

| | <u> </u> | www.pakcit | ty.org | | | Class 11 th : | Chemist | ry notes |
|-----|---|---|--------------------------|------------------|---|---|----------|--|
| 32. | Which statement about the following equilibrium is correct? $2SO_2 + O_2 \Longrightarrow 2SO_3$ (A) The value of K_p falls with increase in pressure. (B) The value of K_p falls with rise in temperature. (C) The value of K_p is equal to K_c . (D) Adding V_2O_5 catalyst increases the equilibrium yield of SO_3 . For the reaction $2SO_2 + O_2 \Longrightarrow 2SO_3$: | | | | | | | |
| 33. | For | the reaction 2SO ₂ - | $+ O_2 = 2S$ | O ₃ : | | | | |
| | A | $K_c = K_x$ | | | © | $K_c < K_p$ | (D) | $K_c = K_p$ |
| 34. | The | units for K _w of H ₂ O | are: | | | | | |
| | A | Mol+2 dm-6 | [®] Mol⁻² dn | n+6 | © | Mol ⁻² dm ⁻³ | (D) | Mol dm ⁻³ |
| 35. | 3 <u>2.</u> 300.00 | ction of BiCl ₃ with ppears by: Adding HCI Decreasing tempe | | vhile ppt. | of B D | BiOCl and HCl i Adding BiCl ₃ Increase tempe | | ed. The white ppt |
| 36. | The | value of K _w at 25°C | lis: | | • | | | |
| | (A) | 0.30×10^{-14} | (B) 3 × 10 ⁻¹ | 4 | (c) | 0.11×10^{-14} | (D) | 1×10^{-14} |
| 37 | The | optimum tempera | ture for the s | onthesis o | | | ncess is | • |
| | | 500 °C | B 300 °C | y Hericolo C | | 400 °C | | 200 °C |
| 38 | | llyst used in prepar | | from No 2 | <u> </u> | ~~ (O) | | |
| 50. | (A) | V ₂ O ₅ | B Fe | II OIII INZ a | (A) | Ni Ni | | Pt |
| 20 | In cr | | | process | Pho | ontimum conditi | on for n | roccuro ici |
| 39. | | nthesis of ammon 200-300atm | | - 10 O | | _ | | |
| | | | V V | 3.) | | | | |
| 40. | | en KCI is added to a Forward Direction | | | | - 1 | | |
| | | | | A Simple | X THE | | | ackward Direction |
| 41. | | which system does 2HF | • | | | $H_2 + I_2 \stackrel{\text{Unit (o)}}{=} 2H$ | | rationj-1: |
| | | $2NO_2 \longrightarrow N_2O_4$ | | A-IIII | | $N_2 + 3H_2 =$ | | |
| 42. | An a | queous solution of | ethanol in w | ater may | hav | e vapour pressur | e: | |
| | A | Less than that of | | | B | More than that | | |
| | (C) | Equal to that of w | | | (D) | Equal to that of | t ethand |)] |
| 43. | A | increase in dilution Decreases the deg Increase the degr | gree of dissoc | | _ | Depends upon t Does not affect | | ree of the solute ree of dissociation |
| 44. | A so | lution with $pH = 0$ | indicates mo | lar conce | ntra | tion of H ⁺ ions: | | |
| | A | 10 ⁺⁷ | | | © | 1.0 | D 1 | 10^{-14} |
| 45. | Mola | arity of pure water | is: | | | | | |
| | A | 6 | B 55.5 | | © | 10 | (D) | 18 |
| 46. | The | nature of milk is: | | | | | | |
| | A | Acidic | Normal | | © | Basic | (D) | Neutral |
| 47 | Sum | of p K_a and p K_b is ϵ | _ | | | | | |
| | | pria ana prip io c | 7 | | | | | |

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|------------|------------|---|--------------------------------------|--------|---|-------------|---------------|---------------------------------------|
| | | | | | | | | .ev |
| | A | 0 | B 1 | © | 7 | D | 14 | |
| 48. | The | pH of 10^{-4} moles d | m^{-3} of Ba (OH) ₂ is: | | | | | |
| | A | 6.4 | ® 7.5 | 0 | 10.3 | D | 4.5 | |
| 49. | The | pH of 10^{-3} moles d | m ⁻³ of an aqueous so | lutio | on of H ₂ SO ₄ is: | | | |
| | A | 3.0 | B 2.7 | © | 2.0 | D | 1.5 | |
| 50. | PH | of buffer can be calc | ulated by using: | | | | | |
| | (A) (C) | Bohrs equation Moseley's equation | n | (D) | Henderson's equa De-Broglie's equa | | | |
| 5 1 | | | | 0 | De Brogne 3 equi | acioi. | <u> </u> | |
| | A | pH of the gastric jui | (B) 2.5 | (C) | 3 | (D) | 5.5 | |
| 52 | Whi | ch aqueous solution | | | | | | |
| 32. | (A) | 0.1M HCI | B 0.2M HNO ₃ | (c) | 0.1M H ₂ SO ₄ | | 0.1M NaOH | |
| 53 | The | | COONa and CH ₃ COOH | ic | 01111112004 | | OIIII IIII | |
|]33. | (A) | 1 | B 7 | (3) | < 7 | (D) | > 7 | |
| 54. | The | term pH was introd | ucod by: | | | | | |
| J.T. | (A) | Thomson | Sorenson | (C) | Goldstein | (D) | Henderson | |
| 55 | The | pH of human blood | | ^ | | | nenderbon | |
| | A | 7.35 | B 8.0 | | 7.63 | (D) | 7.53 | |
| 56 | The | | n water is suppresse | d hv | adding | | | ••••••• |
| | (A) | KMnO ₄ | (B) NaCl@3 | (C) | NaCl | (D) | KCI | |
| 57. | In th | e presence of comp | non ion, the ionization | n of | an electrolyte will: | | | |
| | A | Moderate change | | | decrease | D | Increase | |
| 58. | The | | eous solution of H ₂ S t | the s | olubility will be: | *********** | | |
| | A | Decreases | | В | Increase | | | |
| | (c) | First decreases the | en increases | (D) | Remain constant | | | |
| 59. | 98. | en HCI is added to H First increases the | 2SO ₄ aqueous solution | n, its | Remain constant | | | |
| | © | Increases | II decreases | (D) | Decreases | | | |
| 60. | An e | excess of aqueous si | lver nitrate is added | to a | queous barium chl | orid | e and precipi | tate is |
| | | | Vhat are the main ion | 04:20 | New Y | 1 (1 | _ | |
| | 5-5-5-A | Ag ⁺ and NO only Ag ⁺ and Ba ²⁺ NO $_3^{-2}$ | 1, | | Ba ²⁺ and NO ₃ ⁻¹ ar Ba ²⁺ and NO ₃ ⁻¹ o | | | |
| 61. | | | of AgCl is 2.0×10^{-10} | | | | oncentration | of Ag ⁺ |
| | | in the solution is: | rigorio zio / To | | | | | · · · · · · · · · · · · · · · · · · · |
| | | 1.0×10^{-10} moldn | | | 2.0×10^{-10} mold | | | |
| | | 4.0×10^{-2} moldm | ار د | (U | 1.4×10^{-5} mold | II-3 | | |
| | | | | | | | | |



