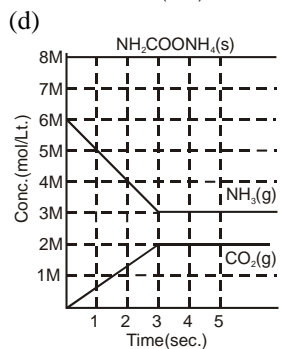
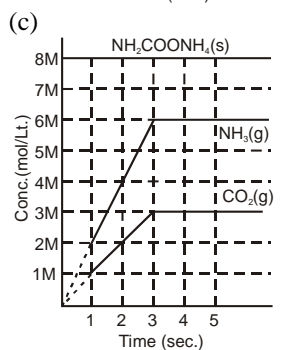
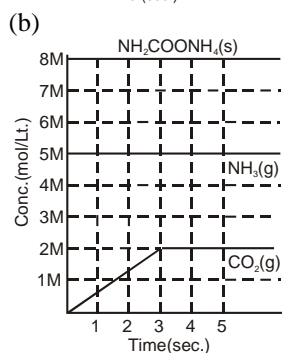
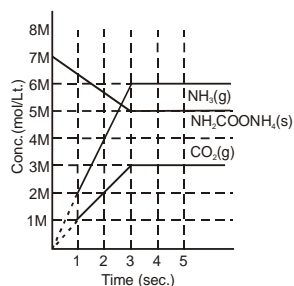


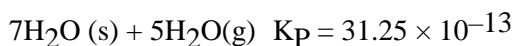
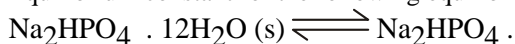
1. Starting with 2 moles of  $\text{PCl}_5$  in 1 litre container, 0.5 moles of  $\text{PCl}_5$  dissociates till equilibrium at  $25^\circ\text{C}$ . Equilibrium constant  $K_{\text{cis}}$  is :
- (a) 6 (b) 4.5 (c)  $\frac{1}{6}$  (d) None
2. The r.m.s. speed of molecules of a gas of density  $4.0 \text{ kg m}^{-3}$  and pressure  $1.2 \times 10^5 \text{ Nm}^{-2}$  is -
- (a)  $300 \text{ m s}^{-1}$  (b)  $900 \text{ m s}^{-1}$  (c)  $600 \text{ m s}^{-1}$  (d)  $400 \text{ m s}^{-1}$
3. In a chemical reaction, the rate constant for the backward reaction is  $7.5 \times 10^{-4}$  and the equilibrium constant is 1.5. The rate constant for the forward reaction is -
- (a)  $5 \times 10^{-4}$  (b)  $2 \times 10^{-3}$   
(c)  $1.125 \times 10^{-3}$  (d)  $9.0 \times 10^{-4}$
4. For the equilibrium  $\text{NH}_4\text{HS(s)} \rightleftharpoons \text{NH}_3\text{(g)} + \text{H}_2\text{S(g)}$   
 $K_c = 1.8 \times 10^{-4}$  at 298 K. The value of  $K_p$  at 298 K would be -
- (a) 0.108 (b)  $4.4 \times 10^{-3}$  (c)  $1.8 \times 10^{-4}$  (d)  $4.4 \times 10^{-4}$
5. If  $K$  for the reaction  $2\text{HI(g)} \rightleftharpoons \text{H}_2\text{(g)} + \text{I}_2\text{(g)}$ , is 0.025 at a certain temperature, the value of  $K$  for the reaction  $\text{H}_2\text{(g)} + \text{I}_2\text{(g)} \rightleftharpoons 2\text{HI(g)}$  at the same temperature would be -
- (a) 10 (b) 40 (c)  $\sqrt{30}$  (d)  $\sqrt{40}$
6. The dissociation constants of two acids  $\text{HA}_1$  and  $\text{HA}_2$  are  $3.0 \times 10^{-4}$  and  $1.8 \times 10^{-5}$  respectively. The relative strengths of the acids will be approximately
- (a) 1 : 4 (b) 4 : 1 (c) 1 : 16 (d) 16 : 1
7. The equilibrium constant,  $K_p$  for the reaction
- $$2\text{SO}_2\text{(g)} + \text{O}_2\text{(g)} \rightleftharpoons 2\text{SO}_3\text{(g)}$$
- is  $4.0 \text{ atm}^{-1}$  at 1000 K. What would be the partial pressure of  $\text{O}_2$  if at equilibrium the amount of  $\text{SO}_2$  and  $\text{SO}_3$  is the same ?
- (a) 16.0 atm (b) 0.25 atm  
(c) 1 atm (d) 0.75 atm
8. For the reaction  $\text{A}_2\text{(g)} + 2\text{B}_2 \rightleftharpoons 2\text{C}_2\text{(g)}$   
the partial pressure of  $\text{A}_2$ ,  $\text{B}_2$  at equilibrium are 0.80 atm and 0.40 atm respectively. The pressure of the system is 2.80 atm. The equilibrium constant  $K_p$  will be
- (a) 20 (b) 5.0 (c) 0.02 (d) 0.2
9. If the anions (a) form hexagonal closest packing and cations (c) occupy only  $\frac{2}{3}$  octahedral voids in it, then the general formula of the compound is
- (a) CA (b)  $\text{CA}_2$  (c)  $\text{C}_2\text{A}_3$  (d)  $\text{C}_3\text{A}_2$
10. For the reaction  $\text{N}_2\text{O}_4\text{(g)} \rightleftharpoons 2\text{NO}_2\text{(g)}$ , if percentage dissociation of  $\text{N}_2\text{O}_4$  are 20%, 45%, 65% & 80%, then the sequence of observed vapour densities will be :
- (a)  $d_{20} > d_{45} > d_{65} > d_{80}$  (b)  $d_{80} > d_{65} > d_{45} > d_{20}$   
(c)  $d_{20} = d_{45} = d_{65} = d_{80}$  (d)  $(d_{20} = d_{45}) > (d_{65} = d_{80})$
11. Solid ammonium carbonate dissociate to give ammonia and carbon dioxide as follows
- $$\text{NH}_2\text{COONH}_4\text{(s)} \rightleftharpoons 2\text{NH}_3\text{(g)} + \text{CO}_2\text{(g)}$$
- which of the following graph correctly represents the equilibrium.
- (a)



