

WEEKLY TEST TYM TEST 16 UNISION
SOLUTION Date 08-12-2019

[PHYSICS]

1. The effective acceleration of a bob in water = $g' = g \left[1 - \frac{\sigma}{\rho} \right]$

where σ and ρ are the densities of water and the bob respectively. Since, the periods of oscillation of the bob in air and water are given as

$$T = 2\pi\sqrt{\frac{l}{g}} \text{ and } T' = 2\pi\sqrt{\frac{l}{g'}}$$

$$\begin{aligned} \therefore \frac{T}{T'} &= \sqrt{\frac{g'}{g}} = \sqrt{\frac{g \left(1 - \frac{\sigma}{\rho} \right)}{g}} \\ &= \sqrt{1 - \frac{\sigma}{\rho}} = \sqrt{1 - \frac{1}{\rho}} \quad [\because \sigma = 1] \end{aligned}$$

Putting $\frac{T}{T'} = \frac{1}{\sqrt{2}}$

We obtain, $\frac{1}{2} = 1 - \frac{1}{\rho} \Rightarrow \rho = 2$

2. When the elevator is at rest, its time period is given by

$$T = 2\pi\sqrt{\frac{l}{g}} = 2\pi\sqrt{\frac{l}{10}}$$

When the elevator accelerates upwards, its time period becomes

$$\begin{aligned} T' &= 2\pi\sqrt{\frac{l}{g+a}} = 2\pi\sqrt{\frac{l}{10+10}} \\ &= 2\pi\sqrt{\frac{l}{20}} \\ &= 2\pi\sqrt{\frac{l}{10}} \times \frac{1}{\sqrt{2}} \\ &= \frac{T}{\sqrt{2}} \end{aligned}$$

3. Velocity of bob $v = \sqrt{2gl(1 - \cos\theta)}$
 $= \sqrt{2 \times 9.8 \times 2 \times (1 - \cos 60^\circ)}$
 $v = \sqrt{2 \times 9.8} \text{ m/s}$

4.
5.

6. Let displacement equation of particle executing SHM is

$$y = a \sin \omega t$$

As particle travels half of the amplitude from the equilibrium position, so

$$y = \frac{a}{2}$$

Therefore, $\frac{a}{2} = a \sin \omega t$

or $\sin \omega t = \frac{1}{2} = \sin \frac{\pi}{6}$

or $\omega t = \frac{\pi}{6}$

or $t = \frac{\pi}{6\omega}$

or $t = \frac{\pi}{6 \left(\frac{2\pi}{T} \right)}$ (as $\omega = \frac{2\pi}{T}$)

60. Given,

and

or $t = \frac{T}{12}$

Hence, the particle travels half of the amplitude from equilibrium in $\frac{T}{12}$ s.

7.
8.
9.
10.

11. Time period of a simple pendulum

$$T = 2\pi \sqrt{\frac{l}{g}}$$

It is independent of the mass of the bob. Therefore time period of the pendulum will remain T .

12.

13. Time period of simple pendulum,

$$T = 2\pi \sqrt{\frac{l}{g}} \Rightarrow \frac{\Delta T}{T} = \frac{1}{2} \frac{\Delta l}{l}$$

But at temperature $\theta^\circ \text{C}$, increase in length of pendulum,

$$\frac{\Delta l}{l} = \alpha \Delta \theta$$

$\therefore \frac{\Delta T}{T} = \frac{1}{2} \alpha \Delta \theta$

or $\frac{\Delta T}{T} = \frac{1}{2} \times 9 \times 10^{-7} \times (30 - 20)$

$$= \frac{1}{2} \times 9 \times 10^{-7} \times 10$$

$$= 4.5 \times 10^{-6}$$

or
$$\Delta T = 4.5 \times 10^{-6} \times 0.5$$

$$= 2.25 \times 10^{-6} \text{ s}$$

14.
15.

