Equilibrium

1.	For a weak acid H. dissociation is nearly the concentration of a the correct option for temperature is: (a) 1×10^{-4} (b) 1×10^{-6} (c) 1×10^{-5} (d) 1×10^{-3}	br a weak acid HA, the percentage of association is nearly 1% at equilibrium. If the concentration of acid is 0.1 mol L ⁻¹ , then the correct option for its K _a at the same emperature is: (2023) 1×10^{-4} 1×10^{-6} 1×10^{-5} 1×10^{-3} the acidic buffer is prepared by mixing: (2023) $1 \times 10^{$		(a) 3.6 (b) 0.36 (c) 3.6 × 10 ⁻² (d) 3.6 × 10 ⁻³ The pH of the solution containing 50 mL each of 0.10 M sodium acetate and 0.01 M acetic acid is (2022) [Given: pK _a of CH ₃ COOH = 4.57] (a) 5.57 (b) 3.57 (c) 4.57 (d) 2.57 $3O_2(g) \neq 2O_3(g)$ For the above reaction at 298 K, K _c is found to be 3.0 × 10 ⁻⁵⁹ . If the concentration of O ₂ at equilibrium is 0.040 M then concentration of O ₃ in M is (2022) (a) 4.38 × 10 ⁻³² (b) 1.9 × 10 ⁻⁶³ (c) 2.4 × 10 ³¹ (d) 1.2 × 10 ²¹ The pK _b of dimethylamine and pK _a of acetic acid are 3.27 and 4.77 respectively at T(K). The correct option for the pH of dimethylammonium acetate solution is:
2.	 An acidic buffer is pre (a) weak acid and it's (b) equal volumes of e weak acid and weak (c) strong acid and it's (d) strong acid and it 			
3.	pK _a of acid = pK _b of the base) 0.01 M acetic acid solution is 1% ionised, then pH of this acetic acid solution is: (2022) (a) 1 (b) 3 (c) 2 (d) 4		8.	
4.	K_H value for some temperature 'T' are giGasAr CO_2 HCHOCH4Where K_H is Henry's I The order of their solu (a) HCHO < CH4 < CO	gases at the same ven: (2022) K_H/k bar 40.3 1.67 1.83 × 10 ⁻⁵ 0.413 wave constant in water. ubility in water is: $D_2 < Ar$	9.	(2021) (a) 5.50 (b) 7.75 (c) 6.25 (d) 8.50 Find out the solubility of Ni(OH) ₂ in 0.1 M NaOH. Given that the ionic product of Ni(OH) ₂ is 2×10^{-15} . (2020) (a) 2×10^{-8} M (b) 1×10^{-13} M (c) 1×10^{8} M (d) 2×10^{-13} M
5.	(a) Herio $< CH_4 < CO_2 < H_1$ (b) Ar $< CO_2 < CH_4 < HCHO$ (c) Ar $< CH_4 < CO_2 < HCHO$ (d) HCHO $< CO_2 < CH_4 < Ar$ 5. K _p for the following reaction is 3.0 at 1000 K. $CO_2(g) + C(s) \rightleftharpoons 2CO(g)$ What will be the value of K _c for the reaction at the same temperature? (2022) (Given: R = 0.083 L bar K ⁻¹ mol ⁻¹)		10.	 (d) 2 × 10⁻¹³ M HCl was passed through a solution of CaCl₂, MgCl₂ and NaCl. Which of the following compound(s) crystallies(s)? (2020) (a) Only NaCl (b) Only MgCl₂ (c) NaCl, MgCl₂ and CaCl₂ (d) Both MgCl₂ and CaCl₂

4.