12. The correct relationship between free energy change in a reaction and the corresponding equilibrium constant  $K$  is

- (a)  $-\Delta G^0 = RT \ln K$                       (b)  $\Delta G = RT \ln K$   
 (c)  $-\Delta G = RT \ln K$                       (d)  $\Delta G^0 = RT \ln K$

13. Equilibrium constant for the following equilibrium is given at  $0^\circ\text{C}$ .



At equilibrium what will be partial pressure of water vapour :

- (a)  $\frac{1}{5} \times 10^{-3}$  atm                      (b)  $0.5 \times 10^{-3}$  atm  
 (c)  $5 \times 10^{-2}$  atm                      (d)  $5 \times 10^{-3}$  atm.

14. A liquid is in equilibrium with its vapour at its boiling point. On the average the molecules in the two phases have equal :

- (a) Inter molecular forces                      (b) Potential energy  
 (c) Kinetic energy                      (d) None of these .

15. Given the following reaction at equilibrium

$\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$ . Some inert gas at constant pressure is added to the system. Predict which of the following facts will be affected

- (a) More  $\text{NH}_3(\text{g})$  is produced
- (b) Less  $\text{NH}_3(\text{g})$  is produced
- (c) No affect on the equilibrium
- (d)  $K_p$  of the reaction is decreased

16. Introduction of inert gas (at the same temperature) will affect the equilibrium if :

- (a) Volume is constant and  $\Delta n_g \neq 0$
- (b) Pressure is constant and  $\Delta n_g \neq 0$
- (c) volume is constant and  $\Delta n_g = 0$
- (d) Pressure is constant and  $\Delta n_g = 0$

17. In the preceding problem, if  $[\text{A}^+]$  and  $[\text{AB}_2^{\square}]$  are  $y$  and  $x$  respectively, under equilibrium produced by adding the substance AB to the solvents, then  $K_1/K_2$  is equal to

- (a)  $\frac{y}{x}(y-x)^2$
- (b)  $\frac{y^2(x+y)}{x}$
- (c)  $\frac{y^2(x+y)}{x}$
- (d)  $\frac{y}{x}(x-y)$

18. The equilibrium constant for the reaction,  $\text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g})$  at temperature T is  $4 \times 10^{-4}$ . The value of  $K_c$  for the reaction,  $\text{NO}(\text{g}) \rightleftharpoons \frac{1}{2}\text{N}_2(\text{g}) + \frac{1}{2}\text{O}_2(\text{g})$  at the same temperature is

- (a)  $2.5 \times 10^2$
- (b) 0.02
- (c)  $4 \times 10^{-4}$
- (d) 50

19. For the reaction,  $2\text{NO}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g}) + \text{O}_2(\text{g})$ ,

( $K_C = 1.8 \times 10^{-6}$  at  $184^\circ\text{C}$ )