16. (b) Acceleration of simple harmonic motion is

$$a_{\max} = -\omega^2 A$$

or
$$\frac{(a_{\max})_1}{(a_{\max})_2} = \frac{\omega_1^2}{\omega_2^2} \quad (\text{as } A \text{ remains the same})$$

or
$$\frac{(a_{\max})_1}{(a_{\max})_2} = \frac{(100)^2}{(1000)^2} = \left(\frac{1}{10}\right)^2 = 1:10^2$$

17. (d)
$$v = \frac{dy}{dt} = A\omega \cos \omega t = A\omega \sqrt{1 - \sin^2 \omega t}$$

$$= \omega \sqrt{A^2 - y^2}$$

Here, $y = \frac{a}{2}$

$$\therefore v = \omega \sqrt{a^2 - \frac{a^2}{4}} = \omega \sqrt{\frac{3a^2}{4}} = \frac{2\pi a\sqrt{3}}{T} \cdot \frac{1}{2} = \frac{\pi a\sqrt{3}}{T}$$

18. (b) Acceleration $\propto -$ (displacement).

$$A \propto -y$$

$$A = -\omega^2 y$$

$$A = -\frac{k}{m} y$$

$$A = -ky$$

Here, $y = x + a$

$$\therefore \text{acceleration} = -k(x + a)$$

19. (b) Use the law of conservation of energy. Let x be the extension in the spring.

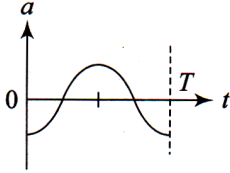
Applying conservation of energy

$$mgx - \frac{1}{2}kx^2 = 0 - 0 \Rightarrow x = \frac{2mg}{k}$$

20. (c) Displacement, $x = A \cos(\omega t)$ (given)

$$\text{Velocity, } v = \frac{dx}{dt} = -A\omega \sin(\omega t)$$

$$\text{Acceleration, } a = \frac{dv}{dt} = -A\omega^2 \cos(\omega t)$$



Hence graph (c) correctly depicts the variation of a with t .

21. (c) The two displacement equations are $y_1 = a \sin(\omega t)$

$$\text{and } y_2 = b \cos(\omega t) = b \sin\left(\omega t + \frac{\pi}{2}\right)$$

$$\begin{aligned} y_{\text{eq}} &= y_1 + y_2 \\ &= a \sin \omega t + b \cos \omega t \\ &= a \sin \omega t + b \sin\left(\omega t + \frac{\pi}{2}\right) \end{aligned}$$

Since the frequencies for both SHMs are same, resultant motion will be SHM.

$$\text{Now } A_{\text{eq}} = \sqrt{a^2 + b^2 + 2ab \cos \frac{\pi}{2}}$$

$$\Rightarrow A_{\text{eq}} = \sqrt{a^2 + b^2}$$

22. (a) Maximum velocity $V_{\text{max}} = A\omega = \beta$ (i)
 maximum acceleration $\alpha_{\text{max}} = A\omega^2 = \alpha$ (ii)

$$\text{Equation (ii) divided by (i) } \omega = \frac{\omega}{\beta} \Rightarrow \frac{2\pi}{T} = \frac{\omega}{\beta}$$

$$T = \frac{2\pi\beta}{\alpha}$$

23. (a) If initial length $l_1 = 100$ then $l_2 = 121$

$$\text{By using } T = 2\pi\sqrt{\frac{l}{g}} \Rightarrow \frac{T_1}{T_2} = \sqrt{\frac{l_1}{l_2}}$$

$$\text{Hence, } \frac{T_1}{T_2} = \sqrt{\frac{100}{121}} \Rightarrow T_2 = 1.1T_1$$

$$\% \text{ increase} = \frac{T_2 - T_1}{T_1} \times 100 = 10\%$$

Alternative: Time period of simple pendulum

$$T = 2\pi\sqrt{\frac{l}{g}} \Rightarrow T \propto \sqrt{l}$$

$$\therefore \frac{\Delta T}{T} = \frac{1}{2} \frac{\Delta l}{l}$$

$$\text{Since, } \frac{\Delta l}{l} = 21\%$$

$$\therefore \frac{\Delta T}{T} = \frac{1}{2} \times 21\% \approx 10\%$$

24. (d) As springs are connected in series, effective force constant

$$\frac{1}{k} = \frac{1}{k_1} + \frac{1}{k_2} \Rightarrow k = \frac{k_1 k_2}{k_1 + k_2}$$

