



Topics to be covered



- Revision of Last Class, Medics test
- Non ideal solution
- 3 Numericals
- 4 Home work from modules



Rules to Attend Class



- 1. Always sit in a peaceful environment with headphone and be ready with your copy and pen.
- Never ever attend a class from in between or don't join a live class in the middle of the chapter.
- 3. Make sure to revise the last class before attending the next class & always complete your Magarmach Practice Questions.
- 4. Never ever engage in chat whether live or recorded on the topic which is not being discussed in current class as by doing so u can be blocked by the admin team or your subscription can be cancelled.

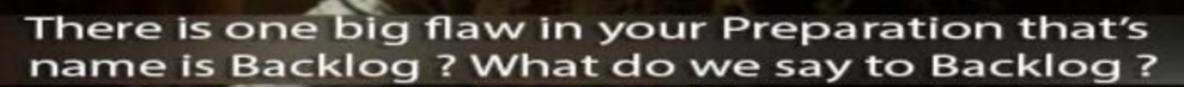


Rules to Attend Class



- Try to make maximum notes during the class if something is left then u can use the notes pdf after the class to complete the remaining class.
- Always ask your doubts in doubt section to get answer from faculty. Before asking any doubt please check whether same doubt has been asked by someone or not.









MEDICS

Mastery

Checks your grasp over NEET-level concepts

Evaluation

Judging both knowledge and test-smartness

Decision Making

Testing your speed + accuracy under pressure

Intuition

Some answers need gut + logic - can you spot the trick?

Concepts

It's all about strong basics no shortcuts here

Strategy

The MEDICS test – built for those who heal, hustle, and hope.

find Molality if MC6H6 = 789/mol

(d) None of these

91 mole fraction of Ta in CoH6 is 0.2.

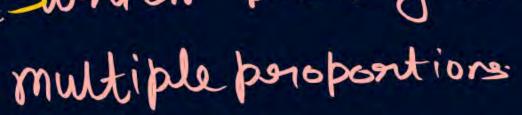


92 which statement is connect? miæture of 649 ta 2 649 og is ignited @ Hais L. R. 64=32 64=2 Umoles

(B) Ca is L. R. 32=16 2=2. UX18=729 160

(B) Ca is L. R. 32=16 My unreacted. 0-32-4-28 mass to unsteated -28×2-569 on mix. has 72g Hgo 2 56g unneacted Hg. (d) 11 11 569 80 & 789 ~~~X

@ 3which pain of Compound do not illustrate law of



(a) Sog & Soz

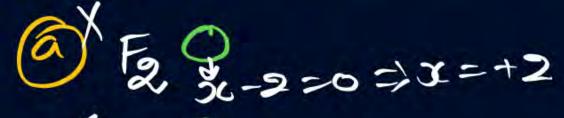
(b) Ng & Ngo

MgO & Mg (OH)

NO & 1205



94 which moleules orepresented by green Colown are in lowest oxidation 8tt ?





oxian number of S in Sg, $\frac{5}{2}$ Fg & Hg S $\frac{3}{2}$ O $\frac{3}{2}$ $\frac{2}{2}$ $\frac{2}{2}$ $\frac{2}{2}$ $\frac{2}{2}$ $\frac{2}{2}$



Test Syllabus zome Vodenate & Tough





Revision of Last class



Ideal sol Ps=PA%A+PB%B AGMix = (-)ve J& spontaneous ASMix = (+)ve J& spontaneous ASMix = (+)ve J& spontaneous RS-PAXA. DVmix = 0 Atmix = 0 AU inix = 0 AP = 0



P°A and P°B are the vapor pressure of pure liquid components A and B respectively of an ideal binary solution. If χA represents the mole fraction of component A, the total pressure of the solution will be

$$(A) p_A^\circ + \chi_A (p_B^\circ - p_A^\circ)$$

$$\mathbf{B} \quad \mathbf{p}_{A}^{\circ} + \chi_{A} \left(\mathbf{p}_{A}^{\circ} - \mathbf{p}_{B}^{\circ} \right)$$

$$p_B^* + \chi_A (p_B^* - p_A^*)$$

$$p^{\circ}_{B} + \chi_{A} (p^{\circ}_{A} - p^{\circ}_{B})$$

QUESTION - (NEET 2019)



For an ideal solution, the correct option is:

- Δ_{mix} S = 0 at constant T and P
- \triangle Δ_{mix} V \neq 0 at constant T and P
- Δ_{mix} H = 0 at constant T and P
- $\Delta_{\text{mix}} G = 0$ at constant T and P



Which one of the following is incorrect for ideal solution?



$$\triangle U_{mix} = 0$$

$$\Delta P = P_{obs} - P_{calculated by Raoult's law} = 0$$

$$\Delta G_{\text{mix}} = 0$$

QUESTION - (NEET 2015)



Which one is not equal to zero for an ideal solution?