( $R = 0.0831 \text{ kJ}/(\text{mol}\cdot\text{K})$ )

When  $K_p$  and  $K_c$  are compared at  $184^\circ\text{C}$  it is found that

- (a) Whether  $K_p$  is greater than, less than or equal to  $K_c$  depends upon the total gas pressure
- (b)  $K_p = K_c$
- (c)  $K_p$  is less than  $K_c$
- (d)  $K_p$  is greater than  $K_c$

20. For the following three reactions a, b and c, equilibrium constants are given:

(a)  $\text{CO}(\text{g}) + \text{H}_2\text{O}(\text{g}) \rightleftharpoons \text{CO}_2(\text{g}) + \text{H}_2(\text{g}); K_1$

(b)  $\text{CH}_4(\text{g}) + \text{H}_2\text{O}(\text{g}) \rightleftharpoons \text{CO}(\text{g}) + 3\text{H}_2(\text{g}); K_2$

(c)  $\text{CH}_4(\text{g}) + 2\text{H}_2\text{O}(\text{g}) \rightleftharpoons \text{CO}_2(\text{g}) + 4\text{H}_2(\text{g}); K_3$

Which of the following relations is correct ?

- (a)  $K_2 K_3 = K_1$
- (b)  $K_3 = K_1 K_2$
- (c)  $K_3 K_2^3 = K_1^2$
- (d)  $K_1 \sqrt{K_2} = K_3$

21. A vessel at 1000 K contains  $\text{CO}_2$  with a pressure of 0.5 atm. Some of the  $\text{CO}_2$  is converted into CO on the addition of graphite. If the total pressure at equilibrium is 0.8 atm, the value of K is :

- (a) 1.8 atm
- (b) 3 atm
- (c) 0.3 atm
- (d) 0.18

22. Ratio of active masses of 22 g  $\text{CO}_2$ , 3g  $\text{H}_2$  and 7g  $\text{N}_2$  in a gaseous mixture :

- (a) 22 : 3 : 7
- (b) 0.5 : 3 : 7
- (c) 1 : 3 : 1
- (d) 1 : 3 : 0.5

23.  $\text{PCl}_5 \rightleftharpoons \text{PCl}_3 + \text{Cl}_2$  in the reversible reaction the moles of  $\text{PCl}_5$ ,  $\text{PCl}_3$  and  $\text{Cl}_2$  are a, b and c respectively and total pressure is P then value of  $K_p$  is :

- (a)  $\frac{bc}{a} \cdot RT$  (b)  $\frac{b}{(a+b+c)} \cdot P$   
 (c)  $\frac{bc \cdot P}{a(a+b+c)}$  (d)  $\frac{c}{(a+b+c)} \cdot P$

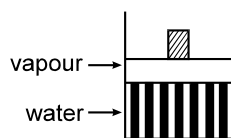
24. The degree of dissociation of  $\text{PCl}_5$  (a) obeying the equilibrium,  $\text{PCl}_5 \rightleftharpoons \text{PCl}_3 + \text{Cl}_2$ , is approximately related to the pressure at equilibrium by (given  $a \ll 1$ ) :

- (a)  $\alpha \propto P$  (b)  $\alpha \propto \frac{1}{\sqrt{P}}$  (c)  $\alpha \propto \frac{1}{P^2}$  (d)  $\alpha \propto \frac{1}{P^4}$

25. In the system,  $\text{LaCl}_3(\text{s}) + \text{H}_2\text{O}(\text{g}) + \text{heat} \rightleftharpoons \text{LaClO}(\text{s}) + 2\text{HCl}(\text{g})$ , equilibrium is established. More water vapour is added to reestablish the equilibrium. The pressure of water vapour is doubled. The factor by which pressure of HCl is changed is:

- (a) 2 (b)  $\sqrt{2}$  (c)  $\sqrt{3}$  (d)  $\sqrt{5}$

26. Some quantity of water is contained in a container as shown in figure. As neon is added to this system at constant pressure, the amount of liquid water in the vessel



- (a) Increases (b) Decreases  
 (c) Remains same (d) Changes unpredictably

27. When a bottle of cold drink is opened, the gas comes out with a fizz due to :

- (a) Decrease in temperature  
 (b) Increase in pressure  
 (c) Decrease in pressure suddenly which results in decrease of solubility of  $\text{CO}_2$  gas in water  
 (d) None

28. In the Haber process for the industrial manufacturing of ammonia involving the reaction,

$\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$  at 200 atm pressure in the presence of a catalyst, a temperature of about  $500^\circ\text{C}$  is used. This is considered as optimum temperature for the process because

- (a) Yield is maximum at this temperature  
 (b) Catalyst is active only at this temperature  
 (c) Energy needed for the reaction is easily obtained at this temperature  
 (d) Rate of the catalytic reaction is fast enough while the yield is also appreciable for this exothermic reaction at this temperature.