Hence, frequency of oscillation is

$$n = \frac{1}{2\pi} \sqrt{\frac{k}{m}} = \frac{1}{2\pi} \sqrt{\frac{k_1 k_2}{(k_1 + k_2)m}}$$

25. (c) $n = \frac{1}{2\pi} \sqrt{\frac{K_{\text{effective}}}{m}}$

Springs are connected in parallel

$$K_{\text{eff}} = K_1 + K_2 = K + 2K = 3K$$

$$\Rightarrow n = \frac{1}{2\pi} \sqrt{\frac{(K + 2K)}{m}} = \frac{1}{2\pi} \sqrt{\frac{3K}{m}}$$

26. (a) As springs are connected in series, effective force constant

$$\frac{1}{k_{\text{eff}}} = \frac{1}{k} + \frac{1}{k} = \frac{2}{k} \Rightarrow k_{\text{eff}} = \frac{k}{2}$$

Hence, frequency of oscillation is

$$n = \frac{1}{2\pi} \sqrt{\frac{k_{\text{eff}}}{m}} = \frac{1}{2\pi} \sqrt{\frac{k}{2M}}$$



27. For a simple harmonic motion

$$\frac{d^2y}{dt^2} \propto -y$$

Hence, equation $y = \sin \omega t - \cos \omega t$

and $y = 5 \cos\left(\frac{3\pi}{4} - 3\omega t\right)$ are satisfying this

condition and equation $y = 1 + \omega t + \omega^2 t^2$ is not periodic and

$y = \sin^3 \omega t$ is periodic but not SHM.

28. The motion of planets around the sun is periodic but not simple harmonic motion.

29. For freely falling case the effective g is zero, so that frequency of oscillation will be zero.

As
$$f = \frac{1}{2\pi} \sqrt{\frac{g_{\text{eff}}}{\lambda}}$$

$$f = \frac{1}{2\pi} \sqrt{\frac{0}{\lambda}}$$

$$f = 0$$

30. $x(t) = A \cos(\omega t + \phi)$

where, ϕ is the phase constant.

31. $y_1 = 5 [\sin 2\pi t + \sqrt{3} \cos 2\pi t]$

$$= 10 \left[\frac{1}{2} \sin 2\pi t + \frac{\sqrt{3}}{2} \cos 2\pi t \right]$$

$$= 10 \left[\cos \frac{\pi}{3} \sin 2\pi t + \sin \frac{\pi}{3} \cos 2\pi t \right]$$

$$= 10 \left[\sin \left(2\pi t + \frac{\pi}{2} \right) \right] \Rightarrow A_1 = 10$$

Similarly, $y_2 = 5 \sin \left(2\pi t + \frac{\pi}{4} \right)$

$$\Rightarrow A_2 = 5$$

Hence, $\frac{A_1}{A_2} = \frac{10}{5} = 2$

32. The potential energy, $U = \frac{1}{2} kx^2$

$$2U = kx^2$$

$$2U = -Fx \quad (\because F = -kx)$$

or $\frac{2U}{F} = -x$

or $\frac{2U}{F} + x = 0$

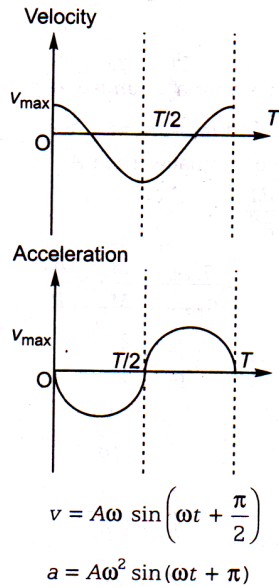
33. Phase difference $\Delta\phi = \phi_1 - \phi_2$