11. Which among the following salt solutions is basic in nature? (2020 covid Re-NEET) (a) Ammonium sulphate (b) Ammonium nitrate (c) Sodium acetate (d) Ammonium chloride 12. The solubility product for a salt of the type AB is 4×10^{-8} . What is the molarity of its standard solution? (2020 Covid Re-NEET) (a) $16 \times 10^{-16} \text{ mol/L}$ (b) $2 \times 10^{-16} \text{ mol/L}$ (c) $4 \times 10^{-4} \text{ mol/L}$ (d) $2 \times 10^{-4} \text{ mol/L}$ 13. Hydrolysis of sucrose is given by the following reaction. Sucrose + $H_20 \rightleftharpoons$ Glucose + Fructose If the equilibrium constant (K_c) is 2×10^{13} at 300 K, the value of $\Delta_r G^\circ$ at the same temperature will be : (a) $8.314 J mol^{-1}K^{-1} \times 300 K \times \ln(2 \times 10^{13})$ (b) 8.314 $J mol^{-1}K^{-1} \times 300 K \times \ln(3 \times 10^{13})$ (c) $-8.314 J mol^{-1} K^{-1} \times 300 K \times \ln (4 \times 10^{13})$ (d) $-8.314 \ I \ mol^{-1} \ K^{-1} \times 300 \ K \times lb(2 \times 10^{13})$ 14. pH of a saturated solution of $Ca(OH_2)$ is 9. The solubility product (K_{sp}) of Ca(OH)₂ is: (2019)(a) 0.5×10^{-15} (b) 0.25×10^{-10} (c) 0.125×10^{-15} (d) 0.5×10^{-10} 15. Conjugate base for Bronsted acids H₂O and HF are : (2019)(a) OH^- and H_2F^+ , respectively (b) H_3O^+ and F^- , respectively (c) OH- and F-, respectively (d) H_3O^+ and H_2F^+ , respectively 16. Which will make basic buffer? (2019)(a) 50 mL of 0.1 M NaOH + 25 mL of 0.1 M CH₃COOH (b) 100 mL of 0.1 M CH₃COOH + 100 mL of 0.1 M NaOH (c) 100 mL of 0.1 M HCl + 200 mL of 0.1 M NH₄OH (d) 100 mL of 0.1 M HCl + 100 mL of 0.1 M NaOH 17. Which one of the following conditions will favour maximum formation of the product in the reaction, $A_2(g) + B_2(g) \rightleftharpoons X_2(g) \ \Delta_r H = -X \ kJ$ (2018)(a) Low temperature and high pressure

(b) Low temperature and low pressure (c) High temperature and low pressure (d) High temperature and high pressure 18. The solubility of BaSO₄ in water is $2.42 \times$ 10⁻³ gL⁻¹ at 298 K. The value of its solubility product (K_{sp}) will be: (2018)(Given molar mass of $BaSO_4 = 233 \text{ g mol}^{-1}$) (a) 1.08 ×10⁻¹⁰ mol² L⁻² (b) 1.08 ×10⁻¹² mol² L⁻² (c) $1.08 \times 10^{-8} \text{ mol}^2 \text{ L}^{-2}$ (d) $1.08 \times 10^{-14} \text{ mol}^2 \text{ L}^{-2}$ 19. Following solutions were prepared by mixing different volumes of NaOH and HCl of different concentrations : (2018)(A) $60mL\frac{M}{10}HCl + 40mL\frac{M}{10}NaOH$ (B) $55mL\frac{M}{10}HCl + 45mL\frac{M}{10}NaOH$ (c) $75mL\frac{M}{5}HCl + 25mL\frac{M}{5}NaOH$ (d) $100mL\frac{M}{10}HCl + 100mL\frac{M}{10}NaOH$ pH of which one of them will be equal to 1? (a) B (b) A (c) C (d) D 20. The equilibrium constants of the following (2017-Delhi) are : $N_2 + 3H_2 \rightleftharpoons 2NH_3$ K_1 $N_2 + O_2 \rightleftharpoons 2NO$ K_2 $H_2 + \frac{1}{2}O_2 \rightarrow H_2O$ K3

- The equilibrium constant (K) of the reaction: $2NH_3 + \frac{5}{2}O_2 \stackrel{K}{\rightleftharpoons} 2NO + 3H_2O$, will be:
- (a) $K_2^3 K_3 / K_1$
- (b) $K_1 K_3^3 / K_2$
- (c) $K_2 K_3^3 / K_1$
- (d) $K_2 K_3 / K_1$
- 21. Concentration of Ag⁺ ions in a saturated solution of $Ag_2C_2O_4$ is 2.2×10^{-4} mol L⁻¹. Solubility product of Ag_2CO_4 is:

(2017-Delhi)

(a) 5.3×10^{-12} (b) 2.42×10^{-8}

- (c) 2.66×10^{-12}
- (d) 4.5×10^{-11}
- 22. Which one of the following statements is not correct? (2017-Delhi)
 - (a) Coenzymes increase the catalytic activity of enzyme
 - (b) Catalyst does not initiate any reaction

- (c) The value of equilibrium constant is changed in the presence of a catalyst in the reaction at equilibrium
- (d) Enzymes catalyse mainly bio-chemical reactions
- 23. The standard equilibrium constant K_P at 298 K for the reaction, N₂(g) + 3H₂(g) ≓ 2NH₃(g) is 5.8 × 10⁵. The value of standard equilibrium constant, if the concentration of gases is expressed in terms of mol L⁻¹, will be: (2017-Gujarat) [Given : R = 0.08314 L bar K⁻¹ mol⁻¹]
 (a) 3.99 × 10⁹
 (b) 3.51 × 10⁶
 (c) 3.84 × 10⁷
 - (d) 3.56×10^8
- 24. Consider the following reaction for which the change in enthalpy is positive $2A(g) + B(g) \rightleftharpoons C(g) + D(g)$ Which of the following will not affect the equilibrium? (2017-Gujarat)
 - (a) Presence of catalyst
 - (b) Change in concentration of reactants
 - (c) Change in pressure
 - (d) Change in temperature
- 25. For the reaction $CO(g) + Cl(g) \rightleftharpoons COCl_2(g)\frac{K_p}{K_c}$
 - is equal to:
 - (a) $(RT)^2$
 - (b) $\frac{1}{RT}$
 - (c) *RT*
 - (d) \sqrt{RT}
- 26. Which of the following fluoro-compounds is most likely to behave as a Lewis base?
 - (2016-II)

(2017-Gujarat)

- (a) CF₄
- (b) SiF_4
- (c) BF_3
- (d) PF₃
- 27. The solubility of AgCl(s) with solubility product 1.6×10^{-10} in 0.1 M NaCl solution would be: (2016-II) (a) 1.6×10^{-11} M
 - (b) Zero
 - (c) 1.26×10^{-5} M
 - (d) 1.6×10^{-9} M
- 28. The percentage of pyridine (C_5H_5N) that forms pyrimidine ion $(C_5H_5N^+H)$ in a 0.10 M aqueous pyridine solution $(K_b \text{ for } C_5H_5N =$ $1.7 \times 10^{-9})$ is : (2016-II) (a) 0.77%

- (b) 1.6%
- (c) 0.0060%
- (d) 0.013%
- 29. Consider the nitration of benzene using mixed conc. H₂SO₄ and HNO₃. If a larger amount of KHSO₄ is added to the mixture, the rate of nitration will be: (2016-I)
 (a) Doubled
 - (b) Increase
 - (c) Decrease
 - (d) Unchanged
- 30. MY and NY₃, two nearly insoluble salts, have the same K_{sp} values of 6.2×10^{-13} at room temperature, which statements would be true in regard to MY and NY₃?

(2016-I)

- (a) The addition of the salt of KY to solution of MY and NY_3 will have no effect on their solubilities
- (b) The molar solubilities of MY and NY_3 in water are identical
- (c) The molar solubility of MY in water is less than of NY_3
- (d) The salts MY and NY_3 are more soluble in 0.5 M KY than in pure water

31. If the equilibrium constant for $N_2(g) + O_2(g) \rightleftharpoons 2NO(g)$ is K, the equilibrium constant for $\frac{1}{2}N_2(g) + \frac{1}{2}O_2(g) \rightleftharpoons NO(g)$ will be: (2015 Re) (a) K^2

