



## CHEMISTRY

### BOOKS - IIT-JEE PREVIOUS YEAR (CHEMISTRY)

#### CHEMICAL AND IONIC EQUILIBRIUM

Jee Main And Advanced

1. The equilibrium constant at  $298K$  for a reaction,  $A + B \rightleftharpoons C + D$  is 100. If the initial concentrations of all the four species were 1M each, then equilibrium concentration of  $D$  (in  $\text{molL}^{-1}$ ) will be

A. 0.818

B. 1.818

C. 1.182

D. 0.182

**Answer: B**



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2. The standard Gibbs energy change at  $300K$  for the reaction  $2A \rightleftharpoons B + C$  is  $2494.2J$ . At a given time, the composition of the reaction mixture is  $[A] = \frac{1}{2}$ ,  $[B] = 2$  and  $[C] = \frac{1}{2}$ . The reaction proceeds in the

( $R = 8.314JK/mole = 2.718$ )

A. forward direction because  $Q > K_e$

B. reverse direction because  $Q > K_e$

C. forward direction because  $Q < K_e$

D. reverse direction because  $Q < K_e$

**Answer: B**

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3. For the reaction,  $SO_2(g) + \frac{1}{2}O_2(g) \rightleftharpoons SO_3(g)$  if  $K_p = K_C(RT)^x$  where, the symbols have usual meaning, then the value of  $x$  is (assuming ideality)

A.  $-1$

B.  $-\frac{1}{2}$

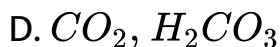
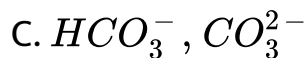
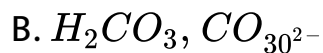
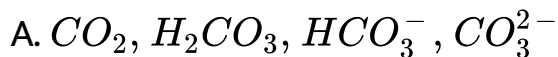
C.  $\frac{1}{2}$

D. 1

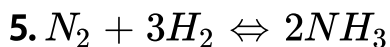
**Answer: B**

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4. The species present in solution when  $CO_2$  is dissolved in water



**Answer: A**



Which is correct statement if  $N_2$  is added at equilibrium condition?

A. The equilibrium will shift to forward direction

because according to II<sup>nd</sup> law of thermodynamics, the entropy must increases in the direction of spontaneous reaction

B. The condition for equilibrium is

$$G(N_2) + 3G(H_2) = 2G(NH_3) \text{ where } G \text{ is Gibbs}$$

free energy per mole of the gaseous species

measured at that partial pressure. The condition of

equilibrium is unaffected by the use of catalyst, which increases the rate of both the forward and backward reactions to the same extent

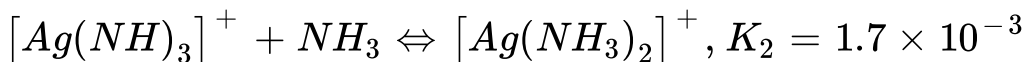
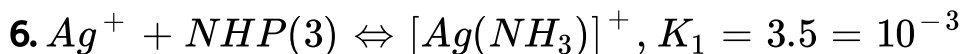
C. The catalyst will increase the rate of forward reaction by  $\alpha$  and that of backward reaction by  $\beta$

D. Catalyst will not alter the rate of either of the reaction

**Answer: B**



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then the formation constant of  $[Ag(NH_3)_2]^+$  is

A.  $6.08 \times 10^{-6}$

B.  $6.08 \times 10^6$

C.  $6.08 \times 10^{-9}$

D. None of these

**Answer: A**



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7. Consider the following equilibrium in a closed container



At a fixed temperature, the volume of the reaction container is halved. For this change, which of the following

statements hold true regarding the equilibrium constant ( $K_p$ ) and degree of dissociation ( $\alpha$ )?

- A. Neither  $K_p$  nor  $\alpha$  changes
- B. Both  $K_p$  and  $\alpha$  change
- C.  $K_p$  changes but  $\alpha$  does not change
- D.  $K_p$  does not change but  $\alpha$  changes

**Answer: D**



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8. At constant temperature, the equilibrium constant  $K_p$  for by  $K_p = \frac{4x^2p}{(1-x^2)}$ , where  $p$  = pressure  $x$  = extent of



decomposition. Which one of the following statement is true?

- A.  $K_p$  increases with increase of  $p$
- B.  $K_p$  increases with increase of  $x$
- C.  $K_p$  increases with decreases of  $x$
- D.  $K_p$  remains constant with change in  $p$  and  $x$

**Answer: D**



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9. When two reactants  $A$  and  $B$  are mixed to give products,  $C$  and  $D$ , the reaction quotient ( $Q$ ) at the initial stages of the reaction

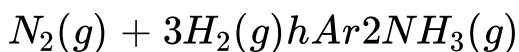
- A. is zero
- B. decreases with time
- C. is independent of time
- D. increases with time

**Answer: D**



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**10.** For the reversible reaction



at  $500^\circ C$  the value of  $K_p$  is  $1.44 \times 10^{-5}$  when partial pressure is measured in atmosphere. The corresponding value of  $K_c$  with concentration in mol/L is

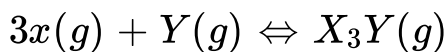
- A.  $\frac{1.44 \times 10^{-5}}{(0.082 \times 500)^{-2}}$
- B.  $\frac{1.44 \times 10^{-5}}{(8.314 \times 73)^{-2}}$
- C.  $\frac{1.44 \times 10^{-5}}{(0.082 \times 773)^2}$
- D.  $\frac{1.44 \times 10^{-5}}{(0.082 \times 773)^{-2}}$

**Answer: D**



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**11.** For the chemical reaction,



the amount of  $X_3Y$  at equilibrium is affected by

A. temperature and pressure

B. temperature only

C. pressure only

D. temperature, pressure and catalyst

**Answer: A**



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**12.** For the reaction,

$CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g)$ , at a given temperature, the equilibrium amount of  $CO_2(g)$  can be increased by

A. adding a suitable catalyst

B. adding an inert gas

C. decreasing the volume of the container

D. increasing the amount of  $CO(g)$

**Answer: D**

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13. One mole of  $N_2O(g)$  at  $300K$  is kept in a closed container under one atmosphere. It is heated to  $600K$  when 20 % by mass of  $N_2O_4(g)$  decomposes of  $NO_2(g)$ .

The resultant pressure

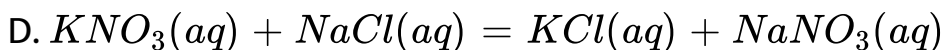
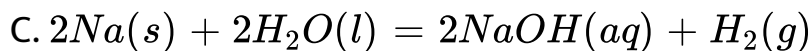
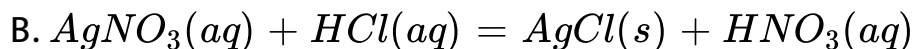
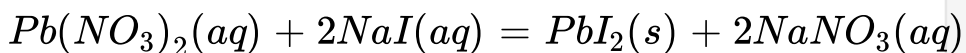
- A. 1.2 atm
- B. 2.4 atm
- C. 2.0 atm
- D. 1.0 atm

**Answer: B**

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**14.** An example of a reversible reaction is

A.



