KD EDUCATION ACADEMY (9582701166

Time: 3 Hour 20 Minute

11.

STD 11 Science Chemistry kd 90+ ch-6 equilibrium

Total Marks: 230

*	Choose The Righ	nt Answer From The Gi	iven Options.[1 Marks	Each]	[69]
1.	0.10M CH ₃ COOH	is 1.34% ionised, calcula	ate its K _a :		
	(A) 1.8×10^{-5}	(B) 1.8×10^{-4}	(C) 5×10^{-4}	(D) 4×10^{-5}	
2.	substances capal	stance that is capable of ole of accepting a hydrog nent is justified by	donating a hydrogen ion gen ion, H ⁺ ".	H ⁺ and bases are	
	(A) Arrhenius con	cept.	(B) Bronsted-Low	ry theory.	
	(C) Lewis concept		(D) All of the abo	ve.	
3.	What will be the $ m GHF, H_2SO_4 \ and$		following Bronsted acids?		
	(A) $\mathrm{F^-,SO_4^{2-}}$ an	$ m d~CO_3^{2-}$	(B) $\mathrm{F}^-, \mathrm{SO}_4^{2-}$ an	$ m d~H_2CO_3$	
			(C) F^- , HSO_4^- and	${ m id}\ { m H}_{2}{ m CO}_{3}$	
	(D) $\mathrm{F^-,HSO_4^-}$ a	${ m nd}~{ m CO}_3^{2-}$			
4.	What will be the r	nolar solubility S of a sol	id salt with general form	ula $ m M_{x}^{p+} m X_{y}^{q}$?	
			(C) $\left(\frac{K_{\rm sp}}{x^x \cdot y^y}\right)^{\frac{1}{x+y}}$	v	
5.	The pH of boiling	water is 6.4. This implie:	s that boiling water is:		
	(A) Slightly basic.		(B) Slightly acidic		
	(C) Neutral.		(D) Amphoteric.		
6.		constant of a reaction at μ . The $\triangle H$ for the reaction	298K and 1000K is 5×1 on is:	0^{-3} and 2 \times	
	(A) Positive.		(B) Negative.		
	(C) Either positive	or negative.	(D) Zero.		
7.	The mass of acet $COOH = 1.8 \times 10$		of solution in which it is	1% ionised (Ka of CH ₃	
	(A) 5.4g	(B) 12.6g	(C) 6.4g	(D) 10.8g	
8.	Which among the (A) Change in ten		es the value of ionic pro (B) Addition of ac		
	(C) Addition of ba	se.	(D) Addition of ei	ther acid and base.	
9.	In which condition	n, the reaction proceeds	in the forward direction?	?	
	(A) $\mathrm{Q}_{\mathrm{C}}=\mathrm{K}_{\mathrm{C}}$	(B) $ m Q_{C} > m K_{C}$	(C) $ m Q_{C} < m K_{C}$	(D) $ m Q_{ m C} eq m K_{ m C}$	
10.	The strength of a	cid is highest in:			
	(A) $pK_a = 6$	(B) $pK_a = 5$	(C) $pK_a = 10$	(D) $pK_a = 1$	

 $0.1M\ CH_3COOH$ and $1.01CH_3COONa$ are mixed togethere, what will be pH of buffer

solution if $pK_a=4.75[\log 10-1=-1]$

	(4) 2.75	(D) 4.75	(C) F 7F	(D) 6.75			
	(A) 3.75	(B) 4.75	(C) 5.75	(D) 6.75			
12.							
	(A) Hardy-Schulze rul	e.					
	(B) Distribution law.						
	(C) Le Chatelier's prin	•					
	(D) Neutralization of o	charge on the colloidal	particles.				
13.	A solution which mair	ntains constant pH whe	n small amounts of ac	id or alkali are added			
	is known as						
	(A) Indicator		(B) Buffer				
	(C) Amphoteric		(D) Neutral				
14.	The pH value of blood base, because the blo	d does not appreciably bod:	change by a small add	dition of an acid or a			
	(A) Is a body fluid.						
	(B) Can be easily coa	gulated.					
	(C) Contains iron as a	part of the molecule.					
	(D) Contains serum p	rotein which acts as but	ffer.				
15.	Addition of HCl will no	ot suppress the ionization	on of:				
	(A) Acetic acid		(B) Benzoic acid				
	(C) H ₂ S		(D) Sulphuric acid	d			
16.	-	t K of the sparingly so					
10.	solubility of the salt is	t K _{sp} of the sparingly so s:					
	(A) $1.0 \times 10^{-4} \text{mol L}^{-1}$		(B) 2 × 10 ⁻⁶ mol L				
	(C) 1.0×10^{-5} mol L ⁻¹		(D) 2×10^{-12} mol	L ⁻¹			
17.	Which of the following	g species is amphoteric	in nature.				
	(A) $\mathrm{H_3O}^+$	(B) Cl ⁻	(C) HSO_4^-	(D) CO_3^{2-}			
18.	The ionic product of v	water if a few dro	ops of acid or base are	e added to it.			
	(A) Increases.		(B) Decreases.				
	(C) Remains the same	3.	(D) Can not predi	ct.			
19.	A buffer solution is a	solution whose pH valu	e on keeping in the ai	r:			
	(A) Increases rapidly.		(B) Decreases rap	oidly.			
	(C) May increase or d	ecrease.	(D) Does not char	nge.			
20.	In the reaction,						
	$N_2(g) + O_2(g) \rightleftharpoons 2NO($	g) - 180.7 kJ,					
	on increasing the tem	nperature, the production	on of NO:				
	(A) Increases.		(B) Decreases.				
	(C) Remains same.		(D) Cannot be pre	edicted.			
21.	Calculate the molar s	solubility (S) of a salt like	e zirconium phosphate	e of molecular			
	formula $(\mathrm{Zr}^{4+})_3(\mathrm{PC})$	$(p_4^{3-})_4$.					
	(A) $\left(\frac{\mathrm{K_{sp}}}{9612}\right)^{\frac{1}{8}}$	(B) $\left(\frac{\mathrm{K_{sp}}}{6912}\right)^{\frac{1}{7}}$	(C) $\left(\frac{\mathrm{K_{sp}}}{5348}\right)^{\frac{1}{6}}$	(D) $\left(rac{ ext{K}_{ ext{sp}}}{8435} ight)^{rac{1}{7}}$			
22.	Strong electrolyte of	,	(0040)	(0400)			
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(B) 0.1M HCl