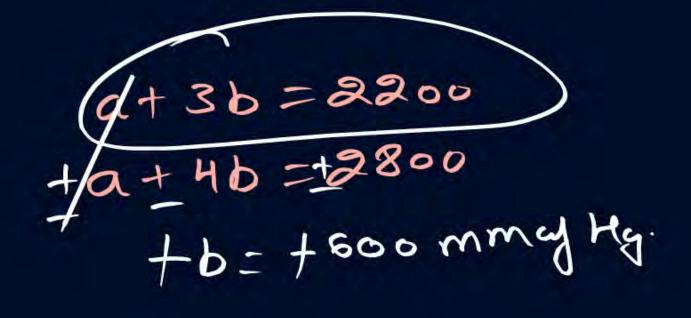


- $oxed{B}$ ΔV_{mix}
- $\Delta P = P_{observed} P_{Raoult}$
- D ΔH_{mix}



Two liquids X and Y form an ideal solution. At 300 K, vapour pressure of the solution containing 1 mol of X and 3 mol of Y is 550 mm Hg. At the same temperature, if 1 mol of Y is further added to this solution, vapour pressure of the solution increases by 10 mm Hg. Vapour pressure (in mm Hg) of X and Y in

- 200 and 300
- 300 and 400
- 400 and 600
- 500 and 600



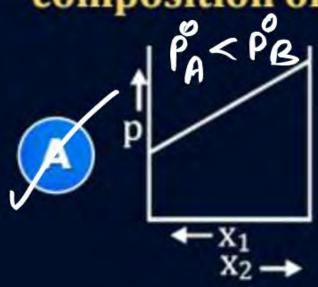


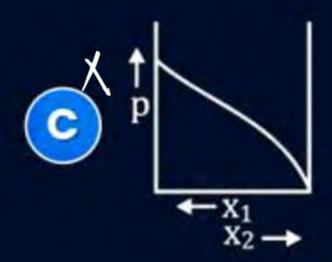
QUESTION* - (NCERT Exemplar)

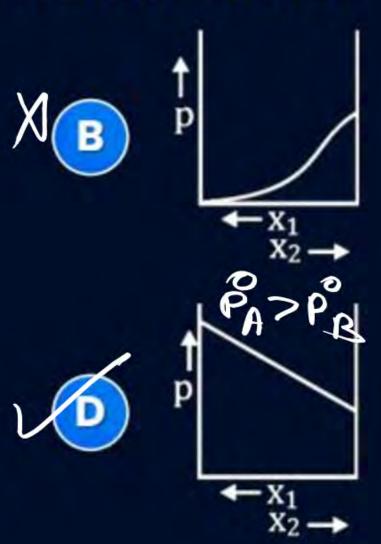


For a binary ideal liquid solution, the variation in total vapour pressure versus

composition of solution is given by which of the curves?











Formation of a solution from two components can be considered as

- (i) Pure solvent \rightarrow to separate solvent molecules, ΔH_1
- (ii) Pure solute \rightarrow to separate solute molecules, ΔH_2
- (iii) Separated solvent & solute molecules \rightarrow Solution, ΔH_3 Solution so formed will be ideal if
- $\Delta H_{\text{soln}} = \Delta H_3 \Delta H_1 \Delta H_2$
- $\Delta H_{\text{soln}} = \Delta H_1 + \Delta H_2 \Delta H_3$
- $\Delta H_{soln} = \Delta H_1 \Delta H_2 \Delta H_3$





Think without Ink for Ideal Solution numerical

#ITT

6) if PA > PB then you masc V. P.

9CA=1\$ \$150

Ps=PA

if PR > PA

then max V-P.

\$68=12 18A=0

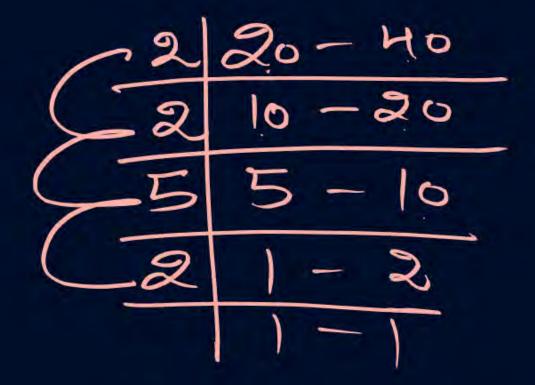
Ps = PR

Titwa=WB=Xg. Lic.mg Mae MB=Xg

(8) if nA = nB Vapowers urste jyeada banege jikka pohigh hai

9 24 nA=nB & half of moles Vaponised then Ps=JPAPR

L.C.M. of 20240





QUESTION - (AIPMT 2012)



 P_A and P_B are the vapour pressure of pure liquid components, A and B respectively of an ideal binary solution. If X_A represents the mole fraction of component A, the total pressure of the solution will be:

- $(A) p_A + x_A (p_B p_A)$

- $p_B + x_A (p_A p_B)$



If $P_A^0 = 200$ mm of Hg, $P_B^0 = 300$ mm of Hg and Which of these cannot be P_S for two miscible liquids following ideal behaviour?