**Answer: D**

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15. Pure ammonia is placed in a vessel at a temperature where its dissociation constant ( $\alpha$ ) is appreciable. At equilibrium,  $N_2 + 3H_2 \rightleftharpoons 2NH_3$

A.  $K_p$  does not change significantly with pressure

B.  $\alpha$  does not change with pressure

C. concentration of  $NH_3$  does not change with pressure

D. concentration of hydrogen is less than that of nitrogen

**Answer: A**



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16. For the reaction  $H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$

the equilibrium constant  $K_p$  changes with

A. total pressure

B. catalyst

C. the amount of  $H_2$  and  $I_2$  present

D. temperature

**Answer: D**



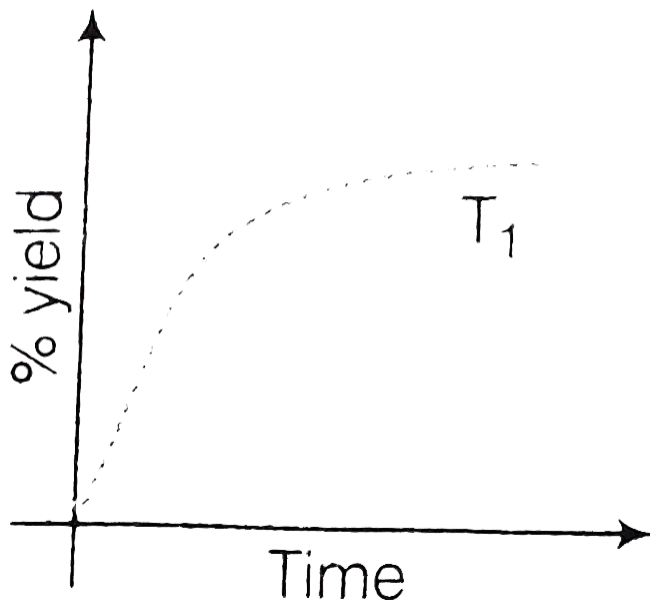
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17. The % yield of ammonia as a function of time in the

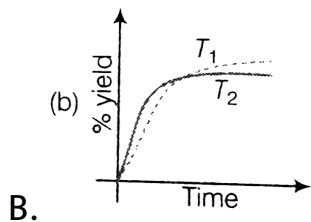
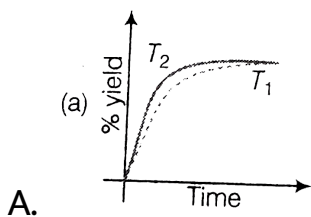
reaction,  $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ ,  $\Delta H < 0$

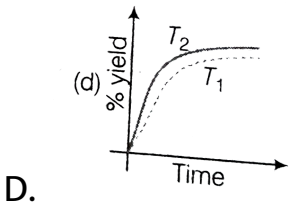
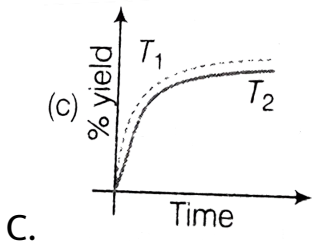
at  $(p, T_1)$  is given below





If this reaction is conducted at  $(p, T_2)$ , with  $T_2 > T_1$  the % yield by of ammonia as a function of time is represented by





**Answer: B**

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**18.** The thermal dissociation of equilibrium of  $CaCO_3(s)$  is studied under different conditions



For this equilibrium, the correct statement (s) is/are

- A.  $\Delta H$  is dependent on  $T$
- B.  $K$  is independent of the initial amount of  $CaCO_3$
- C.  $K$  is dependent of the pressure of  $CO_2$  at a given  $T$
- D.  $\Delta H$  is independent of the catalyst, if any

**Answer: A::B::D**



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**19.** The initial rate of hydrolysis of methyl acetate (1M) by a weak acid ( $HA$ , 1M) is 1/100th of that of a strong acid ( $HX$ , 1M), at  $25^\circ C$ . The  $K_a(HA)$  is

A.  $1 \times 10^{-4}$

B.  $1 \times 10^{-5}$

C.  $1 \times 10^{-6}$

D.  $1 \times 10^{-3}$

**Answer: A**



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**20.** The equilibrium



in aqueous medium at  $25^\circ C$  shifts towards the left in the presence of



D.  $CN^-$

**Answer: B::C::D**

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21. For the reaction  $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$  the forward reaction at constant temperature is favoured by

- A. introducing an inert gas at constant volume
- B. introducing chlorine gas at constant volume
- C. increasing the volume of the container
- D. introducing  $PCl_5$  at constant volume

**Answer: C::D**

22. The equilibrium  $SO_2Cl_2(g) \rightleftharpoons SO_2(g) + Cl_2(g)$  is attained at  $25^\circ C$  in a closed container and an inert gas, helium is introduced. Which of the following statements are correct?

- A. Concentration of  $SO_2$ ,  $Cl_2$  and  $SO_2Cl_2$  change
- B. More chlorine is formed
- C. Concentration of  $SO_2$  is reduced
- D. None of the above

**Answer: D**

23. When  $NaNO_3$  is heated in a closed vessel, oxygen is liberated and  $NaNO_2$  is left behind. At equilibrium

- A. addition of  $NaNO_2$  favours reverse reaction
- B. addition of  $NaNO_3$  favours forward reaction
- C. increasing temperature favours forward reaction
- D. increasing pressure favours reverse reaction

**Answer: C::D**

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24. For the gas phase reaction,

$C_2H_4 + H_2 \rightleftharpoons C_2H_6$  ( $\Delta H = -32.7$  kcal) carried out in a

vessel, the equilibrium concentration of  $C_2H_4$  can be increased by

- A. increasing the temperature
- B. decreasing the pressure
- C. removing some  $H_2$
- D. adding some  $C_2H_6$

**Answer: A::B::C::D**

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25. For a gaseous reaction  $2B \rightarrow A$ , the equilibrium constant  $K_p$  is .....to/than  $K_c$

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26. A ten-fold increase in pressure on the reaction,  $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$  at equilibrium, results in .....in  $K_p$

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27. For a given reversible reaction at a fixed temperature, equilibrium constants  $K_p$  and  $K_c$  are related by .....

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28. The rate of an exothermic reaction increases with increasing temperature. True or False?



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29. Catalyst makes a reaction more exothermic. True or False?

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30. If equilibrium constant for the reaction

$A_2 + B_2 \rightleftharpoons 2AB$  is  $k$ , then for the backward reaction

$AB \rightleftharpoons 1/2A_2 + 1/2B_2$  the equilibrium constant  $k'$  is

$1/K$ .

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**31.** When a liquid and its vapour are at equilibrium and the pressure is suddenly decreased, cooling occurs. True or false?

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**32.** (a) In the following equilibrium  $N_2O_4(g) \rightleftharpoons 2NO_2(g)$  when 5 moles of each are taken, the temperature is kept at  $298K$  the total pressure was found to be 20 bar. Given that  $\Delta G_f^\circ(N_2O_4) = 100kJ$ ,  $\Delta G_f^\circ(NO_2) = 50kJ$

(i) Find  $\Delta G$  of the reaction.

(ii) The direction of the reaction in which the equilibrium shifts.

(b) A graph is plotted for a real gas which follows van der

Waals' equation with  $pV_m$  taken on  $Y$ -axis and  $p$  on  $X$ -axis.

Find the intercept of the line where  $V_m$  is molar volume.