29. For the reaction,  $\text{CO}(\text{g}) + \text{Cl}_2(\text{g}) \rightleftharpoons \text{COCl}_2(\text{g})$  then

$K_p/K_c$  is equal to :

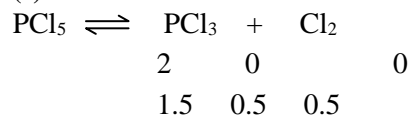
- (a)  $1/RT$  (b) 1.0 (c)  $\sqrt{RT}$  (d)  $RT$

30. Consider the following reversible gaseous reactions (at 298 K) :

- (a)  $\text{N}_2\text{O}_4 \rightleftharpoons 2\text{NO}_2$  (a)  $\text{N}_2\text{O}_4 \rightleftharpoons 2\text{NO}_2$   
 (b)  $2\text{SO}_2 + \text{O}_2 \rightleftharpoons 2\text{SO}_3$   
 (c)  $2\text{HI} \rightleftharpoons \text{H}_2 + \text{I}_2$   
 (d)  $\text{X} + \text{Y} \rightleftharpoons 4\text{Z}$



1. (c)



$$K_c = \frac{(0.5)(0.5)}{1.5}$$

2. (a)

$$u = \sqrt{\frac{3P}{d}}$$

3. (c)

$$K = \frac{K_f}{K_b}$$

4. (a)

$$K_p = K_c (RT)^{\Delta n_g}$$

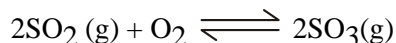
5. (b)

$$\frac{1}{K} = \frac{1}{0.025} = 40$$

6. (b)

$$\frac{\text{st. of HA}_1}{\text{st. of HA}_2} = \sqrt{\frac{K_{a_1}}{K_{a_2}}}$$

7. (b)



$$K_p = 4.0 \text{ atm}^{-2}$$

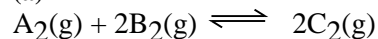
$$K_p = \frac{(\text{SO}_3)^3}{(\text{SO}_2)^2(\text{O}_2)}$$

Given that at equilibrium the amount of  $\text{SO}_2$  and  $\text{SO}_3$  is the same so

$$\frac{(\text{SO}_3)^2}{(\text{SO}_2)^2(\text{O}_2)} = 4$$

$$\Rightarrow \square [\text{O}_2] = \frac{1}{4} = 0.25 \text{ atm.}$$

8. (a)



$$P_{\text{A}_2} = 0.80 \text{ atm.}, \quad P_{\text{B}_2} = 0.4 \text{ atm.}$$

Total pressure of the system = 2.8 atm.

$$\therefore P_{\text{C}_2} = 2.8 - 0.8 - 0.4 = 1.6$$

$$K_p = \frac{P_{\text{C}_2}^2}{P_{\text{A}_2} \times P_{\text{B}_2}^2} = \frac{(1.6)^2}{0.8 \times (0.4)^2} = 20$$

9. (c)

No. of A atoms = 6.

$$\text{No. of C atoms} = 6 \times \frac{2}{3} = 4.$$

$$\therefore \text{Formula} = \text{C}_4\text{A}_6 \text{ or } \text{C}_2\text{A}_3.$$

10. (a)

$$(\text{VD})_{\text{mix}} \frac{M_{\text{mix}}}{2} = \frac{M}{2(1+\alpha)}$$

so, as  $\alpha$  increases  $(\text{VD})_{\text{mix}}$  decreases.

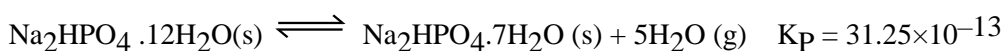
11. (c)

Gas product concentration increases while that of solid reactant is constant.

12. (a)

From thermodynamics.

13. (d)



$$K_p = (P_{\text{H}_2\text{O}})^5$$

$$(P_{\text{H}_2\text{O}})^5 = 31.25 \times 10^{-3}$$

$$(P_{\text{H}_2\text{O}}) = (3125)^{1/5} \times (10^{-15})^{1/5}$$

$$(P_{\text{H}_2\text{O}}) = 5 \times 10^{-3}$$

14. (d)

At equilibrium between liquid and vapour, molecules have same kinetic energy.