- Q1: Law of mass action states that the at which a reaction proceeds, is directly proportional to the product of the active masses of the
- Q2: In an exothermic reversible reaction temperature will shift the equilibrium towards the forward direction.
- Q3: The equilibrium constant for the reaction $20_3 \iff 30_2$ is 10^{55} at 25° C, it tells that ozone is at room temperature.
- Q4: In a gas phase reaction, if the number of moles of reactants are equal to the number of moles of the products, K_C of the reaction is to the K_P .
- Q5: Buffer solution is prepared by mixing together a weak base and its salt with or a weak acid and its salt with

Answers

| 1. | rate, reactants | 2. | decrease | 3. | unstable | 4. | equal |
|----|--------------------------------|----|----------|----|----------|----|-------|
| | Church and all advances leaves | | | | 4 | | |

5. Strong acid, strong base

Chapter: 08

Chemical Equilibrium





- Q1: Define State of Dynamic Equilibrium.
- Ans: Dynamic Equilibrium:

 When the forward and reverse reactions are taking place exactly at the same rate the equilibrium established is called dynamic equilibrium.
- Q2: Difference between reversible and irreversible reactions.
- Ans: The difference between reversible and irreversible reactions is:

| Reversible reaction | Irreversible reaction |
|--|--|
| The reaction which can precede both in | The reaction which takes place only in one |
| forward as well as backward direction is | direction is called irreversible reaction. |
| called reversible reaction. | |
| $N_2 + 3H_2 = 2NH_3$ | $2Na + 2H_2O \longrightarrow 2NaOH + H_2$ |

- Q3: The direction of reaction is determined by K_C . OR How K_C predict the direction of a reaction.
- Ans: For this purpose K_C is compared with ratio of [Product]/ [reactants]:
 - \blacktriangleright If ratio is equal to K_C it means reaction is at equilibrium.
 - \succ If ratio is less than K_C the reaction will proceed in forward direction.
 - \triangleright If ratio is greater than K_C the reaction will proceed in backward direction.
- Q4: State law of mass action.



The rate at which substances react is directly proportional to their active masses.

Q5: Using Law of Mass action, how can you derive following expression for

$$N_2O_{4(g)} = 2NO_{2(g)} K_C = \frac{4x^2}{(a-x)V}$$
 (Indicate what a, x,V stands for).

Ans: $N_2O_{4(g)}$, the expression of K_C involves the factor of volume.

$$N_2O_{4(g)}$$
 = $2NO_{2(g)}$
 N_2O_4 = $2NO_2$

$$at t = 0sec 'a' moles$$
 $0 moles$
 $at t = t_{eq} 'a-x' moles$ $2x moles$

$$\frac{a-x}{v}$$
 $\frac{2x}{v}$

$$K_{C} = \frac{[NO_{2}]^{2}}{[N_{2}O_{4}]}$$

$$K_{C} = \frac{\left[\frac{2x}{v}\right]^{2}}{\left[\frac{a-x}{v}\right]} = \frac{\frac{4x^{2}}{v^{2}}}{\frac{a-x}{v}}$$

$$K_C = \frac{4x^2}{(a-x)v}$$

'a' is the initial number of moles of N_2O_4 , 'x' is number of moles of N_2O_4 decomposed and V is total volume of N_2O_4 and NO_2 at equilibrium stage.

Q6: Give equilibrium constant Kc expression for

$$N_2 + 3H_2 = 2NH_3$$

$$N_2 + 3H_2$$
 $2NH_3$

$$a-x$$
 $b-3x$ $2x$

$$\frac{a-x}{v}$$
 $\frac{b-3x}{v}$ $\frac{2x}{v}$

So,
$$K_{C} = \frac{[NH_{3}]^{2}}{[N_{2}][H_{2}]^{3}}$$
 pakci

$$K_{C} = \frac{\left[\frac{2x}{v}\right]^{2}}{\left[\frac{a-x}{v}\right]\left[\frac{b-3x}{v}\right]^{3}} = \frac{\left(\frac{4x^{2}}{v^{2}}\right)}{\left(\frac{a-x}{v}\right)\left[\frac{(b-3x)^{3}}{v^{3}}\right]}$$

$$K_C = \frac{4x^2v^2}{(a-x)(b-3x)^3}$$

Q7: State law of Mass Action. Give equilibrium constant expression K_c for the following reaction.

$$N_2 + 3H_2 = 2NH_3$$

Ans: Law of mass action:

Ans:

The rate at which substance reacts is directly proportional to its active mass and the rate of reaction is directly proportional to the product of active masses of the reactants.

$$N_2 + 3H_2 = 2NH_3$$

$$K_C = \frac{[NH_3]^2}{[N_2][H_2]^3}$$

Q8: Write two applications of equilibrium constant.

Ans: There are following applications of equilibrium constant:

- Prediction of direction of reaction.
- Extent of a chemical reaction.
- Effect of various factors on chemical equilibrium.
- Q9: Why the equilibrium constant value has its units for some of the reversible reactions but has not units for some other reactions?

Ans: Units of $K_{\mathbb{C}}$ depend upon the number of moles of reactants and products involved in the reaction.

- If number of moles of reactant and products are same it has no unit.
- If number of moles of reactants is different from products then K_C has units, related to the concentration or pressure.
- Q10: Reversible reaction attains the position of equilibrium which is dynamic in nature and not static. Explain it.
- Ans: At equilibrium state the reaction is not stopped. Only the rate of forward reaction becomes equal to reverse reaction. Since reaction is in progress in both the directions. Therefore equilibrium is dynamics in nature not static.
- Q11: The change of temperature disturbs both the equilibrium position and equilibrium constant of a reaction. Explain with reason?
- Ans: According to Le-Chateliers principle an increase in temperature will favour the endothermic reaction and decrease in temperature will favour the exothermic reaction. Therefore, change of temperature will disturb equilibrium position. The equilibrium constant is temperature dependent therefore with the change of temperature a new equilibrium position will be established.
- Q12: How K_C predicts the extent of chemical reaction?
- Ans: The value of K_C also helps us to predict extent of chemical reaction.

