

SAFE HANDS & IIT-ian's PACE
BPT-03 (NEET) SOLUTIONS

1. (b)

Sol. Apparent coefficient of Volume expansion

$$\gamma_{app.} = \gamma_L - \gamma_s = 7\gamma_s - \gamma_s = 6\gamma_s \text{ (given } \gamma_L = 7\gamma_s \text{)}$$

Ratio of absolute and apparent expansion of liquid

$$\frac{\gamma_L}{\gamma_{app.}} = \frac{7\gamma_s}{6\gamma_s} = \frac{7}{6}$$

2. (b)

Sol. Use heat required \equiv heat released.

3. (a)

Sol. Heat Lost by water = Heat gain by ice cold brass.

4. (b)

$$\text{Sol. } \frac{H_1}{H_2} = \frac{m_1 c_1}{m_2 c_2} = \frac{\frac{4}{3}\pi r_1^3 \cdot \rho_1 \cdot C_1}{\frac{4}{3}\pi r_2^3 \cdot \rho_2 \cdot C_2} = \frac{r_1^3}{r_2^3} \times \frac{\rho_1}{\rho_2} \times \frac{C_1}{C_2}$$

5. (b)

Sol. Heat gain by ice = Heat Lost by water.

6. (b)

$$\text{Sol. } P = \frac{1}{64} = \frac{mL}{t} \quad \therefore L = \frac{Pt}{m}$$

7. (a)

Sol. 5 kg

water \rightarrow water
[20°C] [100°C]

$$\Delta Q = mC_{gm} \Delta T$$

8. (c)

Sol. Let T be the same numerical value of temperature on Celsius scales and Farenheit scale,

$$\text{Then using the relations. } \frac{^{\circ}\text{C}}{100} = \frac{^{\circ}\text{F} - 32}{180} \text{ According}$$

to questions

$$^{\circ}\text{C} = ^{\circ}\text{F} = T \quad \therefore \frac{T}{100} = \frac{T - 32}{180} \quad \therefore T = -4^{\circ}\text{C} = -40^{\circ}\text{F}$$

9. (c)

$$\text{Sol. } \frac{\Delta L}{L} = 2\% = \frac{2}{100}$$

$$\alpha = 0.00002^{\circ}\text{C}^{-1}$$

$$\text{As } \Delta L = \alpha L \Delta T$$

$$\text{Or } \Delta T = \frac{\Delta L}{\alpha L} = \frac{2}{100 \times 0.00002} = \frac{1}{0.001}$$

$$= 10^3 = 1000^{\circ}\text{C}$$

10. (b)

$$\text{Sol. As } \beta = 2\alpha \text{ and } \gamma = 3\alpha \quad \therefore \frac{\beta}{\gamma} = \frac{2\alpha}{3\alpha} = \frac{2}{3}$$

11. (d)

Sol. According to Wien's law

$$\lambda_m T = b = \text{constant}$$

12. (c)

Sol. The circular plate maximum surface area and as such it cools the fastest. For a given volume (mass in this case), a sphere has the least surface area and accordingly cools the slowest.

13. (c)

$$\text{Sol. } P_0 V_0 = nRT$$

If temperature is changed to T without changing the pressure then

$$P_0 V = nRT$$

$$\Rightarrow P_0(V - V_0) = nRT(T - T_0) \Rightarrow P_0 \Delta V = nR \Delta T$$

$$\text{using } nR = \frac{P_0 V_0}{T_0}$$

$$\frac{\Delta V}{\Delta T} = \frac{V_0}{T_0}$$

Now thermal coefficient of volume expansion at constant pressure

$$\frac{1}{V_0} \frac{\Delta V}{\Delta T} = \frac{1}{T_0}$$

14. (d)

Sol. 150 J of heat has been added to the gas.

15. (d)

Sol. According to an ideal gas equation

$$PV = nRT$$

$$\text{Or } V = \frac{nRT}{P}$$

$$\therefore P = \frac{a}{T} \text{ (Given) } \dots (i)$$

$$\therefore V = \frac{nRT^2}{a}$$

$$\Rightarrow dV = \frac{2nRT}{a} dT \dots (ii)$$

Work done by the gas, $dW = PdV$

$$\text{Or } W = \int_T^{4T} \frac{a}{T} \frac{2nRT}{a} dT \text{ (using (i) and (ii))}$$

$$= [2nRT]_T^{4T} = 6nRT$$

16. (d)

