# UNIT: 6 SOLUTIONS

## **Important Points**

When two or more than two substance mix and from a uniform or homogeneous mixture, Such a mixture is called solution.

### Type of solutions

The solutions can be found in three states; Solid, Liquid and Gas. The solute and solvent can also be in three states. The physical state of the resulting solution can be decided on the basis of physical state of solute and solvent.

Sr No.	Type of solution	Physical state		Examples
		Solute	Solvent	
1	Solid solution	Solid	Solid	Alloy formed from copper and zinc (Brass).
		Liquid	Solid	Zinc amalgam-Zinc
		Gas	Solid	dissolved in mercury (Zn/Hg adsorption of $H_2$ gas on Pd.
2	Liquid solution	Solid	Liquid	Homogeneous mixture of
2	Liquid solution	Solid	Liquid	sugar and water.
		Liquid S	Liquid	Homogeneous mixture of
				water and ethanol.
		Gas	Liquid	Homogeneous mixture of
				$CO_2$ gas in water.
3	Gaseous solution	Solid	Gas	Homogeneous mixture of
		Liquid	Gas	camphor in $N_2$ gas. Air containing moisture
		Gas	Gas	Mixture of $H_2$ and $O_2$ gas.

$$Formality(F) = \frac{1000 \times mass of solute(gram)}{Formula mass of solute \times volume of solution(ml)}$$

 $% V/V = \frac{100 \times \text{volume of solute}}{\text{volume of solute + volume of solvent}} = \frac{100 \times \text{volume of solute(ml)}}{\text{volume of solvent(ml)}}$  $% W/V = \frac{100 \times \text{mass of solute(gram)}}{\text{volume of solution(ml)}}$ 

parts per million by mass to volume =  $\frac{\text{amount of solute(mg)}}{\text{amount of solution(litre)}}$ 

Factors which effect the solubility of gaseous solute in liquid solution formed by homogeneous Mixture of gaseous solute and are given as under.

Nature of gaseous solute and the solvent (ii) Effect of temperature (iii) Effect of pressure (i)

Henry's law :  $p = K_H X$  where,  $K_H$  is Henry's constant.

When solid solute is dissolved in solid solvent is gives solid Solution. The molecules are arranged in two ways:

#### Substituted solid solution (2) Interstitial solid solution (1)

#### **Solution-Colligative Properties**

When solute substances are dissolved in pure solvent, the solutions are obtained. Some properties of solvent change viz. the vapour pressure of a solution prepared from a solvent is less than that of pure solvent, while the boiling point increases and freezing point decreases. The osmotic pressure also changes. The change in these properties depend in number of molecules of solute but not on nature of solute. Such properties are called colligative properties of solution.

#### Raoult's Law (For Non-volatile Solute)

"If dilute and ideal solution is prepared by dissolving non-volatile solute in a volatile solvent, the relative lowering of vapour pressure of the solution is equal to the mole fraction of the dissolved solute." shikshe

$$\frac{\frac{P_1^0 - P1}{p_1^0}}{\frac{1}{p_1^0}} = \frac{n_2}{n_1 + n_2^0}$$

i.e. mole fraction of solute. Where, n, and n, are the moles of solvent and solute respectively. |For Very dilute solution  $n_2 \ll n_1$  hence putting  $n_1 + n_2$  in equation

$$\frac{P_1^0 - P1}{P1} = \frac{n_2}{n_1} \text{ but } = n_2 = \frac{w_1}{M_2} \text{ and } n_1 = \frac{w_1}{M_1}, \text{ putting this values in equation}$$

$$\frac{P_1^0 - P1}{P1} = \frac{W_2 \times M_1}{M_2 \times W_1}$$
 where,  $W_1 = \text{mass of solvent}$ ,  $M_1 = \text{molecular mass of solvent}$ 

 $W_2$  = mass of solute,  $M_1$  = molecular mass of solute

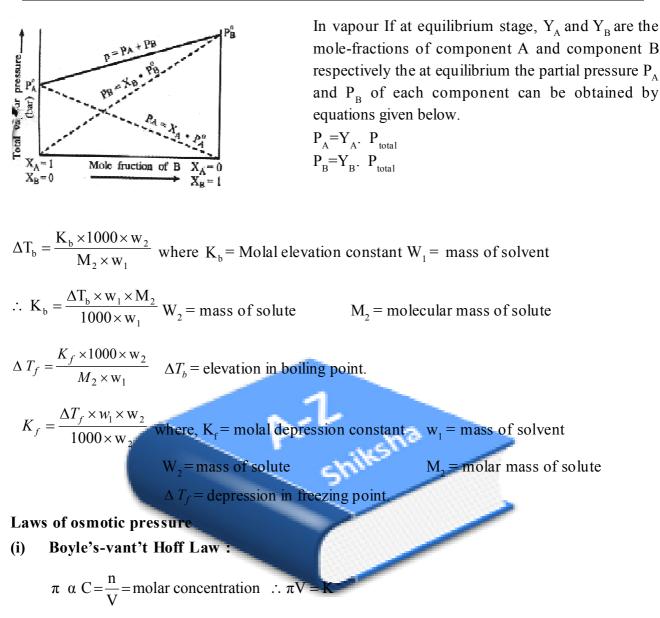
#### Raoult's Law (For Volatile Solute and Volatile Solvent)

Suppose in a binary solution X<sub>A</sub> is the mole-fraction of solute A and its partial pressure is p and XB is the mole fraction of the solvent B and its partial pressure is PB then according to Raoult law  $P_A \alpha X_A$  and  $P_B \alpha X_B$ .

According to experimental observations Raoul's law can be proved that if the vapour pressure of solute is  $P_A^0$  and vapour pressure of pure solvent is  $P_B^0$  then,

 $P = p_A^0 X_A$  and  $p_B^0 X_B$ 

Total Pressure  $P = p_A + p_B$  Total pressure  $p = p_A^0 X_A + p_B^0 X_B$ 



(ii) Gay-Lussac van't Hoff Law :

 $\pi = KT$  where, K = proportionality conctant

#### (iii) Avogadro's-vant't-Hoff law

$$\pi \alpha n$$
  $\pi = \frac{nRT}{V}$  where, R is proportionality constant and its value is equation

that of gas Constant

 $\pi \alpha T$  (Boyle-van't-Hoff Law).  $\therefore \pi V = nRT$ 

 $\pi \alpha T$  (Gay-Lussac-van't-Hoff Law) where  $\pi$  = Osmotic pressure in bar

V = volume of solution in life

n = number of moles of solute

R = gas constant

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T = absolute temperature in Kelvin.

Then, solution whose osmotic pressure is less it is known as hypotonic with comparison to higher pressure solution.

A solution whose osmotic pressure is higher, in comparison to solution having less osmotic pressure is called hypertonic solution.

Less than 0.91% W/V pure NaCl solution is hypotonic compared to fluid inside human blood NaCl solution having concentration more than 0.91% W/V is hypertonic with respect to fluid inside human body.

Van't Hoff factor (i) 
$$= \frac{\text{Normal molar mass of solute}}{\text{Abnormal moral mass of solute}}$$
$$= \frac{\text{Theorical molar mass of solute}}{\text{Experimental moral mass of solute}}$$
$$= \frac{\text{Observed colligative property}}{\text{Colligative property}}$$

Theoritical collingative property

By introducing van't Hoff factor (i) the formula to obtain molar mass can be written as follows

Raoult's law : 
$$\frac{p_1^0 - p_1}{p_1^0} = i \frac{n_2}{n_1 + n_2} \approx i \frac{n_2}{n_1}$$
  
Molal elevation :  $\Delta T_b = i K_b , m$   
Molal depression :  $\Delta T_f = i K_f , m$   
Osmotic pressure of solution :  $\pi = i \frac{nRT}{V}$   
Molality =  $\frac{1000 \times Molarity}{(1000 \times density) - (molar mass of solute \times molarity)}$   
M.V =  $M_1 \cdot V_1 + M_2 \cdot V_2 + M_3 \cdot V_3$ ,  $V = V_1 + V_2 + V_3$   
 $K_a = Molarity \times \left(\frac{a^2}{1-\alpha}\right) \alpha = \frac{i-1}{n-1}$   $i = \frac{\Delta T_{ob}}{\Delta T_{cal}} = \frac{\pi_{ob}}{\pi_{cal}} = \frac{M}{M_{ob}} = \text{Theoretical molar mass} \left[\frac{\Delta Tf}{\Delta Tb} = \frac{Kf}{Kb}\right]$   
Degree of association (X) = (1-i)  $\frac{n}{n-1}$   
Degree of dissociation ( $\alpha$ ) =  $\frac{i-1}{n-1}$   
 $I = \frac{(i) p = p_1^0 \cdot X + p_B^0 \cdot X_B}{VR = QY \cdot (VW - QY)QU}$   
 $(ii) p = p_A^0 + (p_B^0 - p_A^0) \cdot X_B$   
 $UR = QY + (VW - QY)QU$   
 $(iii) p = p_B^0 + (p_A^0 - p_B^0) \cdot X_A$   
 $UR = VW + [(QY - VW)VU]$ 