(A) 01M HAc

	(C) 0.1M KCI	(D) 0.1M NaCl			
23.	In the presence of a common ion (incapable of forming complex ion), the solubility of salt in solution.				
	(A) Increases.	(B) Decreases.			
	(C) Remains the same.	(D) Cannot predict.			
24.	In a reversible reaction $H_2 + I_2 \leftrightharpoons 2HI$, if the concentration of H_2 and I_2 are increased, the value of Kc:				
	(A) Decreases.	(B) Increases.			
	(C) Remains the same.	(D) Changes exponentially.			
25.	Given the chemical equilibrium, $A \rightleftharpoons B + C$, where ΔH_{rxn} is negative, what effect increasing the temperature (at constant pressure) would have on the system at equilibrium?				
	(A) No change.				
	(B) Shift to the left.				
	(C) Shift to the right.				
	(D) Shift to the left for K<1 and to the r	ight for K>1.			
26.	$\rm K_a$ for CH $_3$ COOH is 1.8×10^{-5} and $\rm K_b$ for NH $_4$ OH is 1.8×10^{-5} . The pH of ammonium acetate will be				
	(A) 7.005	(B) 4.75			
	(C) 7.0	(D) Between 6 and 7			
27.	Which of the following will produce a buffer solution when mixed in equal volumes?				
	(A) $0.1 \text{mol dm}^{-3} \text{ NH}_4 \text{OH and } 0.1 \text{mol dm}^{-3} \text{ HCl}$.				
	(B) 0.05 mol dm ⁻³ NH ₄ OH and 0.1 mol dm ⁻³ HCl.				
	(C) 0.1mol dm ⁻³ NH ₄ OH and 0.05mol dm ⁻³ HCl.				
	(D) 0.1mol dm ⁻³ CH ₄ COONa and 0.1mol dm ⁻³ NaOH.				
28.	Which one does not give a buffer solution?				
_0.	(A) Ammonia and sodium hydroxide in water.				
	(B) Sodium acetate and acetic acid in water.				
	(C) Ammonia and ammonium chloride in water.				
	(D) Sodium acetate and hydrochloric acid in water.				
29.	Acidity of BF ₃ can be explained on the basis of which of the following concepts?				
	(A) Arrhenius concept.				
	(B) Bronsted Lowry concept.				
	(C) Lewis concept.				
	(D) Bronsted Lowry as well as Lewis concept.				
30.	Strong acid dissociates completely in water, the resulting base formed would be very weak. The reason is that:				
	(A) Strong acids have strong conjugate bases.				
	(B) Strong acids have strong conjugate acids.				
	(C) Strong acids have very weak conjugate bases.				
	(D) Strong acids have very weak conjugation	gate acids.			
31.					

		${ m D.}$ If the concentration ${ m ED}$ will be twice that of ${ m AD}$		at equilibrium and e equilibrium constant of		
	(A) 4	(B) 6	(C) $\frac{4}{5}$	(D) $\frac{6}{7}$		
32.	9					
	(pink)		(blue)			
	(A) $\Delta { m H} > 0$ for t	the reaction.				
	(B) $\Delta \mathrm{H} < 0$ for t					
	(C) $\Delta H = 0$ for t					
		$\Delta \mathrm{H}$ cannot be predicted		itormation.		
33.		$2SO_2(g) + O_2(g) \rightleftharpoons 2SO_2$	3(g) shifts forward if:			
	(A) Catalyst is us		0			
		t is used to remove SO_3 at of products are used.	as soon as it is forme	a.		
	•	nt of products are used.				
34.				Jecanse.		
J-7.	The addition of NaCl to AgCl decreases the solubility of AgCl because: (A) Solubility product decreases.					
		duct remains constant.	221			
		omes unsaturated.				
	(D) Solution beco	omes supersaturated.				
35.	. In the equilibrium, $AB ightleftharpoons A+B$ if the equilibrium concentration of A is double, then equilibrium concentration of B will be:					
	(A) Half	(B) twice	(C) $\frac{1}{4}$ th	(D) $\frac{1}{8}$ th		
36.	precipitation occ	cur? $[K_{SP} \text{ of } CaF_2 = 1.7]$		which of the Solutions will		
	• •	$^{+}+10^{-5} { m MF}^{-}$				
		$^+~{ m and}~10^{-3}{ m MF}^-$				
	(C) $10^{-2} \mathrm{M~Ca^{2-}}$					
	(D) $10^{-3} \mathrm{M~Ca}^2$	$^{+} { m \ and \ } 10^{-5} { m MF}^{-}$				
37.	$SO_2 + O_2 \rightleftharpoons 2SO$					
	•	reaction proceeds in fo	-			
	(A) Addition of O		(B) Removal	=		
	(C) Addition of in	_	(D) Cannot p	roceed		
38.		lution of NaNO ₂ and HN		dala sak		
	(A) Strong reduction		(B) Strong ox (D) None of t			
20	(C) Buffer solution					
39.	(A) Low P, High T	quilibrium A + 2B ⇌ C +				
	(A) LOW F, FIGIL I		(B) Low P, Lo	VV I		

