### SAMPLE CONTENT MHT-CET 2021 TRIUMPH CHEENSSERV

## MULTIPLE CHOICE QUESTIONS 3049 MCQS

A chameleon basks in the sun. As its body temperature increases, the chemical reactions of its metabolism speed up!

BASED ON STD. XII SYLLABUS 2020-21



#### MHT-CET TRIUMPH CHEMISTRY MULTIPLE CHOICE Based on New Syllabus

#### **Salient Features**

- ☞ Includes chapters of Std. XII as per textbook of 2020.
- Exhaustive subtopic wise coverage of MCQs.
- 3049 MCQs including questions from v<sup>-1</sup> Jous comp<sup>-1</sup> tive e<sup>-1</sup> ams.
- Quick Review provided for all the chapte .
- Important Formulae provided for relevant vpters
- Includes MCQs from JEE (Main) (8<sup>th</sup> April, shn, 1), NEET (UG), NEET (Odisha), MHT-CET (6<sup>th</sup> May, Afternoon) 2019 2<sup>rd</sup> TEE (Main) (7<sup>th</sup> January, shift 1) 2020.
- Includes MCQs from 'EE (M n),  $\mathbb{M}_{E^{T}}$  and MHT-CET upto 2018.
- Solution Various competitive exal. 'nation nuest insupdated till the latest year.
- Evaluation test protent with the nd of each chapter.
- Tinclusions: 'Rea vorld applications' and 'Compilation of organic reaction based MCQs'.

Scan the adj. ont Qk de or visit www.targetpublications.org/tp1629 to download Hints free revealed and Evaluation Test in PDF format.



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**TEID: 1629** 

#### PREFACE

"Don't follow your dreams; chase them!"- a quote by Richard Dumbrill is perhaps the most pertinent for one who is aiming to crack entrance examinations held after std. XII. We are aware of an aggressive competition a student appearing for such career defining examinations experiences and hence wanted to create books that develop the necessary knowledge, tools and skills required to excel in these examinations.

For the syllabus of MHT-CET 2020, 80% of the weightage is given to the syllabus for XII<sup>th</sup> standard while only 20% is given to the syllabus for XI<sup>th</sup> standard (with inclusion of only selected chapters). Since there is no clarity on the syllabus for MHT-CET 2021 till the time when this book was going to be printed and taking the fact into consideration that the entire syllabus for std. XII<sup>th</sup> Science has always been an integral part of MH<sub>1</sub> CET syllabus, this book includes all the topics of std. XII<sup>th</sup> Chemistry.

We believe that although the syllabus for Std. XII and MHT-CET is aligned, the outlook to stable should be altered based on the nature of the examination. To score in MHT-CET, a student as to be not just good with the concepts but also quick to complete the test successfully. Such in enuity in 'b de loped through sincere learning and dedicated practice.

Having thorough knowledge of basic principles, laws, concepts and their applications . a prerequisite for beginning with MCQs on a given chapter in Chemistry. For physical chemistry tuo, 's must tow formulae, conversion factors, units and dimensions of physical quantities involved in the chap. For inorganic and organic chemistry, students need to focus on chemical behaviour of elements and compounds and understand the mechanism of chemical reactions. It should be kept in mind that every ingle the of text has potential of generating several MCQs.

As a first step to MCQ solving, students should start with the neuronatory mestion. Once a momentum is gained, complex MCQs with higher level of difficulty should be practised. Que, the from previous years as well as from other similar competitive exams should be solved to obtain an in "ght about plausible questions."

The competitive exams challenge understanding of  $\xi$  dents above subject by combining concepts from different chapters in a single question. To figure these q. for out, cognitive understanding of subject is required. Therefore, students should put in extra effort to practise such questions.

Such a holistic preparation is the key to such a different such a holistic preparation is the key to such a different such as the sum in the s

Our **Triumph Chemistry** book has tonde and a chieve the above objectives. Commencing from basic MCQs the book proceeds to develop concentence to solve complex MCQs. It offers ample practice of recent questions from various competiated in the solve to min. In It also includes hints that provide explanations and solutions to help students learn how solve t. MCQs. Each chapter ends with an Evaluation test to allow self-assessment.

Features of the book resen. <sup>1</sup> on ... next page will explicate more about the same!

We hope the boc per. <sup>3</sup>ts the le\_rner us we have envisioned.

The journey to rate a complete book is strewn with triumphs, failures and near misses. If you think we've nearly m<sup>2</sup> 1 son. Using or want to applaud us for our triumphs, we'd love to hear from you. Please vrite 1's on. hail@targetpublications.org

<sup>1</sup> boo. af cts e mity; one can never tell where its influence stops.

Best of luck to all the aspirants!

From Isher

Edition: First

#### **FEATURES**

Quick Review		
	Iron	
Cast iron	• Wrought iron	Steel
Contains 4% C	Contains less than 0.2% C	Contains 0.2 - 2% C
Hard and brittle	Very soft	Neither too hard nor too soft
Uses: making pipes, manufacturing automotive parts, pots, pans, utensils, etc.	Uses: making pipes, bars for stay bolts, engine bolts and rivets, etc.	Uses: in buildings infrastructure, tools, ships, automobiles, weapons, et

#### **Quick Review**

Quick Review includes tables, arts to summarize the key prime tant chemical reactions in ' e ch? .er This is our att.  $\gamma p^*$  to  $\gamma t$  stude s to reinforce key construction reinforce key constru

#### Formulae

Formulae includes all of the k. formulae in the chapter. This is our attempt to mak tools formulae accessible for studies while solving problems and ... sing ' .t minute at a glance

#### 1.

**Density** ( $\rho$ ) =  $\frac{Mn}{a^3 N_A}$ 

Formunae

Where M = molar mass of substance (g/mol), n = number of particles in a cubic unit cell, a = edge length (cm), $N_A = Avogadro number (6.022 \times 10^{23} mol^{-1})$ 

2. **Packing efficiency** 

#### Volume occupied by particles in unit cell ×100 Total volume of unit cell

#### **Classical Thinking**

(@\* 9.1 Introduction

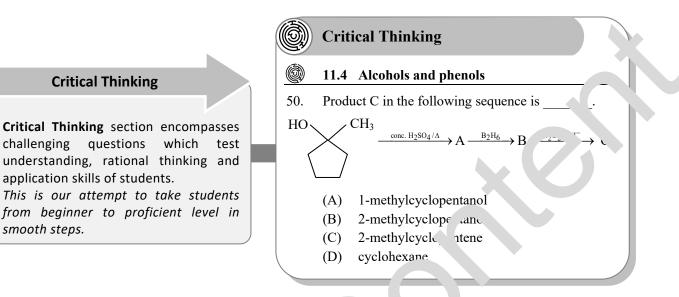
- Coordination compounds contain ligands attached to 1. central metal atom/ion through bonds. (A) covalent (B) ionic (C)
  - coordinate (D) metallic

#### **Classical Thinking**

Classical Thinking section encompasses straight forward questions including knowledge based questions.

This is our attempt to revise chapter in its basic form and warm up students to deal with complex MCQs.

#### **FEATURES**



#### **Competitive Thinking**

**Critical Thinking** 

application skills of students.

challenging

smooth steps.

#### 4.8 Enthalpy (H)

- $A_{(g)}+2B_{(g)} \longrightarrow 2C_{(g)}+3D_{(g)}$  for the sover action 13. the value of  $\Delta$ H is 19.0 kcal + 27°C. The value of  $\Delta U$  in kcal is (Given  $R = 2.0 \text{ cal } \text{K}^{-1} \text{ mol}^{-1}$ ) [M. T CET 2019]
  - 19.8 kcal (A)
  - (C) 18.8 kcal
- $2\sqrt{8}$  kcal
- (B) J)
  - 1' 8 kcal

#### **Competitive Thinking**

Competitive Thinking section encompasses questions from various competitive examinations like MHT CET, JEE, AIPMT/NEET-UG, etc.

This is our attempt to give students practice of competitive questions and advance them to acquire knack essential to solve such questions.