	М.С.	Q.			
1.	Which of the following possesses physical s respectively ?	tate of solute and solven	t are liquid and solid		
	(a) solution of sugar in water	(b) zinc amalgam			
	(c) solution of Naphthelene in benzene	(d) brass			
2.	Which of the following alternative is correct for of camphor in $N_2$ gas ?	physical state of solute and	d solvent for a solution		
	(a) solid, gas (b) solid, liquid	(c) gas, solid	(d) gas, gas		
3.	Which of the following is an example of solid	solution ?			
	(a) interstitial compound of hydrogen and Pd	(b) WC			
	(c) Zn / Hg	(d) Given all			
4.	Which of the following pair of solution having	same physical state of so	lute ?		
	(a) homogenous mixture of chloroform in $N_2$ g	gas, solution of $CO_2$ in wa	ter		
	(b) brass, sodium amalgam				
	(c) homogenous mixture of camphor in $N_2$ gas, solution of $H_2$ gas in Pd metal				
	(d) moist air, solution of ethanol in water				
5.	Which of the following pair of solution having				
	(a) homogenous mixture of chloroform in N <sub>2</sub> §	gas, solution of $CO_2$ in wa	ter		
	(b) brass, solution of salt in water				
	(c) homogenous mixture of camphor in $N_2$ gas				
	(d) solution of $H_2$ gas in Pd metal, solution of				
6.	Which of the following pair of solution having	different physical state of	solute ?		
	(a) homogenous mixture of chloroform in $N_2$ g	gas, solution of $CO_2$ in wa	ter		
	(b) brass, homogenous mixture of camphor in	N <sub>2</sub> gas			
	(c) sodium amalgam, moist air				
	(d) solution of $H_2$ gas in Pd metal mixture of	$N_2$ and $O_2$			
7.	Which of the following pair of solution having	different physical state of	solvent?		
	(a) homogenous mixture of $N_2$ gas in chlorofo	orm, mixture of $N_2$ and $O_2$	1		
	(b) brass, solution of $H_2$ gas in Pd metal				
	(c) sodium amalgam, solution of urea in water				
	(d) solution of $CO_2$ in water, solution of Naph	thelene in benzene			
8.	Which of the following compound possesses m	aximum solubility in wate	er?		
	(a) pentan $-1$ - ol	(b) Pentane – 2, 3 – di	ol		
	(c) pentane - 1, 2, 3 - triol	(d) Pentane – 1, 2, 3, 4	4 – tetraol		

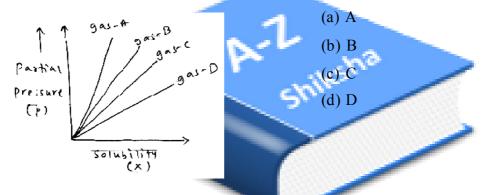
9.	Which of the followin	g given compound is lea	ast soluble in water?	
	(a) Hexan – 1 – ol		(b) glycerol	
	(c) Propane – 1, 3 –	diol	(d) ethylene glycol	
10.	To convert molarity in of the solution ?	to which of the followir	ng unit of concentration, de	oes not require density
	(a) molality	(b) normality	(c) mole-fraction	(d) % w/w
11.	To convert molality in solution ?	nto which of the follow	ing unit of concentration	require density of the
	(a) percentage weight	by weight	(b) percentage by volum	me
	(c) mole-fraction		(d) given all	
12.	Which of the followin	g unit of concentration	does not depend on temp	erature ?
	(a) formality	(b) molarity	(c) molality	(d) normality
13.	Which of the followin	g unit of concentration	depends on temperature?	
	(a) molality	(b) normality	(c) mole-fraction	(d) given all
14.		mality of solution prepa W of Potash-alum = 94	red by dissolving 9.48 gm. 48 gm/mole]	Potash-alum dissolved
	(a) 0.04 F	(b) 0.02 F	(c) 0.002 F	(d) 0.004 F
15.	What quantity of pota	sh-alum is required to p	prepare 500ml solution ha	ving strength 1.5 F
	(a) 711gm	(b) 355.5 gm	(c) 35.55 gm	(d) 71.1 gm
16.	What would be the c dissolved in 4 lt. Solu		of aqueous solution in w	hich 80 ml ethanol is
	(a) 4% v/v	(b) 10% v/v	(c) 2% v/v	(d) 8% v/v
17.	What would be the ve Solution ?	olume of acetone requi	red to prepare 10% v/v ac	cetone solution in 5 lt.
	(a) 50 ml	(b) 5 ml	(c) 100 ml	(d) 500 ml
18.	15 % w/v solution of s be the volume of the s	0 1 1 5	olving 1200 gm sugar in w	vater, then, what would
	(a) 8000 ml	(b) 4 lt	(c) 800 ml	(d) 5000 ml
19.	0.004 gm $O_2$ is dissolution by weight-vol		tion of 50 litre, then, wha	t would be the ppm of
	(a) 0.04	(b) 0.008	(c) 0.004	(d) 0.08
20.	The concentration of	Ca <sup>2+</sup> ion in a sample of	of water is 0.0002 M the me? (Atomic wt. of ca =	n what would be the
	(a) 4	(b) 8	(c) 0.08	(d) 0.4

21.	The concentration of solution in % w/v is	F-ion in a sample of	water is 10 ppm; then, c	oncentration of F <sup>-</sup> ion in a
	(a) 10 <sup>-3</sup>	(b) 10 <sup>-2</sup>	(c) 10	(d) $10^{-4}$
22.	What would be the w strength 5 ppm ?	weight of $O_2$ gas in gran	n dissolved in an aqueor	as solution of 500ml having
	(a) 0.025	(b) $2.5 \times 10^{-4}$	(c) $2.8 \times 10^{-3}$	(d) $2.8 \times 10^{-5}$
23.		nolarity of solution pr 1 1.2 M aqueous solu		ture of 1400 ml 0.3 M, 700
	(a) 0.5 M	(b) 0.8 M	(c) 0.6 M	(d) 0.7 M
24.	What quantity of KC ?	OH is required to prepa	re 10 % w/w KOH solut	ion having weight 1000 gm
	(a) 50 gm	(b) 25 gm	(c) 100 gm	(d) 150 gm
25.	What amount of wat to prepare 0.5 M sol	-	ous solution of 5000 ml l	naving concentration 1.5 M
	(a) 15 litre	(b) 5 litre	(c) 10 litre	(d) 20 litre
26.	On Which factors, th	e solubility of gaseous	solute in liquid depend	s ?
	(a) temperature	(b) Pressure of the	e gas	
	(c) Nature of gaseou	s solute and solvent	(d) Given all	/
27.		, if partial pressure of a um solubility in water		hen, which of the following
	(a) He	(b) N <sub>2</sub>	(c) H <sub>2</sub>	(d) O <sub>2</sub>
28.	At 298 K temperature gas possesses least s		ll given gases are same, t	hen, which of the following
	(a) carbon dioxide	(b) formaldehyde	(c) methane	(d) vinyl chloride
29.	is the correct ascend (a) Ar < HCHO < C (b) Ar < $CH_2 = CH$ (c) Ar < $CH_4 < HC$	c, if partial pressure of a ing order of solubility $CH_4 < CH_2 = CH - C$ $- Cl < CH_4 < HCH$ $HO < CH_2 = CH - C$ $CH_2 = CH - C$	of gases in water ? 11 10 Cl	hen, which of the following
30.	At 293 K temperature is the correct descen (a) $H_2 > N_2 > O_2 >$	2	ll given gases are same, t y of gases in water ? > He	hen, which of the following
31.	2 2 2			n gas possesses higher value
	(a) He	(b) N <sub>2</sub>	(c) H <sub>2</sub>	(d) O <sub>2</sub>

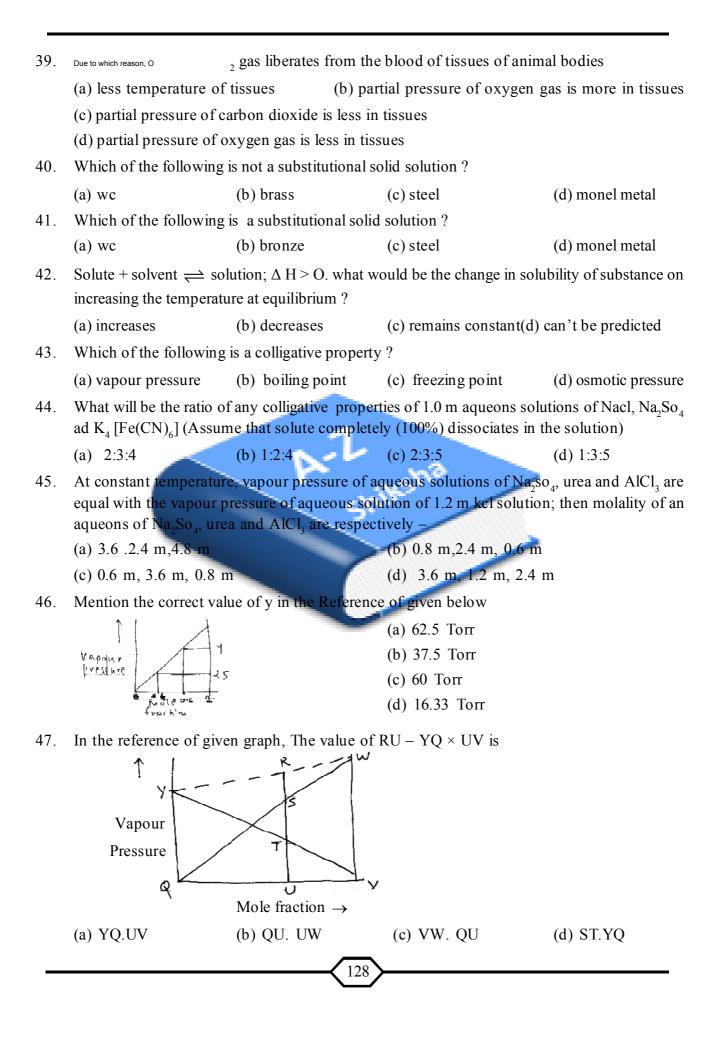
- 32. At 293 K temperature, for solubility of all given gases, in water, which gas possesses lower value of KH ?
  - (a) Carbon dioxide (b) formaldehyde (c) methane
- (d) vinyl chloride
- 33. At 298 K temperature, for solubility of all given gases, which of the following is the correct ascending order of values of Henry's constant ?