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**33.** When  $3.06\text{g}$  of solid  $\text{NH}_4\text{HS}$  is introduced into a two-litre evacuated flask at  $27^\circ\text{C}$ ,  $30\%$  of the solid decomposes into gaseous ammonia and hydrogen sulphide. (i) Calculate  $K_c$  and  $K_p$  for the reaction at  $27^\circ\text{C}$ . (ii) What would happen to the equilibrium when more solid  $\text{NH}_4\text{HS}$  is introduced into the flask?

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**34.** The degree of dissociation is 0.4 at  $400K$  and 1.0 atm for the gaseous reaction

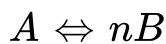


assuming ideal behaviour of all gases, calculate the density of equilibrium mixture at  $400K$  and 1.0 atm (relative atomic mass of P is 31.0 and of Cl is 35.5).

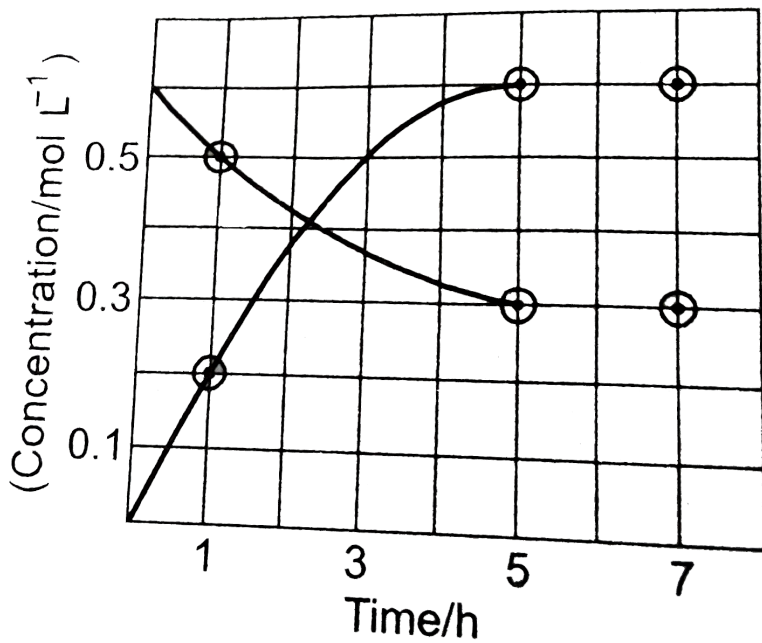


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**35.** The progress of reaction



with time, is represented in fig use given below.



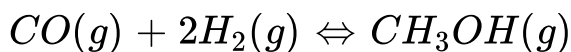
Determine:

- (i) the value of  $n$
- (ii) the equilibrium constant,  $K$  and
- (iii) the initial rate of conversion of  $A$



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**36.** When 0.15 mol of CO taken in a 2.5L flask is maintained at 750K along with a catalyst, the following reaction takes place



Hydrogen is introduced until the total pressure of the system is 8.5 atm at equilibrium and 0.08 mol of methanol is formed.

Calculate

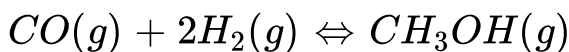
a.  $K_p$  and  $K_c$

b. The final pressure, if the same amount of CO and  $H_2$  as before are used, but with no catalyst so that the reaction does not take place.



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**37.** For the reaction



Hydrogen gas is introduced into a five-litre flask at  $327^\circ C$ , containing 0.2 mol of  $CO(g)$  and a catalyst, until the pressure is  $4.92\text{atm}$ . At this point, 0.1 mol of  $CH_3OH(g)$  is formed. Calculate the equilibrium constants  $K_p$  and  $K_c$ .

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**38.** The equilibrium constant  $K_p$  of the reaction:



containing  $SO_3$  and  $O_2$  having initial pressure of 1 atm and 2 atm respectively, is heated at constant volume to equilibrate. Calculate the partial pressure of each gas at  $800K$  at equilibrium.





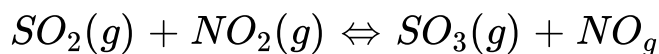
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39.  $N_2(g)$  is 25% dissociated at  $37^\circ C$  and one atmosphere pressure. Calculate (i)  $K_p$  and (ii) the percentage dissociation at 0.1 atm and  $37^\circ C$



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40. At a certain temperature, equilibrium constant ( $K_c$ ) is 16 for the reaction,



If we take one mole each of all the four gases in a one litre container, what would be the equilibrium concentrations of  $NO$  and  $NO_2$ ?





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**41.** The equilibrium constant of the reaction  $A_2(g) + B_2(g) \rightleftharpoons 2AB(g)$  at  $100^\circ C$  is 50. If a one litre flask containing one mole of  $A_2$  is connected to a two litre flask containing two moles of  $B_2$ , how many moles of  $AB$  will be formed at  $373K$ ?



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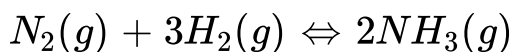
**42.** 1mol of  $Cl_2$  and 3 mol of  $PCl_5$  are placed in a 100L vessel heated to  $227^\circ C$ . The equilibrium pressure is 2.05 atm. Assuming ideal behaviour, calculate the degree of dissociation for  $PCl_5$  and  $K_p$  for the reaction.



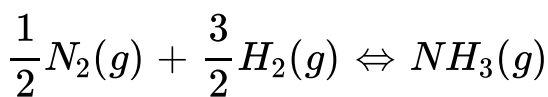


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43. 1 mol of nitrogen is mixed with 3 mol of hydrogen in a 4L container. If 0.25 % of nitrogen is converted to ammonia by the following reaction

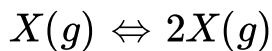


then calculate the equilibrium constant  $K_c$  in concentration units. What will be the value of  $K_c$  for the following equilibrium?



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**44.** Thermal decomposition of gaseous  $X_2$  to gaseous  $X$  at  $298K$  takes place according to the following equation:



The standard reaction Gibbs energy  $\Delta_r G^\circ$ , of this reaction is positive. At the start of the reaction, there is one mole of  $X_2$  and no  $X$ . As the reaction proceeds, the number of moles of  $X$  formed is given by  $\beta$ . Thus  $\beta_{\text{equilibrium}}$  is the number of moles of  $X$  formed at equilibrium. The reaction is carried out at a constant total pressure of 2 bar. Consider the gases to behave ideally.

[Given,  $R = 0.083L \text{ bar } K^{-1} \text{ mol}^{-1}$ ]

The equilibrium constant  $K_p$  for this reaction at  $298K$ , in terms of  $\beta_{\text{equilibrium}}$  is

A. 
$$\frac{8\beta^2_{\text{equilibrium}}}{2 - \beta_{\text{equilibrium}}}$$

- B.  $\frac{8\beta^2 - (\text{equilibrium})}{4 - \beta_{\text{equilibrium}}}$
- C.  $\frac{4\beta^2 - (\text{equilibrium})}{2 - \beta_{\text{equilibrium}}}$
- D.  $\frac{4\beta^2 - (\text{equilibrium})}{4 - \beta_{\text{equilibrium}}}$

**Answer: D**

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**45.** Thermal decomposition of gaseous  $X_2$  to gaseous  $X$  at  $298K$  takes place according to the following equation:



The standard reaction Gibbs energy  $\Delta_r G^\circ$ , of this reaction is positive. At the start of the reaction, there is one mole of  $X_2$  and no  $X$ . As the reaction proceeds, the number of moles of  $X$  formed is given by  $\beta$ . Thus  $\beta_{\text{equilibrium}}$  is the

number of moles of  $X$  formed at equilibrium. The reaction is carried out at a constant total pressure of 2 bar. Consider the gases to behave ideally.