#### Sub wir wise segregation

Every section is segregated sub-topic ··· ·

This is our attempt to cater to individualistic pace and preferences of studying a chapter in students and enable easy assimilation of questions based on the specific concept.

#### Subtopics

- 1.1 Introduction
- 1.2 Types of solids
- 1.3 Classification of crystalline solids
- 1.4 Crystal structure
- Cubic system 1.5

#### **FEATURES**

#### Ò) Miscellaneous

76. The end product in the following sequence of reaction is

 $HC \equiv CH \xrightarrow{1\% HgSO_4} A \xrightarrow{CH_3MgX} B \xrightarrow{[O]} ?$ 

(B)

(D)

- (A) acetic acid
- (C) acetone

#### Miscellaneous

Every section, in general, ends with a sub-topic; miscellaneous.

Miscellaneous incorporates MCQs whose solutions require knowledge of cor covered in different sub-topics  $\gamma$  same chapter or from different chapters.

This is our attempt to dev \_\_\_\_\_ rogi, ve thinking in the students  $\epsilon$  senting to solve questions involving fusion of aultiper key concepts.

#### **Evaluation test**

Evaluation Test covers questions from chapter for self-evaluation purpose. This is our attempt to provide the students with a practice test and help them assess their range of preparation of the chapter.



1.

isopropyl alcohol

ethanol

#### Evaluari in Tes

Which of t' ... 'own cell the chemicals consumed d ring cur. it generation CANNOT be ated'. re

Lead students (B)(A)

(C)

- Merct y cell
- Dry cell (D) NICAD cell

Hair is primarily composed of keratin, a protein, which grows from a sac called the follicle. Cells in the hair follicle generate keratin, and various other proteins, which become a part of the hair shaft. These proteins contain sulfur atoms, and when two of these suy. atoms pair up and bodd, they form a disulfide on. If the two sulfur atc in the ne protein are ai distanc and joir rm the *isulfide* bond, ie pri 'n will end.

#### Straight or curly?



The state the mber of links, the curlier the hair, and the fewer the umbe of links, the straighter the hair.

#### **Real-world applications**

Each includes chapter real-world applications or examples related to the concept discussed.

This is our attempt to link learning to the life and make students conscious of how Chemistry is related to everything we see, feel, touch and taste.

22. Predict the products in the following reactions.

$$\underbrace{\bigcirc}_{\text{CO,HCl}} \xrightarrow{\text{CO,HCl}} \text{'B'} \xrightarrow{\text{HCHO}} \text{'C'} + \underbrace{\bigcirc}_{\text{CuCl}} \xrightarrow{\text{CO, MCl}} \text{'B'} \xrightarrow{\text{HCHO}} \text{'C'} + \underbrace{\bigcirc}_{\text{CuCl}} \xrightarrow{\text{CO, HCl}} \text{'B'} \xrightarrow{\text{HCHO}} \text{'C'} + \underbrace{\bigcirc}_{\text{CuCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \text{'B'} \xrightarrow{\text{HCHO}} \text{'C'} + \underbrace{\bigcirc}_{\text{CuCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{HCHO}} \text{'C'} + \underbrace{\bigcirc}_{\text{CuCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{HCHO}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{HCHO}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{HCHO}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text{HCHO}} \xrightarrow{\text{CO, HCl}} \xrightarrow{\text$$

- (A) Benzoic acid, benzyl acohol, sodium formate
- Benzaldehyde, sodium benzoate, methanol (B)
- Benzoic acid, sodium benzoate, methanol (C)
- (D) Benzaldehyde, benzyl alcohol, sodium formate

#### **Organic Reactions**

**Organic Reactions** – is a compilation of questions based on various organic chemistry concepts and reactions.

This is our attempt to help the students develop a strong understanding of organic chemistry.



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#### Disclaimer

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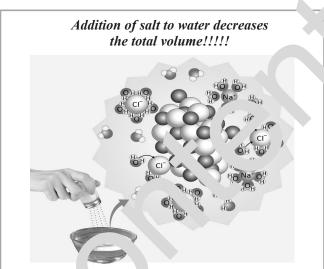
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#### Textbook Chapter No.

## 02 Solutions

#### Subtopics

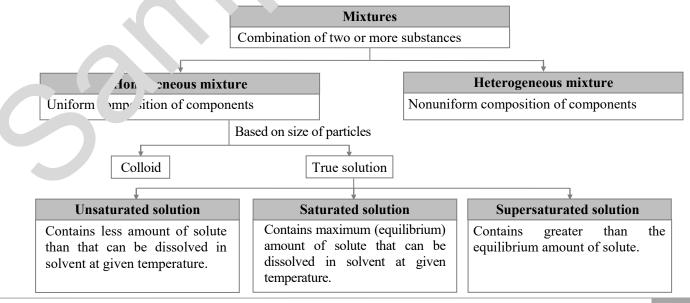
- 2.1 Introduction
- 2.2 Types of solutions
- 2.3 Capacity of solutions to dissolve solute
- 2.4 Solubility
- 2.5 Vapour pressure of solutions of liquids in liquids
- 2.6 Colligative properties of nonelectrolyte solutions
- 2.7 Vapour pressure lowering
- 2.8 Boiling point elevation
- 2.9 Depression in freezing point
- 2.10 Osmotic pressure
- 2.11 Colligative properties of electrolytes



*V* It is  $c^{-led}$  to the water there is an increase in the volume before . "c" dissolves in it. However, as the dissolution of "urs, there is a decrease in the total volume! This is due to be breaking of crystal structure of NaCl into "e free ior of Na<sup>+</sup> and Cl<sup>-</sup>. Moreover, the water n. "cules" and to the free ions i.e., the negative oxygen ends of water molecules surround the positive sodium ions (Na<sup>+</sup>) and the positive hydrogen ends surround the negative chlorine ions. Salt ions (being smaller than the water molecules) enter the voids between them and hence, cause shrinkage of the total volume.

#### Ouick Review

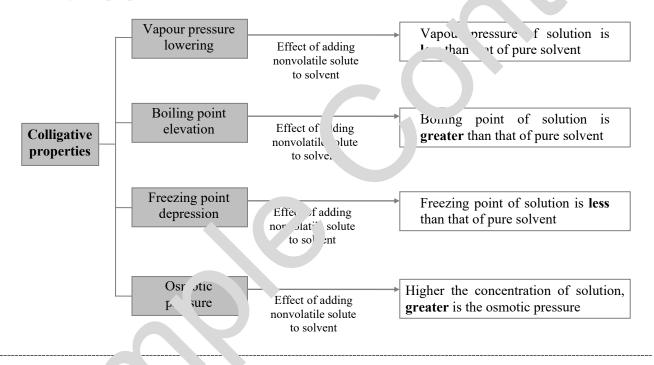
Classification f mix. "es:



#### **Types of solutions:**

State of solute	State of solvent	Examples
Solid	Liquid	Sea water, benzoic acid in benzene, sugar in water
Solid	Solid	Metal alloys such as brass, bronze.
Solid	Gas	Iodine in air
Liquid	Liquid	Gasoline, ethanol in water
Liquid	Solid	Amalgams of mercury with metals i.e., mercury in silver
Liquid	Gas	Chloroform in nitrogen
Gas	Liquid	Carbonated water (CO <sub>2</sub> in water), oxygen in water.
Gas	Solid	H <sub>2</sub> in palladium
Gas	Gas	Air (O <sub>2</sub> , N <sub>2</sub> , Ar and other gases)

#### **Colligative properties:**



#### > Types of so ation pending on the osmotic pressure of two solutions:

No.	Type of Solu .con	Definition	Example
i	• `tonìc	Two or more solutions having the	0.1 M urea solution and 0.1 M
	isc eans equal)	same osmotic pressure are said to be isotonic solutions.	sucrose solution are isotonic.
Ч,	pertonic	If two solutions have unequal	If osmotic pressure of sucrose
	(Hyper means	osmotic pressures, the more	solution is higher than that of urea
	higher)	concentrated solution with higher	solution, the sucrose solution is
		osmotic pressure is said to be	hypertonic to urea solution.
		hypertonic solution.	
iii.	Hypotonic	A solution having an osmotic	If osmotic pressure of sucrose
	(Hypo means	pressure lower than that of another	solution is higher than that of urea
	lower)	solution owing to lower	solution, the urea solution is
		concentration of solute is called	hypotonic to sucrose solution.
		hypotonic solution.	