PALPS<PB

- A 230 mm of Hg
- B 270 mm of Hg
- 150 mm of Hg
- Cannot be solved



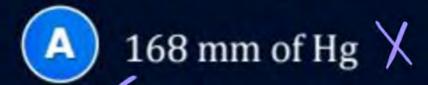
- <u>156</u>
- B 145
- C 150
- 108

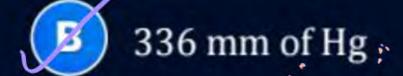


The correct option for the value of vapour pressure of a solution at 45° C with benzene to octane in molar ratio 3:2 is:

[At 45°C vapour pressure of benzene is 280 mm Hg and that of octane is 420 mm Hg.

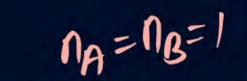
Assume Ideal gas]





- 350 mm of Hg.
- 160 mm of Hg \

QUESTION - (NEET 2016-I)





Which of the following statements about the composition of the vapour over an ideal 1:1 molar mixture of benzene and toluene is correct? Assume that the temperature is at 25°C. (Given, vapour pressure data at 25°C, benzene = 12.8 kPa, toluene = 3.85 kPa)

- The vapour will contain equal amounts of benzene and toluene. A $\frac{\sqrt{A}}{R^2}$
- B Not enough information is given to make a prediction.
- The vapour will contain a higher percentage of benzene.
- The vapour will contain a higher percentage of toluene.



If $P_A^0 = 200$ mm of Hg, $P_B^0 = 300$ mm of Hg and Which of these cannot be P_S for two miscible liquids following ideal behaviour?

- A 200 mm of Hg
- B 250 mm of Hg
- C 240 mm of Hg
- 700 mm of Hg



A and B form a ideal solution and V.P. of pure A and B are 160 mm of Hg and 60 mm of Hg. Calculate partial pressure of benzene and toluene and total pressure

also Pa=160mn of Hg, PB=60mm of Hg

- Containing equal mass of both A and B $= \omega_A = \omega_B = 409$
- Containing equal molecules of both A and B
- Containing 1 mole of A and 4 moles of B
- Also calculate mole fraction of A and B in vapour phase if equal moles of A and B mixed
- UP=UB => XY=XB Ps = PA + PB = 160 + 60 = 100 mm g/lg.

PA = 40 = 2) NB = 40 = 1 NA = 23 = 320 PA = 160×23 = 320 PB = 60×13 = 20 PB = 120 PB =

$$\begin{array}{lll}
\text{D} & n_{A} = n_{B} \\
Y_{A} = \frac{c_{A} x_{A}}{P_{S}} & = \frac{160 \times \frac{1}{2}}{14D} = \frac{80}{14D} = \frac{16}{20} = \frac{8}{11} \\
Y_{B} = 1 - \frac{8}{11} = \frac{3}{11}
\end{array}$$



At a given temperature, the vapor pressure in mm of Hg of a solution of two volatile liquids A and B is given by equation

 $P = 120 - 80\chi_B$ Calculate V.P. of pure A and B at same temperature



If $P_A^0 = 200$ mm of Hg, $P_B^0 = 300$ mm of Hg and solution both have same mass and $M_A = 20$ g and $M_B = 200$ g. Find vapour pressure of solution. Ps -200×10 + 300×1

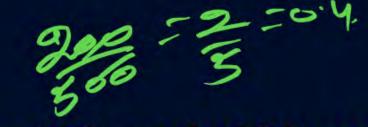
- **∀325 mm of Hg**
- 2009 NA = 200 = 10
- 146.35 mm of Hg
- NB = 200 = 1
- 209 mm of Hg
- 285 mm of Hg



If $P_A^0 = 100$ mm of Hg, $P_B^0 = 500$ mm of Hg and have same no. of moles in solution. Find vapor pressure of solution.