- (b) $K^{1/2}$
- (c) $\frac{1}{2}K$
- (d) *K*
- 32. Which one of the following pairs of solution is not an acidic buffer? (2015 Re)
 - (a) H_3PO_4 and Na_3PO_4
 - (b) HClO₄ and NaClO₄
 - (c) CH₃COOH and CH₃COONa
 - (d) H_2CO_3 and Na_2CO_3
- 33. If the value of an equilibrium constant for a particular reaction is 1.6×10^{12} , then at equilibrium the system will contain: (2015)
 - (a) Mostly reactants
 - (b) Mostly products
 - (c) Similar amounts of reactants and products
 - (d) All reactants
- 34. The K_{sp} of Ag_2CrO_4 , AgBr and AgI are respectively, 1.1×10^{-12} , 1.8×10^{-10} , 5.0×10^{-13} , 8.3×10^{-17} . Which one of the following salts will precipitate last if AgNO₃ solution

is added to the solution containing equal 40. For the reversible reaction, moles of NaCl, NaBr, NaI and Na₂CrO₄? $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g) + Heat.$ (2015)The equilibrium shifts in forward direction: (a) AgCl (2014)(b) AgBr (a) By decreasing the pressure (b) By decreasing the concentration of $N_2(g)$ (c) Ag₂CrO₄ (d) AgI and $H_2(g)$ (c) By Increasing pressure and decreasing 35. What is the pH of the resulting solution temperature when equal volumes of 0.1 M NaOH and (d) By increasing the concentration of 0.01 M HCl are mixed? (2015 Re) (a) 2.0 $NH_3(g)$ (b) 7.0 41. Which of these is least likely to act as a (c) 1.04 Lewis base? (2013)(d) 12.65 (a) CO 36. Aqueous solution of which of the following (b) Fcompounds is the best conductor of electric (c) BF_3 (2015 Re) current? (d) PF_3 (a) Hydrochloric acid, HCl 42. $KMnO_4$ can be prepared from K_2MnO_4 as (b) Ammonia, NH₃ per the reaction: (c) Fructose, $C_6H_{12}O_6$ $3MnO_4^{2-} + 2H_2O \rightleftharpoons 2MnO_4^- + MnO_2 + 4OH^-$ (d) Acetic acid, $C_2H_4O_2$ The reaction can go to completion by 37. Which of the following statements is correct removing OH- ions by adding: (2013)for a reversible process in a state of (a) HCl equilibrium? (b) KOH (2015)(a) $\Delta G^{\circ} = -2.303 \text{ RT} \log K$ (c) CO_2 (b) $\Delta G^{\circ} = 2.303 \text{ RT} \log K$ (d) SO_2 (c) $\Delta G = -2.303 \text{ RT} \log K$ (d) $\Delta G = 2.303 \text{ RT} \log K$ 38. Which of the following salts will give highest pH in water? (2014)(a) NaCl (b) Na_2CO_3 (c) CuSO₄ (d) 4KC1 39. Using the Gibb's energy change, ΔG° = +63.3 kJ, for the following reaction, $Ag_2CO_3(s) \rightleftharpoons 2Ag^+(aq) + CO_3^-(aq)$ The K_{sp} of $Ag_2CO_3(s)$ in water at 25°C is (R = 5.314 JK⁻¹ mol⁻¹) (2014) (a) 3.2×10^{-26} (b) 8.0×10^{-12} (c) 2.9×10^{-3} (d) 7.9×10^{-2}

S1. Ans. (c)		S30. Ans. (c)
S2. Ans. (a)		S31. Ans. (b)
S3. Ans. (d)		S32. Ans. (b)
S4. Ans. (b)		S33. Ans. (b)
S5. Ans. (c)		S34. Ans. (c)
S6. Ans. (a)		S35. Ans. (d)
S7. Ans. (a)		S36. Ans. (a)
S8. Ans. (b)		S37. Ans. (a)
S9. Ans. (d)		S38. Ans. (b)
S10. Ans. (a)		S39. Ans. (b)
S11. Ans. (c)		S40. Ans. (c)
S12. Ans. (d)		S41. Ans. (c)
S13. Ans. (d)		S42. Ans. (c)
S14. Ans. (a)		
S15. Ans. (c)		
S16. Ans. (c)		
S17. Ans. (a)		
S18. Ans. (a)		
S19. Ans. (c)		
S20. Ans. (c)		
S21. Ans. (a)		
S22. Ans. (c)		
S23. Ans. (d)		
S24. Ans. (a)		
S25. Ans. (b)		
S26. Ans. (d)		
S27. Ans. (c)		
S28. Ans. (d)		
S29. Ans. (c)		
	•	

Solutions

S6.