Sol. Here, $\eta_1 = 1 - \frac{T_2}{T_1}$ Or $0.25 = 1 - \frac{T_2}{T_1} \Rightarrow \frac{1}{4} = 1 - \frac{T_2}{T_1}$

$$\frac{T_2}{T_1} = 1 - \frac{1}{4} = \frac{3}{4}$$

According to question

$$\eta_2 = 2\eta_1 \text{ and } T_2 = T_2 - 58^\circ\text{C}$$

$$\therefore 2 \times \frac{1}{4} = 1 - \frac{(T_2 - 58^\circ\text{C})}{T_1} \Rightarrow 1 - \frac{1}{2} = \frac{T_2 - 58^\circ\text{C}}{T_1}$$

$$\frac{1}{2} = \frac{T_1}{T_1} - \frac{58^\circ}{T_1} \Rightarrow \frac{3}{4} - \frac{1}{2} = \frac{58}{T_1} \Rightarrow T_1 = 232^\circ\text{C}$$

17. (b)

Sol. Here, coefficient of performance (β) = 5

$$T_1 = 27^\circ\text{C}$$

$$= (27 + 273)\text{K}$$

$$= 300\text{K}$$

$$\text{As } \beta = \frac{T_2}{T_1 - T_2} \Rightarrow 5 = \frac{T_2}{300 - T_2}$$

$$\text{Or } 1500 - 5T_2 = T_2 \text{ or } 6T_2 = 1500$$

$$\therefore T_2 = \frac{1500}{6} = 250\text{K}$$

18. (b)

Sol. As constant volume,

$$dW = PdV = P \times 0 = 0$$

$$dQ = mC_v \Delta T = 5 \times 0.172 \times 4\text{cal}$$

$$= 5 \times 0.172 \times 4 \times 4.2\text{J}$$

$$= 14.4\text{J}$$

$$\therefore dU = dQ - dW = 14.4 - 0$$

$$= 14.4\text{J}$$

19. (c)

Sol. $\frac{n_1 + n_2}{1.5 - 1} = \frac{n_1}{\frac{5}{3} - 1} + \frac{n_2}{\frac{7}{5} - 1}$

20. (c)

Sol. $1 - \frac{Q_2}{Q_1} = 1 - \frac{T_2}{T_1}$

$$\frac{W}{Q_1} = 1 - \frac{300}{600}$$

$$\Rightarrow \frac{800}{Q_1} = \frac{1}{2}$$

$$Q_1 = 1600\text{J}$$

21. (c)

Sol. $\frac{\Delta W}{(\Delta Q)_P} = \frac{(\Delta Q)_P - (\Delta U)_P}{(\Delta Q)_P} = 1 - \frac{(\Delta U)_P}{(\Delta Q)_P}$

$$= 1 - \frac{\frac{5}{2}R}{\frac{7}{2}R} = 1 - \frac{5}{7} = \frac{2}{7}$$

22. (a)

Sol. $P_1 V_1^\gamma = P_2 V_2^\gamma$ ($\gamma = 7/5$), $V_2 = 1.05 V_1$

$$P_1 V_1^\gamma = P_2 (1.05 V_1)^\gamma$$

$$P_2 = \frac{P_1}{(1.05)^\gamma} \text{ so \% change } \frac{P_2 - P_1}{P_1} \times 100$$

$$= \frac{\frac{P_1}{(1.05)^\gamma} - P_1}{P_1} \times 100 = \frac{1 - (1.05)^\gamma}{(1.05)^\gamma} \times 100 = 7\%$$

23. (b)

Sol. $VP^3 = \text{constant}$, $P = \frac{nRT}{V}$

$$\therefore V \frac{T^3}{V^3} = \text{constant} \Rightarrow \boxed{T^3 \propto V^2}$$

24. (b)

Sol. $\Delta Q = \Delta U + \Delta W$; $110\text{J} = 40\text{J} + \Delta W$; $\Delta W = 70\text{J}$

25. (c)

Sol. $V = \text{constant}$. $\Rightarrow P \propto T \Rightarrow T_2 = \frac{P_2}{P_1} T_1$

$$= \frac{2}{1} \times 293\text{K} = 586\text{K} = 313^\circ\text{C}$$

26. (c)

Sol. $nC_p \Delta T = 100\text{J}$

$$n \left(\frac{\gamma R}{\gamma - 1} \right) \Delta T = 100\text{J}, nR \Delta T = \left(100 \times \frac{\gamma - 1}{\gamma} \right)$$

$$W = nR \Delta T = 100 \times \frac{0.4}{1.4}$$

27. (c)