There are three possibilities:

Large K_C value:

If K_C value is large it means reaction is almost completed.

Small Kc value:

If $K_{\mathbb{C}}$ value is small it means reaction does not proceed appreciably in forward direction. Small amount of products will be formed.

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K_C is in fractions:

If K_C is in fractions. It means little forward reaction.

- Q13: What happen to the direction of reaction of a reversible reaction? OR When the ratio of concentration is less than actual K_c ?
- Ans: In reversible reaction, when the ratio of concentration is less than K_c , it indicates that more of the product is required to attain the equilibrium; therefore, the reaction will proceed in forward direction.

Qc < Kc

- Q14: Define K_f and K_b for reversible reactions.
- Ans: K_f and K_b :



 K_f is rate constant for forward reaction and K_b is rate constant for reverse or backward reaction. At equilibrium, rate of forward reaction becomes equal to reverse reaction.

Q15: Give the physical significance of K_f and K_b .

Ans: The rate of forward and reverse reaction tells us the condition on which a reaction will depend. It also tells about:

- Extent of a reaction.
- Direction of a reaction.

Q16: Why do rates of forward reactions slow down when the reversible reaction approaches the equilibrium stage?

Ans: The rate of forward reaction is directly proportional to molar conc. of reactants. Near the equilibrium stage, the concentration of reactant becomes small. Therefore, the rates of forward reaction slow down due to decrease in concentration.

Q17: Derive value of K_C for the reaction

$$PCl_5 \longrightarrow PCl_3 + Cl_2$$
?

Ans:
$$PCl_5 \longrightarrow PCl_3 + Cl_2$$
:

$$K_{C} = \frac{[PCl_{3}]3[Cl_{2}]}{[PCl_{5}]}$$

Q18: Increasing pressure increase the oxidation of SO_2 to SO_3 . Explain why?



Ans:
$$SO_2 + \frac{1}{2}O_2 = SO_3$$

The number of moles of products are smaller than number of moles of reactants. Hence according to Le-Chatelier's principle, an increase in pressure will favour the forward reaction. During this process the pressure is kept at one atmosphere. Hence by increasing the pressure the conc. of Oxygen increases, which results in greater yield of SO₃.

Q19: What are K_c and K_p and how these are related?

Ans: \mathbf{K}_{C} : When the concentrations of reactants or products are expressed in moles dm⁻³ the equilibrium expression called \mathbf{K}_{C} .

 $\mathbf{K_p}$: When the reactants or products are gases then the conc. terms may be replaced by partial pressure called K_p .

$$K_p = K_C (RT)^{\Delta n}$$

Where $R = Universal\ gas\ constant$.

 $\Delta n = Number of moles of products - number of moles of reactants.$

Q20: State Le-Chatelier's Principle. Discuss the effect of change in pressure on equilibrium position.

Ans: If a stress is applied to a system at equilibrium, the system acts in such a way so as to nullify, as far as possible, the effect of that stress.

The change in pressure or volume is important only for the reversible gaseous reactions where the number of moles of reactants and products are not equal Le-Chatelier's principle plays an important role, to predict the position and direction of the reaction.

Take the example of formation of SO_3 gas from SO_2 gas and O_2 gas.

Q21: How some reactions are effected by change in pressure?

Ans: The change in pressure or volume is important only for the reversible gaseous reaction where number of moles of reactants and products are not equal.



Le-Chatelier's principle plays an important role, to predict the position and direction of reaction,

For example:

$$2SO_2 + O_2 = 2SO_3$$

$$\Delta H = -194 \text{ KJmol}^{-1}$$

$$N_2 + 3H_2 = 2NH_3$$

$$\Delta H = -92.46 \text{ KJmol}^{-1}$$

In both reactions, numbers of moles of reactants are greater so increase in pressure will favor the reaction in forward direction.

Q22: Why during the synthesis of NH_3 temperature is kept low?

$$N_2 + 3H_2 = 2NH_3$$

This is an exothermic reaction. Hence decrease in temperature will favour the forward reaction. That's why during NH₃ preparation temperature kept low. Actually optimum temperature is 400 to 450°C.

This graph is just for your information.

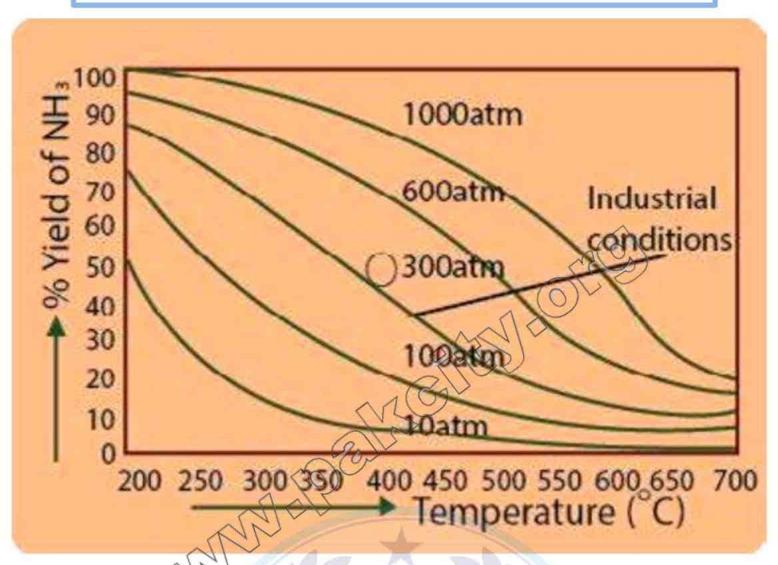


Fig Graphical representation of temperature and pressure for NH, synthesis.

Q23: How does the change of pressure shift the equilibrium position in the synthesis of ammonia?

$$N_{2(g)} + 3H_{2(g)}$$
 = 2NH_{3(g)}

Ans: When we increase or decrease the pressure of a gaseous system the equilibrium position is disturbed. If moles of products and reactants are same in reaction no disturbance in equilibrium position takes place. If moles of products and reactants are unequal, then reaction will move toward less number of moles by increasing pressure. In the formation of NH₃ more products is formed by increasing pressure.

$$N_2 + 3H_2 = 2NH_3$$

Q24: What are optimum conditions for the synthesis of NH₃? OR What are favourable and what are condition to maximize yield of NH₃?

Ans: At high pressure, low temperature and continuous removal of NH_3 will give best yield of NH_3 .

The optimum conditions are:

- 200-300 atm pressure.
- ➤ 400°C temperature
- \triangleright Piece of iron as catalyst embaded in fused mixture of MgO, Al₂O₃ and SiO₂.