Ans.(a)

S1.	Ans.(c) $K_a = C\alpha^2$
	$K_a = (0.1) \times (0.01)^2$
	$K_a = 1 \times 10^{-5}$
S2.	Ans.(a) Acidic buffer is prepared by mixing weak acid and its salt with strong base.
S3.	Ans.(d)
	For weak acid (i.e., CH ₃ COOH)
	$[H^+] = C\alpha$
	= 0.01 × $\frac{1}{100}$ = 10 ⁻⁴ M
	$pH = -\log H^+ = -\log 10^{-4} = 4$
S4.	Ans.(b)
	According to Henry's Law,
	$p = K_H x$
	Where 'p' is partial pressure of gas in
	vapour phase.
	'K _H ' is Henry's Law constant.
	'x' is mole fraction of gas in liquid.
	Higher the value of K_H at a given pressure, lower is the solubility of the gas in the liquid.
	\therefore Solubility: Ar < CO ₂ < CH ₄ < HCHO
S5.	Ans.(c) CO ₂ (g) + C(s) \rightleftharpoons 2CO(g)
	$\Delta n_{g} = 2 - 1 = 1$
	$K_p = K_c (RT)^{\Delta n_g}$
	$K_p = K_c(RT)$
	$[:: K_p = 3]$
	$K_{c} = \frac{K_{p}}{RT} = \frac{3}{0.083 \times 1000}$
	= 0.036
	$= 3.6 \times 10^{-2}$

	It is a mixture of weak acid and salt of
	its conjugate base. Hence, it is acidic
	buffer.
	$pH = pK_a + \log \frac{[Salt]}{[Acid]}$
	$= 4.57 + \log\left(\frac{0.1}{0.01}\right)$
	= 4.57 + 1
	= 4.57
S7.	Ans.(a) $3O_2(g) \rightleftharpoons 2O_3(g)$
	$K_{c} = \frac{[O_{3}]^{2}}{[O_{2}]^{3}}$
	$[O_3]^2 = K_c[O_2]^3 = 3 \times 10^{-59} \times (0.04)^3$
	$[O_3]^2 = 1.9 \times 10^{-63} = 19 \times 10^{-64}$
	$[O_3] = 4.38 \times 10^{-32}$
	Concentration of O_3 at equilibrium = $4.38 \times 10^{-32} \text{ M}$
S8.	Ans.(b)
	$pH = \frac{1}{2} [P^{k_w} + P^{k_a} - P^{k_b}]$
	$= 7 + \frac{1}{2}P^{k_a} - \frac{1}{2}p^{k_b}$
	$= 7 + \frac{1}{2} \times 4.77 - \frac{1}{2} \times 3.27$
	= 7.75
S9.	Ans.(d)
	Let the solubility of $Ni(OH)_2$ is s
	$Ni(OH)_2 \rightleftharpoons Ni_s^{2+} + 2OH_{2s}^{-}$
	$NaOH \rightarrow Na + OH^{-}$
	0.1 0.1 0.1
	As K_{sp} is small 2s <<0.10
	Therefore $(0.10 + 2s) \approx 0.10$
	So total [OH]- = 0.10
	Ionic product = [Ni ²⁺] [OH-] ²
	$2 \times 10^{-15} = s(0.10)^2$
	$s = 2 \times 10^{-13} M$
S10.	Ans.(a)

When HCl is passed through the mixture of CaCl₂, MgCl₂ and NaCl, Clion concentration increases. As CaCl₂ and MgCl₂ are more stable than NaCl.