$$= \frac{3\pi}{6} - \frac{\pi}{6}$$

$$= \frac{2\pi}{6} = \frac{\pi}{3}$$



34. In SHM, the acceleration is ahead of velocity by a phase angle $\frac{\pi}{2}$.



35. The total energy a particle executing SHM

$$= \frac{1}{2} m\omega^2 A^2$$

The PE of the particle at a distance x from the equilibrium position

$$= \frac{1}{2} m\omega^2 x^2$$

From the question, $\frac{1}{2} m\omega^2 x^2 = \frac{1}{2} \left(\frac{1}{2} m\omega^2 A^2 \right)$

$$\Rightarrow x^2 = \frac{A^2}{2} \Rightarrow x = \frac{A}{\sqrt{2}}$$

36. The average acceleration of a particle performing SHM over one complete oscillation is zero.

37. Let x be the point where $KE = PE$

Hence $\frac{1}{2} m\omega^2 (a^2 - x^2) = \frac{1}{2} m\omega^2 x^2$

$$2x^2 = a^2, x = \frac{a}{\sqrt{2}}$$

$$x = \frac{4}{\sqrt{2}} = 2\sqrt{2} \text{ cm}$$

38. By using $k \propto \frac{1}{l}$

Since, one-fourth length is cut away so remaining length is $\frac{3}{4}$ th, hence k becomes $\frac{4}{3}$ times ie, $k' = \frac{4}{3} k$.

39. Maximum velocity $v_{\max} = A\omega$

$$\omega = \frac{2\pi}{T}$$

$$\therefore v_{\max} = \frac{2\pi A}{T}$$

$$v \propto \frac{A}{T}$$

$$\therefore \frac{v_1}{v_2} = \frac{A_1}{A_2} \times \frac{T_2}{T_1} = \frac{1}{2} \times \frac{1}{3} = \frac{1}{6}$$

$$v_2 = 6v_1 = 6v$$

40. For the given figure,

$$f = \frac{1}{2\pi} \sqrt{\frac{k_{\text{eq}}}{m}} \quad \dots(i)$$

$$= \frac{1}{2\pi} \sqrt{\frac{2k}{m}}$$

If one spring is removed, then $k_{\text{eq}} = k$ and

$$f' = \frac{1}{2\pi} \sqrt{\frac{k}{m}} \quad \dots(ii)$$

From Eqs. (i) and (ii), we get

$$\frac{f}{f'} = \sqrt{2}$$

$$f' = \frac{1}{\sqrt{2}} f$$

41. $\frac{d^2x}{dt^2} + 16x = 0$

$$\therefore \omega^2 = 16 \Rightarrow \omega = 4$$

$$T = \frac{2\pi}{\omega} = \frac{2\pi}{4} = \frac{\pi}{2}$$

42. In a complete cycle of SHM, potential energy varies for half the cycle and kinetic energy varies for the other half of the cycle. Thus, for a time period T , the potential energy varies for $\frac{T}{2}$ time.

43.

44.

45.

[CHEMISTRY]

46.

The corresponding acids are HI, HCl, HNO_2 and HCN. Their acid strength follows the order $\text{HI} > \text{HCl} > \text{HNO}_2 > \text{HCN}$. Hence, their conjugate base follows the reverse order.