20



1.

- Henry's law:  $S = K_H P$ where, S = solubility, P = pressure of the gas  $K_H =$  Henry's constant
- 2. Raoult's law: For a binary solution of two volatile components:

 $\mathbf{P}_1 = \mathbf{P}_1^0 \mathbf{x}_1$ 

 $P_2 = P_2^0 x_2$ 

where,

 $P_1^0$  is the vapour pressure of pure component 1 and  $P_1$  is the partial vapour pressure of component 1 in solution.

 $x_1$  is the mole fraction of component 1 in solution.

 $P_2^0$  is the vapour pressure of pure component 2 and  $P_2$  is the partial vapour pressure of component 2.

 $x_2$  is the mole fraction of component 2 in solution.

#### 3. Dalton's law of partial pressures:

 $P = P_1 + P_2$  OR  $P = P_1^0 x_1 + P_2^0 x_2$ where, P is the total pressure of solution.

#### 4. Partial pressures in vapour phase:

 $\mathbf{P}_1 = \mathbf{y}_1 \mathbf{P}$ 

 $P_2 = y_2 P$ where,  $y_1$  and  $y_2$  as the mole fination. If two components in the vapour.  $P_1$  and  $P_2$  are the part  $p_1$  sure. If two components in the vapor  $P_1$  is the total vapour press.  $P_2$ .

5. For a solution con vining a non-volatile solute:

 $P_1 = P_1^0 x_1$ 

where,  $P_1$  is 're vapor pressure of the solution,  $P_1^0$ ; vapor pressure of pure solvent and x is its in fraction in solution.

**Rela ve lo ering of vapour pressure:**   $^{\wedge}P_{+} = \frac{^{\circ} P_{1}}{P_{1}^{\circ}}$ w ere,  $P_{1}^{\circ}$  = vapour pressure of pure solvent and

 $r_1$  = vapour pressure of solution

7. Molecular mass determination from lowering of vapour pressure:

i. 
$$\frac{\Delta P}{P_1^0} = x_2 = \frac{P_1^0 - P_1}{P_1^0} = \frac{n_2}{n_1 + n_2}$$
  
where  $P_1^0 = Vanour pressure of$ 

where,  $P_1^0$  = Vapour pressure of pure solvent and  $P_1$  = Vapour pressure of solution

**Chapter 02: Solutions**  $x_2$  = Mole fraction of solute  $n_1 = Moles of solvent$  $n_2 = Moles of solute$  $n_2 = \frac{W_2}{M_2}$  and  $n_1 = \frac{W_1}{M_1}$ ii. where,  $W_2 = Mass$  of solute,  $W_1 = Mass of solvent,$  $M_2 = Molar mass of solute,$  $M_1 = Molar mass of solvent$ Relative lowering of vapour pressure. iii.  $\frac{P_1^0 - P_1}{P_1^0} = \frac{\triangle P}{P_1^0} = \frac{W_2 M_1}{W_1 M_2}$ 8. Elevation of boiling at:  $\Delta T_{\rm b} = T_{\rm b} - T_{\rm b}^0$ i. where,  $\Delta T_b = F$  svatic in bound point,  $T_b = Boiling$ , nt of soli on and  $T_{h}^{0} = Bc^{m}$  point of pure solvent.  $\Delta T_{\rm b} = K_{\rm b} m$ ii. wher m = Mc lity of solution  $K_b = bo$ . Jint elevation constant  $1000 \text{K}_{\text{b}} \text{W}_2$  $\Delta T_{h}$ iii.  $M_2W_1$ ...,  $\Delta T_b$  = Elevation in boiling point,  $K_b = Molal$  elevation constant,  $W_1 = Mass of solvent,$  $W_2 = Mass of solute,$  $M_2 = Molar mass of solute$ Molecular mass determination from elevation iv. of boiling point: Molecular mass of solute,  $M_2 = \frac{1000 \text{ K}_b \text{W}_2}{\Delta T_b \text{W}_1}$ 9. **Depression of freezing point:**  $\Delta T_f = T_f^0 - T_f$ i. where,  $\Delta T_f$  = Depression in freezing point,  $T_f =$  Freezing point of solution and  $T_{f}^{0}$  = Freezing point of pure solvent. ii.  $\Delta T_f = K_f m$ where, m = Molality of solution  $K_f =$  Freezing point depression constant  $\Delta T_{\rm f} = \frac{1000 K_{\rm f} W_2}{M_2 W_1}$ iii. where,  $\Delta T_f =$  Depression in freezing point,

 $K_f =$  Molal depression constant,

 $W_2 = Mass of solute,$ 

 $W_1 = Mass of solvent,$ 

 $M_2 = Molar mass of solute$ 

- iv. Molecular mass determination from depression of freezing point: Molecular mass of solute,  $M_2 = \frac{1000 \text{ K}_f W_2}{\Lambda \text{ T} W}$  $\Delta T_{\rm f} W_{\rm I}$ van't Hoff equation for osmotic pressure: 10.  $\pi = MRT = CRT$ where,  $\pi = Osmotic$  pressure, M = C = Concentration of solution inmol  $L^{-1}$ , R = Gas constantT = Temperature in Kelvin
  - 11. Molecular mass from osmotic pressure:  $\pi = \frac{W_2 RT}{M_2 V} \quad OR \qquad M_2 = \frac{W_2 RT}{\pi V}$ where,  $\pi = O$ smotic pressure, R = Gas constant (0.08205 atm dm<sup>-3</sup> K<sup>-1</sup> mol<sup>-1</sup>),  $M_2 = M$ olecular mass of solute,  $W_2 = M$ ass of solute,
    - T = Temperature in Kelvin,
    - n = number of moles of solute,
    - $V = volume in dm^3$

#### 12. van't Hoff factor (i):

- $i = \frac{\text{Colligative property of electrolyte solution}}{\text{Colligative property of nonelectrolyte solution}}$ of the same concentration
  - Actual moles of particles in solution

after dissociation

- Moles of formula units dissolve 'n sol, 'n Formula mass of substance
- Observed molar mass of s \_\_\_\_\_
- $= \frac{M_{\text{Theoretical}}}{M_{\text{Observed}}}$
- 13. Modified equations for colling five properties by inclusion . n't Hor. 'rctor.

i. 
$$\Delta P = i P_1^{0} \lambda_2 - i \frac{W_2 M_1}{M_2 W_1}$$
  
ii.  $\Delta^{r} \lambda_3 = i k = i - \frac{00 K_b W_2}{M_2 W_1}$   
iii.  $\Delta T_f = i K_f r = i \frac{1000 K_f W_2}{M_2 W_1}$   
iv.  $\pi = i MRT = i \frac{W_2 RT}{M_2 V}$   
14. Degree of dissociation ( $\alpha$ ):

 $\alpha = \frac{i-1}{n-1}$ where,  $\alpha$  = Degree of dissociation, i = van't Hoff factor, n = Moles of ions obtained from dissociation of 1 mole of electrolyte Classical Thinking