(a) 
$$CH_4 < Ar < HCHO < CH_2 = CH - Cl$$

- (b) HCHO <  $CH_2 = CH Cl < CH_4 < Ar$
- (c) HCHO <  $CH_4 < CH_2 = CH Cl < Ar$
- (d)  $CH_2 = CH Cl < HCHO < CH_4 < Ar$
- 34. At 293 K temperature, for solubility of all given gases, which of the following is the correct descending order of values of Henry's constant ?
  - (a)  $H_2 > N_2 > O_2 > He$  (b)  $N_2 > H_2 > O_2 > He$
  - (c) He >  $N_2$  >  $H_2$  >  $O_2$  (d)  $O_2$  >  $H_2$  >  $N_2$  > He
- 35. At constant temperature, on the basis of the given graph, which gas possesses higher solubility?



- 36. In which of the following specific condition,  $CO_2$  gas is filled in cold drinks, and in soda water?
  - (a) at high temperature and high pressure
- (b) at low temperature and high pressure
- (c) at low temperature and low pressure (d) at high temperature and low pressure
- 37. In which condition, Henry's law is applicable?
  - (a) ideal behaviour of gaseous solute at high pressure and low temperature
  - (b) gaseous solute neither associate nor dissociate in solution
  - (c) gaseous solute react with solvent
  - (d) applicable in given all conditions
- 38. Now a days, divers uses the cylinder having gaseous mixture contains -
  - (a) 2 %  $O_2$  and 98 % He (b) 11.7 % He, 56.2 %  $N_2$  and 32.1 %  $O_2$
  - (c) 11.7 %  $N_2$ , 56.2% He and 32.1%  $O_2$  (d) 11.7 % He, 56.2 %  $O_2$  and 32.1 %  $N_2$



48.	Ionic substances are co solutions possesses hig		the given solutions, then	which of the following
	(a) 0.01m Urea		(b) 0.01m NaCl	
	(c) 0.01m BaCl <sub>2</sub>		(d) $0.01 \text{ m Al}_2(\text{SO}_4)_3$	
49.	•	4 and its partial vapour	B, at constant temperature, pressure is 400 mm, then	-
	(a) 600 mm	(b) 300 mm	(c) 500 mm	(d) 200 mm
50.	-		poils at 100.55 °C ; then, g - mole <sup>-1</sup> and Kb = $1.8$	
	(a) 272 K	(b) 271 K	(c) 375 K	(d) 274 K
51.	Ionic substances are c substances having leas		hen aqueous solution of	which of the following
	(a) glucose	(b) NaCl	(c) $Al_2(SO_4)_3$	(d) $CaCl_2$
52.	0.2 M aqueous solutio	n of NH <sub>4</sub> Cl is isotonic v	vith which of the followin	ng aqueous solution ?
	(a) 0.1 M Na <sub>3</sub> PO <sub>4</sub>	(b) 0.2 M $K_2SO_4$	(c) 0.1 M $Al_2(SO_4)_3$	(d) none of thses
53.	At constant temperature $Al_2(SO_4)_3$ are equal, the dissociates in the solution	$n$ mention the value of $\Sigma$	f an aqueous solution of f X. (Assume that ionic solid	1.5 M NH <sub>4</sub> NO <sub>3</sub> and <i>x</i> N substances completely
	(a) 0.1	(b) 3.6	(c) 1.2	(d) 0.6
54.	$Al_2(SO_4)_3$ are equal; the		an aqueous solution of 1 y of an aqueous solution tes in the solution)	. 5
	(a) 0.3 m	(b) 2.1 m	(c) 3.75 m	(d) 0.6 m
55.	water at 1 atmosphere		d by dissolving 1.5 mole n which of the following a	
	(a) i = 1	(b) 1 < i < 2	(c) $i < 1$	(d) i > 2
56.		g aqueous solution is isonces completely dissocia	otonic with 0.2 m Na <sub>4</sub> [Collites in the solution)	$F_6$ ] solution ? (Assume
	(a) 0.2 m urea	(b) 0.25 m AlCl <sub>3</sub>	(c) 0.15m CaCl <sub>2</sub>	(d) 0.2 m CuSo <sub>4</sub>
57.	Which type of solution	n, moist air is ?		
	(a) gas	(b) liquid	(c) solid	(d) colloidal
58.	-	num osmotic pressure ?	meable membrane. For w (Assume that ionic solid	
	(a) 0.5 m NaCl   0.1	m Na <sub>2</sub> SO <sub>4</sub>	(b) 0.3 m NaCl   0.1 n	m Na <sub>2</sub> SO <sub>4</sub>
	(c) 0.5 m NaCl   0.1	m FeCl <sub>3</sub>	(d) 0.5 m NaCl   0.1 n	n sugar
		129	<b>}</b>	

- 59. At constant temperature, the vapour pressure of an aqueous solution of Na, SO<sub>4</sub> and 0.3 m Na<sub>3</sub>PO<sub>4</sub> are approximately equal; then, what would be the molality of an aqueous solution of  $Na_2SO_4$ ? (Assume that ionic solid substances completely dissociates in the solution) (a) 0.5 m (b) 0.6 m (c) 0.4 m(d) 1.2 m What will be the ratio of elevation in boiling point of aqueous solution of 1 m sugar, 1 C<sub>s</sub>Cl, and 60. 1 m Na<sub>2</sub>SO<sub>4</sub>? (Assume that ionic solid substances completely dissociates in the solution) (a) 3 : 2 : 1 (c) 1 : 2 : 3 (d) 3 : 1 : 2 (b) 1 : 3 : 2 61. At constant temperature, solubility of which of the following substances decreases with increase in temperature? (a) aqueous solution of sugar (b) aqueous solution of salt (c) aqueous solution of  $CO_2$ (d) aqueous solution of KNO<sub>3</sub> At constant temperature, in a closed vessel, an ideal solution is formed by liquid - A and liquid 62. - B; and mole-fraction of A and B are 0.6 and 0.4 respectively. If vapour pressure of pure liquids are 125.0 and 62.5 mm respectively, then their mole-fraction in vapour state are respectively – (In vessel, no other component is in gaseous form) (a) 0.6 and 0.4 (b) 0.4 and 0.6 (c) 0.25 and 0.75 (d) 0.75 and 0.25 At constant temperature, two liquids having osmotic pressure  $\pi_1$  and  $\pi_2$  are seperated by 63. semipermeable membrane, then, what will be the osmotic pressure of the system ? (d)  $\frac{\pi_1 - \pi_2}{2}$ (a)  $\pi_1 + \pi_2$ Which of the following pair of solutions forms ideal solution? 64. (a) Chloro benzene, chloro ethane (b) benzene-toluene (d) water, HCL (c) acetone-chloroform Which of the following pair forms true solutions? 65. (a) Hexane, heptane (b) chloro benzen, bromo benzen (c) chloro ethane, bromo ethane (d) phenol, aniline 66. What state does point s indicate ?
  - (a) Mole-fraction and partial vapour pressure of both the liquids are same
  - (b) Mole-fraction of the both the liquids are same, but their partial vapour pressures are different

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(c) Mole-fraction and partial vapour pressures of both the liquids are different

Mole-Incion

(d) Mole-fraction of both the liquids are different, but their partial pressures are same.

67. What would be the elevation in boiling point of 0.1 m NaCl solution ? (Assume that Nacl dissociates completely)

(a) kb/10 (b) 10 kb (c) kb/5 (d) k b/20

68. Which of the following semipermeable membrane is best one?

(a) parchment paper (b) copper frocyanide

- (c) butter paper (d) cello phane
- 69. At constant temperature, binary ideal solution is formed by two liquids A and B. At equilibrium, mole-fraction of liquid A is 0.7 and in vapour state mole-fraction of A is 0.4 Poa + Pob =90 mm then at the same temperature, what will be the vapour pressure of pure liquid A and B?

(a) 40 mm, 50 mm (b) 30 mm, 60 mm (c) 50 mm, 40 mm (d) 20 mm, 70 mm

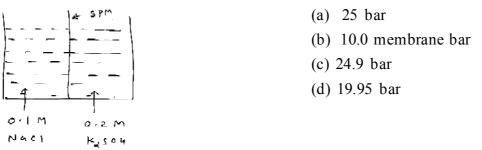
- 70. At constant temperature, binary ideal solution is formed by two liquids A and B. At equilibrium, mole-fraction of liquid B is 0.4 and vapour state mole-fraction of B is 0.25. P°B=40 mm, then at the same temperature, what will be the vapour pressure of pure liquid 'A'?
  (a) 80 mm
  (b) 60 mm
  (c) 40 mm
  (d) 50 mm
- 71. Choose correct alternative for True and False statements for given diagram. (For correct statement T and for wrong statement F) (Assume that ionic solid substances completely dissociates in the solution) (For Correct- Stutement T and for Wrory worry Srut ment F)