	(C) High P, Low T		(D) High P, Hig	hΤ
40.	The solubility produ	uct of CaSO $_4$ is 6.4 $ imes$	x 10 ⁻⁵ . The solubility of sa	alt in mol L ⁻¹ is:
(A) 8.10 ⁻¹⁶	(B) 8.10 ⁻²	(C) 8.10 ⁻³	(D) 1.6 ⁻³
41.	Buffer Solution is p	repared by mixing _	·	
	(A) Weak acid and i	ts salt of strong base	e.	
	(B) Strong acid + its	_		
	(C) Weak acid + its			
	(D) Strong base + it	_		
42.		owing reactions, the gon at constant volu	equilibrium remains una ıme?	affected on addition of
	(A) $H_2(g) + I_2(g) =$	$ ightharpoonup 2 \mathrm{HI}(\mathrm{g})$		
	(B) $\operatorname{PCl}_5(g) \rightleftharpoons \operatorname{PCl}_5(g)$	$\mathrm{cl}_3(\mathrm{g})+\mathrm{Cl}_2(\mathrm{g})$		
	(C) $N_2(g) + 3H_2(g)$	- (-/		
			ted in all the three cases	
43.			500K in a closed contain	
			$1.2 imes 10^{-3}$ mol L $^{-1}$ and 1 action $ ext{PCl}_5 ext{g} ightharpoons ext{PCl}_3$ (§	
	(A) $1.8 \times 10^3 \text{mol L}^{-1}$		(B) 1.8×10^{-3}	5) C128 Will BC.
	(C) 1.8×10^{-3} L mol		(D) 0.55×10^4	
44.	• •			energy is $\Delta \mathrm{G}^\ominus>0$ The
	equilibrium constar			energy is $\Delta \alpha > 0$ the
(A) $K = 0$	(B) K > 1	(C) K = 1	(D) K < 1
45.			$egin{aligned} K_c & and \ K_p & is \ K_p & = K_c(I) \ \mathcal{C} I(s) & & \mathcal{C} I(s) & \mathcal{C} I(s) \end{aligned}$	
(A) 1	(B) 0.5	(C) 1.5	(D) 2
46.	1mL of $rac{ m N}{100} m HCl$ is a	added to 1L of buffer	r having pH = 5. The pH	of the solution will be:
	(A) Become 7		(B) Become 3	
	(C) Become 6		(D) Remain un	affected.
47.	-		pheric pressure, the soli	
	(A) Normal melting	point.	(B) Equilibrium	temperature.
	(C) Boiling point.		(D) Freezing po	oint.
48.	The chemical equil	ibrium of reversible	reaction is not influence	ed by:
	(A) Pressure.	5	(B) Catalyst.	
	(C) Concentration o	f the reactants.	(D) Temperatu	re.
49.			presented by $K=rac{[H_3O]}{[H_3O]}$	$^+][\mathrm{OH}^-] \ \mathrm{H_2O]}$
	$\mathrm{or} = [\mathrm{H}^+][\mathrm{OH}^-]\mathrm{H}$			_
	(A) Ionic product of		(B) Ionic produ	
	(C) Ionisation const	ant of water.	(D) Ionisation (constant of acid and base.

50.	The equibrium constant K_c for $H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$ at	700K is 49			
	The 'K' for reaction $HI(g) \rightleftharpoons G$ (A) 49 (B) 0.02		(D) 1.43		
	. ,	(C) 7	(5) 1.15		
51.	Production of ammonia accord $ m N_2(g) + 3H_2(g)2NH_3(g);$				
	$\Delta ext{H} = -92.38 ext{kJ mo}$		116		
	is an exothermic process. At lo	w temperature, the reaction s	nifts in:		
	(A) Forward direction. (B) Backward direction.				
	(C) Either forward or backward	direction			
	(D) None of the above.	an ection.			
52.	Which buffer solution comprisi	ng of the following has its pH v	value greater than 7?		
	(A) CH ₃ COOH + CH ₃ COONa	(B) HCOOH -	\ _ '		
	(C) CH ₃ COONH ₄	(D) NH ₄ OH -	l'		
53.	1M NaCl and 1M HCl are present in an aqueous solution . The solution is:				
	(A) Not a buffer solution with p	H < 7 (B) Not a but	ffer solution with pH > 7		
	(C) A buffer solution with pH $<$	7 (D) A buffer	solution with pH > 7		
54.	It accepts a proton. It is called	as:			
	(A) A Bronsted acid.	(B) A Bronst	ed base.		
	(C) A strong acid.	(D) A weak k	oase.		
55.	The addition of HCl will not suppress the ionisation of:				
	(A) Acetic acid	(B) Sulphurio			
	(C) H ₂ S	(D) Benzoic	acid		
56.	100mL of a solution contains 0 change on adding:	$0.1 {\sf MNH_4OH}$ and $0.1 {\sf MNH_4CI}$. Th	ne pH of the solution will not		
	(A) 20mL of 0.1M NH ₄ OH soluti		0.1M NH ₄ Cl solution.		
	(C) 10mL of 0.1M NaOH solutio		distilled water.		
57.	A 0.2 molar solution of formic				
	(A) 9.6×10^{-3}	(B) 2.1×10^{-1}			
	(C) 1.25×10^{-6}	(D) 4.8 × 10	-5		
58.	What will be pH of $0.01M \text{ CH}_3\text{C}$	COOH?			
	$(K_a = 1.80 \times 10^{-5})$	(0)	(5)		
	(A) 3.4 (B) 3.6	(C) 3.9	(D) 3.0		
59.	Buffer solution can be obtaine		of		
	(A) Sodium acetate and excess(B) Sodium acetate and acetic				
	(C) Sodium chloride and HCl.	aciu.			
	(D) Acetic acid and excess of N	laOH.			
60.	The concentration of hydroger pH?		s 3.8 $ imes$ 10 ⁻³ M. What is its		

	(A) 4.32	(B) 5.12	(C) 3.31	(D) 2.42	
				(D) 2.42	
61.		nis solution will not chang			
	(A) Chemical pH indica		(B) Acid/ base buffer.		
	(C) Anhydrous solution		(D) Hypotonic solution		
62.			n that in pure water beca	use:	
	(A) Agl forms complex				
	(B) Of common ion eff		Nal		
		of AgI is less than that of of the solution decreases			
62	·				
63.	its molar solubility is:	of a sparingly soluble sa	It AB at room temperatu	re is 1.21 × 10 ⁻⁰ ,	
	(A) $1.21 \times 100M$.		(B) 1.1×10^{-4} M.		
	(C) 1.1×10^{-3} M.		(D) None of these.		
64.			will the gas phase reacti the Le Chatelier's princip	•	
	$ m N_2(g) + 3H_2(g) ightleftharpoons 2$	$2NH_3(g)$			
	Which of the following	- (9)	essure at which the equil temperature?	librium is	
	(A) K will remain same				
	(B) K will decrease.		427		
	(C) K will increase.				
	(D) K will increase initially and decrease when pressure is very high.				
65.	55. The solubility of CO ₂ in water increases with:				
	(A) Increase in temper	rature.	(B) Reduction of gas p	ressure.	
	(C) Increase in gas pre	essure.	(D) Increase in volume	2.	
66.	Which is not a buffer s	solution:			
	(A) $NH_4CI + NH_4OH$		(B) $CH_3COOH + CH_3COOH$	OONa	
	(C) CH ₃ COONH ₄		(D) NH_4NO_3		
67.		ne concentration of H ⁺ io	emperature increases, ic ons and OH [–] ions are equ		
	(A) Equal to 7.0		(B) Greater than 7.0		
	(C) Less than 7.0		(D) Equal to zero.		
68.	What is pH of resulting 0.01M HCl are mixed?		lume when equal of 0.1M	I NaOH and	
	(A) 7	(B) 1.04	(C) 12.65	(D) 2.0	
69.	Which of the following	is the example of a reve	ersible reaction?		
	(A) $Pb(NO_3)_2(aq) +$	$(\mathrm{s}) + 2\mathrm{NaNO}_3(\mathrm{aq})$			
	(0, (-,	$) \longrightarrow 2 NaOH(aq) +$, -, -,		
	` '	$\overset{,}{\operatorname{Cl}}(\operatorname{aq}) \longrightarrow \operatorname{AgCl}(\overset{,}{\operatorname{s}}) +$	(-)		
	(D) $\mathrm{KNO_3(aq)} + \mathrm{NaCl(aq)} \longrightarrow \mathrm{KCl(aq)} + \mathrm{NaNO_3(aq)}$				
	- \ -/	· · ·	- \ -/		