Sol. $\Delta T = \frac{Q}{m \cdot s} = \frac{420/4.2}{10} = 10^\circ\text{C}$

28. (b)

Sol. $80 \times m = 80 \times 30 \times 1$
 $m = 30\text{gm}$

29. (c)

$$\text{Sol. } Q = 1 \times 80 + 1 \times 1 \times 100 + 1 \times 540 \\ = 720 \text{ cal}$$

30. (b)

$$\text{Sol. Here, } C_V = \frac{3}{2}R$$

$$\text{Since } C_p - C_V = R$$

$$\therefore C_p = C_V + R = \frac{3}{2}R + R = \frac{5}{2}R$$

31. (c)

Sol. According to Mayer's relation

$$C_p - C_V = R \text{ or } 1 - \frac{C_V}{C_p} = \frac{R}{C_p}$$

$$1 - \frac{1}{\gamma} = \frac{R}{C_p} \quad \left(\because \gamma = \frac{C_p}{C_V} \right)$$

Or

$$\text{or } \frac{\gamma - 1}{\gamma} = \frac{R}{C_p} \text{ or } C_p = \frac{\gamma R}{\gamma - 1}$$

$$\text{Specific heat capacity} = \frac{\text{molar heat capacity}}{\text{molecular weight}}$$

Specific heat capacity at constant pressure

$$= \frac{\gamma R}{M(\gamma - 1)}$$

32. (b)

Sol. As 32 gm O_2 means 1 mole therefore 8 gm

$$O_2 \text{ means } 1/4 \text{ mole i.e. } \mu = \frac{1}{4}$$

So from $PV = \mu RT$ we get $PV = \frac{1}{4}RT$ or

$$PV = \frac{RT}{4}$$

33. (a)

$$\text{Sol. } \mu = \frac{\text{Mass of water}}{\text{Molecular wt. of water}} = \frac{4.5 \text{ kg}}{18 \times 10^{-3} \text{ kg}} = 250,$$

$$T = 273 \text{ K and } P = 10^5 \text{ N/m}^2 \text{ (STP)}$$

From $PV = \mu RT$

$$\Rightarrow V = \frac{\mu RT}{P} = \frac{250 \times 8.3 \times 273}{10^5} = 5.66 \text{ m}^3.$$

34. (a)

Sol. Specific gas constant

$$r = \frac{\text{Universal gas constant } (R)}{\text{Molecular weight of gas } (M)} = \frac{8.3}{2} = 4.15$$

Joule/mole-K.

35. (c)

Sol. From ideal gas equation $PV = \mu RT$ we get

$$\frac{T_2}{T_1} = \left(\frac{P_2}{P_1} \right) \left(\frac{V_2}{V_1} \right) = \left(\frac{2P_1}{P_1} \right) \left(\frac{3V_1}{V_1} \right) = 6$$

$$\therefore T_2 = 6T_1 = 6 \times 300 = 1800 \text{ K} = 1527^\circ\text{C}.$$

36. (c)

Sol. From $PV = \mu RT$ we get

$$\frac{P_2}{P_1} = \left(\frac{T_2}{T_1} \right) \left(\frac{V_1}{V_2} \right) = \left(\frac{T_1/2}{T_1} \right) \left(\frac{V_1}{2V_1} \right) = \frac{1}{4} \Rightarrow P_2 = \frac{P_1}{4}$$

37. (a)

Sol. Root means square velocity

$$v_{rms} = \sqrt{\frac{3RT}{M}} = 1930 \text{ m/s (given)}$$

$$\therefore M = \frac{3RT}{(1930)^2} = \frac{3 \times 8.31 \times 300}{1930 \times 1930} = 2 \times 10^{-3} \text{ kg} = 2 \text{ gm}$$

i.e. the gas is hydrogen.

38. (c)

Sol. Root mean square velocity does not depend upon the quantity of gas. For a given gas and at constant temperature it always remains same.

39. (c)

$$\text{Sol. } E \propto T \quad \therefore \frac{E_2}{E_1} = \frac{T_2}{T_1} \Rightarrow \frac{2E_1}{E_1} = \frac{T_2}{(20 + 273)}$$

$$\Rightarrow T_2 = 293 \times 2 = 586 \text{ K} = 313^\circ\text{C}.$$

40. (d)

$$\text{Sol. } E = \frac{3}{2}RT = \frac{3}{2} \times 8.31 \times 273 = 3.4 \times 10^3 \text{ Joule}$$

41. (a)

Sol. $P_1 = P, T_1 = T,$

$$P_2 = P + (0.4\% \text{ of } P) = P + \frac{0.4}{100}P = P + \frac{P}{250}$$

$$T_2 = T + 1$$

$$\text{From Gay Lussac's law } \frac{P_1}{P_2} = \frac{T_1}{T_2}$$

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$$\Rightarrow \frac{P}{P + \frac{P}{250}} = \frac{T}{T+1} \quad [\text{As } V = \text{constant for closed}$$

vessel]

By solving we get $T = 250 \text{ K}$.