#### \* 21 Introduction

	2.1	Introduction		
1.	diame	solutions contain		nm.
	(A) (C)	0.1 to 2 10 to 200		1 to 20 100 to 2000
2.	compo (A)	solution, the larg onent is known as _ solution solvent	(B)	-
	2.2	Types of solu 'on	×	
3.		nding on the phy in es, there are five nine	(B)	vents and s of solutions. veven twelve
4.	Sugar solut <sup>;</sup>	discolvea wate	r is a	type of
	(A)		(B) (D)	solid in gas gas in solid
5.	type	by 15 a nomogeneo solution. solid in solid solid in liquid	(B)	solid in gas
6.		example for solid		
	(A) (C)	bronze H <sub>2</sub> in palladium		iodine in air gasoline
	2.3	Capacity of soluti	ons to	dissolve solute
7.	amour some be (A)	continue the additi nt of solvent, the time. The solution supersaturated unsaturated	dissolution dissolution dissolution dissolution dissolution distribution distribution distribution dissolution distribution di distribu	ution stops after
<b>@</b> *	2.4	Solubility		
8.	Solub unit(s) (A) (C)	ility is expressed ) of mol L <sup>-1</sup> mole fraction		me concentration mol $g^{-1}$ mol $kg^{-1}$
9.		ility of which of slightly with temper NaBr KBr	ature? (B)	llowing changes KNO3 NaNO3
10.	The s $\frac{(A)}{(B)}$ (C)	olubility of a gas increase in tempera reduction in gas pr decrease in temper	ature essure	er increases with

(D) amount of liquid taken

11. Henry's law constant is expressed in the unit(s) of \_\_\_\_\_.

 $(A) \quad mol^{-1} L^{-1} bar^{-1}$ 

- (B) mol  $L^{-1}$  bar
- (C)  $L \mod^{-1} \operatorname{bar}^{-1}$
- (D) mol  $L^{-1}$  bar<sup>-1</sup>

#### 2.5 Vapour pressure of solutions of liquids in liquids

- 12. Partial vapour pressure of any volatile component in a solution is \_\_\_\_\_.
  - (A) directly proportional to its mole fraction in the solution
  - (B) inversely proportional to its mole fraction in the solution
  - (C) inversely proportional to the mole fraction of the solvent
  - (D) directly proportional to its molarity
- 13. An ideal solution is that which \_\_\_\_\_ over the entire range of concentration.
  - (A) shows positive deviation from Raoult's law
  - (B) shows negative deviation from Raoult's law
  - (C) obeys Raoult's law
  - (D) Both (A) and (B)
- 14. Which of the following behaves as an ideal solution?
  - (A) Ethanol + acetone
  - (B) Carbon disulphide + ace.
  - (C) Benzene + toluene
  - (D) Chloroform + acetor
- 15. Which of the folloing show negative deviation from Raoult's law?
  - (A) Benzene + Juc.
  - (B) Phenol iline
  - (C) Ethar 1 + a, 'one
  - (D) Cart disulpt. '-+ acetone

#### 2.6 C 'ligat, properties of nonelectrolyte

- 16. Which of t' 2 following is a colligative property?
  - (B Surface tension of a solution
  - (' , Boiling point elevation
  - (D) Radioactivity of a substance
  - 17. While dealing with colligative properties of nonelectrolyte solutions, the relatively dilute solutions with concentrations \_\_\_\_\_\_ or less are considered.
    (A) 0.2 M
    (B) 1 M
    (C) 2 M
    (D) 0.5 M

18. V

0

2.7

18. Vapour pressure of the solution, of a non-volatile solute is always \_\_\_\_\_.

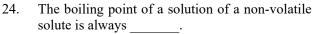
Vapour pressure lowering

- (A) equal to the vapour pressure of pure solvent
- (B) higher than vapour pressure of pure s vert
- (C) lower than vapour pressure of p solvent
- (D) constant
- 19. The lowering of vapour pressure of . Ivent by the addition of a non-volatile solute it, is equal to .
  - (A) the product of vapor presure of pure solvent and solvent free of he nonvolatile solute
  - (B) the product of var in pressure of pure solvent in the fraction of the pure solven
  - (C) the sun of vapour pressure of pure and mole fraction of the nonvolatile lute
  - (D) 'he sum of vapour pressure of pure 'vent and mole fraction of the pure solvent
- 20. At 30 K, when a solute is added to a solvent, its pour pressure over mercury reduces from mm to 45 mm. The value of mole fraction of solute will be

21. The mathematical expression for relative lowering of vapour pressure is

(A) 
$$\frac{P_1^0 - P_1}{P_1^0} = \frac{W_2 M_1}{W_1 M_2}$$
 (B)  $\frac{P_1 - P_1^0}{P_1} = \frac{W_2 M_1}{W_1 M_2}$   
(C)  $\frac{P_1 - P_1^0}{P_1^0} = \frac{W_2 M_1}{W_1 M_2}$  (D)  $\frac{P_1^0 - P_1}{P_1^0} = \frac{W_1 M_1}{W_2 M_2}$ 

- **2.8** Boiling point elevation
- 22. Boiling point of water is defined as the temperature at which \_\_\_\_\_.
  - (A) vapour pressure of water equals to that of atmospheric pressure
  - (B) bubbles are formed
  - (C) steam comes out
  - (D) vapour pressure of water is higher than that of atmospheric pressure
- 23. Which of the following statement is CORRECT for the boiling point of solvent, containing a dissolved solid substance?
  - (A) Boiling point of the liquid is lowered.
  - (B) Boiling point of the liquid is elevated.
  - (C) There is no effect on the boiling point.
  - (D) Boiling point of the liquid becomes equal to the boiling point of water.



- lower than the boiling point of the solvent (A)
- higher than the boiling point of the (B) solvent
- equal to the boiling point of the solvent (C)
- (D) independent of the boiling point of the solvent
- 25. The molal elevation constant is the ratio of the elevation of boiling point to
  - molarity of the solution (A)
  - (B) molality of the solution
  - (C) mole fraction of the solute
  - (D) mole fraction of the solvent
- Unit of boiling point elevation constant (K<sub>b</sub>) is 26.

$\overline{(A)}$	kg mol <sup>-1</sup>	(B)	K $mol^{-1}$
(C)	g mol <sup>-1</sup>	(D)	K kg $mol^{-1}$

The boiling point of 0.15 molal aqueous 27. solution of an unknown solute is 373.23 K at 1 atm. The molal elevation constant of water is K kg  $mol^{-1}$ .

> (A) 0.53 (B) 0.88 (C) 1.8 (D) 5.3

K<sub>b</sub> is given by \_\_\_\_\_. 28.

(A) 
$$\frac{\Delta T_{b} \times W_{2} \times M_{2}}{1000 \times W_{1}}$$
 (B) 
$$\frac{W_{2} \times 1000}{\Delta T_{b} \times W_{1} \times M_{2}}$$
  
(C) 
$$\frac{\Delta T_{b} \times W_{1} \times M_{2}}{1000 \times W_{2}}$$
 (D) 
$$\frac{W_{1} \times 1000}{\Delta T_{b} \times W_{2} \times W_{2}}$$

int

- 29. The temperature at which the vapo ores. a solid is equal to the vapor ssur of liquid is called
  - elevation of boilin, voint (A)
  - (B) freezing poir\*
  - (C) boiling poi *c*
  - (D) depres of freez. poin.
- Solute why dissolv in water 30.
  - (A) increa. s the vat our pressure of water
  - (B) .. rease. he boiling point of water
  - (f , d , ses the freezing point of water
  - Acres the freezing point of water (1)
- Depr. : , in freezing point in any dilute 31. sc tion containing a non-volatile solute is di ctly proportional to
  - molarity of the solution 15
  - **(B)** molality of the solution
  - (C) mass of solvent
  - (D) mole fraction of solvent
  - 32. Relationship between  $K_f$ , m and  $\Delta T_f$  can be written as

(A) 
$$\Delta T_f = K_f / m$$
 (B)  $\Delta T_f = K_f m$   
(C)  $\Delta T_f = K_f + m$  (D)  $\Delta T_f = m / K_f$ 

- 33. What is the molality of solution of a certain solute in a solvent, if there is a freezing point depression of 0.184 °C, and if the freezing point depression constant is  $18.4 \text{ K kg mol}^{-1}$ ?
  - (A) 0.01 m (B) 1 m
  - 0.001 m (D) 100 m (C)
- 34. The molar mass of the solute using depress of freezing point may be calculated using the formula, .