- A 100 mm of Hg
- B 150 mm of Hg
- 300 mm of Hg
- 500 mm of Hg

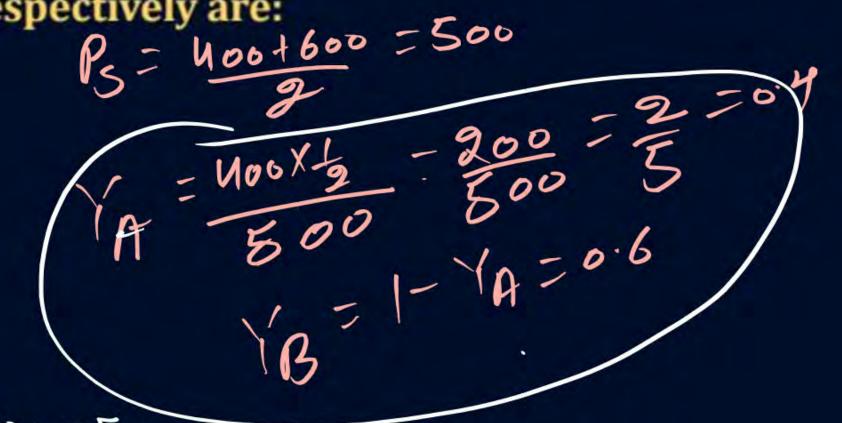
QUESTION (8th April, 1st shift-2019)





The vapour pressures of pure liquids A and B are 400 and 600 mm Hg, respectively at 298 K. On mixing the two liquids, the sum of their initial volumes is equal to the volume of the final mixture. The mole fraction of liquid B is 0.5 in the mixture. The vapour pressure of the final solution, the mole fractions of components A and B in vapour phase, respectively are:

- A 450 mm Hg, 0.5, 0.5 P = PA XA
- B 450 mm Hg, 0.4, 0.6
- 500 mm Hg, 0.5, 0.5 Pg = 600 XL
- D 500 mm Hg, 0.4, 0.6 R = 200 + 300 = 500.





Find Mole Fraction of A and B if V.P. of solution is 160 torr. If V.P. of pure A and B are 100 mm of Hg and 200 mm of Hg.

$$160 = (260 - 100) \% B + 100$$

$$160 - 100 = 60 = 100 \% B$$

$$\% B = \frac{60}{100} = \frac{3}{5}$$

$$\% A = \frac{2}{5}$$



If in a solution 1 mole each of A and B are mixed. Find vapour pressure of solution if half of the total moles are vaporised ($P_A^0 = 500 \text{ mm}$ of Hg and $P_B^0 = 20 \text{ mm}$ of Hg)

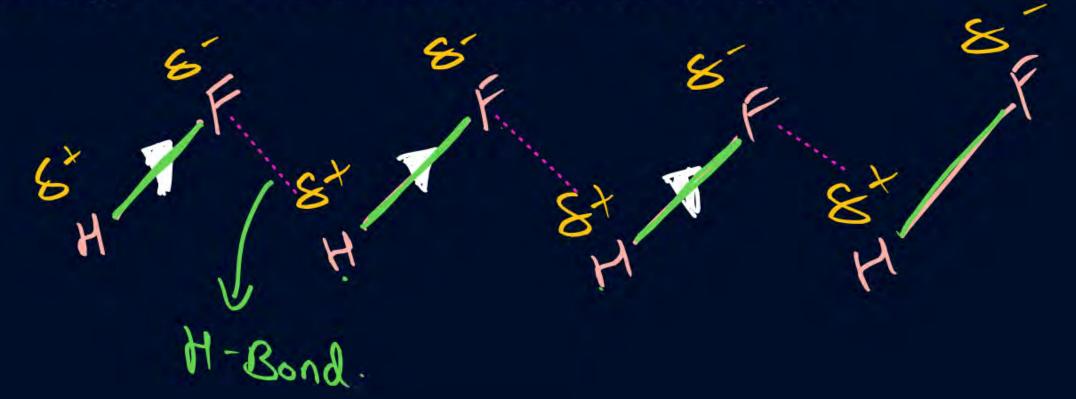


Don't attempt today's dept Jo it tomorrow.

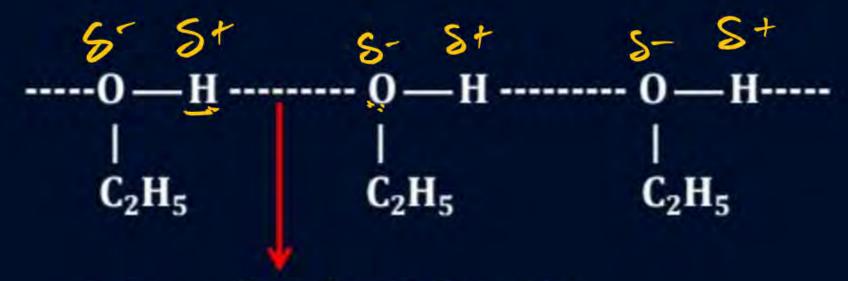




Strong force of attraction between H & F, O, N







Intermolecular H-Bonding



Home work from modules



Solve all questions of Ideal Solution.