* a statement of Assertion (A) is followed by a statement of Reason (R). Choose the correct option.

[2]

70. **Note:** In the following questions a statement of Assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given\ below each question.

Assertion (A): A solution containing a mixture of acetic acid and sodium acetate maintains a constant value of pH on addition of small amounts of acid or alkali.

Reason (R): A solution containing a mixture of acetic acid and sodium acetate acts as a buffer solution around pH 4.75.

- i. Both A and R are true and R is correct explanation of A.
- ii. Both A and R are true but R is not the correct explanation of A.
- iii. A is true but R is false.
- iv. Both A and R are false.
- 71. **Note:** In the following questions a statement of Assertion (A) followed by a statement of Reason (R) is given. Choose the correct option out of the choices given\ below each question.

Assertion (A): The ionisation of hydrogen sulphide in water is low in the presence of hydrochloric acid.

Reason (R): Hydrogen sulphide is a weak acid.

- i. Both A and R are true and R is correct explanation of A.
- ii. Both A and R are true but R is not correct explanation of A.
- iii. A is true but R is false.
- iv. Both A and R are false.

* Answer The Following Questions In One Sentence.[1 Marks Each]

[25]

72. What is Kc for the following equilibrium when the equilibrium concentration of each substance is:

$$[SO_2] = 0.60M$$
, $[O_2] = 0.82M$ and $[SO_3] = 1.90M$?

$$2SO_{2(g)} + O_{2(g)} \rightleftharpoons 2SO_{3(g)}$$

73. Calculate the hydrogen ion concentration in the following biological fluids whose pH are given below:

Human blood, 7.38

74. The equilibrium constant expression for a gas reaction is,

$$K_{c} = \frac{\left[NH_{3}\right]^{4}\left[O_{2}\right]^{5}}{\left[NO\right]^{4}\left[H_{2}O\right]^{6}}$$

Write the balanced chemical equation corresponding to this expression.

- 75. Write the conjugate acids for the following Brönsted bases: NH_2^- , NH_3 and HCOO-.
- 76. Which of the following reactions will get affected by increasing the pressure? Also, mention whether change will cause the reaction to go into forward or backward direction.

$$\mathrm{CH_{4}\ (g)} + 2\mathrm{S_{2}\ (g)} \rightleftharpoons \mathrm{CS_{2}\ (g)} + 2\mathrm{H_{2}S\ (g)}$$

77. Write the expression for the equilibrium constant, K_C for each of the following reactions:

$$Fe^{3+}(aq) + 3OH^{-}(aq) \rightleftharpoons Fe(OH)_3(S)$$

78. At 473K, equilibrium constant K_c for decomposition of phosphorus pentachloride, PCI_5 is 8.3 $\times 10^{-3}$. If decomposition is depicted as,

 $\mathrm{PCl}_5 \; (\mathrm{g})
ightleftharpoons \mathrm{PCl}_3 \; (\mathrm{g}) + \mathrm{Cl}_2 \; (\mathrm{g}) \; \Delta_\mathrm{r} \mathrm{H}^\ominus = 124.0 \; \mathrm{kJ} \; \mathrm{mol}^{-1}$

what is the value of K_c for the reverse reaction at the same temperature?

79. At 473K, equilibrium constant K_c for decomposition of phosphorus pentachloride, PCI_5 is 8.3×10^{-3} . If decomposition is depicted as,

$$PCl_{5}~(g) \rightleftharpoons PCl_{3}~(g) + Cl_{2}~(g)~\Delta_{r}H^{\ominus} = 124.0~kJ~mol^{-1}$$

what would be the effect on Kc if

- i. more PCI5 is added
- ii. pressure is increased
- iii. the temperature is increased?
- 80. Describe the effect of:

addition of H2

on the equilibrium of the reaction:

$$2H_2(g) + CO(g) \rightleftharpoons CH_3OH(g)$$

81. Which of the following reactions will get affected by increasing the pressure? Also, mention whether change will cause the reaction to go into forward or backward direction.

$$CO_2$$
 (g) + C (S) \rightleftharpoons 2CO (g)

82. Does the number of moles of reaction products increase, decrease or remain same when each of the following equilibria is subjected to a decrease in pressure by increasing the volume?

$$CaO(S) + CO_2(g) \rightleftharpoons CaCO_3(S)$$

- 83. Predict if the solutions of the following salts are neutral, acidic or basic: ${\rm KB}_{\rm r}$
- 84. Calculate the hydrogen ion concentration in the following biological fluids whose pH are given below:

Human stomach fluid, 1.2

85. Describe the effect of:

removal of CH₃OH

on the equilibrium of the reaction:

$$2H_2(g) + CO(g) \rightleftharpoons CH_3OH(g)$$

- 86. What will be the conjugate bases for the Brönsted acids: HF, ${
 m H_2SO_4}$ and ${
 m HCO_3^-}$?
- 87. For the following equilibrium, $K_c = 6.3 imes 10^{14}~at~1000 K$

$$NO(g) + O_3(g) \rightleftharpoons NO_2(g) + O_2(g)$$

Both the forward and reverse reactions in the equilibrium are elementary bimolecular reactions. What is K_c , for the reverse reaction?