42. (b)

$$\text{Sol. } \gamma = 1 + \frac{2}{f} \Rightarrow \gamma - 1 = \frac{2}{f} \Rightarrow \frac{f}{2} = \frac{1}{\gamma - 1} \text{ or } f = \frac{2}{\gamma - 1}$$

43. (b)

$$\text{Sol. } \bar{V}_m = \frac{(1)^2 + (2)^2 + (3)^2 + (4)^2}{4}$$

44. (c)

$$\text{Sol. } v_{rms} = \sqrt{\frac{3RT}{M}} \quad \therefore \frac{v_{N_2}}{v_{O_2}} = \sqrt{\frac{M_{O_2}}{M_{N_2}}} = \sqrt{\frac{32}{28}} = \sqrt{\frac{8}{7}}$$

45. (b)

$$\text{Sol. } V = \sqrt{\gamma P / \rho}$$

$$330 = \sqrt{\gamma \times \frac{1 \times 10^5}{1.3}}$$

$$\frac{(33)^2 \times 100 \times 1.3}{1 \times 10^5} = \gamma$$

$$\frac{1.089 \times 10^3 \times 10^2 \times 1.3}{1 \times 10^5} = \gamma$$

$$\frac{2}{f} + 1 = \gamma = 1.4 = 7/5$$

$$\frac{2}{f} = 2/5$$

$$f = 5$$

46. (a)

$$\text{Sol. } K = \frac{\alpha^2 C}{1-\alpha}; \alpha = \frac{0.01}{100} \approx 1 \therefore K = \alpha^2 C = \left[\frac{0.01}{100} \right]^2 \times 1$$

$$= 1 \times 10^{-8}.$$

47. (b)

Sol. Mathematical form of Ostwald's dilution law.

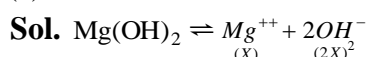
48. (b)

Sol. The value of pK_a for strong acid is less.

49. (c)

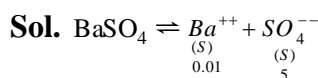
Sol. Due to common ion effect.

50. (a)



$$K_{sp} = 4X^3$$

51. (c)



$$K_{sp} = S^2 = S \times S = 0.01 \times S$$

$$S_{(\text{SO}_4^{--})} = \frac{K_{sp}}{S_{(\text{Ba}^{++})}} = \frac{1 \times 10^{-9}}{0.01} = 10^{-7} \text{ mole/litre.}$$

52. (d)

$$\text{Sol. } K_{sp} \text{ for } \text{CaF}_2 = 4s^3 = 4 \times [2 \times 10^{-4}]^3 = 3.2 \times 10^{-11}.$$

53. (c)

Sol. For the precipitation of an electrolyte, it is necessary that the ionic product must exceed its solubility product.

54. (d)

$$\text{Sol. } K_{sp} = [\text{Ag}^+]^2 [\text{CrO}_4^{--}] = [2S]^2 [0.01]$$

$$= 4S^2 [0.01] = 4[2 \times 10^{-8}]^2 \times 0.01 = 16 \times 10^{-18}.$$

55. (c)

Sol. Complex salts contain two different metallic elements but give test for only one of them. e.g.

$K_4\text{Fe}(\text{CN})_6$ does not give test for Fe^{3+} ions.

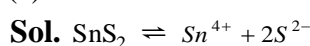
56. (b)

Sol. NH_4CN is a salt of weak acid and weak base and thus for it.

57. (d)

Sol. Because CCl_4 is an organic solvent and AgNO_3 is insoluble in organic solvent.

58. (a)



$$\therefore K_{sp} = [\text{Sn}^{4+}] [\text{S}^{2-}]^2$$

59. (a)

Sol. pH of blood does not change because it is a buffer solution.