Q25: What will be the effect of increase of pressure and temperature on the following reactions?



Ans:
$$N_2 + 3H_2 = 2NH_3$$

In this case number of moles of products is smaller than reactant. Therefore by increase in pressure will favour the forward reaction. Hence, NH₃ concentration will increase. But as we know that it is an exothermic reaction. So, by increasing temperature yield of NH₃ become lower.

Q26: What is the effect of temperature change on the direction of the reaction?

$$N_{2(g)} + 3H_{2(g)} = 2NH_{3(g)}$$

Ans: According to Le- Chatelier's principle "Decreasing the temperature will shift it to the forward direction".

Q27: What will be the effect of volume change on the following system at equilibrium state?

(A)
$$PCl_{5(g)} \longrightarrow PCl_{3(g)} + Cl_{2(g)}$$
 (B) $2SO_{2(g)} + O_{2(g)} \longrightarrow 2SO_{3(g)}$

Ans:
$$PCl_{5(g)} \longrightarrow PCl_{3(g)} + Cl_{2(g)}$$

The change in volume will affect the equilibrium position only. When volume is decreased reaction moves in backward direction to establish equilibrium again. But equilibrium constant is not affected.

$$2SO_{2(g)} + O_{2(g)} = 2SO_{3(g)}$$

When we decrease in volume for this reaction equilibrium position will shift in forward direction.

Q28: Why change of volume disturbs the equilibrium position for some of the gaseous phase reactions but not the equilibrium constant?

Ans:
$$2SO_{2(g)} + O_{2(g)} = 2SO_{3(g)}$$

This gas phase reaction proceeds with the decrease in the number of moles and hence decreases in volume at equilibrium stage. When the reaction approaches the equilibrium stage, the volume of the equilibrium mixture is less than the volume of reactants taken initially.

If one decrease the volume further at equilibrium state, the reaction is established a new equilibrium position while K_C remains constant. The reverse happens when the volume is increased or pressure is decreased at equilibrium stage.

Q29: What is the effect of catalyst of equilibrium constant?

Ans: A catalyst does not change the equilibrium position. A catalyst only increases the rate of forward and reverse reaction and attains equilibrium earlier.

Q30: Why catalyst does not affect the equilibrium position? OR A catalyst does not affect the equilibrium position and K_c of a reversible reaction. Explain.

Ans: In most of the reversible reactions, the equilibrium is not always reached within a suitable short time. So, an appropriate catalyst is added. A catalyst does not affect the equilibrium position of the reaction. It increases the rates of both forward and backward reactions and it reduces the time to attain the state of equilibrium. Actually a catalyst lowers the energy of activation of both forward and reverse steps by giving new path to the reaction.

Q31: What is the effect of increase in temperature on the yield of the product for the reaction?



Ans:
$$2SO_{2(g)} + O_{2(g)} = 2SO_{3(g)} + Heat$$

It is an exothermic reaction in the forward direction. It means that by increasing temperature, the reaction at equilibrium will shift the reaction to the backward direction by decreasing the yield of product.

What will be the effect of increase of pressure on the decomposition of PCl₅?

Ans:
$$PCl_{5(g)} \longrightarrow PCl_{3(g)} + Cl_{2(g)}$$

Since the number of moles of reactant are smaller than products. Therefore by increasing pressure equilibrium will shift towards left side and more amount of PCl₅ will produce.

Define pK_a and pK_b.

The values of Ka and Kb for weak acids and bases are small numbers usually expressed in Ans: exponential form. It is convenient to convert them into whole numbers by taking their negative log. Thus we obtain pKa and pkb Values of acids and bases.

$$pK_a = -\log K_a$$

$$pK_b = -\log K_b$$

Q34: Prove that
$$pK_a + pK_b = 14$$
 at 25 °C.

Ans: For an acid $K_a \times K_b = K_W$ at 25 °C the value of K_W is 1×10^{-14} . By taking the log of above equation we get

$$\log K_a + \log K_b = \log K_w$$

Multiply above equation by -1, we get

$$-\log K_a + (-1) \log K_b = -\log K_w$$

$$pK_a + pK_b = pK_w$$

$$-\log K_a + (-1) \log K_b = -\log K_w$$

$$pK_a + pK_b = pK_w$$

$$pK_a + pK_b = 14 \qquad \{pK_w = -\log (1 \times 10^{-14}) = 14\}$$

What is the justification for the increase of ionic product of water with temperature?

Ans:
$$H_2O \longrightarrow H^+ + OH^-$$

Water is a very weak electrolyte. The ionization of H₂O is an endothermic process which is increased with the rise in temperature.

Is it true that value of K_w increases 75 times when temperature is increased from 0 °C to 100 °C?

lonic product of water is increase with rise in temperature because at high temperature ionization becomes easy.

For example:

$$K_W at 0$$
 °C is 0.10×10^{-14} . At 100 °C 7.5×10^{-14}

Thus the value from 0 °C-100 °C is:

$$\frac{7.5 \times 10^{-14}}{0.10 \times 10^{-14}} = 75 \text{ times}$$
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What is ionic product of water?

How does this value change by change in temperature?

Water undergoes self-ionization as follows and reaction is reversible:

$$H_2O + H_2O \longrightarrow H_3O^+ + OH^-$$

$$H_2O \longrightarrow H^+ + OH^-$$

$$K_C = \frac{[H^+][OH^-]}{[H_2O]}$$
 $K_C = 1.8 \times 10^{-16} \text{ moles dm}^{-3}$

Since the water is present in very large excess and very few of its molecules undergo ionization, so its concentration remains effectively constant. Constant concentration of water is taken on L.H.S. and multiplied with K_C to get another constant K_w .

$$1.8 \times 10^{-16} \times 55.5 = 1.01 \times 10^{-14} = [H^+][OH^-]$$

This 1.01×10^{-14} is called K_w of water at 25 °C

$$K_C[H_2O] = [H^+][OH^-]$$

 $K_w = [H^+][OH^-] = 10^{-14} \text{ at } 25 \text{ °C}$

Kw is called ionic product of water.

The value of K_w increases almost 75 times when temperature is increased from 0 °C to 100 °C. Anyhow, the increase in K_w is not regular.

Q38: Is it true that the sum of pK_a and pK_b is always equal to 14 at all temperature for any acid?

Ans: It is not true because pK_a and pK_b values are temperature dependent. The degree of ionization of any acid increase as the temperature increases. Hence the value of pK_a and pK_b change with change in temperature.

Q39: What will be nature of solution when: (a) pH = 3.0 (b) pH = 8.0?

Ans: The value of pH varies between 0-14. A solution having pH value 0-7 are acidic in nature while a solution having pH value 7-14 are basic in nature.