- 88. If $Q_c < K_c$, in which direction reaction will proceed?
- 89. What happens to ionic product of water if some acid is added to it?
- 90. Classify the following as Lewis acid or Lewis base:

 $\mathrm{NH_4^+}$ and $\mathrm{NH_3}$

- 91. SO_3^{2-} is Bronsted base or acid and why?
- 92. Why pH of our blood remains constant at 7.4 through we quite often eat spicy food?

- 93. What is the effect of temperature on solubility product (K_{sp}) ?
- 94. Write K_p in terms of K_c for the following chemical reaction: $2SO_2(g) + O_2(g) \rightleftharpoons 2SO_3(g)$
- 95. Which of the following is weakest acid? $HClO_4, HClO_3, HCl_2, HClO$
- 96. For Tribasic acid $K_{a1} > K_{a2} > K_{a3}$ what will happen to the acid strength of polyprotic acids if protons are lost?

* Given Section consists of questions of 2 marks each.

[66]

- 97. A liquid is in equilibrium with its vapour in a sealed container at a fixed temperature.The volume of the container is suddenly increased.How do rates of evaporation and condensation change initially?
- 98. Assuming complete dissociation, calculate the pH of the following solutions: 0.002 M KOH
- 99. Assuming complete dissociation, calculate the pH of the following solutions: 0.003M HCl
- 100. Assuming complete dissociation, calculate the pH of the following solutions: 0.002 M HBr
- 101. Calculate the pH of the resultant mixtures: 10mL of 0.01M $H_2SO_4 + 10mL$ of 0.01M $Ca(OH)_2$
- 102. A liquid is in equilibrium with its vapour in a sealed container at a fixed temperature. The volume of the container is suddenly increased.
 What happens when equilibrium is restored finally and what will be the final vapour pressure?
- 103. Assuming complete dissociation, calculate the pH of the following solutions: 0.005 M NaOH
- 104. Calculate the pH of the resultant mixtures: 10mL of 0.1M $H_2SO_4 + 10mL$ of 0.1M KOH
- 105. $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g);$ K = 0.50 at 673K.

Write the equilibrium expression and equilibrium constant for reverse reaction.

- 106. On the basis of the equation $pH = -log [H^+]$, the pH of 10^{-8} mol dm⁻³ solution of HCl should be 8. However, it is observed to be less than 7.0. Explain the reason.
- 107. Which of the following is strongest acid? $HCl, HClO_3, HNO_3, H_2SO_4, HClO_4$
- 108. The K_{sp} of Ag_2CrO_4 AgCl, 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 first AgNO₃ solution is adding to a solution containing equal mole of NaCl, NaBs, NaI and Na₂CrO₄.
- 109. At what temperature the solid and liquid are in equilibrium under 1 atm pressure?
- 110. What will be effect on boiling point of liquid if pressure is increased?
- 111. How does a catalyst affect the equilibrium constant? Explain.

- 112. Mention the conditions of temperature and pressure when gas will dissolve in liquid to maximum extent with decrease in volume and absorption of heat.
- 113. The value of Kc for the reaction $2\mathrm{HI}(\mathrm{g}) \rightleftharpoons \mathrm{H}_2(\mathrm{g}) + \mathrm{I}_2(\mathrm{g})$ is 1×10^{-4} At a given time, the composition of reaction mixture is [HI] = 2×10^{-5} mol, [H₂] = 1×10^{-5} mol and [I₂] = 1×10^{-5} mol In which direction will the reaction proceed?
- 114. $CaCl_2(s) + aq \rightleftharpoons CaCl_2(aq) + Heat$ Discuss the solubility if temperature is increased.
- 115. One millilitre solution of 0.01M HCl is added to 1L of sodium chloride solution. What will be the pH of the resulting solutions?
- 116. Consider the following equilibrium:

$$CO_2(g) + C \text{ (graphite)} \Rightarrow 2CO(g)$$

Write the equilibrium expression for K_c and calculate its units.

- 117. Will AgCl be more soluble in aqueous solution or NaCl solution and why?
- 118. What is the effect of reducing volume on the following system?

$$2C(s) + O_2 \rightleftharpoons 2CO(g)$$

119. $C(diamond) \rightleftharpoons C(graphite)$

$$d = 3.5 g cm^{-3}$$
 $d = 2.3 g cm^{-3}$

What will be effect of increasing pressure in this equilibrium?

120. Conjugate acid of a weak base is always stronger. What will be the decreasing order of basic strength of the following conjugate bases?

- 121. pK_a value of acids A, B, C, D are 1.5, 3.5, 2.0 and 5.0. Which of them is strongest acid?
- 122. What could be temperature 15°C or 100°C for $K_w = 7.5 \times 10^{-14}$. What happens to ionic product if some acid is added to it?
- 123. What will be the pH of $1M \text{ Na}_2SO_4$ solution?
- 124. What is the relation between K_p and K_c ?
- 125. What is the effect of increasing pressure on the equilibrium?

$$N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$$

126. Ionisation constant of a weak base MOH, is given by the expression.

$$K_b = \frac{[M^+][OH^-]}{[MOH]}$$

Values of ionisation constant of some weak bases at a particular temperature are given below:

Base	Dimethylamine	Urea	Pyridine	Ammonia
K_b	5.4×10^{-4}	1.3×10^{-14}	1.77×10^{-9}	1.77×10^{-5}

Arrange the bases in decreasing order of the extent of their ionisation at equilibrium. Which of the above base is the strongest?

127. At 0°C, ice and water are present in equilibrium.

What will happen on increasing the pressure?

- 128. Why do we sweat more on humid day?
- 129. Write conjugate acid and conjugate base of H_2O .

Given Section consists of questions of 3 marks each.