(A) 
$$M_2 = \frac{K_f W_2 1000}{\Delta T_f m}$$
 (B)  $M_2 = \int \frac{1}{r} \frac{W_1 10 v}{V_T W_2}$   
(C)  $M_2 = \frac{\Delta T_f W_2 1000}{K_f W_1}$  (D)  $M_2 = \frac{K_f v}{\Delta T_f W_1} \frac{1000}{\Delta T_f W_1}$ 

#### (@ 2.10 Osmotic press

- 35. A membrane which allow solvent molecules but NOT the scate n. 'ecules to pass through it is called as
  - (A) semipern ble membrane
  - Jerm 'len mbrane (B)
  - (C) filter mu brane
  - (D)porous r embrane
- 36. During comosis, flow of water through a semi rmeable membrane is
  - unidirectional bidirectional (A) **(B)**
  - (C)multidirectional unpredictable (D)
- 37. Osmosis is a process in which
  - solvent molecules flow through a semi-(A) permeable membrane from a solution of lower concentration to a solution of higher concentration
  - solute molecules flow through a semi-(B) permeable membrane from a solution of lower concentration to a solution of higher concentration
  - solvent molecules flow through a semi-(C) permeable membrane from a solution of higher concentration to a solution of lower concentration
  - solute molecules flow through a semi-(D) permeable membrane from a solution of higher concentration to a solution of lower concentration
- 38. A solution having a higher osmotic pressure than another solution is called a .
  - hypotonic solution (A) (B) isotopic solution
  - (C) isotonic solution
  - hypertonic solution (D)
- 39. If two solutions separated by a semipermeable membrane have the same osmotic pressure, they are called solutions.
  - hypertonic (A) (B) hypotonic (C)
    - isotonic (D) saturated

#### **Chapter 02: Solutions**

- 40. At constant temperature, the osmotic pressure of a solution is
  - (A) directly proportional to the concentration
  - inversely proportional to the (B) concentration
  - directly proportional to the square of the (C) concentration
  - directly proportional to the square root of (D) the concentration
- 41. If mole fraction of the solvent in a solution decreases, then .
  - (A) vapour pressure of solution increases
  - (B) boiling point decreases
  - (C) osmotic pressure increases
  - osmotic pressure decreases (D)
- 42. Which statement is INCORRECT about osmotic pressure  $(\pi)$ , volume (V) and temperature (T)?
  - (A)  $\pi \propto 1/V$  if T is constant.
  - $\pi \propto T$  if V and n are constant. (B)
  - $\pi \propto V$  if T is constant. (C)
  - (D)  $\pi V$  is constant if T and n are constant.
- 1 M and 2 M solutions of glucose are prepared 43. in water. Hence,
  - the osmotic pressure of both the solutions (A) will be the same at the same temperature
  - 2 M solution will have higher (B) tic pressure
  - (C) 1 M solution will hav, highe or notic pressure
  - osmotic pressure will be ina, endent of (D) the concentration
- 44. The molar mass  $(M_2)$  or  $V_2$  g sc ite and the osmotic pressure ., the state in prepared in V litres by the solute a smpth fure T has the following re' .ac. hip •

(A) 
$$M_2 = \frac{W_2 RT}{V}$$
 (B)  $M_2 = \frac{W_2 R}{\pi T}$   
(C  $M_2 = mR_2$  (D)  $M_2 RT = \pi$ 

$$M_0 = mR_1 = \pi$$

The rectin of osmosis can be reversed by apply or pressure larger than the - SSURC.

(A	absolute	(B)	atmospheric
1	osmotic	(D)	vapour

#### 2.11 Colligative properties of electrolytes

- 46. The van't Hoff factor will be highest for
  - sodium chloride (A)
  - (B) magnesium chloride
  - (C) sodium phosphate
  - (D) urea

- 47. Which of the following has the highest boiling point?
  - (A) 1 m glucose solution
  - 1 m CH<sub>3</sub>COOH solution (B)
  - 1 m Na<sub>2</sub>SO<sub>4</sub> solution (C)
  - 1 m urea solution (D)
- 48. van't Hoff factor is
  - less than one in case of dissociation (A)
    - (B) always more than one
    - (C) always less than one
    - greater than one in case of dissoc 'ion (D)
- 49. Which of the following is NOT .ne cc. ect modified equation (f collig 'iv' prop ties for electrolyte solutions?

(A) 
$$\Delta P = i \frac{W_2 M_1}{M_1}$$
  $\Delta T_b = i K_b m$ 

(C) 
$$\Delta T_f = \frac{1}{1} \frac{\partial O K_f W_2}{\partial W_1}$$
 (C)  $\pi = i \frac{M_2 RT}{W_2 V}$ 

Misc \_aneo.

5 cm of methy alcohol is added to 100 cm<sup>3</sup> of water, 'n t' vapour pressure of the solution

- (A) vill be equal to the vapour pressure of pure water
- will be less than the vapour pressure of pure water
- will be greater than the vapour pressure of (C) pure water
- will be very large (D)
- Select the CORRECT option for a solution 51. containing non-volatile solute in it.

(A) 
$$T_{f}^{0} < T_{f}$$
 (B)  $T_{b}^{0} < T_{b}$   
(C)  $\Delta P = P_{l}^{0} x_{l}$  (D)  $\Delta T_{f} = T_{f} - T_{f}^{0}$ 

- 52. When swimming for a long time in salt water, the skin of one's fingertips wrinkles. Which one of the following properties is responsible for this observation?
  - (A) Osmosis (B) Dialysis
  - Electrodialysis (D) Coagulation (C)
- When a substance is dissolved in a solvent, the 53. vapour pressure of the solvent is decreased. This results in
  - an increase in the boiling point of the (A) solution
  - a decrease in the boiling point of the **(B)** solution
  - no change in the boiling point of the (C) solution
  - an initial decrease in the boiling point (D) followed by a sharp increase in the boiling point of the solution

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To see complete chapter buy Target Notes or Target E-Notes

919]

72. On dissolving 3.24 g of sulfur in 40 g of benzene, boiling point of solution was higher than that of benzene by 0.81 K. K<sub>b</sub> value of benzene is 2.53 K kg mol<sup>-1</sup>. Atomic mass of sulfur is 32 g mol<sup>-1</sup>. The molecular formula of sulfur is \_\_\_\_\_.

(A)  $S_6$  (B)  $S_7$  (C)  $S_8$  (D)  $S_9$ 

- 73. Pressure cooker reduces cooking time for food because \_\_\_\_\_.
  - (A) heat is more evenly distributed in the cooking space
  - (B) boiling point of water involved in cooking is increased
  - (C) the higher pressure inside the cooker crushes the food material
  - (D) cooking involves chemical changes helped by a rise in temperature

#### **Competitive Thinking**

#### 2.4 Solubility

- The relation between solubility of a gas in liquid at constant temperature and external pressure is stated by which law? [MHT CET 2016]
   (A) Beauk's law?
  - (A) Raoult's law
  - (B) van't Hoff Boyle's law
  - (C) van't Hoff Charles' law
  - (D) Henry's law
  - 2.5 Vapour pressure of selutions f lie .ds in liquids
- 2. The vapour pressures of pure liquid A and B are 400 and 600 mm lig, specifiely at 298 K. On mixing the poliquid, the sum of their initial volumes is equal to the volume of the final mixture in mole final of liquid B is 0.5 in the mixture. The vapour pressure of the final solution in mole fractions components A and F n vapor phase, respectively are [JEE (Main) 2019]

 (A)
 ) mm
 '7, 0.5, 0.5

 (I)
 4'
 m Hg, 0.4, 0.6

 (I)
 J0 r
 Hg, 0.5, 0.5

 (D)
 500 m Hg, 0.4, 0.6

1. mole fractions of benzene and toluene vapurs in equilibrium with the ideal solution of azene in toluene at 300 K are 0.61 and 0.39 respectively. The total vapour pressure of the solution is 41 mm Hg. If the vapour pressures of pure benzene and toluene at 300 K are 50 and 32 mm Hg respectively, the mole fractions of benzene and toluene in solution respectively are

	•	AP LAMC	EI (Mea.) 2019
(A)	0.25, 0.75	(B)	0.75, 0.25
(C)	0.30, 0.70	(D)	0.50, 0.50

4. The vapour pressures of pure heptane and pure octane are 92 and 31 torr, respectively at 40 °C. The total vapour pressure (in torr) of a solution containing 1.00 mole of heptane and 4.00 moles of octane is

		TS EAMO	CET (Med.) 2
(A)	18.4	(B)	24.8
(C)	43.2	(D)	51.2

5. Which of the following state ents CORRECT regarding a solution of two components A and B exb<sup>:1.:</sup>ing sitive deviation from ideal behavior.?