[Given,  $R = 0.083L \text{ bar } K^{-1} \text{ mol}^{-1}$ )]

The incorrect statement among the following for this reaction, is

A. Decrease in the total pressure will result in the formation of more moles of gaseous  $X$

B. At the start of the reaction, dissociation of gaseous  $X_2$  takes place spontaneously

C.  $\beta_{\text{equilibrium}} = 0.7$

D.  $K_C < 1$

**Answer: D**

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46.  $pK_a$  of a weak acid ( $HA$ ) and  $pB_b$  of a weak base ( $BOH$ ) are 3.2 and 3.4 respectively. The  $pH$  of their salt ( $AB$ ) solution is

A. 7.2

B. 6.9

C. 7.0

D. 1.0

**Answer: B**

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47. How many litres of water must be added to 1L of an aqueous solution of  $HCl$  with a  $pH$  of 1 to create an aqueous solution with  $pH$  of 2?

A. 0.1L

B. 0.9L

C. 2.0L

D. 9.0L

**Answer: D**



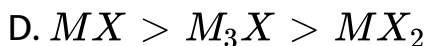
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48. The solubility product constant ( $K_{sp}$ ) of salts of types  $MX$ ,  $MX_2$ , and  $M_3X$  at temperature  $T$  are



$4.0 \times 10^{-8}$ ,  $3.2 \times 10^{-14}$ , and  $2.7 \times 10^{-15}$ , respectively.

The solubilities of the salts at temperature  $T$  are in the order



**Answer: D**



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49.  $2.5\text{mL}$  of  $\frac{2}{5}$  weak monoacidic base ( $K_b = 1 \times 10^{-12}$  at  $25^\circ\text{C}$ ) is titrated with  $\frac{2}{15}MHCl$  in water at  $25^\circ\text{C}$ . The

concentration of  $H^+$  at equivalence point is:

$$(K_w = 1 \times 10^{-14} \text{ at } 25^\circ C)$$

A.  $3.7 \times 10^{-13} M$

B.  $3.2 \times 10^{-7} M$

C.  $3.2 \times 10^{-2} M$

D.  $2.7 \times 10^{-2} M$

**Answer: D**



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50. 0.1 mole of  $CH_3NH_2$  ( $K_b = 5 \times 10^{-4}$ ) is mixed with 0.08 mole of  $HCl$  and diluted to one litre. The  $[H^+]$  in solution is

A.  $1.6 \times 10^{-11}$

B.  $8 \times 10^{-11}$

C.  $5 \times 10^{-5}$

D.  $8 \times 10^{-2}$

**Answer: B**



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51.  $HX$  is a weak acid ( $K_a = 10^{-5}$ ). It forms a salt  $NaX(0.1M)$  on reacting with caustic soda. The degree of hydrolysis of  $NaX$  is

A. 0.01 %

B. 0.0001 %

C. 0.1 %

D. 0.5 %

**Answer: A**



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52. A solution which is  $10^{-3}M$  each in  $Mn^{2+}$ ,  $Fe^{2+}$ ,  $Zn^{2+}$ , and  $Hg^{2+}$  is treated with  $10^{-16}M$  sulphide ion. If the  $K_{sp}$  of  $MnS$ ,  $FeS$ ,  $ZnS$  and  $HgS$  are  $10^{-15}$ ,  $10^{-23}$ ,  $10^{-20}$ , and  $10^{-54}$ , respectively, which one will precipitate first?

A.  $FeS$

B.  $MgS$

C.  $HgS$

D.  $ZnS$

**Answer: C::D**



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**53.** Identify the correct order of solubility in aqueous medium

A.  $CuS > ZnS > Na_2S$

B.  $ZnS > Na_2S > CuS$

C.  $Na_2S > CuS > ZnS$

D.  $Na_2S > ZnS > CuS$

**Answer: D**

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**54.** For a sparingly soluble salt  $A_pB_q$ , the relationship of its solubility product ( $L_s$ ) with its solubility ( $S$ ) is

A.  $L_s = S^{p+q} \cdot p^p \cdot q^q$

B.  $L_s = S^{p+q} \cdot p^q \cdot q^p$

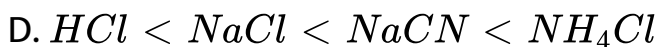
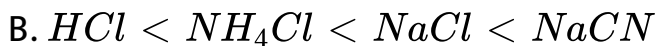
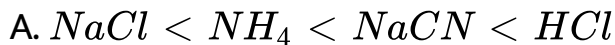
C.  $L_s = S^{pq} \cdot p^p \cdot q^q$

D.  $L_s = S^{pq} \cdot (p \cdot q)^{(p+q)}$

**Answer: A**

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55. The  $pH$  of  $0.1M$  solution of the following salts decreases in the order



**Answer: B**

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56. Which solution will have  $pH$  closer to 1.0 ?



B.  $55\text{mL}$  of  $(M/10)\text{HCl}$  +  $45\text{mL}$  of  $(M/10)\text{NaOH}$

C.  $10\text{mL}$  of  $(M/10)\text{HCl}$  +  $90\text{mL}$  of  $(M/10)\text{NaOH}$

D.  $75\text{mL}$  of  $(M/5)\text{HCl}$  +  $25\text{mL}$  of  $(M/5)\text{NaOH}$

**Answer: D**

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57. Amongst the following hydroxides, the one which has the lowest value of  $K_{sp}$  is:

A.  $\text{Mg}(\text{OH})_2$

B.  $\text{Ca}(\text{OH})_2$

C.  $\text{Ba}(\text{OH})_2$

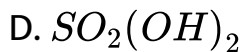
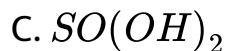
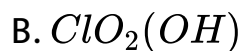
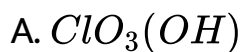
D.  $\text{Be}(\text{OH})_2$



**Answer: D**

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**58.** Which of the following is the strongest acid?



**Answer: A**

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59. When equal volumes of following solution are mixed, precipitation of  $AgCl$  ?

( $K_{sp} = 1.8 \times 10^{-10}$ ) will occur only with

A.  $10^{-4}M(Ag^+)$  and  $10^{-4}M(Cl^-)$

B.  $10^{-5}M(Ag^+)$  and  $10^{-6}M(Cl^-)$

C.  $10^{-6}M(Ag^+)$  and  $10^{-6}M(Cl^-)$

D.  $10^{-10}M(Ag^+)$  and  $10^{-10}M(Cl^-)$

**Answer: A**



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60. The  $pK_a$  of acetylsalicylic acid (aspirin) is 3.5. The pH of gastric juice in human stomach is about 2 – 3 and the pH

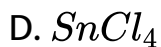
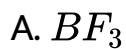
in the small intestine is about 8. Aspirin will be:

- A. unionised in the small intestine and in the stomach
- B. completely ionised in the small intestine and in the stomach
- C. ionised in the stomach and almost unionised in the small intestine
- D. ionised in the small intestine and almost unionised in the stomach

**Answer: D**

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**61.** The compound that is not a Lewis acids is



**Answer: C::D**



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**62.** The conjugate acid of amide ion ( $NH_2^-$ ) is



D.  $N_2H_4$

**Answer: A**

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**63.** The best indicator for detection of end point in titration of a weak acid and a strong base is