60. (b)

$$\text{Sol. } [\text{H}^+] = [\text{OH}^-]$$

$$K_w = [\text{H}^+] [\text{OH}^-] = 10^{-14}$$

$$\therefore [\text{H}^+] = 10^{-7}, \text{pH} = -\log[\text{H}^+] = 7.$$

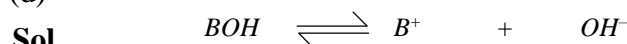
61. (d)

$$\text{Sol. } \text{pH} = 5 \text{ means } [\text{H}^+] = 10^{-5}$$

$$p\text{OH} = 14 - \text{pH} = 14 - 5 = 9$$

$$[\text{OH}^-] = 10^{-p\text{OH}} = 10^{-9}$$

62. (d)



	C		0		
Initial					
0					

	C - C α				C α
At eq.					
C α					

$$K_b = \frac{C^2 \alpha^2}{C(1-\alpha)} = C\alpha^2 \text{ assuming } \alpha \ll 1; 1-\alpha \approx 1$$

$$10^{-12} = 10^{-2} \times \alpha^2; \alpha^2 = 10^{-10}; \alpha = 10^{-5}$$

$$[\text{OH}^-] = C\alpha = .01 \times 10^{-5} = 10^{-7}$$

63. (c)

$$\text{Sol. } \text{pH} = 4 \text{ means; } [\text{H}^+] = 10^{-4} \text{ mol}$$

64. (a)

Sol. Buffer solution is a mixture of weak acid and its conjugate base.

65. (d)

Sol. As the solution is acidic, $\text{pH} < 7$. This is because $[\text{H}^+]$ from H_2O [10^{-7} M] cannot be neglected in comparison to 10^{-10} M

66. (a)

$$\text{Sol. } [\text{OH}^-] = 10^{-2} \text{ M}; \text{pOH} = 2$$

$$\text{pH} + \text{pOH} = 14, \text{pH} = 14 - \text{pOH}$$

$$\text{pH} = 14 - 2 = 12$$

67. (b)

$$\text{Sol. } \text{pH of the solution A} = 3$$

$$[\text{H}^+]_A = 10^{-3} \text{ M.}$$

$$\text{pH of the solution B} = 2$$

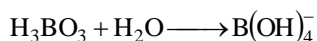
$$[\text{H}^+]_B = 10^{-2} \text{ M}$$

$$[\text{H}^+] = 10^{-3} + 10^{-2} = 10^{-3} + 10 \times 10^{-3} = 11 \times 10^{-3}.$$

$$\text{pH} = -\log(11 \times 10^{-3}) = 3 - \log 11$$

$$= 3 - 1.04 = 1.95$$

68. (a)

Sol. H_3BO_3 is monobasic acid

69. (a)

Sol. Old pH = 7

$$\text{New } [\text{OH}^-] = 10^{-2} \times \frac{1}{10} = 10^{-3}$$

New pH = 11

Change in pH = 4

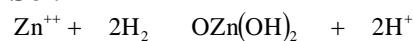
70. (b)

Sol. For colour characteristic of H indicator

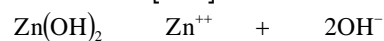
$$\text{pH} = \text{p}K_{\text{in}} - \log \frac{[\text{H}_{\text{in}}]}{[\text{In}^-]}$$

$$\text{pH} = \text{p}K_{\text{in}} \pm 1$$

71. (b)

Sol.

$$\therefore K_{\text{h}} = \frac{[\text{Zn}(\text{OH})_2 + 2\text{H}^+]}{[\text{Zn}^{2+}]} \quad \dots 1$$

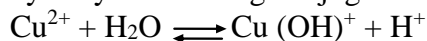


$$\therefore K_{\text{b}} = \frac{[\text{Zn}^{2+}][\text{OH}^-]^2}{[\text{Zn}(\text{OH})_2]}, \quad K_{\text{w}} = [\text{H}^+][\text{OH}^-] \therefore \frac{K_{\text{w}}^2}{K_{\text{b}}} = K_{\text{h}}$$

72. (c)

Sol. H_3O^+ can not take up proton ; H_2PO_2^- can not give up proton, SO_4^{2-} can not give proton

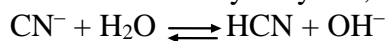
73. (a)

Sol. A solution of CuSO_4 is acidic due to hydrolysis of strong conjugate acid Cu^{2+} ions,

74. (c)

Sol. A solution of NaCl is neutral as both Na^+ and Cl^- ions are weak conjugate acid and base respectively.