**INE T** Jdisl ) 2019]

- (A) Intermolecular ractiv force between A–A and B–B are vial to mose between A–B.
- (B) Interm<sup>1</sup> cular a, ctive forces between A-A an, B-B are stronger than those <sup>1</sup> ... <sup>1</sup> A-
- (C)  $\Delta_{\text{mix}}$  H = at constant T and P.
- (D)  $_{\text{mix}}$ V = at constant T and P.

#### 2.7 Vapour pressure lowering

ő

7.

- 9 g
   g lucose (mol wt = 180) is dissolved in

   G' g of H<sub>2</sub>O. Relative lowering of vapour

   pressure is \_\_\_\_\_.
   [MH CET 2011]

   (A)
   0.99
   (B)
   0.099

   (C)
   0.0099
   (D)
   0.00099
- Relative lowering of vapour pressure of a dilute solution of glucose dissolved in 1 kg of water is 0.002.

The molality of the solution is \_\_\_\_\_

			[KCET 2019]
(A)	0.111	(B)	0.021
(C)	0.004	(D)	0.222

- 8. On dissolving 18 g solid in 100 g H<sub>2</sub>O at 20 °C, water vapour pressure decreases from 17.53 mm to 17.22 mm. The molecular weight of solid is \_\_\_\_\_. [BCECE 2014] (A)  $18 \text{ g mol}^{-1}$  (B)  $183 \text{ g mol}^{-1}$ 
  - (C)  $27 \text{ g mol}^{-1}$  (D)  $274 \text{ g mol}^{-1}$
- 9. The vapour pressure of acetone at 20 °C is 185 torr. When 1.2 g of a non-volatile substance was dissolved in 100 g of acetone at 20 °C, its vapour pressure was 183 torr. The molar mass (g mol<sup>-1</sup>) of the substance is \_\_\_\_\_.

	,	[,	JEE (Main) 2015]
(A)	32	(B)	64
(C)	128	(D)	488

10. Vapour pressure of pure solvent and its solution at certain temperature are 660 mm and 600 mm of Hg respectively. If  $3.6 \times 10^{-3}$  kg of solute is added into  $40 \times 10^{-3}$  kg of solvent, what is the molar mass of solute? (solvent = Benzene, C = 12, H = 1)

[MH CET 2013]

- (A)  $78.0 \text{ g mol}^{-1}$  (B)  $58.5 \text{ g mol}^{-1}$
- (C)  $72.0 \text{ g mol}^{-1}$  (D)  $156 \text{ g mol}^{-1}$
- 11. The vapour pressure of a solution of 6.0 g of non-volatile solute in 390 g of benzene at 298 K is 3.00 kPa. If 78 g of benzene is added to this solution the vapour pressure becomes 3.02 kPa at the same temperature. The molar mass of solute in g mol<sup>-1</sup> is

	-	[AP EAMCET (Med.) 2019]
(A)	60.8	(B) 50.4
(C)	31.2	(D) 21.2

12. 18 g glucose  $(C_6H_{12}O_6)$  is added to 178.2 g water. The vapour pressure of water (in torr) for this aqueous solution is

		[ <b>JEE</b> (Main) 2016]
(A)	7.6	(B) 76.0
(C)	752.4	(D) 759.0

#### 2.8 Boiling point elevation

13. An aqueous dilute solution containing nonvolatile solute boils at 100.052 °C. What is the molality of solution? ( $K_b = 0.52 \text{ kg r}$ Boiling temperature of water = 100 °C [TS EAMCL `(En. 2015]

(B)

0. m

1.0 n.

(A) 0.1 m

- (C) 0.001 m (D)
- 14. The CORRECT relation between elevation of boiling point and multimeters of solute is \_\_\_\_\_. CET 2018]
  - (A)  $M_2 = \frac{K_b \cdot V_2}{r_b}$  (C)  $M_2 = \frac{\Delta T_b \cdot K_b}{r_c}$  (D)  $M_2 = \frac{\Delta T_b \cdot K_1}{K_b \cdot W_2}$
- 15. If ne e<sup>1</sup>, 'ion n boiling point of a solution of 5 g sol (molar mass = 100) in 500 g of wate is  $\Delta^r_{b}$ . The ebullioscopic constant  $K_b$  of ter \_\_\_\_ual to . [MHT CET 2019]  $\Delta T_{b}$ (B) (A) $100 \Delta T_{\rm h}$ 50  $10 \Delta T_{\rm h}$ (D)  $\Delta T_{h}$ . Ľ) 2.9 **Depression in freezing point**
- 16. If molality of the dilute solution is doubled, the value of molal depression constant (K<sub>f</sub>) will be

   [NEET (UG) 2017]
   (A) halved
   (B) tripled
   (C) unchanged
   (D) doubled

17. Find the K<sub>f</sub> if 6 g of urea is dissolved in 0.1 dm<sup>3</sup> of water and it corresponds to 0.15 °C in  $\Delta T_{f}$ . (Molecular weight of urea = 60 g mol<sup>-1</sup>)

[MH CET 2010]

- (A) 0.015 (B) 0.15 (C) 0.30 (D) 0.030
- 18. After adding non-volatile solute, freezing p<sup>'</sup> of water decreases to  $-0.186 \,^{\circ}\text{C}$ . Calculate  $\Delta T_b$  if  $K_f = 1.86 \,\text{K}$  kg r  $51^{-1} \,^{\circ}$  4  $K_b = 0.521 \,\text{K}$  kg mol<sup>-1</sup>. [K T 2015] (A) 0.521 K (B) 0.0521
  - (C) 1.86 K (D) 0.075 K
- 19. 6 g of a mixture  $(n \text{ pht}, 1 \text{ re} (C H_8) \text{ and}$ anthracene  $(C_{14}H_{10})$  disson 1 is 500 g ofbenzene. If the depressic in freezing point is 0.70 K, the composition completion completion and anthracene is the mixtor respectively in g are

(mola <sup>1</sup> , ss	ic. constant of benzene is
5.1 F кg mol	[A: EAMCET (Engg.) 2019]
(A) <sup>2</sup> .60, 3.4	(B) 3.40, 2.60
$(C) \ 2 \ 2 \ J$	(D) 3.10, 2.90
0.10 0 /	

- 2.10 <u>Osmotic press</u>ure
- 0. 0.0€ % (W/V) aqueous solution of urea is conic with \_\_\_\_\_.

   (1)
   0.06 % (W/Z) = .
  - (A) 0.06 % (W/V) glucose solution(B) 0.6 % (W/V) glucose solution
  - (C) 0.01 M glucose solution
  - (D) 0.1 M glucose solution
- 21. If M, W and V represent molar mass of solute, mass of solute and volume of solution in litres respectively, which among following equations is TRUE? [MH CET 2015]

22. The osmotic pressure of solution containing  $34.2 \text{ g of cane sugar (molar mass} = 342 \text{ g mol}^{-1})$ in 1 L of solution at 20 °C is (Given: R = 0.082 L atm K<sup>-1</sup> mol<sup>-1</sup>) [MHT CET 2017] (A) 2.40 atm (B) 3.6 atm (C) 24 atm (D) 0.0024 atm

23. The osmotic pressure of solution at 0 °C is 4 atm. What will be the osmotic pressure at 546 K under similar condition?