47.

pH of a weak acid is given by

$$(I) \quad 2\text{pH} = \frac{1}{2}[pK_a - \log C] \text{ at } C = 0.1 \text{ M}$$

$$(ii) \quad \text{pH} = \frac{1}{2}[pK_a - \log C'] \text{ at } C' = ?$$

$$\therefore \quad 4\text{pH} = pK_a - \log C'$$

$$2\text{pH} = pK_a = \log C$$

$$2\text{pH} = \log C - \log C' = \log \frac{0.1}{C'}$$

$$\text{From I,} \quad \text{pH} = \frac{1}{2}[4.74 - \log 0.1] = \frac{1}{2}[4.74 + 1.0] = 2.87$$

$$\therefore \quad 2 \times 2.87 = \log \frac{0.1}{C'} \Rightarrow 5.74 = \log \frac{0.1}{C'}$$

$$\therefore \quad \frac{0.1}{C'} = 5.5 \times 10^5$$

$$\text{Thus, dilution } \frac{1}{C'} = \frac{5.55 \times 10^5}{0.1} = 5.55 \times 10^6 \text{ times}$$

48.

The conjugate acid-base pairs are (HCl, Cl⁻) and (CH₃COOH₂⁺, CH₃COOH).

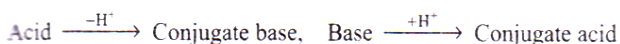
49.

The conjugate acids are H₂O, NH₃, HC ≡ CH and CH₃CH₃. Their order of acid strength is CH₃CH₃ < NH₃ < HC ≡ CH < H₂O. Their conjugate base follows the reverse order.

50.

NH₃ donates pair of electrons while BF₃, Cu²⁺ and AlCl₃ accept lone pair of electrons.

51.



52.

H₃O⁺ (acid), H₂O (conjugate base) and not OH⁻.

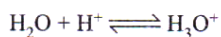
53.

54.

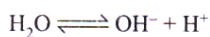
For each weak polyprotic acid $K_{a_1} > K_{a_2} > K_{a_3}$

55.

H₂O is the weaker base, hence, its conjugate acid is the stronger acid,



H₂O is the weakest acid, hence, its conjugate base is the strongest base.



56.

$$\begin{aligned} \text{pH [HCl]} &= 2.0 \\ \therefore [\text{H}^+] &= 10^{-2} \text{ M} \\ [\text{HCl}] &= 10^{-2} \text{ M} \\ \text{Volume} &= 200 \text{ mL} \\ \text{pH [NaOH]} &= 12.0 \\ \text{pOH} &= 2.0 \\ [\text{OH}^-] &= 10^{-2} \text{ M} \\ [\text{NaOH}] &= 10^{-2} \text{ M} \\ \text{Volume} &= 300 \text{ mL} \\ N_1 V_1 (\text{acid}) &= 200 \times 10^{-2} = 2 \\ N_1 V_2 (\text{base}) &= 300 \times 10^{-2} = 3 \\ N_2 V_2 &> N_1 V_1 \\ \text{Thus, resultant mixture basic.} \\ N(\text{OH}^-) &= \frac{N_2 V_2 - N_1 V_1}{V_1 + V_2} = \frac{3 - 2}{500} = 2 \times 10^{-3} \text{ M} \\ \text{pOH} &= -\log(2 \times 10^{-3}) = 2.7 \\ \therefore \text{pH} &= 14 - \text{pOH} = 14 - 2.7 = 11.3 \end{aligned}$$

57.

$$\begin{aligned} [\text{H}^+] \text{ after mixing} &= \frac{10^{-2} \times 10 + 10^{-4} \times 990}{1000} = \frac{0.1 + 0.0990}{1000} \\ &= \frac{0.1990}{1000} = 1.99 \times 10^{-4} \\ \text{pH} &= (\log 1.99 \times 10^{-4}) \\ \therefore \text{pH} &= 4 - 0.3 = 3.7 \end{aligned}$$

58.

$$\begin{aligned} [\text{H}^+] &= \frac{50 \times 10^{-1} + 50 \times 10^{-2}}{100} = 5.5 \times 10^{-2} \text{ M} \\ \text{pH} &= \log(1.99 \times 10^{-4}) \\ \therefore \text{pH} &= 2 - 0.74 = 1.26 \end{aligned}$$

59.

