium at 25 °C. Calculate E^{C^-} . $(Hg_2^{2^+}|Hg)$ assuming that the only reaction that occurs is

 $2Hg + 2Fe^{3+} \rightarrow Hg_2^{2+} + 2Fe^{2+}$ Given $: E^{c-} \cdot (Fe^{3+} | Fe^{2+}) = 0.77V$ **24.** The standard reduction potential for Cu^{2+} | Cu is +0.34V. Calculate the reduction potential al pH = 14 for the above couple . K_{sp} of $Cu(OH)_2$ is 1.0×10^{-19}

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25. How many grams of silver could be plated out on a serving tray by the electrolysis of a solution containing silver in +1 oxidation state of a period of 8.0*h* at a current of 8.46*A* ? What is the area of the ray, if the thickness of the silver plating is 0.0254cm? The density of silver is 10.5 g cm⁽⁻³⁾.

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26. Calculate the equilibrium constant for the reaction : $Fe^{2^+} + Ce^{4^+} \Leftrightarrow Fe^{3^+} + Ce^{3^+}$ Given, $E_{Ca^{4^+}/Ce^{3^+}}^{\circ} = 1.44V$ and

$$E_{Fe^{3+}/Fe^{2+}}^{\circ} = 0.68V$$



27. Calculate the equilibrium constant for the reaction :

$$2Fe^{3+} + 3I^{\Theta} \Leftrightarrow 2Fe^{2+} + I_3^{\Theta}$$

The standard reduction potential in acidic conditions is 0.78V and 0.54V,

respectively, for Fe^{3+} | Fe^{2+} and I_3^{c-} | I^{c-} couples

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28. Find the solubility product of a saturated solution of Ag_2CrO_4 in

water at 298K, if the *EMF* of the cell :

$$Ag \left| Ag \oplus \left(satAg_2 CrO_4 sol \right) \right| |Ag(0.1M)| Agis 0.164V \text{ at } 298K.$$

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29. A cell, $Ag | Ag^{\oplus} | | Cu^{2+} | Cu$, initially contains $1MAg^{\oplus}$ and $1MCu^{2+}$ ions. Calculate the change in the cell the potential after the passage of 9.65A of current for 1h.



30. Copper sulphate solution (250*ML*) was electrolyzed using a platinum anode and a copper cathode. A constant current of 2mA was passed for 16*min*. It was found that after electrolysis the absorbance of the solution was reducted to 50 % of its original value . Calculate the concentration of copper sulphate in the solution to begin with.

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31. The following electrochemical cell has been set up :

$$Pt(l) \left| Fe^{3+}, Fe(a = 1) \right| Ce^{4+}, Ce^{3+}(a = 1) \mid Pt(2)$$
$$E^{C-} \cdot \left(Fe^{3+}, Fe^{2+} \right) = 0.77V$$

and
$E^{C} \cdot (Ce^{4+} Ce^{3+}) = 1.61V$

If an ammeter is connected between two platinum electrodes, predict the direction of the flow of current. Will the current increase or decreases with time ?

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32. The standard potential of the following cell is 0.23 V at $15 \degree C$ and 0.21V at $35 \degree C.Pt |H_2|$ |HCl(aq)|Agcl(s)|g(s)

(i) write the cell reaction .

(ii) Calculate $\Delta H\,^\circ\,$ and $\Delta S\,^\circ\,$ for the cell reaction by assuming that these

quantities remain unchanged in the range $15 \degree C$ to $35 \degree C$.

(iii) calculate the solubility of AgCl in water at 25 ° C.

Give , the standard reduction potential of the $\left(Ag^+(aq)/Ag(s)\right)$ is 0.80 V at 25 ° C.



33. Two students use same stock solution of $ZnSO_4$ and a solution of $CuSO_4$. The *EMF* of one cell is 0.03 higher than the other. The concentration of $CuSO_4$ in the cell with higher *EMF* value is 0.5*M*. Find the concentration of $CuSO_4$ in the other cell.

(Take 2.303RT/F = 0.06)

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34. Find the equilibrium constant for the reaction :

$$Cu^{2+} + In^{2+} \Leftrightarrow Cu^{\oplus} + In^{3+}$$

Given that $E^{c-} \cdot \cdot_{Cu^{2+} \mid Cu^{\oplus}} = 0.15V, E^{c-} \cdot_{\ln^{2+} \mid \ln^{\oplus}} = -0.4V,$
$$E^{c-} \cdot_{\ln^{3+} \mid \ln^{\oplus}} = -0.42V$$

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35. We have taken a saturated solution of AgBr, whose K_{sp} is 12×10^{-14} . If $10^{-7}M$ of $AgNO_3$ are added to 1L of this solutino, find the conductivity (specific conductance) of the solution in terms of $10^{-7}Sm^{-1}$ units. Given :

$$\lambda^{\circ} \cdot \left(Ag^{\oplus}\right) = 6 \times 10^{-3} Sm^2 mol^{-1}$$
$$\lambda^{\circ} \cdot \left(Br^{c}\right) = 8 \times 10^{-3} Sm^2 mol^{-1}$$
$$\lambda^{\circ} \cdot \left(NO_3^{C}\right) = 7XX10^{-3} Sm^2 mol^{-1}$$

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