#### [MHT CET 2019]

(A)	2 atm	(B)	8 atm
(C)	4 atm	(D)	0.5 atm

			Chapter 02: Solutions
24.	$30 \times 10^{-4}$ kg of urea dissolved in water to make 500 mL aqueous solution and this solution is isotonic with cane-sugar solution. How much mass of cane sugar in its one litre solution? (H=1, N=14, O=16, C=12) [MH CET 2013] (A) 17.1 g (B) 171.0 g (C) 3.42 g (D) 34.2 g	33.	van't Hoff factor of centimolal solution of $K_3[Fe(CN)_6]$ is 3.333. Calculate the percent dissociation of $K_3[Fe(CN)_6]$ .[MH CET 2015](A) 33.33(B) 0.78(C) 78(D) 23.33
25.	Pure water can be obtained from sea water by(A) centrifugation(B) plasmolysis(C) reverse osmosis(D) sedimentation	34.	Miscellaneous In water saturated air, the mole fraction of w. hr vapour is 0.02. If the total pres. to of the saturated air is 1.2 atm, the partial produce of
8	2.11 Colligative properties of electrolytes		dry air is [NF : T (' dis' ۱) 2019]
26.	Of the following 0.10 m aqueous solutions, which one will exhibit the largest freezing point depression?[AIPMT 2014](A) KCl(B) $C_6H_{12}O_6$ (C) $Al_2(SO_4)_3$ (D) $K_2SO_4$	35.	(A) $0.98 \text{ atm}$ (B) $(.18 \text{ at}^{-1})$ (C) $1.76 \text{ atm}$ (C) $1.76 \text{ atm}$ (C) $1.76 \text{ atm}$ (C) $1.70 \text{ atm}$ For an ideal solution, $\circ$ CORRECT option is <b>NEET (UG) 2019</b>
27.	Identify the compound amongst the following of which 0.1 M aqueous solution has highest boiling point.(A) Glucose(B) Sodium chloride(C) Calcium chloride(D) Ferric chloride		(A) $4 = 40$ a onstant T and P (B) $\Delta_{mix} H = 4$ constant T and P (C) $4 = 4$ constant T and P (D) $\Delta_{mix} 0$ at constant T and P
28.	The van't Hoff factor (i) for a dilute aqueous solution of the strong electrolyte barium hydroxide is [NEET P-II 2016] (A) 3 (B) 0 (C) 1 ( $\Gamma$ )	36.	From he given graph at constant temperature, wh: I gas has the least solubility? [GUJ CET 2014]
29.	For which among the fol. ving ui slar aqueous solutions, van't Hoff ctor is the lowest value? [MH1 'ET 2000 (A) (A) Aluminium chloride (B) Potassium sulpha' (C) Ammonium chlorios (D) Urea		(A) Gas - D (B) Gas - B
30.	Which one $(1, followin_{12})$ elect. Lytes has the same valy $(1)$ of van Hoff's factor (i) as that of Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub> (1) II are 10 % ionised)? [AIPMT 2015] (() K 5 (B) K <sub>3</sub> [Fe(CN) <sub>6</sub> ] (() Al(N $_{3/3}$ (D) K <sub>4</sub> [Fe(CN) <sub>6</sub> ]	37.	(C) Gas - A (D) Gas – C A non-volatile solute, 'A' tetramerises in water to the extent of 80%. 2.5 g of 'A' in 100 g of water, lowers the freezing point by 0.3 °C. The molar mass of A in g mol <sup>-1</sup> is (K <sub>f</sub> for water = $1.86$ K kg mol <sup>-1</sup> )
31	't factor for aqueous monofluoroacetic		[KCET 2019]
	ac is [MH CET 2013] () $i = 1 + 2 \alpha$ (B) $i = 1 - 2\alpha$ (C) $i = 1 + \alpha$ (D) $i = 1 - \alpha$		(A)155(B)354(C)62(D)221
32.	vant Hoff's factor for 0.1 m Ba(NO <sub>3</sub> ) <sub>2</sub> solution is 2.74. The percentage dissociation of it is [MHT CET 2019; Similar in Assam CEE 2015]	38.	To observe an elevation of boiling point of 0.05 °C, the amount of solute (Mol. Wt. = 100) to be added to 100 g of water ( $K_b = 0.5$ ) is [WB JEEM 2014]
	(A) 90% (B) 100%		(A) 2 g (B) 0.5 g
	(C) 87% (D) 75%		(C) 1 g (D) 0.75 g

39. The freezing point of benzene decreases by 0.45 °C when 0.2 g of acetic acid is added to 20 g of benzene. If acetic acid associates to form a dimer in benzene, percentage association of acetic acid in benzene will be (K<sub>f</sub> for benzene = 5.12 K kg/mol)

[**JEE** (Main) 2017]

- (A) 61.2 % (B) 75.6 % 94.6 % (C) 82.7 %
  - (D)
- 40. At 100 °C, the vapour pressure of a solution of 6.5 g of a solute in 100 g water is 732 mm. If  $K_b = 0.52$ , the boiling point of this solution will be [NEET P-I 2016] 102 °C (B) 103 °C (A) (C) 101 °C (D) 100 °C
- 41. The freezing point of solution containing 10 mL of non-volatile and non-electrolyte liquid "A" in 500 g of water is -0.413 °C. If K<sub>f</sub> of water is 1.86 K kg mol<sup>-1</sup> and the molecular weight of A = 60 g mol<sup>-1</sup>, what is the density of the solution in g mL<sup>-1</sup>? (Assume  $\Delta_{mix} V = 0$ )

[TS EAMCET (Engg.) 2019] 1.13 1.3 **(B)** 0.993 (D)

0.90 (C)

(A)

Why salt is applied on leeches?

When leeches are stuck to any bod the e. est way to get rid of them is to  $ap_1 \lor sal^t \land n$  them. It involves simple phenome on of c nc s res 'ing in the drying up of the org. m wh enour salt is applied. Some worms or pests 'e the s..., seech have no protective barrier ween i. cell walls and the outside world. On a Jicai. of sai. A their body, the high concentration of the solution of the outside of the organism initiates process of osmosis wherein water from "s of panism moves out of its body and the c ganism c 'es up

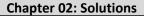
Answei *zev* 

	Cla	ssica	l Thi	nkin	g														
1.	(A)	2.	(C)	3.	(C)	4.	(C)	5	(A	6.	(A)	7.	(B)	8.	(A)	9.	(A)	10.	(C)
11.	(D)	12.	(A)	13.	(C)	14.	(Ċ)	<b>.</b>	(F	16.	(C)	17.	(A)	18.	(C)	19.	(A)	20.	(C)
21.	(A)	22.	(A)	23.	(B)	24.	(т	25.	-(B)	26.	(D)	27.	(A)	28.	(C)	29.	(B)	30.	(C)
31.	(B)	32.	(B)	33.	(A)	34.	<b>D</b> )	٦5.	(A)	36.	(A)	37.	(A)	38.	(D)	39.	(C)	40.	(A)
41.	(C)	42.	(C)	43.	("	44.		45.	(C)	46.	(C)	47.	(C)	48.	(D)	49.	(D)	50.	(C)
51.	(B)	52.	(A)	53.	(A,	54.	( )	55.	(A)										

Critical In ing

1.	(C)	2.	$\langle v \rangle$	3.	<b>B</b> )	4.	(D)	5.	(A)	6.	(C)	7.	(B)	8.	(D)	9.	(B)	10.	(C)
11.	(B)		(Ċ,	13.	(C)	14.	(C)	15.	(C)	16.	(D)	17.	(A)	18.	(A)	19.	(D)	20.	(C)
21.	(A)	22	<b>B</b> )	3.	(A)	24.	(A)	25.	(A)	26.	(A)	27.	(A)	28.	(C)	29.	(C)	30.	(B)
31.	(/	3.	$\sum$	33.	(C)	34.	(C)	35.	(B)	36.	(B)	37.	(B)	38.	(D)	39.	(C)	40.	(B)
41.	(C)	2.	/ A)	43.	(C)	44.	(B)	45.	(B)	46.	(D)	47.	(D)	48.	(D)	49.	(A)	50.	(A)
51.	( <b>L</b>		(B)	53.	(A)	54.	(B)	55.	(D)	56.	(C)	57.	(B)	58.	(B)	59.	(A)	60.	(C)
	(B)	62.	(B)	63.	(C)	64.	(A)	65.	(D)	66.	(D)	67.	(C)	68.	(D)	69.	(A)	70.	(C)
71.	(D	72.	(C)	73.	(B)														