On heating pure water the value of ionic product of water increases i.e., $K_w = 10^{-14}$ at 25°C and at 100°C , $K_w = 10^{-12}$. Thus pH and pOH both become 6 at 100°C (pH and pOH = 7 at 25°C).

60.

- (a) At 25°C , $[\text{H}^+]$ in a solution of $10^{-8} \text{ M HCl} > 10^{-7} \text{ M}$.
 (b) $[\text{H}^+] = 10^{-8} \text{ M}$.
 (c) $[\text{OH}^-] = 4 \times 10^{-6} \text{ M} \Rightarrow [\text{H}^+] = 2.5 \times 10^{-9} \text{ M}$
 (d) $[\text{H}^+] = 10^{-9} \text{ M}$

61.

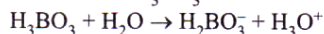
K_w changes with temperature. As temperature increases, $[\text{OH}^-]$ and $[\text{H}^+]$ decrease.

62.

Meq. of HCl = $10 \times 10^{-1} = 1$
 Meq. of NaOH = $10 \times 10^{-1} = 1$
 Thus both are neutralised and 1 Meq. of NaCl (a salt of strong acid and strong base) which does not hydrolyse and thus pH = 7.

63.

The dissociation of H_3BO_3 is



$$K_1 = \frac{[\text{H}_2\text{BO}_3^-][\text{H}_3\text{O}^+]}{[\text{H}_3\text{BO}_3]} = \frac{(0.18).x}{(01.0)} = 7.3 \times 10^{-10}$$

or $x = [\text{H}_3\text{O}^+] = 4.1 \times 10^{-10}$

or $\text{pH} = -\log x = -\log (4.1 \times 10^{-10}) = 9.39$

64.

- | | |
|--|------------------------------------|
| (a) HCl | NaOH |
| No. of milli eq. = $\frac{1}{10} \times 100 = 10$ | $\frac{1}{10} \times 100 = 10$ |
| So solution is neutral | |
| (b) $\frac{1}{10} \times 55 = 5.5$ | $\frac{1}{10} \times 45 = 4.5$ |
| $[\text{H}^+] = \frac{1}{100} = 10^{-2} \text{ M}$, $\text{pH} = 2$ | |
| (c) $\frac{1}{10} \times 10 = 1$ | $\frac{1}{10} \times 90 = 9$ Basic |
| (d) $\frac{1}{5} \times 75 = 15$ | $\frac{1}{5} \times 25 = 5$ |
| $[\text{H}^+] = 0.1 \text{ M}$, $\text{pH} = 1$ | |

65.

- | | |
|---|-------------------------------------|
| Initial | Final |
| $\text{pH} = 12$ | $\text{pH} = 11$ |
| $[\text{H}^+] = 10^{-12} \text{ M}$ | $[\text{H}^+] = 10^{-11} \text{ M}$ |
| $[\text{OH}^-] = 10^{-2} \text{ M}$ | $[\text{OH}^-] = 10^{-3} \text{ M}$ |
| Initial no. of mole of $\text{OH}^- = 10^{-2}$ | |
| Final no. of mole of $\text{OH}^- = 10^{-3}$ | |
| So no. of mole of OH^- removed = $[0.1 - 0.001] = 0.009$ | |

66.

$$pK_w = -\log K_w = -\log 1 \times 10^{-12} = 12.$$

$$K_w = [\text{H}^+][\text{OH}^-] = 10^{-12}$$

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

$$\Rightarrow [\text{H}^+]^2 = 10^{-12}; [\text{H}^+] = 10^{-6}; \text{pH} = -\log [\text{H}^+] = -\log 10^{-6} = 6.$$

H_2O is neutral because $[\text{H}^+] = [\text{OH}^-]$ at 373 K even when $\text{pH} = 6$.
(d) is not correct at 373 K. Water cannot become acidic.