Hence ionic product [Na⁺] [Cl⁻] becomes more than solubility product. Thus, precipitation of NaCl occurs (due to common ion effect)

S11. Ans.(c)

Acetic acid, CH₃COOH, will react with sodium hydroxide, NaOH, to produce sodium acetate, CH₃COONa, and water.

Sodium acetate is a salt of strong base and Weak acid therefore its salt solution is basic in nature.

S12. Ans.(d)

For salt AB

$$K_{sp} = (s)2$$

$$s = \sqrt{K_{sp}}$$
$$s = \sqrt{4 \times 10^{-8}}$$
$$s = 2 \times 10^{-4} M$$

S13. Ans.(d)

Given : $K = 2 \times 10^{13}$ $\Delta G = \Delta G^{\circ} + RT \ln Q$ At equilibrium $\Delta G = 0, Q = K_{eq}$ So $\Delta_r G^{\circ} = -RT \ln K_{eq}$ $\Delta_r G^{\circ} = -8.314 J mol^{-1}K^{-1} \times 300K \times \ln (2 \times 10^{13})$

$$Ca(OH)_{2} \rightleftharpoons Ca^{2+} + 2OH^{-}$$

$$pH + pOH = 14$$

$$pH = 9, \text{ thus pOH} = 14 - 9 = 5$$

$$[OH^{-}] = 10^{-5}M$$
Thus $[Ca^{2+}] = \frac{10^{-5}}{2}$
So, $K_{sp} = [Ca^{2+}][OH^{-}]^{2}$

$$= \frac{10^{-5}}{2}(10^{-5})^{2} = 0.5 \times 10^{-15}$$
S15. Ans.(c)

$$0H^{\bigoplus} \stackrel{-H^+}{\longleftarrow} H_2 0 \stackrel{+H^+}{\longrightarrow} H_3 0^{\bigoplus}$$

Conjugate Conjugate base acid

 $\underset{Acid}{HF} + \underset{Base}{H_2O} \rightleftharpoons \underset{Conjugate \ base}{F^{\ominus}} + \underset{Conjugate \ acid}{H_3O^{\oplus}}$ S16. Ans.(c) (i) $CH_3COOH + NaOH \rightarrow CH_3COONa + H_2O$ Initial 25 mL 50 mL 0 × 0.1 M × 0.1 M = 2.5 mmol = 5 mmol0 2.5 mmol Final 2.5 mmolThis is basic solution due to NaOH. This is not basic buffer. (ii) $CH_3COOH + NaOH \rightarrow CH_3COONa + H_2O$ Initial 100 mL 100 mL 0 $\times 0.1 \text{ M} \times 0.1 \text{ M}$ = 10 mmol = 10 mmol0 0 Final 10 mmol Hydrolysis of salt takes place. This is not basic buffer. (iii) $HCl + NH_4OH \rightarrow NH_4Cl + H_2O$ Initial 100 mL 200 mL 0 × 0.1 M × 0.1 M = 10 mmol = 20 mmolFinal 0 10 mmol 10 mmol This will make basic buffer. (iv) $HCl + NaOH \rightarrow NaCl + H_2O$ Initial 100 mL 100 mL 0 $\times 0.1 \text{ M} \times 0.1 \text{ M}$ = 10 mmol = 10 mmol Final 0 0 10 mmol \Rightarrow This is Neutral solution. S17. Ans.(a) $A_2(g) + B_2(g) \rightleftharpoons X_2(g); \Delta H = -X kJ$

> On increasing pressure, equilibrium shifts in a direction where pressure decreases i.e. forward direction.

On decreasing temperature, equilibrium shifts in forward direction as it is exothermic direction.

So, high pressure and low temperature favours maximum formation of product.