- A. methyl orange (3 to 4)
- B. methyl red (5 to 6)
- C. bromothymol blue (6 to 7.5)
- D. phenolphthalein (8 to 9.6)

**Answer: D**

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**64.** A certain weak acid has a dissociation constant  $1.0 \times 10^{-4}$ . The equilibrium constant for its reaction with a strong base is :

A.  $1.0 \times 10^{-4}$

B.  $1.0 \times 10^{-10}$

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

D.  $1.0 \times 10^{-14}$

**Answer: C::D**

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65. A certain buffer solution contains equal concentration of  $X^{\ominus}$  and  $HX$ . The  $K_b$  for  $X^{\ominus}$  is  $10^{-10}$ . The  $pH$  of the buffer is

A. 4

B. 7

C. 10

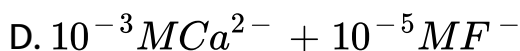
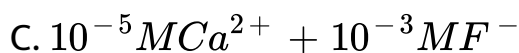
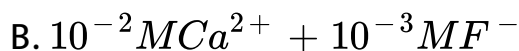
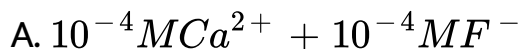
D. 14

**Answer: A**



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66. The precipitate of  $CaF_2$  ( $K_{sp} = 1.7 \times 10^{-10}$ ) is obtained when equal volumes of the following are mixed



**Answer: B**

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67. An acidic buffer solution can be prepared by mixing solution of



- A. acetate and acetic acid
- B. ammonium chloride and ammonium hydroxide
- C. sulphuric acid and sodium sulphate
- D. sodium chloride and sodium hydroxide

**Answer: A**

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**68.** Of the given anions, the strongest base is





**Answer: A**

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69. At  $90^\circ C$  pure water has  $[H_3O^+]$  as  $10^{-6} molL^{-1}$ .

What is the value of  $K_w$  at  $90^\circ C$ ?

A.  $10^{-6}$

B.  $10^{-12}$

C.  $10^{-14}$

D.  $10^{-8}$

**Answer: B**



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70. The  $pH$  of  $10^{-8}M$  solution of  $HCl$  in water is

A. 8

B.  $-8$

C. between 7 and 8

D. between 6 and 7

Answer: D



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71. The  $K_{sp}$  of  $Ag_2CrO_4$  is  $1.1 \times 10^{-12}$  at  $298K$ . The solubility (in mol/L) of  $Ag_2CrO_4$  in a  $0.1M AgNO_3$

solution is

A.  $1.1 \times 10^{-11}$

B.  $1.1 \times 10^{-10}$

C.  $1.1 \times 10^{-12}$

D.  $1.1 \times 10^{-9}$

**Answer: B**



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**72.** Aqueous solution of  $HNO_3$ ,  $CH_3$ ,  $CH_3COOH$ , and  $CH_3COOK$  of identical concentrations are given. The pair (s) of the solution which may form a buffer upon mixing is (are):

A.  $HNO_3$  and  $CH_3COOH$

B.  $KOH$  and  $CH_3COONa$

C.  $HNO_3$  and  $CH_3COONa$

D.  $CH_3COOH$  and  $CH_3COONa$

**Answer: C::D**



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**73.** A buffer solution can be prepared from a mixture of

A. sodium acetate and acetic acid in water

B. sodium acetate and  $HCl$  in water

C. ammonia and ammonium chloride in water

D. ammonia and sodium hydroxide in water

Answer: A::B::C

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74.  $(CH_3(OH)_2^1)$  is..acidic than  $(CH_3NH_3^+)$ .

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75. In the reaction,  $I^- + I_2 \rightarrow I_3^-$  the Lewis acid is....

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76. Silver chloride is sparingly soluble in water because its lattice energy is greater than \_\_\_\_\_

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77. An element which can exist as a positive ion in acidic solution and also as a negative ion in basic solution is said to be.....

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78. The conjugate base of  $HSO_4^-$  in aqueous solution is.

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79. The following species are in increasing order of their acidic property:  $ZnO$ ,  $Na_2O$ ,  $P_2O_5$ ,  $MgO$

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80. The solubility of sodium hydroxide increases with increase of temperature.

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81. Aluminium chloride ( $AlCl_3$ ) is a Lewis acid because it can donate electrons.

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**82.** The molar conductivity of a solution of a weak acid  $HX(0.01M)$  is 10 times smaller than the molar conductivity of a solution of a weak acid  $HY(0.10M)$ . If  $\lambda_{X^-}^\circ = \lambda_{Y^-}^\circ$ , the difference in their  $pK_a$  values,  $pK_a(HX) - pK_a(HY)$ , is (consider degree of ionisation of both acids to be  $< < 1$ ):

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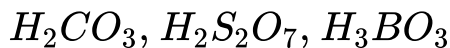
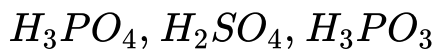
**83.**  $MX_2$  dissociates into  $M^{2+}$  and  $X^-$  ions in an aqueous solution, with a degree dissociation ( $\alpha$ ) of 0.5. The ratio of the observed depression of freezing point of the aqueous solution to the value of the depression of freezing point in the absence of ionic dissociation is

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**84.** In 1L saturated solution of  $AgCl$  [ $K_{SP}(AgCl) = 1.6 \times 10^{-10}$ ], 0.1 mole of  $CuCl$  [ $K_{SP}(CuCl) = 1.0 \times 10^{-6}$ ] is added. The resultant concentration of  $Ag^+$  in the solution is  $1.6 \times 10^{-x}$ . The value of x is:

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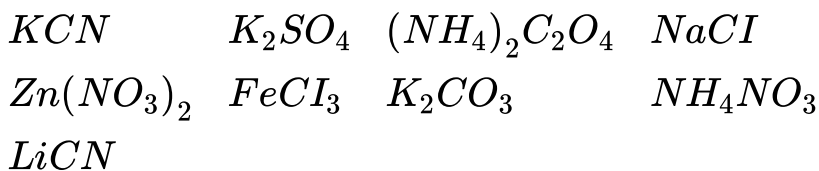
**85.** The total number of diprotic acids among the following is





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**86.** Amongst the following, the total number of compounds whose aqueous solution turns red litmus paper blue is:



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**87.** The dissociation constant of a substituted benzoic acid at  $25^\circ C$  is  $1.0 \times 10^{-4}$ . The  $pH$  of  $0.01M$  solution of its sodium salt is



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**88.**  $0.1M NaOH$  is titrated with  $0.1M HA$  till the end point.  $K_a$  of  $HA$  is  $5.6 \times 10^{-6}$  and degree of hydrolysis is less compared to 1. Calculate  $pH$  of the resulting solution at the end point ?

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**89.**  $500mL$  of  $0.2M$  aqueous solution of acetic acid is mixed with  $500mL$  of  $0.2M HCl$  at  $25^\circ C$ .

a. Calculate the degree of dissociation of acetic acid in the resulting solution and  $pH$  of the solution.

b. If  $6g$  of  $NaOH$  is added to the above solution determine the final  $pH$ . [ $K_a$  of  $CH_3COOH = 2 \times 10^{-5}$ ].

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**90.** The average concentration of  $SO_2$  in the atmosphere over a city on a certain day is 10 ppm, when the average temperature is 298 K. Given that the solubility of  $SO_2$  in water at 298 K is  $1.3653 \text{ mol litre}^{-1}$  and the  $pK_a$  of  $H_2SO_3$  is 1.92, estimate the pH of rain on that day.