- \triangleright pH = 3 *This solution is acidic.*
- \triangleright pH = 8 This solution is basic.

Q40: Define pH and pOH. Give its equation.

Ans: pH: The negative logarithm of H⁺ ions concentration is called pH.

$$pH = -log(H^+)$$

pOH: The negative logarithm of OH⁻ ions concentration is called pOH.

$$pOH = -log(OH^{-})$$

Just for information and important for MCQs

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Table Relationship of [H₃0⁺], [OH⁻], pH and pOH

| ٨ | | [H ₃ O ⁺] | рН | [OH ⁻] | рОН |
|-------------|---------|----------------------------------|------|---------------------|------|
| /\ | | 1x10 ⁻¹⁴ | 14.0 | 1x10 | 0.0 |
| / \ | | 1x10 ⁻¹³ | 13.0 | 1x10 ⁻¹ | 1.0 |
| 4 2 | Basic | 1x10 ⁻¹² | 12.0 | 1x10 ⁻² | 2.0 |
| basic | | 1x10 ⁻¹¹ | 11.0 | 1x10 ⁻³ | 3.0 |
| More basic | | 1x10 ⁻¹⁰ | 10.0 | 1x10 ⁻⁴ | 4.0 |
| Ž | | 1x10 ⁻⁹ | 9.0 | 1x10 ⁻⁵ | 5.0 |
| | | 1x10 ⁻⁸ | 8.0 | 1x10 ⁻⁶ | 6.0 |
| | Neutral | 1x10 ⁻⁷ | 7.0 | 1x10 ⁻⁷ | 7.0 |
| ig G | | 1x10 ⁻⁶ | 6.0 | 1x10 ⁻⁸ | 8.0 |
| a aci | | 1x10 ⁻⁵ | 5.0 | 1x10 ⁻⁹ | 9.0 |
| More acidic | | 1x10 ⁻⁴ | 4.0 | 1x10 ⁻¹⁰ | 10.0 |
| 7 | Acidic | 1x10 ⁻³ | 3.0 | 1x10 ⁻¹¹ | 11.0 |
| \ / | | 1x10 ⁻² | 2.0 | 1x10 ⁻¹² | 12.0 |
| \ / | | 1x10 ⁻¹ | 1.0 | 1x10 ⁻¹³ | 13.0 |
| V | | 1x10 ⁻⁰ | 0.0 | 1x10 ⁻¹⁴ | 14.0 |

Class 11th: Chemistry notes

Approximate pH and pOH of some common materials at 25°C

| | - | | | | <u> </u> |
|---------------|-----|-------|--|---------|----------|
| Material | pН | рОН | Material | pН | pOH |
| 1.0 M HCI | 0.1 | 13.9 | bread | 5.5 | 8.5 |
| 0.1 M HCl | 1.1 | 12.9 | potatoes | 5.8 | 8.2 |
| 0.1 M CH₃COOH | 2.9 | 11.10 | rainwater | 6.2 | 7.8 |
| gastric juice | 2.0 | 12.00 | milk | 6.5 | 7.5 |
| lemons | 2.3 | 11.7 | saliva | 6.5-6.9 | 7.5-7.1 |
| vinegar | 2.8 | 11.2 | pure water | 7.0 | 7.00 |
| soft drinks | 3.0 | 11.00 | eggs | 7.8 | 6.2 |
| apples | 3.1 | 10.9 | 0.1 M NaHCO₃ | 8.4 | 5.6 |
| grapefruit | 3.1 | 10.9 | seawater | 8.5 | 5.5 |
| oranges | 3.5 | 10.5 | milkofmagnesia | 10.5 | 3.5 |
| tomatoes | 4.2 | 9.8 | 0.1 M NH ₃ | 11.1 | 2.9 |
| cherries | 3.6 | 10.4 | 0.05 M Na ₂ CO ₃ | 11.6 | 2.4 |
| bananas | 4.6 | 9.4 | 0.1 M NaOH | 13.0 | 1.00 |

What will be nature of solution when (a) pH is more than 7 (b) pH is smaller than 7?

pH scale generally ranges from 0 to 14. When pH is less than 7 the solution has acidic nature. When pH is greater than 7 then solution is basic.

Define pH and pOH. How are they related with pKw.

$$pK_w = pH + pOH$$

Relation of pH *and* pOH *with* pK_w: Ans:

We know that

$$K_w = (H^+)(OH^-)$$

Taking -ve log on both sides

$$-\log K_w = [-\log(H^+)] + [-\log(OH^-)]$$

Negative log of K_w is called pK_w . So, $pK_w = pH + pOH$

$$pK_w = pH + pOH$$

Why aqueous solution of CH3COONa is basic in nature?

When salt CH₃COONa is dissolved in water, it produces NaOH and CH₃COON. NaOH is a strong base while CH₃COON is a weak acid. So, the solution becomes basic in nature.

Please visit for more data at: www.pakcity.org

Write down Henderson equation for acidic and basic buffers.

Henderson's equation: Ans:

Consider a reaction:

$$NA \longrightarrow H^+ + A^ NaA \longrightarrow Na^+ + A^-$$

The ionization constt. for acid.

$$K_a = \frac{[H^+][A^-]}{[HA]}$$
 $[H^+] = \frac{K_a[HA]}{[A^-]}$

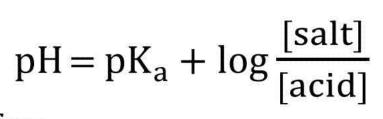
Taking -ve log on both sides

$$-\log[H^{+}] = -\log \frac{K_{a}[HA]}{[A^{-}]}$$

$$-\log[H^{+}] = -\log K_{a} - \log \frac{[HA]}{[A^{-}]}$$

$$pH = pK_{a} - \log \frac{[HA]}{[A^{-}]}$$

$$pH = pK_{a} - \log \frac{[acid]}{[salt]}$$



For basic buffers,

$$pOH = pK_b + log \frac{[salt]}{[base]}$$

Q45: What is the ionization constant of acids?

Ans: Ka Ionization constant of acid:

It is ionization constant of acid. Its value is quantitative measure of the strength of acid.

$$HA + H_2O \longrightarrow H_3O^+ + A^-$$

Equilibrium expression:

$$K_C = \frac{[H_3O^+][A^-]}{[HA][H_2O]}$$

Water being large in excess its concentration remain constant.

$$K_{C}[H_{2}O] = \frac{[H_{3}O^{+}][A^{-}]}{[HA]}$$

 K_C is equilibrium constant and H_2O concentration almost remains constant. So, H_2O is also constant. Thus, these constants are replaced by K_a .

$$K_a = \frac{[H_3O^+][A^-]}{[HA]}$$

Greater the value of Ka stronger will be the acid.