What is meant by the conjugate acid-base pair? Find the conjugate acid/base for the 130. following species:

$$\mathrm{HNO}_2, \mathrm{CN}^-, \mathrm{HClO}_4, \mathrm{F}^-, \mathrm{OH}^-, \mathrm{CO}_3^{2-} \mathrm{and} \ \mathrm{S}^{2-}$$

- The value of K_c for the reaction $3O_2~(g) \rightleftharpoons 2O_3~(g)$ is 2.0 $\times 10^{-50}$ at 25°C. If the 131. equilibrium concentration of O_2 in air at 25°C is 1.6 $\times 10^{-2}$, what is the concentration of 0_{3} ?
- Find out the value of K_c for each of the following equilibria from the value of K_p: 132. $2CaCO_3$ (S) $\rightleftharpoons CaO(S) + CO_2$ (g); $k_p = 167$ at 1073K
- The reaction, $CO(g) + 3H_2(g) \rightleftharpoons CH_4(g) + H_2O(g)$ 133. is at equilibrium at 1300K in a 1L flask. It also contain 0.30mol of CO, 0.10molof H₂ and 0.02mol of H₂O and an unknown amount of CH₄ in the flask. Determine the concentration of CH₄ in the mixture. The equilibrium constant, K_c for the reaction at the given temperature is 3.90.
- The degree of ionization of a 0.1M bromoacetic acid solution is 0.132. Calculate the pH 134. of the solution and the pK_a of bromoacetic acid.
- The pH of a sample of vinegar is 3.76. Calculate the concentration of hydrogen ion in it. 135.
- If $K_w = 49 \times 10^{-14}$, what will be neutral pH of H_2O ? 136.
- Why do we pass H₂S gas in acidic medium in group 2? 137.
- The K_{sp} values of two slightly soluble salts AB and PQ_2 are each equal to 4.0×10^{-18} . 138. Which salt is more soluble?
- For the reaction : $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ 139.

Equilibrium constant
$$K_c = rac{[\mathrm{NH_3}]^2}{[\mathrm{N_2}][\mathrm{H_2}]^3}$$

Some reactions are written below in Column I and their equilibrium constants in terms of K_c are written in Column II. Match the following reactions with the corresponding equilibrium constant.

Column I (Reaction)

Column II (Equilibrium constant)

i.
$$2N_2(g) + 6H_2(g) \rightleftharpoons 4NH_3(g)$$

a.
$$2 {
m K}_{
m c}$$

ii.
$$2NH_3(g) \rightleftharpoons N_2(g) + 3H_2(g)$$

b.
$$K_c^{rac{1}{2}}$$

iii.
$$\frac{1}{2}N_2(g) + \frac{3}{2}H_2(g) \rightleftharpoons NH_3(g)$$

- Write a relation between ΔG and Q and define the meaning of each term and answer 140. the following:
 - Why a reaction proceeds forward when Q < K and no net reaction occurs when a. Q = K.
 - Explain the effect of increase in pressure in terms of reaction quotient Q. for b. the reaction: $CO(g) + 3H_2(g) \rightleftharpoons CH_4(g) + H_2O(g)$
- 141. Urine has a pH of 6.0. If a patient eliminates 1300mL of urine per day, how many gram equivalents of the acid he eliminates per day?

- 142. Calculate the molar solubility of Ni(OH)₂ in 0.10M NaOH. The ionic product of Ni(OH)₂ is 2.0×10^{-15} .
- 143. For the reaction, $N_2(g)+3H_2(g)\rightleftharpoons 2NH_3(g)$, the partial pressures of N_2 and H_2 are 0.80 and 0.40 atmosphere respectively at equilibrium. The total pressure of the system is 2.80 atmosphere. What is K_p for the above reaction?
- i. Why is NH_4Cl added before addition of NH_4OH in qualitative analysis of 3^{rd} group?
 - ii. Which will be added to precipitate soap (RCOONa)? NaCl or KCl and why?'
- Solid Ba(NO₃), is gradually dissolved in a 1.0×10^{-4} M Na₂CO₃ solution. At what concentration of Ba²⁺ will a precipitate begin to form.(K_{sp} for BaCO₃ = 5.1×10^{-9})
- 146. A certain buffer is made by mixing sodium formate and formic acid in water. With the help of equations explain how this buffer neutralizes addition of a small amount of an acid or a base?
- 147. MY and NY₃ two nearly insoluble salts, have same K_{sp} values 6.2 \times 10⁻³ of non temperature. Calculate solubility of each salt. Which has more solubility.
- 148. What is the effect of temperature on the reactions? Give reason.
 - i. $N_2(g)+3H_2(g) \rightleftharpoons 2NH_3(g) + Heat$
 - ii. $N_2(g)+O_2(g) \rightleftharpoons 2NO(g) + Heat$
- i. Give the relationship between K_a , c and $('\alpha')$ where ' K_a ' is acid dissociation constant, 'c' is molar concentration, ' α' is degree of dissociation.
 - ii. If the solubility of $Ca(IO_3)_2$ in water at 18°C is 2.1g/ litre. Calculate the value of solubility product.

[8]

[Molecular mass of $Ca(IO_3)_2 = 390$]

* Case study based questions

150. Read the passage given below and answer the following questions from (i) to (v).

Predicting the Direction of the Reaction- The equilibrium constant helps in predicting the direction in which a given reaction will proceed at any stage. For this purpose, we calculate the reaction quotient Q. The reaction quotient, Q (Qc with molar concentrations and QP with partial pressures) is defined in the same way as the equilibrium constant Kc except that the concentrations in Qc are not necessarily equilibrium values. For a general reaction:

$$aA + bB \rightleftharpoons cC + dD$$

$$Qc = \frac{[C]^c[D]^d}{[A]^a[B]^b}$$

Then

If Qc > Kc, the reaction will proceed in the direction of reactants (reverse reaction).

If Qc < Kc, the reaction will proceed in the direction of the products (forward reaction).

If Qc = Kc, the reaction mixture is already at equilibrium. Consider the gaseous reaction of H_2 with I_2 ,

$$\mathrm{H_{2(g)}} + \mathrm{l_{2(g)}} \rightleftharpoons \mathrm{2Hl_{(g)}}; \mathrm{kc} = 57.0\mathrm{at}700\mathrm{k}.$$

Suppose we have molar concentrations [H $_2$]t = 0.10M, [I $_2$]t = 0.20 M and [HI]t = 0.40 M. (the subscript t on the concentration symbols means that the concentrations were

measured at some arbitrary time t, not necessarily at equilibrium). Thus, the reaction quotient, Qc at this stage of the reaction is given by,

$$\mathrm{Qc} = rac{[\mathrm{Hl}]\mathrm{t}^2}{[\mathrm{H}]^2]_{\mathrm{t}}[\mathrm{l}_2]_{\mathrm{t}}} = rac{(0.40)_2}{(0.10) imes (0.20)} = 8.0$$

Now, in this case, Qc (8.0) does not equal Kc (57.0), so the mixture of $H_{2\ (g)}$, $I_{2\ (g)}$ and $HI_{(g)}$ is not at equilibrium; that is, more $H_{2\ (g)}$ and $I_{2\ (g)}$ will react to form more $HI_{(g)}$ and their concentrations will decrease till Qc = Kc. The reaction quotient, Qc is useful in predicting the direction of reaction by comparing the values of Qc and Kc.Thus, we can make the following generalisations concerning the direction of the reaction

If Qc < Kc, net reaction goes from left to right

If Qc > Kc, net reaction goes from right to left.