67.

$$\text{Relative strength of weak acids} = \sqrt{\frac{K_{a_1} \times C_1}{K_{a_2} \times C_2}}$$

$$\therefore \text{Relative strength} = \sqrt{\frac{K_{a_1}}{K_{a_2}}} \quad (\because C_1 = C_2) = \sqrt{\frac{2 \times 10^{-4}}{2 \times 10^{-5}}}$$

Relative strength for HCOOH to $\text{CH}_3\text{COOH} = \sqrt{10} : 1$

68.

$$\text{pH} = 13$$

$$\therefore [\text{H}^+] = 10^{-13} \text{ M}$$

$$[\text{OH}^-] = 10^{-1} \text{ M} = 0.1 \text{ mol L}^{-1}$$

$$[\text{Ba}(\text{OH})_2] = 0.1 \text{ N},$$

$$= 0.1 \times 100 = 10 \text{ milliequivalents}$$

69.

pH of amphiprotic salts and weak acid-weak base salt is independent of its concentration.

70.

71.

72.

Reaction: $2A + B \rightleftharpoons C + D$

$$K_P = \frac{n_C \times n_D}{n_A^2 \times n_B} \times \left(\frac{P}{\Sigma n} \right)^{\Delta n_g}$$

$$\Delta n_g = 2 - 3 = -1$$

$$K_P = \frac{n_C \times n_D}{n_A^2 \times n_B} \times \left(\frac{\Sigma n}{P} \right)$$

$$PV = \Sigma nRT$$

$$\frac{V}{RT} = \frac{\Sigma n}{P}$$

From equations (i) and (ii),

$$K_P = \frac{n_C \times n_D}{n_A^2 \times n_B} \times \frac{V}{RT}$$

73.

Concentration of $[\text{NO}_2]$ will decrease with increase in concentration $[\text{N}_2\text{O}_4]$.

74.

With passage of time conc. of reactants decreases and products increases.

75.

$$K = 2 = \sqrt{k_1}, K_2 = \frac{1}{K_4}, K_1 = \frac{1}{K_3}$$

$$\therefore K_1 K_3 = 1, \sqrt{K_1} K_4 = 1, \sqrt{K_3} = 1$$

76.

$$\Delta n_g = 4 + 1 - (2 + 2) = 1$$

$$\therefore K_P = k_c (RT)^{\Delta n_g}$$

$$0.03 = K_C (0.082 \times 700)^1$$

$$K_C = 5.23 \times 10^{-4}$$

77.

Required equilibrium is obtained if we operate.

Eq. (III) \times 4 - Eq. (I) \times 2 - Eq. (II) \times 2

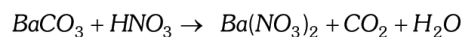
$$K_C = \frac{[\text{N}_2\text{O}_4]^2}{[\text{N}_2\text{O}]^2 [\text{O}_2]^3} = \frac{(4.1 \times 10^{-9})^4}{(2.7 \times 10^{-18})^2 (4.6 \times 10^{-3})^2} = 1.832 \times 10^6$$

78.

At equilibrium rates of backward and forward reactions become equal.

87. (c) HCl is a strong electrolyte since it will produce more H^+ , comparison than that of CH_3COOH . Hence assertion is true but reason false.

88. (a) Barium carbonate is more soluble in HNO_3 than in water because carbonate is a weak base and reacts with the H^+ ion of HNO_3 causing the barium salt to dissociate.



89. (a) The conjugate base of $CHCl_3$ is more stable than conjugate base of $CHF_3(CF_3)$. CCl_3 is stabilized by $-I$ effect of chlorine atoms as well as by the electrons. But conjugate base of $CH_3(CH_3)$ is stabilized only by $-I$ effect of fluorine atoms. Here both assertion and reason are true and reason is correct explanation of assertion.