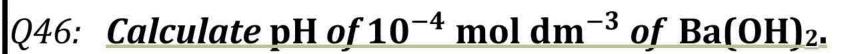
These tables are just for information and imp for MCQs

Table Dissociation constants of some acids at 25°C and their relative strength

| ۸ ما ما | Disconiation | T V | Dolotius atuonath |
|--------------------------------|--|-------------------------------|-------------------|
| Acid | Dissociation | K _a | Relative strength |
| HCI | HCl H++Cl | very large(10 ⁺⁷) | Very strong |
| HNO ₃ | HNO_3 $H^+ + NO_3^-$ | very large(10 ⁺³) | Very strong |
| H ₂ SO ₄ | $H_2SO_4 \square H^+ + HSO_4^-$ | Large(10 ⁺²) | Very strong |
| HSO-4 | $HSO_4^- \square H^+ + SO_4^{-2}$ | 1.3x10 ⁻⁴ | Strong |
| HF | $HF \square H^+ + F^-$ | 6.7x10 ⁻⁵ | Weak |
| CH₃COOH | CH ₃ COOH H ⁺ + CH ₃ COOH | 1.85x10 ⁻⁵ | Weak |
| H ₂ CO ₃ | $H_2CO_3 \square H^+ + HCO_3^-$ | 4.4x10 ⁻⁷ | Weak |
| H ₂ S | $H_2S \square H^+ + HS^-$ | 1.0x10 ⁻⁷ | Weak |
| NH ₄ ⁺ | $NH_4 \square H^+ + NH_3$ | 5.7x10 ⁻¹⁰ | Weak |
| HCO ₃ | $HCO_3^- \square H^+ + CO_3^{2-}$ | 4.7x10 ⁻¹¹ | Weak |
| H ₂ O | $H_2O \square H^+ + OH^-$ | 1.8x10 ⁻¹⁶ | Very weak |

Table K_b of some important bases

| Base | Dissociation | K _b | Relative strength |
|---|---|-----------------------|----------------------|
| NaOH | NaOH Na ⁺ + OH ⁻ | Very high | Very strong |
| КОН | KOH □ K ⁺ + OH ⁻ | Very high | Very strong |
| Ca(OH) ₂ | $Ca(OH)_2 \square Ca^{2+} + 2OH$ | High | Very strong |
| NH ₄ OH | NH₄OH□ NH₄⁺ + OH⁻ | 1.81x10 ⁻⁵ | Weak |
| CH ₃ NH ₂ (Mathyl amine) | $CH_3NH_2 + H_2O \square CH_3NH_3^+ + OH^-$ | 4.38x10 ⁻⁴ | Weak |
| C ₆ H ₅ NH ₂ (Aniline) | $C_6H_5NH_2 + H_2O \square C_6H_5NH_3^+ + OH^-$ | 4.7x10 ⁻¹⁰ | Very weak |



Ans:
$$Ba(OH)_2 = Ba^{2+} + 2OH^{-}$$

$$10^{-4}$$
 2×10^{-4}

$$pOH = -log[OH^-]$$

$$pOH = -[log(2 \times 10^{-4})]$$

$$pOH = 3.69$$

$$pH + pOH = 14$$

$$pH + 3.69 = 14$$

$$pH = 14 - 3.69$$

$$pH = 10.31$$

Q47: Calculate pH of 10^{-4} mol dm⁻³ of HCl solution.



Ans:

$$HCl \longrightarrow H^+ + Cl^-$$

$$10^{-4}$$

$$2 \times 10^{-4}$$

$$pH = -\log[H^+]$$

$$[H^+] = 10^{-4} \text{ mol dm}^{-3}$$

$$pH = -log[10^{-4}]$$

$$pH = -(-4)\log 10$$

$$pH = +4(1)$$

$$pH = +4$$

Q48: Define acid and base by Lowry-Bronsted concept.

Ans: Acid:

An acid is a substance which has ability to donate proton.

Base:

The base is a substance which has ability to accept proton.

$$NH_3 + H_2O \longrightarrow NH_4^+ + OH^-$$

Base Acid

Q49: Why aqueous solution of CuSO₄ is acidic in nature?

Ans: When CuSO₄ is dissolved in water it produce H₂SO₄ which is an acid, so aqueous Solution of CuSO₄ is acidic in nature as shown by following reaction:

$$CuSO_4 + H_2O \longrightarrow Cu(OH)_2 + H_2SO_4$$

Q50: Why Aqueous Solution of NH₄Cl is acidic in Nature?

Ans: When NH4CI is dissolved in water it produces HCI. As HCI is strong acid and NH4OH is weak base. So aqueous Solution of NH4CI is acidic in nature as shown by following reaction:

$$NH_4CI + H_2O \longrightarrow NH_4OH + HCI$$

Q51: Aqueous solution of Na_2CO_3 is alkaline in nature, Why?

Ans: When Na₂CO₃ is dissolved in water it produces weak acid and a strong base. So, aqueous solution of Na₂CO₃ is alkaline in nature as given by following chemical reaction:

$$Na_2CO_3 + H_2O \longrightarrow NaOH + H_2CO_3$$

Q52: Explain the terms buffer and buffer capacity. OR

Define buffer solutions.

Ans: Buffer:



The solutions which resist the change in pH by the addition of small amount of an acid or base is called buffer.

Buffer capacity:

The capacity of a buffer to maintain definite pH is called buffer capacity.

Q53: What happens to the acidic and basic properties of aqueous solutions when pH varies from zero to 14?

Ans: Properties of solutions when pH changes from 0 to 14.

pH = 7 Solution is neutral pH = 0 to 6.9 Solution is acidic pH = 7.1 to 14 Solution is basic

Q54: What will be the nature of solution having pH equal to 12?

Ans: The value of pH varies between 0-14. A solution having pH value 0-7 are acidic in nature while a solution having pH value 7-14 are basic in nature.

pH = 12 This Solution is basic

Q55: How does a buffer act? Explain with an example.

Ans: Let us take the example of an acidic buffer consisting of CH₃COOH and CH₃COONa. Common ion effect helps us to understand how the buffer will work. CH₃COOH being a weak electrolyte undergoes very little dissociation.

 $CH_3COOH + H_2O \longrightarrow H_3COO^- + H_3O$ $CH_3COONa \longrightarrow CH_3COO^- + Na^+$

Actually a buffer mentioned above is a large reservoir of CH_3COOH and CH_3COO^- components. When an acid or H_3O^+ ions are added to this buffer, they will react with CH_3COO to give back acetic acid and hence the pH of the solution will almost remain unchanged.