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**91.** The solubility of  $Pb(OH)_2$  in water is  $6.7 \times 10^{-6} \text{ M}$ . Calculate the solubility of  $Pb(OH)_2$  in a buffer solution of  $pH = 8$ .

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**92.** Find the solubility product of a saturated solution of  $Ag_2CrO_4$  in water at  $298K$ , if the  $EMF$  of the cell :

$$Ag|Ag^{\oplus}(sat\ Ag_2CrO_4\ sol)||Ag(0.1M)|Ag\ is\ 0.164V \quad \text{at}$$

$298K$ .

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**93.** A sample of  $AgCl$  was treated with  $5.00mL$  of  $1.5M$   $Na_2CO_3$  solubility to give  $Ag_2CO_3$  . The remaining solution contained  $0.0026g\ of\ Cl^{-}$  per litre. Calculate the solubility product of  $AgCl$ .

( $K_{SP}$  of  $Ag_2CO_3 = 8.2 \times 10^{-12}$ )

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**94.** An acid type indicator,  $HIn$  differs in colour from its conjugate base ( $In^-$ ). The human eye is sensitive to colour differences only when the ratio  $[In^-]/[HIn]$  is greater than 10 or smaller than 0.1. What should to observe a complete colour change? ( $K_a = 1.0 \times 10^{-5}$ )

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**95.** The ionization constant of  $NH_4^+$  ion in water is  $5.6 \times 10^{-10}$  at  $25^\circ C$ . The rate constant the reaction of  $NH_4^+$  and  $OH^-$  ion to form  $NH_3$  and  $H_2O$  at  $25^\circ C$  is  $3.4 \times 10^{10} Lmol^{-1}s^{-1}$ . Calculate the rate constant for proton transfer form water to  $NH_3$ .

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96. What is the pH of a 0.50M aqueous NaCN solution ?

( $pK_b$  of  $CN^- = 4.70$ )

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97. Calculate the pH of an aqueous solution of 1.0M

ammonium formate assuming complete dissociation. ( $pK_a$

of formic acid = 3.8 and  $pK_b$  of ammonia = 4.8)

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98. For the reaction

$Ag(CN)_2^{\ominus} \rightleftharpoons Ag^{\oplus} + 2CN^{\ominus}$ , the  $K_c$  at  $25^{\circ}C$  is



$4 \times 10^{-19}$  Calculate  $[Ag^{\oplus}]$  in solution which was originally  $0.1M$  in  $KCN$  and  $0.03M$  in  $AgNO_3$ .

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**99.** An aqueous solution of a metal bromide  $MBr_2(0.05M)$  is saturated with  $H_2S$ . What is the minimum pH at which  $MS$  will precipitate ?  $K_{SP}$  for  $MS = 6.0 \times 10^{-21}$  .  
Concentration of saturated  
 $H_2S = 0.1M, K_1 = 10^{-7}$  and  $K_2 = 1.3 \times 10^{-13}$  for  
 $H_2S$  .

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**100.** The pH of blood stream is maintained by a proper balance of  $H_2CO_3$  and  $NaHCO_3$  concentrations. What volume of 5 M  $NaHCO_3$  solution, should be mixed with 10 mL sample of blood, which is 2 M in  $H_2CO_3$  in order to maintain a pH of 7.4 ( $K_a$  of  $H_2CO_3$  in blood =  $7.8 \times 10^{-7}$ )

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**101.** The  $K_{SP}$  of  $Ca(OH)_2$  is  $4.42 \times 10^{-5}$  at  $25^\circ C$ . A 500 mL of saturated solution of  $Ca(OH)_2$  is mixed with equal volume of 0.4 M  $NaOH$ . How much  $Ca(OH)_2$  in mg is precipitated ?

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**102.** A 50 mL solution of weak base BOH is titrated with 0.1N HCl solution. The pH of solution is found to be 10.04 and 9.14 after the addition of 5.0 mL and 20.0 mL of acid respectively. Find out  $K_b$  for weak base.

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**103.** The  $K_{SP}$  of  $Ag_2C_2O_4$  at  $25^\circ C$  is  $1.29 \times 10^{-11} mol^3 L^{-3}$ . A solution of  $K_2C_2O_4$  containing 0.152 mole in 500 mL water is shaken at  $25^\circ C$  with excess of  $Ag_2CO_3$  till the equilibrium is reached.



At equilibrium the solution contains 0.0358 mole of  $K_2CO_3$ . Assuming degree of dissociation of  $K_2C_2O_4$  and  $K_2CO_3$  to be same, calculate  $K_{SP}$  of  $Ag_2CO_3$ .



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**104.** What is the pH of 1 M solution of acetic acid ? To what volume one litre of this solution be diluted so that pH of the resulting solution will be twice of the original value ?

$$(K_a = 1.8 \times 10^{-5})$$



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**105.** Freshly precipitated Al and Mg hydroxides are stirred vigorously in a buffer solution containing 0.25M of  $NH_4Cl$  and 0.05M of  $NH_4OH$ . Calculate  $[Al^{3+}]$  and  $[Mg^{2+}]$  in solution.  $K_b$  for  $NH_4OH = 1.8 \times 10^{-5}$   $K_{SP}$  of

$$Al(OH)_3 = 6 \times 10^{-32}$$

and

$$K_{SP}$$

of

$$Mg(OH)_2 = 8.9 \times 10^{-12}.$$

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**106.** How many mole of HCl will be required to prepare one litre of buffer solution (containing  $NaCN + HCl$ ) of  $pH 8.5$  using  $0.01g$  formula weight of NaCN ?

$$K_{HCN} = 4.1 \times 10^{-10}$$

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**107.** What is the  $pH$  of the solution when  $0.20mol$  of HCl is added to  $1L$  of a solution containing

a.  $1M$  each of acetic acid and acetate ion.

b.  $0.1M$  each of acetic acid and acetate ion.

Assume the total volume is  $1L$ .  $K_a$  for acetic acid is  $1.8 \times 10^{-5}$ .

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**108.** The solubility of  $Mg(OH)_2$  in pure water is  $9.57 \times 10^{-3} \text{ g litre}^{-1}$ . Calculate the pH of its saturated solution. (Assume 100% ionisation)

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**109.** The concentration of hydrogen ions in a  $0.2M$  solution of formic acid is  $6.4 \times 10^{-3} \text{ molL}^{-1}$ . To this solution, sodium formate is added so as to adjust the concentration

of sodium formate to  $1\text{molL}^{-1}$ . What will be the  $pH$  of this solution? The dissociation constant of formic acid is  $2.4 \times 10^{-4}$  and the degree of dissociation for sodium formate is 0.75.

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**110.** A solution contains a mixture of  $Ag^+$  ( $0.10M$ ) and  $Hg_2^{2+}$  ( $0.10M$ ) which are to be separated by selective precipitation. Calculate the maximum concentration of iodide ion at which one of them gets precipitated almost completely. What % of that metal ion is precipitated ?

( $K_{SP}$  of  $AgI = 8.5 \times 10^{-17}$  and  $K_{SP}$  of  $Hg_2I_2 = 2.5 \times 10^{-26}$ )

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**111.** The dissociation constant of weak acid  $HA$  is  $4.9 \times 10^{-8}$ . After making the necessary approximations, calculate  $pH$  in  $0.1M$  acid.