If Qc = Kc, no net reaction occurs.

Calculating Equilibrium Concentrations In case of a problem in which we know the initial concentrations but do not know any of the equilibrium concentrations, the following three steps shall be followed:

Step 1) Write the balanced equation for the reaction.

Step 2) Under the balanced equation, make a table that lists for each substance involved in the reaction: (a) the initial concentration, (b) the change in concentration on going to equilibrium, and (c) the equilibrium concentration. In constructing the table, define x as the concentration (mol/L) of one of the substances that reacts on going to equilibrium, then use the stoichiometry of the reaction to determine the concentrations of the other substances in terms of x.

Step 3) Substitute the equilibrium concentrations into the equilibrium equation for the reaction and solve for x. If you are to solve a quadratic equation choose the mathematical solution that makes chemical sense.

Step 4) Calculate the equilibrium concentrations from the calculated value of x.

Step 5) Check your results by substituting them into the equilibrium equation.

Relationship between equilibrium constant K, reaction quotient Q and gibbs energy G The value of Kc for a reaction does not depend on the rate of the reaction. However, it is directly related to the thermodynamics of the reaction and in particular, to the change in Gibbs energy, $\triangle G$. If,

 $\triangle G$ is negative, then the reaction is spontaneous and proceeds in the forward direction.

 $\triangle G$ is positive, then reaction is considered non-spontaneous. Instead, as reverse reaction would have a negative $\triangle G$, the products of the forward reaction shall be converted to the reactants.

 $\triangle G$ is 0, reaction has achieved equilibrium; at this point, there is no longer any free energy left to drive the reaction. A mathematical expression of this thermodynamic view of equilibrium can be described by the following equation:

$$\triangle G = \triangle G^{\phi} + RTlnQ$$

where, $\triangle G^\phi$ is standard Gibbs energy. At equilibrium, when $\triangle G=0$ and Q = Kc, the equation becomes,

$$riangle \mathrm{G} = \mathrm{G}^\phi + \mathrm{RTlnk} = 0$$

$$\triangle \mathbf{G}^{\phi} = -\mathbf{R}\mathbf{T}\mathbf{l}\mathbf{n}\mathbf{k}$$

$$\mathrm{Ink} = rac{-igtriangle \mathrm{G}^\phi}{\mathrm{RT}}$$

Taking antilog of both sides, we get,

$$K = e - \frac{\triangle G0}{RT}$$

Hence, using the equation, the reaction spontaneity can be interpreted in terms of the value of $\triangle G^{\phi}$.

If $\triangle G^{\phi}>0$ then $\frac{-\triangle G^{\phi}}{RT}$ is positive, and >1, making K > 1, which implies a spontaneous reaction or the reaction which proceeds in the forward direction to such an extent that the products are present predominantly.

If $\triangle G^{\phi}>0$, then $\frac{-\triangle G^{\phi}}{RT}$ is negative, and < 1, that is, K < 1, which implies a non-spontaneous reaction or a reaction which proceeds in the forward direction to such a small degree that only a very minute quantity of product is formed.

Factors affecting equilibria One of the principal goals of chemical synthesis is to maximise the conversion of the reactants to products while minimizing the expenditure of energy. This implies maximum yield of products at mild temperature and pressure conditions. If it does not happen, then the experimental conditions need to be adjusted. For example, in the Haber process for the synthesis of ammonia from N₂ and H₂, the choice of experimental conditions is of real economic importance. Annual world production of ammonia is about hundred million tones, primarily for use as fertilizers. Equilibrium constant, Kc is independent of initial concentrations. But if a system at equilibrium is subjected to a change in the concentration of one or more of the reacting substances, then the system is no longer at equilibrium; and net reaction takes place in some direction until the system returns to equilibrium once again. Similarly, a change in temperature or pressure of the system may also alter the equilibrium. In order to decide what course the reaction adopts and make a qualitative prediction about the effect of a change in conditions on equilibrium we use Le Chatelier's principle. It states that a change in any of the factors that determine the equilibrium conditions of a system will cause the system to change in such a manner so as to reduce or to counteract the effect of the change. This is applicable to all physical and chemical equilibria.

Effect of Concentration Change In general, when equilibrium is disturbed by the addition/removal of any reactant/ products, Le Chatelier's principle predicts that: The concentration stress of an added reactant/product is relieved by net reaction in the direction that consumes the added substance.

The concentration stress of a removed reactant/product is relieved by net reaction in the direction that replenishes the removed substance. or in other words, "When the concentration of any of the reactants or products in a reaction at equilibrium is changed, the composition of the equilibrium mixture changes so as to minimize the effect of concentration changes". Let us take the reaction,

$$H_{2(g)} + l_{2(g)} \rightleftharpoons 2Hl_{(g)}$$

If H_2 is added to the reaction mixture at equilibrium, then the equilibrium of the reaction is disturbed. In order to restore it, the reaction proceeds in a direction wherein H_2 is consumed, i.e., more of H_2 and I_2 react to form HI and finally the equilibrium shifts in right (forward) direction. This is in accordance with the Le Chatelier's principle which implies that in case of addition of a reactant/product, a new equilibrium will be set up in which the concentration of the reactant/product should be less than what it was after the addition but more than what it was in the original mixture. The same point can be explained in terms of the reaction quotient, Qc,

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$$Qc = \frac{[HI]^2}{[H]_2[I]_2}$$

Addition of hydrogen at equilibrium results in value of Qc being less than Kc. Thus, in order to attain equilibrium again reaction moves in the forward direction. Similarly, we can say that removal of a product also boosts the forward reaction and increases the concentration of the products and this has great commercial application in cases of reactions, where the product is a gas or a volatile substance. In case of manufacture of ammonia, ammonia is liquified and removed from the reaction mixture so that reaction keeps moving in forward direction. Similarly, in the large scale production of CaO (used as important building material) from CaCO₃, constant removal of CO₂ from the kiln drives the reaction to completion. It should be remembered that continuous removal of a product maintains Qc at a value less than Kc and reaction continues to move in the forward direction.