- (i) Osmotic pressure of the system increase by adding  $H_2O$  in an aqueous solution of  $C_{\mu}SO_4$
- (ii) The concentration of solution of glucose increases with the passage of time.
- (iii) Osmotic pressure of the system decreases by adding glucose in the solution of glucose.
- (iv) The concentration of solution of  $C_{\mu}SO_{4}$  increases with the passage of time.
- (a) FTTF (b) TFFT (c) FFTT (d) TTFF
- 72. Choose correct alternative for True and False statements for given figure (For correct statement T and for wrong statement F) (Assume that lomic solid substances completely dissociates in the solution)

	emipermable 		concentration of an aqueous sol with the pressure of time.	ution of FeCl <sub>3</sub> increases
		(ii)	aqueous solution of FeCl <sub>3</sub> gradu	ally turns reddish
	KCNS	(iii) concentration of an aqueous solution of KCNS increases with the passage of time		
(χειτοω) (:	ole urless?	(iv)	aqueous solution of KCNS rem	ains colourless.
(a) FTTF		(b) TFFT	(c) FFTT	(d) TTFF

73. What would be the osmotic pressure of the system at 300 k temperature ? (R=8.314 x 10-2 litre-bar-mole-1 k-1) (Assume that lomic solid substances completely dissociates in an aqueous solution)



74. At constant temperature, 2 litres aqueous solution of each 0.2 M kcl and 0.3 M AlCl<sub>3</sub> are in contact with each other by semipermeable membrane. When osmosis stops, then, what mililitre water diffuses from semipermeable membrane to the other side ? (Assume that ionic solids dissociates completely in the aqueous solution)

(a) 500 ml (b) 600 ml (c) 800 ml (d) 1000 ml

75. Choose the correct alternative for the given diagram for correct and wrong statements. (T is for false statement) (Assume that ionic solid substances dissociates completely in the aqueous solution)

(1) concentration of solution of NaCl increases with the passage of time (ii) concentration of solution of urea decreases with the passage of time (iii) concentration of solution of NaCl decreases with the passage of time (iv) concentration of solution of urea increases with the passage of time

(a) FTTF (b) TFFT (c) FFTT (d) TTFF

76. Which of the following solution is hypotonic with fluids in RBC ?(Assume that ionic solid substances completely dissociates in the solution)

(a) 0.2 M NaCl (b) 0.1 M NaCl (c) 0.18 M NaCl (d) given all

- 77. At constant temperature, Which of the following solution is hypotonic in the comparison with fluids in RBC ? (Assume that ionic solid substances completely dissociates in the solution)
  (a) 0.17 M NaCl
  (b) 0.12 M NaCl
  (c) 0.1 M NaCl
  (d) given all
- (a) 0.17 M NaCl
  (b) 0.12 M NaCl
  (c) 0.1 M NaCl
  (d) given all
  78. X M NaCl is isotonic with fluids present in RBC (Red Blood Corpusceles), then what would be the universe for 2 (Marco Of NaCl = 58.5 gram/march) (A summa that is no activity and the times are the times and the times and the times are the time
  - the value of x ? (M.w. Of NaCl =58.5 gm/mole) (Assume that ionic solid substances completely dissociates in the solution)

(a) 0.15 (b) 0.05 (c) 0.18 (d) 0.78

- 79. Which of the following solution is hypotonic in comparison with the solution of 0.4 M glucose?
  - (a)  $0.1 \text{ M CaCl}_2$  (b) 0.2 M NaCl (c)  $0.15 \text{ M FeCl}_3$  (d) 0.3 M urea

80.	Which of the following		-		cCl solu	tion ?(As	sume
	that ionic solid substan						
	(a) 0.1 M $\operatorname{CaCl}_2$	(b) $0.08 \text{ M FeCl}_{3}$	(c) 0.2 M urea			.12 BaCl	2
81.	Which of the following	g solution is isotonic w	ith fluid of RBC	C ? (For N	VaCl, i=	=2)	
	(a) $5.6\%$ w/v glucose		(b) 2.8% w/v g	glucose			
	(c) 1.5% w/v urea		(d) 0.91% w/v	v urea			
82.	Which of the following	g solution is isotonic w	ith fluid of RB	C ? (For 1	VaCl, 2	=2)	
	(a) 2.02 % w/v glucos	e	(b) 4.02% w/v	glucose			
	(c) 8.0% w/v urea		(d) both b and	l c			
83.	In Which of the follow glucose=180 and urea=6 solution)				-	5	
	(a) 1.2% w/v urea glucose	(b) 0.2 MkCl	(c) 0.09% M	FeCl <sub>3</sub>	(d)	6.0%	w/v
84.	In Which of the following lucose=180 and urea=0 solution)				-	2	
	(a) 0.1% w/v CaCl <sub>2</sub> glucose	(b) 2% w/v urea	(c) 1.0% w/v	FeCl <sub>3</sub>	(d)	5.0%	w/v
85.	FeCl <sub>3</sub> ionizes 80% in th	eir aqueous solution, th	en what will be th	he value o	f Vant	'Hoff fac	tor i ?
	(a) 4	(b) 2.7	(c) 3.4	~	(d) 3	.1	
86.	CaCl, ionices 80% in t	heir aqueons solution of	of 0.2 m CaCl <sub>2</sub> , th	nen, mola	lity of	solution	is -
	(a) 0.48 m	(b) 0.52 m	(c) 0.6 m		(d) 2	.6 m	
87.	A substances associate of Van't hoff factor i?		mer (or bimolecu	ule), then	what w	vill be the	value
	(a) 0.2	(b) 0.4	(c) 0.6		(d) gi	ven all	
88.	A substance associates factor i ?	as trimer in their solution	on; then, what we	ould be th	e value	e of Van't	Hoff
	(a) 0.4		(b) 0.3		(c) 0	.2 (d) 0	.25
89.	A substance associates Hoff factor i ?	as bimolecule in their	solution; then wh	nat would	be the	value of	Van't
	(a) 0.4 d" i < 1	(b) 0.5 d" i < 1	(c) $0 < i < 1$		(d) (	0.6 d" i <	< 1
90.	The solute remains as di the solution is -	mer in the m-molal solu	tion; then elevation	on in boili	ng poin	t irrelevan	t with
	(a) $\frac{\text{mkb}}{2}$	(b) $\frac{3\text{mkb}}{5}$	(c) $\frac{3\text{mkb}}{4}$		(d) <sup><u>r</u></sup>	nkb 3	

- 91. Which of the following ratio is irrelevant formula mass and experimental molecular weight obtained from colligative properties of solution of ionic solid of AB type ?
  - (a) 3 : 2 (b) 5 : 3 (c) 4 : 3 (d) 5 : 2

92. Which of the following ratio of correct molecular wt (formula weight) and experimental molecular weight obtained from colligative properties of solution of ionic solid of AB type is possible ?

(a) 5:3 (b) 4:1 (c) 7:3 (d) 5:2

93. A substance associates as trimer in their solution, then which of the following ratio is irrelevant for real molecular weight and experimental molecular weight obtained from colligative properties of solution ?

94. A substance associates as trimer in their solution, then which of the following alternative is possible for depression in freezing point of m molal solution ?

(a) 
$$\frac{mkf}{2}$$
 (b)  $\frac{mkf}{4}$  (c)  $\frac{mkf}{5}$  (d)  $\frac{mkf}{8}$ 

95. Which of the following unit of concentration is common in the field of pharmacy ?(a) formality(b) molarity(c) molality(d) normality

96. A substance associates as trimer in their solution, then what would be the maximum freezing point of their m molal solution is positive ?

(a) 
$$Tf - \frac{mkf}{2}$$
 (b)  $Tf - \frac{mkf}{3}$  (c)  $Tf - \frac{2mkf}{3}$  (d)  $Tf - 2mkf$ 

97. Boiling point of the 0.2 m aqueons solution of a substance is 100.4 °C; then what would be the freezing point of th solution ? (Kb = 0.513°, K+" = 1.86

(a) -0.372 °C (b) -0.37 °C (c) -1.45 °C (d) -0.5 °C

98. Aqueous solution of 0.5 m H2So4 is more concentrated then 0.5 m H2So4 solution; then what will be the possible density of that solution ?

(a) 
$$1.07 - \frac{9m}{m1}$$
 (b)  $1.06 \ 9m/m1$  (c)  $1.05 \ 9m/m1$  (d)  $1.02 \ 9m/m1$ 

99. Which of the following is irrelevant with the boiling point of an aqueous solution of xm AlCl3 ?

(a) Tb + 3 × kb (b) Tb + 5 × kb (c) Tb 
$$-\frac{7xkb}{2}$$
 (d) Tb  $-\frac{5 \times kb}{2}$ 

100. Which of the following is suitable alternative for density of the solution, when molarity (m) and molality (m) of an aqueous solution of urea is same at fixed temperature ? (molecular wt of urea = 60 gm/mole ?