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**112.** Given reason for the statement that the  $pH$  of an aqueous solution of sodium acetate is more than 7.

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**113.**  $20ml$  of  $0.2MNaOH$  is added to  $50ml$ , of  $0.2MCH_3COOH$  to give  $70ml$ , of the solution. What is



the  $pH$  of the solution? The ionization constant of acetic acid is  $2 \times 10^{-5}$

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**114.** What amount of solution propanoate should be added to one litre of an aqueous solution containing 0.02 mole of propanoic acid ( $K_a = 1.0 \times 10^{-5}$  at  $25^\circ C$ ) to obtain a buffer solution of  $pH6$

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**115.** When  $100mL$  of  $1.0M HCl$  was mixed with  $100mL$  of  $1.0M NaOH$  in an insulated beaker at constant pressure, a temperature increase of  $5.7^\circ C$  was measured for the

beaker and its contents (Expt.1). Because the enthalpy of neutralisation of a strong acid with a strong base is constant ( $-57.0 \text{ kJ mol}^{-1}$ ) this experiment could be used to measure the calorimeter constant. In a second experiment (Expt.2),  $100 \text{ mL}$  of  $2.0 \text{ M}$  acetic acid ( $K_a = 2.0 \times 10^{-5}$ ) was mixed with  $100 \text{ mL}$  of  $1.0 \text{ M NaOH}$  (under identical conditions to Expt. 1) where a temperature rise of  $5.6^\circ \text{ C}$  was measured.

Enthalpy of dissociation (in  $\text{kJ mol}^{-1}$ ) of acetic acid obtained from the Expt. 2 is

- A. 1.0
- B. 10.0
- C. 24.5
- D. 51.4

**Answer: A**

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**116.** When  $100\text{mL}$  of  $1.0\text{M HCl}$  was mixed with  $100\text{mL}$  of  $1.0\text{M NaOH}$  in an insulated beaker at constant pressure, a temperature increase of  $5.7^\circ\text{C}$  was measured for the beaker and its contents (Expt.1). Because the enthalpy of neutralisation of a strong acid with a strong base is constant ( $-57.0\text{kJmol}^{-1}$ ) this experiment could be used to measure the calorimeter constant. In a second experiment (Expt.2),  $100\text{mL}$  of  $2.0\text{M}$  acetic acid ( $K_a = 2.0 \times 10^{-5}$ ) was mixed with  $100\text{mL}$  of  $1.0\text{M NaOH}$  (under identical conditions to Expt. 1) where a temperature

rise of  $5.6^{\circ}C$  was measured.

The  $pH$  of the solution after Expt. 2 is

A. 2.8

B. 4.7

C. 5.0

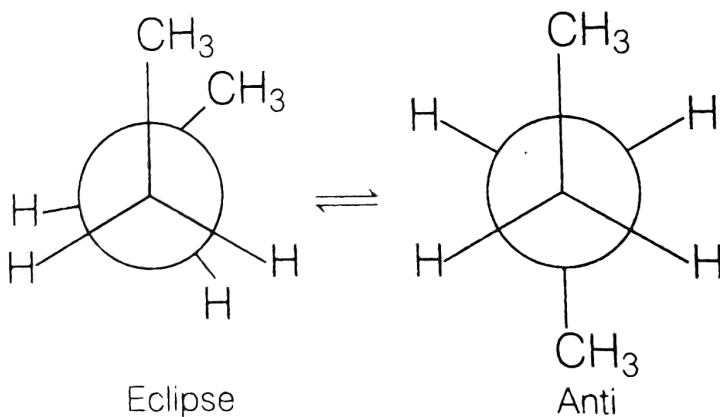
D. 7.0

**Answer: D**



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117. Consider the equilibrium in the right margin.



- A. The change is exothermic and increasing temperature decreases the equilibrium amount of eclipse form
- B. The change is exothermic and increasing temperature increases the equilibrium amount of anti form

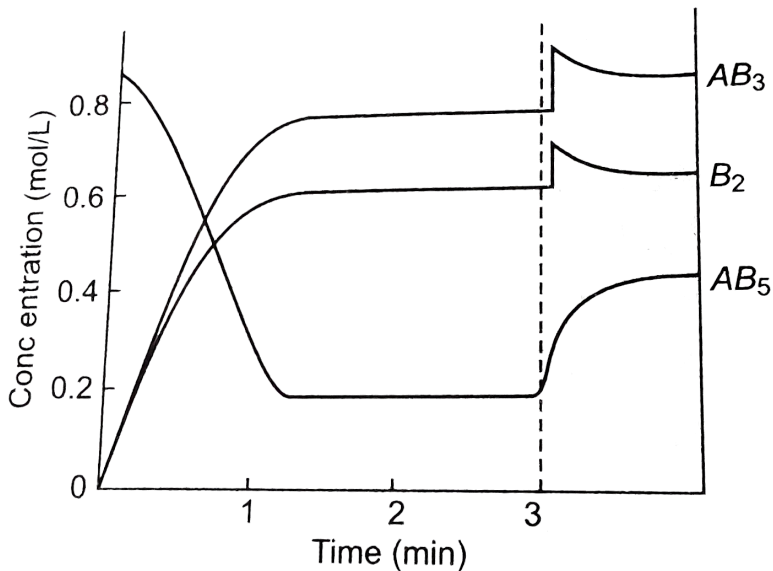
C. The change is exothermic and decreasing temperature decreases the equilibrium amount of eclipse form

D. The change is endothermic and increasing temperature decreases the equilibrium amount of anti form

**Answer: C**

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**118.** The equation for the reaction in the figure given below is



At time three minutes what change was imposed into the equilibrium?

- A. Pressure was increased
- B. Temperature was increased
- C.  $B_2$  was added to the system
- D.  $AB_3$  was added to the system

Answer: A



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119. Which of the following mixtures would result in a buffered solution?

A. Mixing  $100.0\text{mL}$  of  $0.100\text{M HCl}$  with  $100.0\text{mL}$  of

$0.100\text{M NaOH}$

B. Mixing  $100.0\text{mL}$  of  $0.100\text{M NH}_3$  with  $100.0\text{mL}$  of

$0.100\text{M NaOH}$

C. Mixing  $100.0\text{mL}$  of  $0.100\text{M HCl}$  with  $100.0\text{mL}$  of

$0.100\text{M NH}_3$



D. Mixing  $50.0\text{mL}$  of  $0.100\text{M HCl}$  with  $100.0\text{mL}$  of  $0.100\text{M NHP}(3)$

**Answer: D**

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**120.**  $0.03\text{ M}$  aqueous solution of a weak monobasic acid solution ( $K_a = 10^{-12}$ ) is titrated against a  $0.01\text{M NaOH}$  solution. What is the  $[H^+]$  at the end point?

A.  $1.15 \times 10^{-12}\text{M}$

B.  $2 \times 10^{-12}\text{M}$

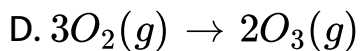
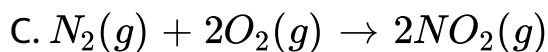
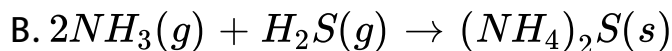
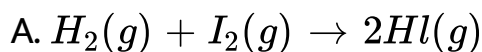
C.  $5 \times 10^{-12}\text{M}$

D.  $4 \times 10^{-12}\text{M}$

**Answer: B**

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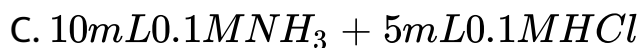
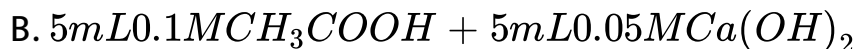
**121.** The reaction for which  $K_C > K_P$  at a given temperature is (are)



**Answer: B::C::D**

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122. Which of the following solution will have  $pH$  greater than 7 at  $25^\circ C$ ?