- i. If ... the reaction will proceed in the direction of reactants (reverse reaction).
 - a. Qc > Kc
 - b. Qc < Kc
 - c. Qc = Kc
 - d. None of above
- ii. If ... the reaction will proceed in the direction of the products (forward reaction).
 - a. Qc > Kc
 - b. Qc < Kc
 - c. Qc = Kc
 - d. None of above
- iii. If ... the reaction mixture is already at equilibrium. Consider the gaseous reaction.
 - a. Qc > Kc
 - b. Qc < Kc
 - c. Qc = Kc
 - d. All of above
- iv. If $\triangle G$ is then the reaction is spontaneous and proceeds in the forward direction.
 - a. Zero
 - b. Positive
 - c. Negative
 - d. None of above
- v. $\triangle G$ is ... reaction has achieved equilibrium; at this point, there is no longer any free energy left to drive the reaction.
 - a. Zero
 - b. Positive
 - c. Negative
 - d. None of above
- 151. Read the passage given below and answer the following questions from (i) to (v).

When a liquid evaporates in a closed container, molecules with relatively higher kinetic energy escape the liquid surface into the vapour phase and number of liquid molecules from the vapour phase strike the liquid surface and are retained in the liquid phase. It gives rise to a constant vapour pressure because of an equilibrium in which the number of molecules leaving the liquid equals the number returning to liquid from the vapour. We say that the system has reached equilibrium state at this stage. However, this is not static equilibrium and there is a lot of activity at the boundary between the liquid and the vapour. Thus, at equilibrium, the rate of evaporation is equal to the rate of condensation. It may be represented by

$$H_2O_{(l)} \rightleftharpoons H_2O_{(vap)}$$

The double half arrows indicate that the processes in both the directions are going on simultaneously. The mixture of reactants and products in the equilibrium state is called an equilibrium mixture.

Equilibrium can be established for both physical processes and chemical reactions. The reaction may be fast or slow depending on the experimental conditions and the nature of the reactants. When the reactants in a closed vessel at a particular temperature react to give products, the concentrations of the reactants keep on decreasing, while those of products keep on increasing for some time after which there is no change in the concentrations of either of the reactants or products. This stage of the system is the dynamic equilibrium

The chemical equilibrium may be classified in three groups.

- 1. The reactions that proceed nearly to completion and only negligible concentrations of the reactants are left. In some cases, it may not be even possible to detect these experimentally.
- 2. The reactions in which only small amounts of products are formed and most of the reactants remain unchanged at equilibrium stage.
- 3. The reactions in which the concentrations of the reactants and products are comparable, when the system is in equilibrium.

The equilibrium involving ions in aqueous solutions which is called as ionic equilibrium. Solid-Liquid Equilibrium Ice and water kept in a perfectly insulated thermos flask (no exchange of heat between its contents and the surroundings) at 273K and the atmospheric pressure are in equilibrium state and the system shows interesting characteristic features. We observe that the mass of ice and water do not change with time and the temperature remains constant. However, the equilibrium is not static. The intense activity can be noticed at the boundary between ice and water. Molecules from the liquid water collide against ice and adhere to it and some molecules of ice escape into liquid phase. There is no change of mass of ice and water, as the rates of transfer of molecules from ice into water and of reverse transfer from water into ice are equal at atmospheric pressure and 273 K. It is obvious that ice and water are in equilibrium only at particular temperature and pressure. For any pure substance at atmospheric pressure, the temperature at which the solid and liquid phases are at equilibrium is called the normal melting point or normal freezing point of the substance. The system here is in dynamic equilibrium and we can infer the following:

- 1. Both the opposing processes occur simultaneously.
- 2. Both the processes occur at the same rate so that the amount of ice and water remains constant.

Solid – Vapour Equilibrium Let us now consider the systems where solids sublime to vapour phase. If we place solid iodine in a closed vessel, after sometime the vessel gets filled up with violet vapour and the intensity of colour increases with time. After certain time the intensity of colour becomes constant and at this stage equilibrium is attained. Hence solid iodine sublimes to give iodine vapour and the iodine vapour condenses to give solid iodine. The equilibrium can be represented as,

$$l_2(solid) \rightleftharpoons l_2(vapour)$$

Other examples showing this kind of equilibrium are,

$$Camphor_{(solid)} \rightleftharpoons Camphor_{(vapour)}$$

$$NH_4CI_{(solid)} \rightleftharpoons NH_4CI_{(vapour)}$$

The equilibrium Involving Dissolution of Solid in Liquids Only a limited amount of salt or sugar can dissolves in a given amount of water at room temperature. If we make a thick sugar syrup solution by dissolving sugar at a higher temperature, sugar crystals separate out if we cool the syrup to the room temperature. We call it a saturated

solution when no more of solute can be dissolved in it at a given temperature. The concentration of the solute in a saturated solution depends upon the temperature. In a saturated solution, a dynamic equilibrium exits between the solute molecules in the solid state and in the solution: Sugar (solution) Sugar (solid), and the rate of dissolution of sugar = rate of crystallisation of sugar. Equality of the two rates and dynamic nature of equilibrium has been confirmed with the help of radioactive sugar. If we drop some radioactive sugar into saturated solution of non-radioactive sugar, then after some time radioactivity is observed both in the solution and in the solid sugar. Initially there were no radioactive sugar molecules in the solution but due to dynamic nature of equilibrium, there is exchange between the radioactive and non-radioactive sugar molecules between the two phases. The ratio of the radioactive to non- radioactive molecules in the solution increases till it attains a constant value.

- i. Which of the following symbol represents equilibrium.
 - a. \rightleftharpoons

 - **c.** ⇔
 - d. ↑↓
- ii. When there is no change in the concentrations of either of the reactants or products, this stage of the system is the ...
 - a. Static equilibrium
 - b. Dynamic equilibrium
 - c. Physical equilibrium
 - d. Chemical equilibrium
- iii. A ... solution means no more of solute can be dissolved in it at a given temperature.
 - a. Unsaturated
 - b. Supersaturated
 - c. Saturated
 - d. None of these.
- iv. The equilibrium involving ions in aqueous solutions which is called as ...
 - a. Static equilibrium
 - b. Dynamic equilibrium
 - c. Physical equilibrium
 - d. Ionic equilibrium
- v. The concentration of the solute in a saturated solution depends upon the ...
 - a. Solvent
 - b. Pressure
 - c. Temperature
 - d. System

---- Stay away from those people who try to disparage your ambitions. ... -----