(a) 
$$1 - \frac{3M}{50}$$
 (b)  $1 - \frac{M}{25}$  (c)  $\frac{50 + 3m}{50}$  (d)  $\frac{25 + 2m}{25}$ 

101.	Choose the correc statement 'F' is us	•	se statement. (For true	statement 'T' and for false	
	(i) solubility of gas in liquid increases with increase in partial pressure of the gas.				
	.,	as in liquid increases wi		-	
		gas in liquid is $K_{\rm H}$ is less	FFF		
		gas in liquid increases, a	s partial pressure of gas	decreases and	
	(a) TTFF	(b) FTTT	(c) TFFF	(d) FFTT	
102.	<b>U</b> 1	aqueous solution of 0.0: tor i ? (Kb = $0.513 \text{ °C}$ -		then, what will be the value	
	(a) 4	(b) 3.4	(c) 2.5	(d) 2.8	
103.		••• ••	-	on of glucose is 104 °C at 1 = 0.513° and K+" = 1.86 °C	
	(a) 2.373 m	(b) 1.05 m	(c) 2.151 m	(d) 1.68 m	
104.	oC; then, state the		ctor i ? (For benzene, K Tf = 5.5 ℃)	prepared in benzene is 75.7 b = $2.65 \text{ OC} - \text{kg} - \text{mo}^{-1}$ ,	
	(a) 1.44	(b) 0.64	(c) 0.83	(d) 0.77	
105.				ion of urea is 100.2372 OC; 3° and K+" = 1.86 °C – kg	
	(a) 59.64	(b) 38.946	(c) 51.65	(d) 40.5	
106.		HCl is prepared by dissoltion ? (Molecular weigh		er. What will be the molarity e)	
	(a) 0.4 M	(b) 0.3 M	(c) 0.8 M	(d) 0.3 M	
107.		molality of the solution ht of NaOH = $40 \text{ gm/m}$		0 gm NaOH in 1.5 kg water	
	(a) 0.5 m	(b) 1.0 m	(c) 0.8 m	(d) 0.4m	
108.	What would be the	e molarity of $3.0 \text{ N H}_2\text{S}$	$O_4$ solution ?		
	(a) 6 M	(b) 1.5 M	(c) 3 M	(d) 1 M	
109.	What quantity of I	NaOH is needed to prep	oare 1.2 m, 800 ml NaO	H solution ?	
	(a) 3.84	(b) 60	(c) 42	(d) 38.4	
110.				$dissolving 19.6 \text{ gm H}_2\text{SO}_4$ to f H <sub>2</sub> SO <sub>4</sub> is 98 gm/mole)	
	(a) 0.5 M, 0.25 N		(b) 0.25 M, 0.125	N	
	(c) 0,25 M, 0.5 N		(d) 0.125 M, 0.25	Ν	
			35		

111.	What amount of $H_2SO_4$ of H2SO4 is 98 gm/mo		tre of 0.5 N $H_2SO_4$ solution	n? (Molecular weight
	(a) 98 gm	(b) 24.5 gm	(c) 49 gm	(d) 73.5 gm
112.	What will be the concen ?	tration of solution prepa	red by dissolving 50 gm gl	ucose in 200 gm water
	(a) 25 % w/w	(b) 20 % w/w	(c) 35 % w/w	(d) 15 % w/w
113.	What quantity of urea	required to prepare 20	% w/w solution having w	eight 150 gm?
	(a) 20 gm	(b) 40 gm	(c) 10 gm	(d) 30 gm
114.	What is the normality of	of an aqueous solution	of $0.5 \text{ Al}_2(\text{SO}_4)_3$ ?	
	(a) 3 N	(b) 1 N	(c) 1.5 N	(d) 2.5 N
115.			lution, prepared by dissolv d ethanol are 18 and 48 g	
	(a) 0.04	(b) 0.004	(c) 0.4	(d) 0.0004
116.	What will be the mole-find 1.8 kg water ? (M.W.		OH respectively, when 260 40)	) gm NaOH dissolved
	(a) 0.96, 0.4	(b) 0.0962, 0.38	(c) 0.0962, .038	(d) 0.962, 0.0038
117.	What would be the most strength 4.5 m?	le-fraction of solute in	an aqueous solution of	a substances heaving
	(a) 0.75	(b) 0.075	(c) 0.45	(d) 0.045
118.	The density of 98% w/v	$W H_2 SO_4$ solution is 1.8	gm/mole then, molarity	of the solution is -
	(a) 20 M	(b) 10 M	(c) 18 M	(d) None of these
119.	Molarity and molality o then, waqht will be the		of $H_2SO_4$ are 1.56 (M) and ?	1.8 (M) respectively;
	(a) 1.835 gm/ml	(b) 1.55 gm/ml	(c) 1.02 gm/ml	(d) 1.725 gm/ml
120.	A solution is prepared respectively then, mole		ole-fraction of A, B and G	C are 0.1, 0.2 and 0.4
	(a) 0.2	(b) 0.1	(c) 0.3	(d) 0.4
121.	The density of 4 M H2S ? (Molecular weight of		(ml then, what will be the n )	nolality of the solution
	(a) 3 M	(b) 3.5 M	(c) 1.2 M	(d) 0.4 M
122.	Molarity of 1.2 N aque	ous solution of AlCl <sub>3</sub> is	s <b>-</b>	
	(a) 3.6 m	(b) 2.4 m	(c) 1.2 m	(d) 0.4 m
123.		• • • •	red using 500 gm of 25 % r weight of NaOH = 40 g	
	(a) 12.74 m	(b) 6.25 m	(c) 9 m	(d) 5 m

124.	24. What will be the molality of solution prepared by taking 25 % w/w NaOH and 15 % w/w NaOH solution ? (Molecular weight of NaOH = 40 gm/mole)					
	(a) 12.74 m	(b) 5.5 m	(c) 9 m	(d) 4 m	l	
125.	The density of 2.5 M Na correct for molarity and	-	n/ml; then, which of the f	ollowing a	alternativ	e is
	(a) M > m predicted	(b) M < m	(c) M = m	(d) (	can't	be
126.	Which of the following	is correct for an ideal s	olution ?			
	(a) " $H = 0$ , " $V = 0$ , " $S$	S = 0	(b) "H " $0$ , "V = $0$ , "S	S = 0		
	(c) " $H = 0$ , " $V = 0$ , " $S$	· `` 0	(d) " $H = 0$ , "V " 0, "S	S = 0		
127.			BC and 0.02 m $X_2Y_3$ are s etor (i) for solute in both t			ure;
	(a) 1.04, 5.1	(b) 1.9, 4.75	(c) 1.2, 3.4	(d) 1.5,	3.7	
128.	higher boiling point? (N	Iolecular weight of Kcl,	entration of aqueous soluti BaCl <sub>2</sub> , glucose and Al <sub>2</sub> (S dissociates completely in	$O_4)^3$ are 74	4.5, 208,	342
	(a) KCl	(b) BaCl <sub>2</sub>	(c) glucose	(d) $Al_2($	$(SO_4)_3$	
129.	Molecular weight of bio	molecules such as prot	ein can be determined by	1	method.	
	(a) osmotic pressure me	asurement	(b) Depression in freezin	ng point m	neasurem	ent
	(c) Elevation in boiling p	ooint measurement (d)	Vapour pressure measurer	ment		
130.	In the references of the	following graph, UR -	Oy=			
	R	- Jw				
	y K		(a) (SU -VW)QU			
	1 S	-	(b) $VU(VW - QY)$			
	1 A		(c) $(VW - QY)QU$ (d) $VW(QV - VU)$			
	q 2 1	¥	$(\mathbf{d}) \mathbf{v} \mathbf{w} (\mathbf{Q} \mathbf{v} - \mathbf{v} 0)$			
131.	What will be the elevat	ion in boiling point of	an aqueous solution of 0	.5 m NaC	21?(i=	1.8)
	(a)2.04, 5.1	(b) 1.9, 4.57	(c) 1.2, 3.4	(d) 1.4,		-
132.	When 2 gm phenol id d	issolve in 100gm benze	ene; then depression in fre	ezing poi	int is 0.69	9 K.

- 132. When 2 gm phenol id dissolve in 100gm benzene; then depression in freezing point is 0.69 K. If its association is dimeric, then calculate its degree of association (X). Molal depression constant for solvent is 5.12 K kg mole-1.
  - (a) 0.0734 (b) 0.374 (c) 0.00734 (d) 0.734
- 133. At 353 K temperature, the Vapour pressure of pure liquids A and B are 600mm and 800 mm respectively. If mixture of liquids A and B boils at 353 K and 1 bar pressure, then mole proportion of B in percent is -

(a) 80%	(b) 60%	(c) 20%	(d) 40%

134. 90 gm glucose and 120 gm urea dissolved in 1.46 kg aqueous solution, then what will be the boiling point of the solution at 1 bar pressure ? ( $Kb = 0.512^{\circ}C - kg - mole^{-1}$ , molecular weight of glucose and urea are 180 and 60 gm/mole respectively) (a) 100.876°C (b) 101.024°C (c)  $100.248^{\circ}C$ (d) 100.007°C 135. pH of 0.2M dibasic acid H<sub>2</sub>A is 1.699; then, what will be its osmotic pressure at T K temperature ? (a) 0.22 RT (b) 0.02 RT (c) 0.4 RT (d) 0.1 RT 136. Boiling point of an aqueous soultion of 0.4m AlCl<sub>3</sub> is 100.7°C; then what would be the pressure of ionization of AlCl<sub>2</sub> ? Kb - 0.512°C - kg - mole-1. (a) 80.67% (b) 60.5% (c) 76.54% (d) 84.75% 137. The vapour pressure of homogenous mixture of 10 mole of liquid X and 30 mole of liquid Y at constant temperature is 550 mm. In this solution, 10 mole of liquid Y increases, hence, increase in vapour pressure is 10 mm. Then, find the vapor pressure of pure liquid X and Y at that temperature. (a)  $P^{o}x = 200 \text{ mm}$ ,  $P^{o}y = 500 \text{ mm}$ (b)  $P^{o}x = 400 \text{ mm}$ ,  $P^{o}y = 600 \text{ mm}$ (c)  $P^{o}x = 600 \text{ mm}$ ,  $P^{o}y = 300 \text{ mm}$ (d)  $P^{o}x = 350 \text{ mm}, P^{o}y = 500 \text{ mm}$ 138. What amount of urea dissolved in 1 kg water at constant temperature, so that vapour pressure of the solution reduced by 2%? (M.W of urea = 60 gm/mole) (a) 68 gm (b) 60 gm (c) 50 gm (d) 75 gm 139. What would be the volume of 15% w/v and 5% w/v NaOH solution required to prepare 1 litre aqueous solution of 2M NaOH? (M.w. of Naoh = 40 gram/mole) (a) 300 ml, 700ml (b) 250 ml, 750ml (c) 400 ml, 600ml (d) 280 ml, 720ml 140. At constant temperature, vapour pressure of an aqueous solution of 1.5 kg glucose decreases to 0.98% in comparision with vapour pressure of pure water then, what quantity of glucose in gram dissolved in the solution ? (Molecular weight of glucose = 180 gm/mole) (a) 148.5 gm (b) 14.85 gm (c) 125 gm (d) 135 gm 141. At constant pressure, 0.5 m NaCl aqueous solution is diluted by adding water in it. Which of the following statement is correct in this reference? (a) Van't Hoff factor (i) and boiling point of the solution both decreases (b) Van't Hoff factor (i) and boiling point of the solution both increases (c) Van't Hoff factor (i) decreases while boiling point of the solution both increases (d) Van't Hoff factor (i) increases while boiling point of the solution both decreases 142. Boiling point of an aqueous solution of 0.5 m ionic solid substance is 100.5OC; then state the value of i? (Kb =  $0.512^{\circ}$ C -kg - mole<sup>-1</sup>) (a) 1.95 (b) 1.85 (c) 1.25 (d) 0.85 143. Aqueous solution of substance boils at 100.5°C at 1 bar pressure; then at what temperature it freezes ? (Kb =  $0.512^{\circ}$ C -kg - mole<sup>-1</sup>, Kf =  $1.86^{\circ}$ C - kg - mole<sup>-1</sup>) (b) 29.84°F (a) 11.2°C (c)  $271.8^{\circ}$ K (d)  $-1.2^{\circ}C$ 138