75. (c)

Sol. Ba^{2+} is not hydrolysed, $\text{CN}^- = 1$ (M)

$$\text{pH} = \frac{1}{2} (\text{p}K_{\text{w}} + \text{p}K_{\text{a}} + \log c)$$

$$\text{pH} = \frac{1}{2} (\text{p}K_{\text{w}} + \text{p}K_{\text{a}}) \quad \text{since } c = 1(\text{M})$$

$$\text{pH} = \frac{1}{2} (2\text{p}K_{\text{w}} - \text{p}K_{\text{b}}) = \frac{1}{2} (28 - 9.3)$$

$$\text{pH} = \frac{1}{2} (28 - 9.3) = 9.35$$

76. (a)

Sol. For forward reaction $[\text{Al}(\text{H}_2\text{O})_6]^{3+}$ is acid and for backward reaction H_2CO_3 is an acid.

77. (d)

Sol. On adding H_2SO_4 , $[\text{H}^+]$ increases. Therefore of keep K_{w} constant, $[\text{OH}^-]$ decreases.

78. (c)

Sol. 10^{-3} M NaOH has $[\text{OH}^-] = 10^{-3}$ or $[\text{H}^+] = 10^{-11}$. Hence pH = 11.

79. (b)

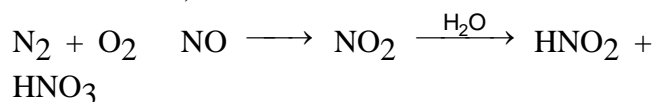
Sol. For pure water, $[\text{H}_3\text{O}^+] = [\text{OH}^-]$ Hence $K_{\text{w}} = 10^{-12}$.

80. (c)

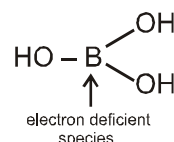
Sol. It is the salt of weak acid and weak base, hence its degree of hydrolysis will be independent of concentration

$$h = \sqrt{\frac{K_{\text{w}}}{K_{\text{a}} \times K_{\text{b}}}}$$

81. (a)

Sol. When rain is accompanied by a thunderstorm,

82. (c)

Sol.

So it will behave as Lewis acid.

83. (c)

Sol. Ostwald's dilution law is valid for weak electrolysis.

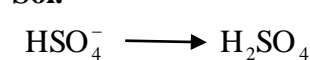
84. (d)

Sol. Factual

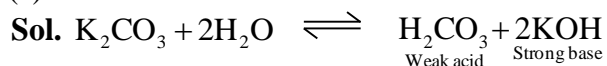
85. (d)

Sol. All other can accept electrons. PH_3 in not electron deficient.

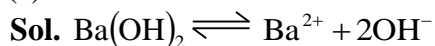
86. (b)

Sol. HSO_4^- can accept or give a proton.

87. (c)



88. (c)

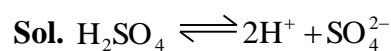


$$[\text{OH}^-] = 2 \times 1 \times 10^{-3} \text{ M}$$

$$\text{pOH} = -\log [\text{OH}^-] = -\log(2 \times 10^{-3}) = 2.7$$

$$\text{pOH} + \text{pH} = 14 \Rightarrow \text{pH} = 14 - 2.7 = 11.3$$

89. (b)



$$[\text{H}^+] = 2 \times 1 \times 10^{-4} \text{ M}$$

$$\text{pH} = -\log(2 \times 10^{-4}) = 3.70$$

90. (a)

Sol. Nucleophiles are Lewis bases while electrophiles are Lewis acids.

BIOLOGY ANSWER KEY

Q.No.	Ans.	Q.No.	Ans.	Q.No.	Ans.	Q.No.	Ans.	Q.No.	Ans.
91	B	109	D	127	D	145	A	163	C
92	C	110	D	128	C	146	D	164	D
93	A	111	C	129	C	147	D	165	A
94	D	112	A	130	B	148	D	166	C
95	C	113	A	131	C	149	A	167	A
96	A	114	B	132	A	150	B	168	A
97	B	115	C	133	C	151	A	169	A
98	A	116	A	134	B	152	B	170	C
99	C	117	A	135	D	153	D	171	A
100	B	118	A	136	A	154	A	172	D
101	C	119	C	137	C	155	D	173	B
102	A	120	D	138	C	156	D	174	C
103	C	121	D	139	B	157	A	175	B
104	B	122	B	140	C	158	A	176	A
105	D	123	C	141	D	159	C	177	C
106	C	124	B	142	B	160	B	178	B
107	D	125	D	143	D	161	B	179	C
108	D	126	B	144	B	162	C	180	B