S18. Ans.(a)

Solubility of BaSO₄, $s = \frac{2.42 \times 10^{-3}}{233} (mol \ L^{-1})$

$$= 1.04 \times 10^{-5} (mol \ L^{-1})$$

$$BaSO_{4}(s) \rightleftharpoons Ba^{2+}(aq) + SO_{4}^{2-}(aq)$$

$$K_{sp} = [Ba^{2+}][SO_{4}^{2-}] = s^{2}$$

$$= (1.04 \times 10^{-5})^{2}$$

$$= 1.08 \times 10^{-10} mol^{2} L^{-2}$$
S19. Ans.(c)

$$M_{eq} \text{ of HCl} = 75 \times \frac{1}{5} \times 1 = 15$$

$$- M_{eq} \text{ of HCl} = 75 \times \frac{1}{5} \times 1 = 5$$

$$- M_{eq} \text{ of HCl} \text{ in resulting solution} = 10$$

$$- Molarity \text{ of [H^+] in resulting mixture}$$

$$= \frac{10}{100} = \frac{1}{10}$$

$$pH = -\log[H^+] = -log\left[\frac{1}{10}\right] = 1.0$$
S20. Ans.(c)

$$2NH_{3} \rightleftharpoons N_{2} + 3H_{2} \qquad \frac{1}{K_{1}}$$

$$N_{2} + O_{2} \rightleftharpoons 2NO \qquad K_{2}$$

$$3H_{2} + \frac{3}{2}O_{2} \rightleftharpoons 3H_{2}O \qquad K_{3}^{3}$$

$$2NH_{3} + \frac{5}{2}O_{2} \rightleftharpoons 2NO + 3H_{2} \qquad K$$

$$K = \frac{K_{2}K_{3}^{3}}{K_{1}}$$

S21. Ans.(a)

$$Ag_2C_2O_4 \rightarrow 2Ag^+ + C_2O_4^{2-}$$

 $K_{sp} = [Ag^+]^2[C_2O_4$
 $= (2s)^2 (s) ; [Ag^+ = 2.2 \times 10^{-4}]$
 $= (2.2 \times 10^{-4})^2 (1.1 \times 10^{-4});$
 $[2s = 2.2 \times 10^{-4}]$
 $= 4.84 \times 10^{-8} \times 1.1 \times 10^{-4}$
 $= 5.3 \times 10^{-12}$

S22. Ans.(c)

The value of equilibrium constant is not changed in the presence of a catalyst in the reaction at equilibrium. The catalyst only increases the rates of reaction, it does not affect the equilibrium constant.

$$K_p = K_c (RT)^{\Delta n}$$
$$K_c = \frac{K_p}{(RT)^{\Delta n}} = \frac{5.8 \times 10^5}{(0.08314 \times 298)^{2-4}}$$
$$= \frac{5.8 \times 10^5}{(0.08314 \times 298)^{-2}}$$
$$= 3.56 \times 10^8$$

S24. Ans.(a)

Presence of catalyst does not affect the equilibrium of the reaction. Catalyst only affect the rate of reaction.

S25. Ans.(b)

$$CO(g) + Cl_2(g) \rightleftharpoons COCl_2(g)$$

 $\Delta n = -1$
 $K_p = K_c (RT)^{\Delta n_g}$
 $\frac{K_p}{K_c} = \frac{1}{RT}$
S26. Ans.(d)
Among the 4 options PF₃ w

Among the 4 options PF_3 will behave as a Lewis base due to presence of lone pair of electrons in p-orbitals.

S27. Ans.(d)

$$\begin{split} NaCl(aq) &\rightleftharpoons Na^{+} \quad Cl^{-} \\ \text{Initially} & 0.1 \text{ M} & 0 & 0 \\ \text{At equilibrium 0} & 0.1 & 0.1 + \text{S} \\ & AgCl &\rightleftharpoons Ag^{+} + Cl^{-} \\ \text{Initially} & a & 0 & 0 \\ \text{At equilibrium 0} & \text{S} & 0.1 + \text{S} \\ & K_{sp} = 1.6 \times 10^{-10} = [Ag^{+}] [Cl^{-}] \\ &= S(0.1 + S) \\ & 1.6 \times 10^{-10} = S \times 0.1 \\ & S = 1.6 \times 10^{-9} M \end{split}$$

S28. Ans.(d)

Pyrimidine is a very weak base so $K_b = c\alpha^2$

$$\alpha = \sqrt{\frac{K_b}{c}}$$
$$= \sqrt{\frac{1.7 \times 10^{-9}}{0.9}}$$
$$= 1.3 \times 10^{-4}$$
$$\% \alpha = 0.013\%$$

S29. Ans.(c)

If large amount of KHSO₄ would be added then concentration of NO_2^+ will decrease, rate of nitration will also decrease.