144. 1.4 m aqueous solution of a weak electrolyte  $AB_2$  ionizes 20%, then, state boiling point and freezing point of the solution respectively.

(a) 100.86 °C, -3.12°C (b) 101°C, -3.65°C (c) 274°C, -3.65°C (d) 374°C, -3.65°C

- 145. Solute substance in a 1.4 m aqueous solution associates by 25%, then, find the boiling point and freezing point of solution; where, solute exists as trimer in the solution; thus, n = 3. (Kb = 0.512°C kg mole<sup>-1</sup>, Kf = 1.86°C kg mole<sup>-1</sup>)
  - (a)  $100.448^{\circ}$ C,  $-2.28^{\circ}$ C (b)  $373.58^{\circ}$ K,  $-3.65^{\circ}$ C
  - (c)  $100.59^{\circ}$ C,  $2.17^{\circ}$ C (d)  $213^{\circ}$ F,  $270.83^{\circ}$ k

(b) 2:1

(b) 0.928 N

- 146. Molecular mass of a weak acid HA is 60gm/mo1 Ifs experimental molecular mass in its 0.7 M aqueous solution obtained from colligative properties is 50gm/mo1. Then calculate ionistion consteint of weak acid HA.
  - (a) 0.023 (b) 0.0085 (c) 0.035 (d) 0.085

147. At constant temperature, the total pressure of a homogeneous mixture of gas-A and gas-B in a closed container collected on water is 2.0 bar. Their ratio of mole frachion is 1.6 If the values of their  $K_{\rm H}$  are  $2.4 \times 10^{+4}$  bur and  $4.8 \times 10^{4}$  bar respectively then calcwate its ration of mole fraction when dissolhed in  $H_2O$ .

(a) 1:2

148. If one of the colligative property of 0.3m aqueous tisdn. of Na cland x m aqueows solution of  $H_2So_4$ then what would be the approximate vulwe of aqueows solution of  $H_2So_4$ ? (Density of × m  $H_2So_4$ solution = 1.185 gm/ml)

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(a) 0.464 N
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(c) 0.232 N

(c) 3:1

(d) 0.53 N

(d) 1:3

- 149. At one bar pressure the value of ratio of mole fraction of  $O_2$  and  $N_2$  gas in air is 1:4 The values of KH of  $O_2$  and  $N_2$  are  $3.3 \times 10^7$  Torr and  $6.60 \times 10^7$  Torr respectively. Then calculate the value of ratio of mole fractions of  $O_2$  and  $N_2$  gases will be. (1 Torr = 1mm)
  - (a) 3:1 (b) 2:1 (c) 1:2 (d) 1:3
- 150. According to Boyle-van't-Hofflaw at a constellnt temperature, osmotic pressure of a solution is directly proportional to its molarity. It means  $\pi \alpha C$ , where C = molarity of solution

 $\therefore \pi = \text{Kc}$  Then calculate the value of k in sI unit at 24° c temperatuve. (R = 8.314 J/mole k)

(a) 24.942 J/mol (b) 2494.2 J/mol (c) 0.024942 J/mol (d) 2.4942 J/mol

- 151. Boiling point of an aqueous solution of urea at one bar pressure is 373.41 k. Then at a constant temperature, calculate the percentage decrease in vapoure pressure of a solution compared to  $(k_b = 0.512 \text{ k.kg. mol}^{-1})$ 
  - (a) 1.42% (b) 2.56% (c) 4.17% (d) 3.44%
- 152. Calculate  $P^{H}$  of a solution prepared by mixing equal volume of an aqueous solution of HCI having  $P^{H} = 2$  and  $P^{H} = 5$  at 298 k temp.

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(a) 3.5 (b) 3.0 (c) 7.0 (d) 2.3

153. Which of the following is the correct formula for Rqoult's law when non-volatile solute is mixed with liquid solveut. where n=mole traction of solute N=mok fraction of solveet, P=vapoure pressure of solution  $P^{o}$ =vapoure pressure of pure sovent and  $\Delta P$ =Decrease in vapoure pressure.

(a) 
$$\frac{\Delta p}{p} = \frac{n}{N}$$
 (b)  $\frac{\Delta p}{p^0} = \frac{n}{N}$  (c)  $\frac{\Delta p}{p} = \frac{n}{n+N}$  (d)  $\frac{\Delta p}{p} = \frac{N}{n+N}$ 

154. Decrease in Freezing point of 75.2 gm pnenol when dissolved in a solvent having  $k_f = 14 \text{ k. } k_g$ . mole<sup>-1</sup>. is 7 k. Calculate the percentege of association of phenol if it is forms a dimev in the solutio. (a) 62.5 % (b) 80.5 % (c) 70 % (d) 75 %

155. Which of the following is correct option when kcl is dissolved in H<sub>2</sub>0.?

(a)  $\Delta H = +ve$ ,  $\Delta S = +ve$ ,  $\Delta G = +ve$  (b)  $\Delta H = +ve$ ,  $\Delta S = -ve$ ,  $\Delta G = -ve$ 

(c) 
$$\Delta H = +ve$$
,  $\Delta S = +ve$ ,  $\Delta G = -ve$  (d)  $\Delta H = -ve$ ,  $\Delta S = -ve$ ,  $\Delta G = +ve$ 

156. Naphthaleneis soluble in ether or benzene because.?

- (a) dipole-dipole attraction is equal (b) London forces are equal
- (c)Hydrogenbond (d)Ionicattraction
- 157. Four Liauids are given.
  - (i) Water: more polar and capacity to torm H-bond.
  - (ii) Hexanol:moderatly pdar and partial capacity to form H-bond.
  - (iii) Chloro form : moderatly polar and does not capable to form H-bond.
  - (iv) Octane: non polar and does not capable to form H-bond.

which of the following pair of liquids mixed with each other in very less proportion.

- (a) I, IV (b) I, II (c) II, III
- 158. Which of the following is applicable for the solubility of gases in liquid.
  - (a) Increases with increase in temperature and pressure.
  - (b) decreases with increase in temp and pressure.
  - (c) Increases with decrease in temp and increase in pressure.
  - (d) Decreases with decrease in temp and increase in pressure.
- 159. Eoncentration of lead metal in a blood of any person is more than that of 10 microgram. dm<sup>-1</sup>, than that person is considered as on effect of poision sectrion. Then calculate its concentration in ppb (parts per billion)

(d) III, IV

- (a) 1 (b) 10 (c) 100 (d) 1000
- 160. The ratio of  $\frac{\text{RT}}{\pi}$  of 6%  $\frac{\text{W}}{\text{v}}$  and 9%  $\frac{\text{W}}{\text{v}}$  is one for both. what would be the value of atomic weight of A and B respectively. (AB<sub>2</sub> and A<sub>2</sub>B are electrolytes)

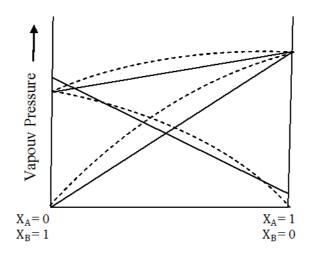
(a) 60, 90	(b) 40, 40	(c) 40, 10	(d) 10, 40

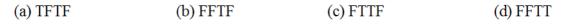
161. Decrease in vapour pressure of an aqueous soln. of an electrolyte is 4% what would be the percentage increase in elevation in Boiling point?  $(k_b = 0.512 \text{ k. } k_g \text{ mol}^{-1})$ 

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(a) 0.55 %(b) 0.02 %(c) 5.5 %(d) 2 %

- 162. Which of the following option is correct for a homogeneous mix tare of liquid A and liquid B which shows positive deviation according to Result's laco.
  - (a) A-B intermolecular attraction is more than intermolecular attraction in A-A and B-B
  - (b) Intermolecular attraction in A-B is less than in termolecular attraction in A-B and B-B.
  - (c) Inter molecular attraction in B-B is less than intermolecular attraction in B-B is less than intermolecular attraction in A-A
  - (d) Intermolecular attraction in A-A is less than in termolecular attraction B-B
- 163. Select the true and false statements with reference to the graph of mole fraction Vapour pressure. T = True statement F = False statement.
  - (i) Azeotropic mixture of liquid A and liquid B has highest boiling point.
  - (ii) When vapour pressure of both liquids in q mixture is same then  $X_A < X_B$ .
  - (iii) It shows positive deviation on the basis of Raoult's law.
  - (iv) When  $X_A = X_B$  then  $P_A < P_B$ .