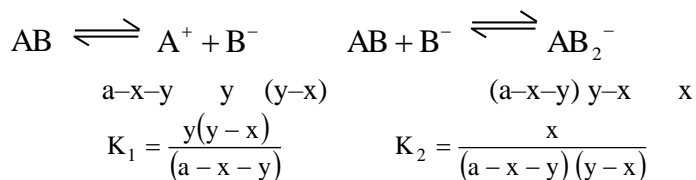
15. (b)

On adding inert gas at constant pressure effect on equilibrium will be similar to as if volume of container has been increased.

16. (b)

For constant volume, reaction quotient (Q) will remain constant.

17. (a)



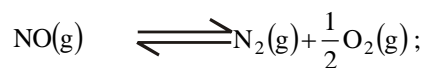
$$\frac{K_1}{K_2} = \frac{\left[ \frac{y(y-x)}{(a-x-y)} \right]}{\frac{x}{(a-x-y)(y-x)}}$$

$$\Rightarrow \frac{K_1}{K_2} = \frac{y}{x} (y-x)^2$$

18. (d)



$$K_c = \frac{[\text{NO}]^2}{[\text{N}_2][\text{O}_2]} = 4 \times 10^{-4}$$



19. (d)

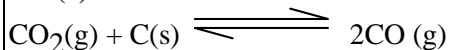
$$K_p = K_c (RT)^{\Delta n} \quad \Delta n = 3 - 2 = 1.$$

$$K_p = K_c (0.0821 \times 457)^1. \quad K_p > K_c.$$

20. (b)

$$c = a + b$$

21. (a)



$$0.5 \text{ atm} \quad \quad 0.5 - p \quad \quad 2p$$

$$\text{Total pressure} = 0.5 - p + 2p = 0.8$$

$$p = 0.3$$

$$K_p \frac{P_{\text{CO}}^2}{P_{\text{CO}_2}} = \frac{(2p)^2}{(0.5 - p)} = \frac{(0.6)^2}{(0.5 - 0.3)}$$

$$K_p = 1.8$$

22. (d)

$$\text{Moles of CO}_2 = \frac{22}{44} = \frac{1}{2}$$

$$\text{Moles of H}_2 = \frac{3}{2}$$

$$\text{Moles of N}_2 = \frac{7}{28} = \frac{1}{4}$$

$$\text{Ratio of active masses} = 1/2 : 3/2 : 1/4 \text{ or } 1 : 3 : 0.5$$

23. (c)

$$K_p = \frac{P'_{\text{PCl}} \times P'_{\text{Cl}_2}}{P'_{\text{PCl}_5}} = \frac{\frac{b}{(a+b+c)} P \times \frac{c}{(a+b+c)} \times P}{\frac{a}{(a+b+c)} P}$$

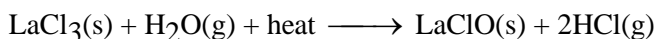
$$K_p = \frac{bc.P}{a(a+b+c)}$$

24. (b)

$$K_p = \frac{\alpha^2}{1 - \alpha^2} P \approx \alpha^2 P.$$

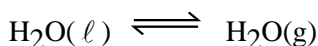
$$\text{so,} \quad \alpha \approx \sqrt{\frac{K_p}{P}}$$

25. (b)



$$K_p = \frac{P_{\text{HCl}}^2}{P_{\text{H}_2\text{O}}} = \frac{P_{\text{HCl}}^2}{2P_{\text{H}_2\text{O}}} \quad P_{\text{HCl}} = \sqrt{2}P_{\text{H}_2\text{O}}$$

26. (b)

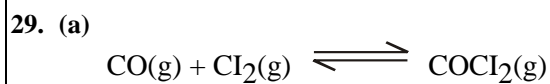




$$K_p = (P_{\text{H}_2\text{O}})$$

When neon is added at constant pressure, we have to increase volume of the container. So more water will evaporate to maintain equilibrium

27. (c) Solubility of gas is directly proportional to the pressure of gas above liquid.
28. (d) On increasing temperature though reaction equilibrium shifts in the backward direction but for rate of reaction to be higher, higher temperature is required and particle  $500^\circ\text{C}$  is found to be optimum temperature.



$$\Delta n = 1 - 2 = -1; K_p = K_c (\text{RT})^{\Delta n} \quad \therefore \quad \frac{K_p}{K_c} = (\text{RT})^{-1} = \frac{1}{\text{RT}}$$

30. (a)
- Use  $K_p = K_c (\text{RT})^{\Delta n}$