For
$$MY \rightleftharpoons M_S^+ + Y_S^-$$

$$K_{sp} = 6.2 \times 10^{-13}$$

$$S = 7.9 \times 10^{-7} mol \ L^{-1}$$

For $NY_3 \rightleftharpoons N^{3+} + 3Y^-$

$$K_{sp} = 27S^4 \implies S = \left(\frac{K_{sp}}{sp}\right)^{1/4}$$

$$K_{sp} = 273 \qquad \Rightarrow \qquad S = \left(\frac{1}{27}\right)$$
$$S = 3.89 \times 10^{-4} mol \ L^{-1}$$

So, solubility of NY_3 is more than solubility of MY in pure water. And addition of KY decrease the solubility because of common ion effect.

S31. Ans.(b)

If the reaction is multiplied by $\frac{1}{2}$, then new equilibrium constant, $K' = K^{1/2}$.

S32. Ans.(b)

 $HClO_4$ and $NaClO_4$ will not make a buffer solution because a buffer solution is formed by the mixture of a weak acid and its conjugate base and $HClO_4$ is not a weak acid.

S33. Ans.(b)

Such big equilibrium constant indicates large concentration of products so equilibrium is mostly towards product.

S34. Ans.(c)

According to solubility law:

$$Ag_{2}CrO_{4} \rightleftharpoons 2Ag^{+} + CrO^{4^{2-}}$$

$$(2S)^{2} \qquad S$$

$$1.1 \times 10^{-12} = 4S^{3}$$

$$Ag_{2}CrO_{4} = S \sim 10^{-3}$$

$$AgCl = S \sim 10^{-5}$$

$$AgBr = S \sim 10^{-6}$$

$$AgI = S \sim 10^{-8}$$
So, $Ag_{2}CrO_{4}$ will be the most soluble.

S35. Ans.(d)

: 1 mole of NaOH is completely neutralized by 1 mole of HCl.

So, 0.01 mole is neutralized by 0.01 mole of HCl.

Left NaOH = 0.09 mole

$$[OH^{-}] = \frac{0.09}{2} = 0.045M$$
$$p^{OH} = -\log[OH^{-}] = 1.35$$
$$pH = 14 - p^{OH} = 14 - 1.35 = 12.65$$

S36. Ans.(a)

HCl is strong acid and dissociated completely into ions in aq. Solution.

S37. Ans.(a) $\therefore \Delta G = \Delta G^{\circ} + 2.303 RT \log Q$ At equilibrium $\Delta G = 0$ $Q = K_{eq}$ So, $\Delta G^{\circ} = (-)2.303 RT \log K_{eq}$ S38. Ans.(b) Highest pH is shown by alkali

So, Na_2CO_3 being salt of a strong base and weak acid will exhibit highest pH.

NaCl \rightarrow neutral pH = 7

 $CuSO_4 \rightarrow Acidic \ pH < 7$

 $KCl \rightarrow Neutral pH = 7$

S39. Ans.(b)

$$\therefore \ \Delta G^{\circ} = -2.303 \ RT \log K_{sp}$$

63.3 × 10³ J = -2.303 × 8.314 × 298 log K_{sp}
 $K_{sp} = 8.128 \times 10^{-12}$

S40. Ans.(c)

On increasing pressure, equilibrium shifts in forward direction, where number of moles decreases, it is an example of exothermic reaction therefore decreasing temperature favours the forward direction.

S41. Ans.(c)

Lewis base species are those which are electron deficient and does require electron to complete their octet. BF_3 is an electron deficient species and is a Lewis acid.

S42. Ans.(c)

HCl and SO₂ are reducing agents so they can reduce MnO_4^- .

CO₂ is neither oxidizing nor reducing agent it will provide only acidic medium and shift the direction in forward reaction towards completion.