ANSWER KEY

1	b	26	d	51	С	76	b	101	С	126	С	151	а
2	а	27	d	52	а	77	а	102	b	127	d	152	d
3	d	28	а	53	b	78	а	103	d	128	а	153	а
4	d	29	b	54	d	79	С	104	d	129	а	154	d
5	С	30	d	55	С	80	С	105	а	130	С	155	С
6	а	31	а	56	b	81	а	106	С	131	а	156	b
7	С	32	b	57	а	82	а	107	b	132	d	157	а
8	d	33	С	58	d	83	а	108	b	133	а	158	С
9	а	34	С	59	С	84	b	109	d	134	b	159	С
10	b	35	d	60	С	85	С	110	С	135	а	160	С
11	b	36	С	61	С	86	b	111	С	136	а	161	а
12	С	37	b	62	d	87	С	112	b	137	b		
13	b	38	b	63	b	88	а	113	d	138	а		
14	d	39	d	64	b	89	b	114	а	139	а		
15	b	40	а	65	d	90	d	115	b	140	d		
16	С	41	а	66	d	91	d	116	C	141	d		
17	d	42	a	67	С	92	а	117	b	142	а		
18	а	43	d	68	d	93	с	118	С	143	С		
19	d	44	С	69	а	94	а	119	U	144	b		
20	b	45	b	70	а	95	а	120	С	145	d		
21	а	46	b	71	а	96	b	121	U	146	С		
22	С	47	С	72	С	97	С	122	d	147	d		
23	а	48	а	73	b	- 98	d	123	b	148	а		
24	С	49	а	74	d	99	b	124	b	149	С		
25	С	50	b	75	d	100	С	125	а	150	b		

### Hints

- (23)  $[M \cdot V = M_1 \cdot V_1 + M_2 \cdot V_2 + M_3 \cdot V_3, \qquad V = V_1 + V_2 + V_3]$  $\therefore M \cdot 2600 = 0.3 \times 1400 + 0.4 \times 700 + 1.2 \times 500$  $\therefore M = 0.5$
- (25)  $M_1 \cdot V_1 = M_2 \cdot V_2$  5000ml = 5 lit.  $1.5 \times 5 = 0.5 \times V_2$   $\therefore V_2 = 15$  lit.  $\therefore$  Amount of water added = 15-15 = 10 lit.
- (35) Slope =  $k_H$ Solubility of gas is more which have less value of  $k_H$ 
  - $\therefore$  gas D has more solubility.
- $(45) \quad n_1 \bullet m_1 = n_2 \bullet m_2$

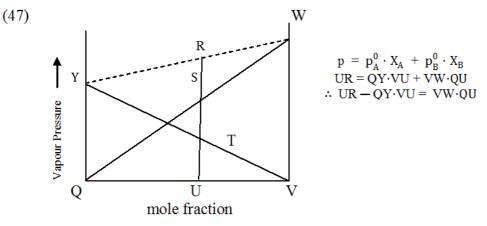
Where  $n_1 =$  number of ions produse when  $1^{st}$  substanle is dis solved in H<sub>2</sub>O

 $m_1 = mulality of 1^{st} substanle.$ 

 $n_2$  = number of ions produced when  $2^{nd}$  substance is dissolved in  $H_2O$ . according to its formula.

The values of  $n_1$  for  $Na_2So_4$ , ureq and  $Alcl_3$  are 3, 1 and 4 respectively.

we get 0.8m, 2.4m and 0.6m for  $Na_2So_4$ , ureq and Alcl<sub>3</sub>.



(49)  $Y_A = 0.4$  Aand  $P_A = 400$  mm

 $P_{\rm B} = 1000 - 400 = 600 {\rm mm}$ 

- $P_A = P_{Total} \times Y_A$
- $\therefore 400 = P \times 0.4$
- $\therefore P_{\text{Total}} = 1000 \text{ mm}$

 $\frac{\Delta Tf}{\Delta Tb} = \frac{Kf}{Kb} \quad \therefore \Delta T_{f} = \frac{Kf}{Kb} \times \Delta T_{b}$  $= \frac{1.86}{0.51} \times 0.55 = 2 \text{ K}$ (50) $\therefore T_f = 273 - 2 = 271 \text{ K}$ (53)  $n_1 M_1 = n_2 M_2$ For NH<sub>4</sub> No<sub>3</sub> values for n<sub>1</sub> and M<sub>1</sub> are 2 and 1.5 resp For Al<sub>2</sub> (so<sub>4</sub>)<sub>3</sub>  $n_2 = 5$  $\therefore M_2 = \frac{2}{5} \times 1.5 = 0.6$ Normality of  $Al_2(SO_4)_3 = 6 \times molarity$  $= 6 \times 0.6$ = 3.6 N(58)Answer = DHighest os motic pressure is for that pair having highest difference between n<sub>1</sub> M<sub>1</sub> and  $n_2 M_2$  $\mathbf{p}_A = \mathbf{p}_A^0 \times \mathbf{X}_A$ (62)  $= 125 \times 0.6$ = 75 mm  $p_B = p_B^0 \times X_B = 62.5 \times 0.4 = 25 \text{ mm}$  $P_{\text{Total}} = p_A + = p_B = 75 + 25 = 100 \text{ mm}$  $p_A = P_{Total} \times Y_A$  $\therefore$  75 = 100 × Y<sub>A</sub>  $\therefore Y_A = 0.75$ Similarly  $\therefore$  Y<sub>B</sub> = 0.25 For Na C1 value of i = 2.0(67) $\therefore \Delta Tb = i \cdot m \cdot Kb = 2 \times 0.1 \times Kb$  $\therefore \Delta Tb = \frac{Kb}{5}$  $X_B = 1 - X_A = 1 - 0.7 = 0.3$   $Y_B = 1 - Y_A = 1 - 0.4 = 0.6$ (69)  $p_A = p_A^0 \times X_A$  Nand  $p_A = P_{Total} \times Y_A$ Noe  $p_A = p_A^{"} \times x_A$  react  $p_A = p_{Total}^{"} \times x_A$   $p_A = p_{Total}^{"} \times 0.4$   $\therefore p_A^0 \times 7 = p_{Total} \times 4$ Similarly  $p_B^0 \times 3 = p_{Total} \times 6$   $\frac{p_A^0 \times 7}{p_B^0 \times 3} = \frac{p_{Total}^{"} \times 4}{p_{Total}^{"} \times 6}$   $\therefore p_A^0 = \frac{p_B^0 \times 2}{7}$   $p_A^0 \times 7 = p_{Total}^{"} \times 6$   $\therefore \frac{p_B^0 \times 2}{7} + p_B^0 = 90$   $\therefore \frac{p_B^0 \times 2}{7} + p_B^0 = 90$   $\therefore 9p_B^0 = 630$   $\therefore p_B^0 = 70 \text{ mm and}$   $p_A^0 = 20 \text{ mm}$ **NNNoe** By dividing

- (70)  $X_A = 1 X_B = 1 0.4 = 0.6$   $Y_A = 1 - Y_B = 1 - 0.25 = 0.75$   $p_B = p_B^0 \times X_B$   $p_B = P_{Total} \times Y_B$   $\therefore p_B = 40 \times 0.4$   $\therefore 16 = P_{Total} \times 0.25$  = 16 mm  $\therefore P_{Total} = 64 \text{ mm}$   $p_A = P_{Total} \times Y_A = 64 \times 0.75 = 48 \text{ mm}$ Now  $p_A = p_A^0 \times X_A$   $\therefore 48 = p_A^0 \times 0.6$  $\therefore p_A^0 = 80 \text{ mm}$
- (73) Whiten  $\times$  Suppose  $\times$  m1 H<sub>2</sub>O is transferred from aqueous solution of KCI to aqueous solution of Alcl<sub>3</sub> then the phenomenon osmosis stops. So morality of soluble particles in both solutions becomes equal at this time.
  - $M_1 = \text{modality of kcl solution}$   $M_2 = \text{modality of kcl solution when x ml water is reduced.}$   $M_1 V_1 = M_2 V_2$   $0.2 \times 2000 = M_2 \times (2000 \text{ x})$  $∴ M_2 = \frac{400}{2000 \text{ x}}$

Similarly volume of Alcl<sub>3</sub> Solution increases by addition of  $x m_1 H_2O$ .

morality of Alcl<sub>3</sub> solution =  $\frac{600}{2000 + X}$ 

Now morality soluble particles in kcl solution = morality of soluble particles in Alcl<sub>3</sub> soln.

- $\therefore n_1 \cdot M_1 = n_2 \cdot M_2$  $\therefore \frac{2 \times 400}{2000 - x} = \frac{4 \times 600}{2000 + x}$  $\therefore x = 1000$
- (74) Liquid present in Red Blood cells is isotonic with 0.91 % W/<sub>V</sub> solution of Nacl
   ∴ morality of soluble particles in 0.91 % W/<sub>V</sub> Nacl solution

$$= 2 \times 0.1555$$
  
= 0.311 M

(75)  $\pi = MRT$  where M = effective molarity of solutions kept in contact with semi-permeable membrane.

 $\therefore \pi = 0.2 \text{ M (molarity of soluble particles in Naclsdn)}$ = 3 × 0.2 - 2 × 0.1 =0.4 M.