Answer: B::C::D

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123. Assertion: To a system at equilibrium addition of inert gas at constant pressure and temperature drive the

reaction to the side where larger number of active species is present.

Reason: Addition of inert gas at constant temperature and pressure increases the equilibrium volume.

- A. Both Assertion and Reason are correct and Reason is the correct explanation on the Assertion.
- B. Both Assertion and Reason are correct but Reason is not the correct explanation of Assertion
- C. Assertion is correct but Reason is incorrect
- D. Assertion is incorrect but Reason is correct

**Answer: D**



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124. A  $50\text{mL} 10^{-3}\text{M HCl}$  solution is mixed with  $50\text{mL} 8 \times 10^{-4}\text{M NaOH}$  solution. What would be the  $pH$  of resulting solution?

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### Objective Type

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

A. The  $pH$  of  $1.0 \times 10^{-8}\text{M}$  solution of  $\text{HCl}$  is 8

B. The conjugate base of  $\text{H}_2\text{PO}_4^-$  is  $\text{HPO}_4^{2-}$

C. Autoprotolysis constant of water increases with temperature

D. When a solution of a weak monoprotic acid is titrated against a strong base, at half-neutralisation point

$$pH = \left(\frac{1}{2}\right)pK_a$$

**Answer: B::C**

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2. The salts  $AgX$ ,  $AgY$  and  $AgZ$  are all equally soluble in water (and none are very soluble at all). When each salt is added to separate beakers of  $100\text{mL}$  of  $1.0\text{M HNO}_3$ , you notice that  $AgY$  is much more soluble than  $AgZ$  in acid. The salt  $AgX$  is no more soluble in strong acid than it is in water. Which of the following best explains this?

A.  $HX$ ,  $HY$  and  $HZ$  are all weak acids

B.  $HX$  is a weak acid and  $HY$  and  $HZ$  are strong acids

C.  $HY$  and  $HZ$  are weak acids and  $HX$  is a strong acid,

$HY$  is a stronger acid than  $HZ$

D.  $HY$  and  $HZ$  are weak acids and  $HX$  is a strong acid

$HZ$  is a stronger acid than  $HY$

**Answer: D**

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3. Solubility of which of the following salts can be increased by adding some  $HNO_3$  solution?

A.  $BaF_2$

B.  $CH_3COOAg$

C.  $BaSO_4$

D.  $PbS$

**Answer: A::B::D**

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## Comprehension Type

1. To determine molar solubility of an unknown metal hydroxide,  $M(OH)_3$  in  $0.00101MNaOH$ , Jane adds  $0.250g$  of  $M(OH)_3$  solid to  $200mL$  of a  $0.00101MNaOH$  solution and stirred for a long time. A cloudy solution is formed. The undissolved hydroxide is separated from the



solution by filtratin  $25.00\text{mL}$  of filtrate is pipetted into a flask  $30\text{mL}$  of water is added adn then the solution is titrated with  $0.00444\text{MHCl}$  solution  $13\text{mL}$  of  $\text{HCl}$  is used for this titration.

The molar concentration of hydroxide ion in filtrate solution is

- A.  $0.0015\text{M}$
- B.  $0.0023\text{M}$
- C.  $0.00333\text{M}$
- D.  $0.00462\text{M}$

**Answer: B**



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2. To determine molar solubility of an unknown metal hydroxide,  $M(OH)_3$  in  $0.00101MNaOH$ , Jane adds  $0.250g$  of  $M(OH)_3$  solid to  $200mL$  of a  $0.00101MNaOH$  solution and stirred for a long time. A cloudy solution is formed. The undissolved hydroxide is separated from the solution by filtration.  $25.00mL$  of filtrate is pipetted into a flask  $30mL$  of water is added and then the solution is titrated with  $0.00444MHCl$  solution  $13mL$  of  $HCl$  is used for this titration.

The solubility product of  $M(OH)_3$  is

A.  $4.8 \times 10^{-9}$

B.  $5.6 \times 10^{-11}$

C.  $2.1 \times 10^{-11}$

D.  $5.34 \times 10^{-12}$

**Answer: D**

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3. To determine molar solubility of an unknown metal hydroxide,  $M(OH)_3$  in  $0.00101MNaOH$ , Jane adds  $0.250g$  of  $M(OH)_3$  solid to  $200mL$  of a  $0.00101MNaOH$  solution and stirred for a long time. A cloudy solution is formed. The undissolved hydroxide is separated from the solution by filtration.  $25.00mL$  of filtrate is pipetted into a flask  $30mL$  of water is added and then the solution is titrated with  $0.00444MHCl$  solution  $13mL$  of  $HCl$  is used for this titration.

If Jane carelessly added  $20mL$  of water to the flask instead of  $30mL$  how would this affect the determination of

solubility product ( $K_{sp}$ ) and molar solubility?

(1) The value of  $K_{sp}$  determined by Jane would be same as true value

(2) The value of  $K_{sp}$  determined by Jane would be higher than the true value.

(3) The value of  $K_{sp}$  determined by Jane would be lower as true value.

(4) The molar solubility of  $M(OH)_3$  determined by Jane would be same as true value.

(5) The molar solubility of  $M(OH)_3$  determined by Jane would be higher than true value.

(6) The molar solubility of  $M(OH)_3$  determined by Jane would be lower than the true value

A. 1 and 4

B. 2 and 5

C. 3 and 6

D. 1 and 6

**Answer: A**



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**Ar Type**

1. Assertion: Addition of  $NaCN$  to a saturated solution of  $AgCN$  increases the solubility of  $AgCN$

Reason:  $NaCN$  produces common ion effect when added to a saturated solution of  $AgCN$

- A. Both Assertion and Reason are correct and Reason is the correct explanation on the Assertion.
- B. Both Assertion and Reason are correct but Reason is not the correct explanation of Assertion
- C. Assertion is correct but Reason is incorrect
- D. Assertion is incorrect but Reason is correct

**Answer: B**

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**Match The Column**

## 1. Match the column I and Column II

Column I	Column II
A. Addition of small amount of HCl decreases the solubility of sparingly soluble salt	p. A saturated solution of AgCl ( $K_{sp} = 2 \times 10^{-10}$ )
B. Addition of small amount of NaOH decreases solubility of sparingly soluble salt.	q. A saturated solution of $PbCl_2$ ( $K_{sp} = 4 \times 10^{-6}$ )
C. Addition of small amount of HCl increases solubility of sparingly soluble salt.	r. HCl solution saturated with AgCl
D. Addition of small amount of $AgNO_3(s)$ increases solubility of sparingly soluble salt.	s. NaOH solution saturated with $Mg(OH)_2$ ( $K_{sp}$ of $Mg(OH)_2 = 4 \times 10^{-7}$ )



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## Integer Type

1. Consider the following reversible system:



At equilibrium there are 1.0 mole of A and 2.0 moles of each

B, C and D present. If 2.0 moles of B is added so that moles of A and D do not change?



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