Q56: Write two uses of buffer solution.

Ans: The uses of buffer solution are:

- Buffers are used in many industrial processes like electroplating, dyes etc.
- The pH of human blood is maintained at 7.35 with the help of buffers. A higher or lower value than this may prove fatal.

Q57: Give preparation of acidic buffer solutions. OR [Ustify mixture of sodium acetate and acetic acid gives us the acidic buffer.]

Ans: Acidic buffer solutions are prepared by mixing a weak acid and its salt with strong base.

<u>For example:</u>

CH₃COOH \longrightarrow CH₃COO⁻ + H⁺
CH₃COONa⁺ \longrightarrow CH₃COO⁻ + Na⁺
CH₃COOH / CH₃COONa

Q58: A mixture of NH4OH and NH4Cl gives a basic buffer. Justify the statement. OF What is Basic Buffer, give example?

Ans: A buffer which consists of a weak base and its salt with strong acid is called basic buffer. Since NH₄OH is a weak base and NH₄CI is its salt with strong acid. Therefore it is basic buffer.

Q59: Why we need buffer solution? OR



Buffers are important in many area of chemistry.

Ans: Buffers are the substances which resist the change in pH. Buffers are very important in many areas of chemistry. Buffers are needed in chemical analysis, pharmaceuticals, electroplating etc.

Q60: What are buffer solutions? OR

How a basic buffer can be prepared?

Ans: Buffer solution:

The solution that resists in pH changes when small amount of an acid or a base is added to it is called buffer solution.

Basic buffers are formed by mixing a weak base and its salt with strong acid. Such solution will give basic buffers with pH more than 7. Mixing of NH₄OH and NH₄CI is one of the best examples of such a basic buffer.

Q61: Differentiate between acidic and basic buffers.

Ans: The difference between acidic and basic buffers is:

| | Acidic buffers | | Basic buffers pakcity.org |
|---|---|-----|---|
| > | The buffer which is prepared by mixing | > | The buffer which is prepared by mixing |
| | a weak acid and a salt of it with a | | a weak base and a salt of it with a |
| | strong base is known as acidic buffer. | | strong acid is known as basic buffer. |
| > | Such solutions give acidic buffers with | > | Such solutions give basic buffers with |
| | pH less than 7, Mixture of acetic acid | | pH more than 7. Mixture of NH ₄ OH and |
| | and sodium acetate is one of the best | ^- | NH4CI is one of the best examples of |
| | examples of such a buffer. | 2/4 | basic buffer. |

Q62: What is the formula to calculate the percentage ionization of weak acid?

Ans: We can calculate the percentage ionization of weak acids and the formula is as follows:

Q63: What are applications of buffer in daily life?

Ans: Buffers are important in many areas of chemistry and allied sciences like molecular biology microbiology, cell biology, soil sciences, nutrition and the clinical analysis.

Q64: What is common ion effect? Give an example.

Ans: Common ion effect:

The decrease in degree of ionization of a weak electrolyte by the addition of another electrolyte having a common ion is called common ion effect.

For example:

NaCl
$$\longrightarrow$$
 Na⁺ + Cl⁻
HCl \longrightarrow H⁺ + Cl⁻

 Cl^- is a common ion.

Q65: Why solid NH₄Cl is added in qualitative analysis of 3rd group basic radical before adding NH₄OH?

Ans:
$$NH_3OH \longrightarrow NH_4^+ + OH^-$$

To provide low concentration of OH⁻ ion NH₄Cl is added which is more soluble.

$$NH_4Cl \longrightarrow NH_4^+ + Cl^-$$

NH₄Cl decrease the OH⁻ ion conc. by suppressing ionization of NH₃OH due to common ion. That's why a mixture of NH₄Cl + NH₃OH is used as reagent for third group analysis.



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- Q66: Define conjugate acid with an example.
- Conjugate acid is a substance which is formed by accepting a proton (H^+) by a base.

$$HCl + H_2O \longrightarrow H_3O^+ + Cl^-$$

conj.base conj.acid Acid Base

How NaCl can be purified by common ion effect?

Give applications of common ion effect.

The impurities like CaCl₂, MgCl₂, Na₂SO₄, can be removed by the use of common ion effect. Ans: HCI gas is passed through saturated solution of NaCl.

$$NaCl \longrightarrow Na_{(aq)}^+ + Cl_{(aq)}^-$$

$$HCl \longrightarrow H_{(aq)}^+ + Cl_{(aq)}^-$$

Concentration of Cl⁻ ions increase by passing HCI. According to Le-Chateliers principle the reaction will move in reverse direction to keep the value of K_C const. Hence NaCl will precipitated out.

- What is the effect of common ion on solubility?
- The solubility of a partially soluble electrolyte is decreased by the addition of a more soluble electrolyte having common ion.

For example:

*Precipitation of KCIO*₃

$$KCIO_3 \longrightarrow K^+ + ClO_3^-$$

$$KCl \longrightarrow K^+ + Cl^-$$

$$KCl \longrightarrow K^+ + Cl^-$$

- Q69: Solubility of glucose increase in water by heating. Give reason.
- Ans: Formation of solution of glucose in H_2O is an endothermic process. The solution becomes cold. Therefore, according to Le-Chateliers principle an increase in temperature will favour the formation of glucose solution.

Thus by increasing temperature more quantity of glucose will dissolve in water.

Q70: Define solubility and solubility product.



The concentration of the solute in the solution when it is in equilibrium with the solid substance at a particular temperature.

Solubility product:

The solubility product is the product of the concentration of ion raised to exponent equal to coefficient of the balanced equation.

It is denoted by K_{sp}.

Example:

$$PbSO_4 \longrightarrow Pb^{2+} + SO_4^{2-}$$

$$K_C = \frac{[Pb^{2+}][SO_4^{2-}]}{[PbSO_4]}$$

Since PbSO₄ is sparingly soluble therefore its concentration will remain constant.

$$K_C[PbSO_4] = [Pb^{2+}][SO_4^{2-}]$$

$$K_{sp} = [Pb^{2+}][SO_4^{2-}]$$

Define solubility product and write down its applications.

Solubility product:



The solubility product is the product of the concentration of ion raised to exponent equal to coefficient of the balanced equation.

It is denoted by K_{sp}.

Applications of K_{sp}:

- > Determination of solubility from K_{sp}.
- \triangleright Determination of K_{sp} from solubility.

Q72: Define solubility product. Derive solubility product expression for Ag₂CrO₄ and PbCl₄.

Ans: Solubility product:

The solubility product is the product of the concentration of ion raised to exponent equal to coefficient of the balanced equation.