(86)  ${}^{\text{o}}\text{l}_{0} \frac{\text{dissociation}}{100} = \frac{i-1}{n-1}$ Fec13 ionises 80% in its aqueous solution  $\therefore \frac{80}{100} = \frac{i-1}{4-1} \quad \therefore i = 3.4$ 

(87) when association of any substance takes place in a solution then degree of association<sup>is</sup>  $\leq 1$ .

$$\therefore 0 < \frac{i-1}{\frac{1}{n}-1} \le 1$$

$$\therefore 0 < \frac{1-i}{1-\frac{1}{n}} \le 1$$

$$\therefore 0 < 1-i \le 1-\frac{1}{n}$$

$$\therefore -1 < -i \le -\frac{1}{n}$$

$$\begin{array}{l} \therefore 1 > i \geq \frac{1}{n} \\ \text{Here n} = 2 (\text{given}) \\ \therefore 1 > i \geq \frac{1}{2} \text{ means } 0.5 \leq i < 1 \\ \end{array}$$

$$(90) \quad \text{For association (given) n=2} \\ \therefore i \geq \frac{1}{2} \\ \therefore \text{ imff} \geq \frac{mKr}{2} \\ \therefore \Delta Tr \geq \frac{mKr}{2} \\ \Rightarrow \text{ imff} \geq \frac{mKr}{2} \\ \Rightarrow \text{ imff} \geq \frac{mKr}{2} \\ \Rightarrow \Delta Tr \geq \frac{mKr}{2} \\ \Rightarrow \frac{mKr}{2} = \frac{mKr}{2} \\ \text{(91)} \quad i = \frac{4\pi}{M_{eff}} = \frac{mK}{M_{eff}} \\ \text{where } M = \text{Actual molecular mass of solute} \\ \text{Mob} = \text{experimental molecular mass of solute} \\ \text{Mob} = \text{experimental molecular mass of solute} \\ \text{For AB Type ionic compound } i \leq 2 \\ \Rightarrow \frac{M}{M_{eff}} \leq 2 \\ 5 : 2 > 2 : 1 \\ \text{(96) The substance forms trimer in a solution due to association.} \\ \Rightarrow \frac{1}{3} \leq i < 1 \\ \Rightarrow \frac{mKr}{3} \leq \text{imKf} < mKr \\ \Rightarrow -mKr = -\Delta Tr \leq -\frac{mKr}{3} \\ \Rightarrow Tr^0 - mKr < -\Delta Tr \leq -\frac{mKr}{3} \\ \Rightarrow Tr^0 - mKr < Tr = \frac{mKr}{3} \\ \text{(98) If x molar solution of any substance is more concentrated than x molul solution so molarity value is less than the molitury value.} \\ \Rightarrow \frac{imMr}{molarity} = \frac{1000 \times W}{M \times V} \times \frac{M \times W_0}{1000 \times W} \\ \Rightarrow \frac{molarity}{molarity} = \frac{1000 \times W}{V} \times \frac{M \times W_0}{1000 \times W} \\ \Rightarrow \frac{molarity}{W} = \frac{W_0 \text{ gm}}{W_0 \text{ mol}} \qquad \Rightarrow \frac{W_0 \text{ W}}{W} = \frac{W}{V} = \frac{W}{V$$

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its

(105) 
$$T_b - T_f = 75.7$$
  
 $\therefore T_b^0 + \Delta T_b - (T_f^0 - \Delta T_f) = 75.7$   
 $\therefore 80 + \Delta T_b - (5.5 - \Delta T_f) = 75.7$   
 $\therefore \Delta T_b + \Delta T_f = 1.2$   
 $\therefore imKb + imKf = 1.2 \quad \therefore i = \frac{1.2}{m(Kb + Kf)} = \frac{1.2}{0.2(2.65 + 5.12)} \quad \therefore i = 0.77$ 

(117) 
$$X = \frac{m}{55.55 + m}$$

$$X = \text{mole fraction of solute}$$

$$m = \text{molality} = 4.5 \text{ m}$$

$$X = \frac{4.4}{55.55 + 4.5} = 0.075$$
(118) Density of 98 % <sup>W</sup>/<sub>W</sub> H<sub>2</sub>SO<sub>4</sub> = 1.8  $\frac{gn}{ml}$ 

$$w = 98 \text{ gm}$$

$$d = \frac{W + W_0}{V} \div V = \frac{W + W_0}{d}$$

$$V = \frac{100}{1.8} \text{ ml molarity} (m) = \frac{1000 \times w}{M \times V}$$

$$= \frac{1000 \times 98 \times 1.8}{98 \times 100}$$

$$= 18 \text{ M}$$
(119) and (121)
$$molality = \frac{100 \times \text{molrity}}{(1000 \times \text{density}) - (\text{mol.mass of solute } \times \text{molrity})}$$
use above formula to solve the question
$$= 5 \times 25 = 125 \text{ gm}.$$
mass of N<sub>a</sub>OH in 500gm 25% <sup>W</sup>/<sub>W</sub> N<sub>a</sub>OH = 5 × 15 = 75 \text{ gm}.
mass of N<sub>a</sub>OH in 500gm 15% <sup>W</sup>/<sub>W</sub> N<sub>a</sub>OH = 5 × 15 = 75 \text{ gm}.
mass of N<sub>a</sub>OH in a mixed solution when both solutions are mixed W = 125 + 75 = 200 \text{ gm}.
Now molality of mixed solution =  $\frac{1000 \times w}{M \times W_0} = \frac{1000 \times 200}{100 \times 800} = 6.25 \text{ m}$ 
(124) molality of 25% <sup>W</sup>/<sub>W</sub> N<sub>a</sub>OH
$$= \frac{1000 \times w}{M \times W_0} = \frac{1000 \times 25}{40 \times 75} = 8.33 \text{ m}$$
molality of 15 % <sup>W</sup>/<sub>W</sub> NaOH =  $\frac{1000 \times w}{M \times W_0} = \frac{1000 \times 15}{40 \times 85} = 4.41 \text{ m}$ 

when two different wncentration containing solutions of same substances are mixed then conc of dil. solution < concentration of mixed solution < conc. of concentration soln.  $\therefore$  4.41 m < conc. (molality) of mixed solution < 8.33 in

(128) From graph

$$p = p_A^0 + (p_B^0 - p_A^0) X_B$$
  
$$\therefore UR = QY + (VW - QY)QU$$

(129)  $\frac{n \times \% W/W}{\text{molecular mass (formula weigh}} = X$ 

If volue of x is hignest than solution hare highest Boiling point.

(n = no. of ions in a formula)

(134) mass of solvent in a solution wo = 
$$1460 - (90 + 120) = 1250$$
 gm.

mole of glucose = 
$$\frac{90}{180} = 0.5$$

mole of urea =  $\frac{120}{60} = 2$ 

Total moles of solute in a solution = 0.5 + 2 = 2.5

molality = 
$$=\frac{1000 \times n}{W_0} = \frac{1000 \times 2.5}{1250} = 2.0$$
 m

$$\Delta Tb = mKb = 2 \times 0.512 = 1.024$$
 °C

$$\therefore$$
 Tb = 100 + 1.024 = 101.024 °C.

(135) 
$$pH = 1.669$$
  $\therefore [H_3O^+] = 0.02 M$ ,  $[H_2A] = 0.2$ ,

Degree of dissociation  $\alpha = \frac{0.02}{0.2} = 0.1 = \frac{i-1}{n-1}$ , (n = 3)

$$\pi = iMRT = 1.2 \times 0.2 \times RT : i = 1.2 = 0.22R1$$

(137) 
$$n_X = 10$$
,  $n_Y = 30$ 

Total mok = 40

$$\therefore X_{\rm X} = 0.25 \ {\rm X}_{\rm Y} = 0.75$$

 $Total \ uapour \ pressure \qquad = \mathbf{P} = \mathbf{p}_{\mathbf{X}} + \mathbf{p}_{\mathbf{X}}$ 

$$\therefore n_X \cdot p_X^0 + n_Y \cdot p_Y^0 = P \qquad \therefore 0.25 p_X^0 + 0.75 p_Y^0 = 550 \dots (1)$$

If mole of liqulid y is in creased by 10 then its uapour pressure is increased by 10 mm.

 $\therefore$  n<sub>X</sub> = 10 , n<sub>Y</sub> = 40  $\therefore$  Total mole = 50

 $X_X = 0.2$ ,  $X_Y = 0.8$  and total vapor pressure  $P = p_X + p_X = 560$  mm

:  $0.2p_X^0 + 0.8p_Y^0 = 560$  .....(2) By soloing eqn. (1) and (2) we get  $p_X^0 = 400$  mm and  $p_Y^0 = 600$  mm (138)  $\Delta p = \frac{2p^0}{100} \therefore \frac{\Delta p}{p^0} = \frac{1}{50} = X$  (mole fraction of urea)  $= \frac{m}{55.55 + m}$  $\therefore m = 1.134$  (molality of urca)

: mass of urea ( $W_2$ ) = 1.134× 60 = 68 gm.

(139) Suppose V<sub>1</sub> liter 15 % W/<sub>V</sub> NaOH and V<sub>1</sub> liter 5 % W/<sub>V</sub> NaOH solution is required to pre paec one litev 2 M NaOH solution.

80 gm. NaOH is required to prepare one liter 2m NaOH solution.

 $\therefore V_1 + V_2 = 1$  liter.....(1).

 $150V_1 + 50V_2 = 80 \text{ gm.} \dots (2)$ 

By soluing (i) and (ii)

are get  $V_1 = 300$  ml. and  $V_2 = 700$  ml