It is denoted by K_{sp}.

PbCl₄:

$$PbCl_{4} = Pb^{2+} + 4Cl^{-}$$

$$K_{C} = \frac{[Pb^{2+}][Cl^{-}]^{4}}{[PbCl_{4}]}$$

$$K_{C} [PbCl_{4}] = [Pb^{2+}][Cl^{-}]^{4}$$

$$K_{sp} = [Pb^{2+}][Cl^{-}]^{4}$$

$$K_{C} [PbCl_{4}] = K_{sp}$$

Ag₂CrO₄:

$$Ag_{2}CrO_{4} = 2Ag^{+} + CrO_{4}^{2}$$

$$K_{C} = \frac{[Ag^{+}]^{2}[CrO_{4}^{-2}]}{[Ag_{2}CrO_{4}]}$$

$$K_{C} [Ag_{2}CrO_{4}] = [Ag^{+}]^{2}[CrO_{4}^{-2}]$$

$$K_{Sp} = [Ag^{+}]^{2}[CrO_{4}^{-2}]$$

So,

This table is just for information and imp for MCQs

Table K_{sp} values for some ionic compounds (compounds are arranged alphabetically).

| Salt | Ion Product | K _{sp} | Salt | Ion Product | K _{sp} |
|---------------------------------|--|-----------------------|--------------------------------|--|-----------------------|
| AgBr | [Ag ⁺][Br ⁻] | 5.0x10 ⁻¹³ | CuS | [Cu ²⁺][S ²⁻] | 8x10 ⁻³⁴ |
| Ag ₂ CO ₃ | [Ag ⁺] ² [CO ₃ ²⁻] | 8.1x10 ⁻¹² | FeS | [Fe ²⁺][S ²⁻] | 6.3x10 ⁻¹⁸ |
| AgCl | [Ag ⁺][Cl ⁻] | 1.8x10 ⁻¹⁰ | Fe ₂ S ₃ | [Fe ³⁺][S ²⁻] ³ | 1.4x10 ⁻⁸⁵ |
| Agl | [Ag ⁺][I ⁻] | 8.3x10 ⁻¹⁷ | Fe(OH) ₃ | [Fe ³⁺][OH ⁻] ³ | 1.6x10 ⁻³⁹ |
| Ag ₂ S | $[Ag^{+}]^{2}[S^{2-}]$ | 8x10 ⁻⁴⁸ | HgS | [Hg ²⁺][S ²⁻] | 2x10 ⁻⁵⁰ |
| Al(OH) ₃ | [Al ³⁺][OH ⁻] ³ | 3x10 ⁻³⁴ | MgCO ₃ | [Mg ²⁺][CO ₃ ²⁻] | 3.5x10 ⁻⁸ |
| BaCO ₃ | [Ba ²⁺][CO ₃ ²⁻] | 2x10 ⁻⁹ | Mg(OH) ₂ | [Mg ²⁺][OH ⁻] ² | 6.3x10 ⁻¹⁰ |
| BaSO ₄ | [Ba ²⁺][SO ₄ ²⁻] | 1.1x10 ⁻¹⁰ | MnS | $[Mn^{2+}][S^{2-}]$ | 3x10 ⁻¹¹ |
| CdS | [Cd ²⁺][S ²⁻] | 8.0x10 ⁻²⁷ | PbCl ₂ | [Pb ²⁺][Cl ⁻] ² | 1.6x10 ⁻⁵ |
| CaCO ₃ | [Ca ²⁺][CO ₃ ²⁻] | 3.3x10 ⁻⁹ | PbCrO ₄ | [Pb ²⁺][CrO ₄ ²⁻] | 2.3x10 ⁻¹³ |
| CaF ₂ | [Ca ²⁺][F ⁻] ² | 3.2x10 ⁻¹¹ | PbSO ₄ | [Pb ²⁺][SO ₄ ²⁻] | 1.6x10 ⁻⁸ |
| Ca(OH) ₂ | [Ca ²⁺][OH ⁻] ² | 6.5x10 ⁻⁶ | PbS | [Pb ²⁺][S ²⁻] | 8.0x10 ⁻²⁸ |



Chapter: 08

Chemical Equilibrium





Q1: The following reaction was allowed to reach the state of equilibrium

$$2A_{(aq)} + B_{(aq)} - C_{(aq)}?$$
 OR

- Q2: Define law of mass action & derive equilibrium constant for a general chemical reaction.
- Q3: $N_{2(g)}$ and $H_{2(g)}$ Combine to give $NH_{3(g)}$. The value of K_C in this reaction at 500°C is 6.0 $\times 10^{-2}$ Calculate the value of K_P for this reaction. (V.Imp)
- Q4: When 1.00 Mole of steam and 1.00 mole of Carbon Monoxide are allowed to reach equilibrium 33.3% of the equilibrium mixture is Hydrogen. Calculate the value of K_p . State the units of K_p .
- Q5: What is the percentage ionization of acetic acid in the solution in which 0.1 moles of it has been dissolved per dm³ of the solution? (V.Imp)
- Q6: Write a note on synthesis of ammonia gas by Haber's process keeping in mind the applications of chemical equilibrium in industry.
- Q7: The equilibrium constant for the reaction between acetic acid and ethyl alcohol is 4.0. A mixture of 3 moles of acetic acid and one mole C₂H₅OH is allowed to come to equilibrium. Calculate the amount of ethyl acetate at equilibrium state in no of moles and grams. Also calculate mass of reactants left behind.
- Q8: Define common ion effect. Give its two applications,
- Q9: Calculate the pH of a buffer solution in which $0.11 \text{ molar CH}_3\text{COONa}$ and 0.09 molar acetic acid solutions are present. K_a for CH_3COONa is 1.85×10^{-5} . (V.Imp)
- Q10: The solubility of PbF₂ at 25°C is 0.64g dm⁻³ Calculate K_{sp} Of PbF₂. (Molecular mass of PbF₂ = 245.2g mol⁻¹) (V.Imp)
- Q11: $Ca(0H)_2$ is a sparingly soluble compound. Its solubility product is 6.5×10^{-6} . Calculate the solubility of $Ca(0H)_2$?
- Q12: The Solubility of CaF₂ in water at 25°C is found to be 2.05×10^{-4} mol dm⁻³. What is the value of K_{sp} at this temperature? (V.Imp)
- Q13: A buffer solution is pre pared by mixing 0.2 M CH₃COONa and 0.5 M CH₃COOH in 1dm³ of solution. Calculate pH of solution. pk_a of acid is 4.74.
- Q14: Define Le. Chaterlieri's principle. Discuss effect of
 - > Change in volume
 - Change in temperature on the formation of ammonia.