	C	omp	etitiv	e Thi	inkin	g													<u> </u>
1.	(D)	2.	(D)	3.	(D)	4.	(C)	5.	(B)	6.	(C)	7.	(A)	8.	(B)	9.	(B)	10.	(A)
11.	(C)	12.	(C)	13.	(A)	14.	(A)	15.	(D)	16.	(C)	17.	(B)	18.	(B)	19.	(B)	20.	(C)
21.	(C)	22.	(A)	23.	(B)	24.	(D)	25.	(C)	26.	(C)	27.	(D)	28.	(A)	29.	(D)	30.	(D)
31.	(C)	32.	(C)	33.	(C)	34.	(D)	35.	(B)	36.	(C)	37.	(C)	38.	(C)	39.	(D)	40.	(C)
41.	(D)																		



#### **Evaluation Test**

- 1. Calculate van't Hoff factor for a 0.2 m aqueous solution of KCl which freezes at -0.680 °C.  $[K_f = 1.86 \text{ K kg mol}^{-1}]$ 
  - $\begin{array}{c} (A) & 1.50 \\ \end{array} \quad (B) & 1.65 \\ \end{array}$
  - (C) 1.83 (D) 2.00
- 2. Which of the following solutions exhibits positive deviation from Raoult's law?
  - (A) Chloroform + Acetone
  - (B) Ethanol + Acetone
  - (C) Aniline + Phenol
  - (D) Benzene + Toluene
- 3. A solution containing 3.56 g of a polymer in 1 litre of a solvent was found to have an osmotic pressure of  $5.2 \times 10^{-4}$  atmosphere at 300 K. The molecular mass of the polymer is \_\_\_\_\_.

 $(R = 0.082 L \text{ atm mol}^{-1} K^{-1})$ 

- (A)  $1.68 \times 10^3 \text{ g mol}^{-1}$
- (B)  $1.68 \times 10^2 \text{ g mol}^{-1}$
- (C)  $1.68 \text{ g mol}^{-1}$
- (D)  $1.68 \times 10^5 \text{ g mol}^{-1}$
- 4. Identify the compound amongst the following of which 0.05 m aqueous solution has highest boiling point.
  - (A) Urea
  - (B) Potassium chloride
  - (C) Sodium sulphate
  - (D) Silver nitrate
- 5. For sodium chloride olved 1 water, the van't Hoff factor (i) accorts for e extent of of the science.
  - (A) solubility (C) disso ation (L, more fraction
- 6. The vapour a ssure of pure benzene at 25 °C is 180 ... Hg. The vapour pressure lowering c used 1 graduation of 25 g of a solute (molar n use 342 a 250 g of benzene is
  - (A) 1.4 <sup>r</sup> m Hg (B) 2.6 mm Hg 4.1 mm Hg (D) 3.2 mm Hg
- 7. W ch one of the following statements is ALSE?
  - (A) The correct order of osmotic pressure for 0.01 M aqueous solution of each compound is

 $BaCl_2 > KCl > CH_3COOH > Sucrose.$ 

(B) The osmotic pressure  $(\pi)$  of a solution is given by the equation  $\pi = MRT$  where M is the molarity of the solution.

- (C) Raoult's law states that the vapour pressure of a component over a solution is proportional to its mole fraction in. he solution.
- (D) Two sucrose solutions of same molality prepared in different solvents at a re the same freezing point depres: n.
- 8. At 300 K, vapour pressure of substan. A is 0.95 atm and vapour pressur of substance? is 0.15 atm. A solution of A at B<sup>+</sup> presser and allowed to equilible with as var ar. The vapour is found to have substance? The vapour is found to have su
  - (A) 0.18 B) 0.14 (C) 0.23 (D) 0.34
- 9. Henr s law pph ble at \_\_\_\_\_. (A) constant oncentration
  - (B) 'nstant ressure
  - (C) communit volume
  - (D) constant temperature
- '0. Coll' ative properties are used for the \_\_\_\_\_.
  - (A) molar mass
  - (B) equivalent weight
  - (C) arrangement of molecules
  - (D) melting point and boiling point
- 11. A solution contains 0.524 g of camphor (molar mass = 152 g/mol dissolved in 36.8 g of ether (boiling point = 34.6 °C). The molal elevation constant of ether is 1.94 K kg mol<sup>-1</sup>. The boiling point of the solution is \_\_\_\_\_.

(A)	317.68 K	(B)	307.78 K
(C)	307.6 K	(D)	317.6 K

- 12. 0.1 m solution each of glucose, sodium chloride, sodium sulphate and sodium phosphate are taken; the ratio of depression in freezing point is
- 13. 18 g of glucose is dissolved in 90 g of  $H_2O$ . Relative lowering of vapour pressure is . [Molar mass of glucose = 180] (A) 0.0196 (B) 0.180 (C) 0.0990 (D) 0.510
- 14. The osmotic pressure of a solution at 276 K is 2.5 atm. Its osmotic pressure at 546 K under similar conditions will be \_\_\_\_\_.
  (A) 0.5 atm (B) 1.0 atm (C) 2.5 atm (D) 5.0 atm

- Two elements X and Y form compounds having 15. molecular formula XY<sub>2</sub> and XY<sub>4</sub>. When dissolved in 20 g of benzene, 1 g of XY<sub>2</sub> lowers the freezing point by 2.3 K, whereas 1.0 g of XY<sub>4</sub> lowers it by 1.3 K. The molal depression constant for benzene is 5.1 K kg  $mol^{-1}$ . The atomic mass of X and Y is respectively.
  - (A) 23.88, 40.31 (B) 25.59, 42.64 (C) 27.13, 44.83 (D) 29.28, 46.73
- 16. Identify the CORRECT statement.
  - Osmosis is a colligative property. (A)
  - Boiling point of solvent is lower than that **(B)** of solution.
  - Osmotic pressure of solution is lower than (C) that of solvent.
  - Vapour pressure of solution is higher than (D) that of pure solvent.
- 17. A solution of glucose containing 9.2 g/litre (molecular weight: 180 g/mol) is isotonic with 3% (W/V) solution of a non-volatile solute. The molecular weight of the solute will be
  - (B)  $267.2 \text{ g mol}^{-1}$  $587 \text{ g mol}^{-1}$ (A)  $567 \text{ g mol}^{-1}$ (D)  $5.87 \text{ g mol}^{-1}$ (C)
- 18. If 10 g of a solute was dissolved in 250 mL of water and osmotic pressure of the solution was found to be 600 mm of Hg at 300 K, then molecular weight of the solute is  $g \text{ mol}^{-1}$ .

(A) 
$$\frac{22800 \times R}{1.5}$$
 (B)  $\frac{22800}{5}$   
(C)  $\frac{1.5 \times R}{22800}$  (D)  $\frac{22c}{1.5}$ 

Henry's law constant r a ga CH<sub>3</sub>Br is 19. 0.159 mol dm<sup>-3</sup>  $e^{+}$  <sup>-1</sup> at <sup>5</sup> °C What is the solubility of CH Br h. water v 25 °C and a partial pressu 0.164 a ?

R .5

- (A) 0.015 M (B)0.164 M (D)
- 0.042 M (C) 0.02
- 20. Th *a*<sub>h</sub> 'r pre. 're will be lowest for () 0  $\square$  sugar solution
  - J.1 N LaCl solution (L)
  - (C) 0.1', Cu(NO<sub>3</sub>)<sub>2</sub> solution
  - 0.1 M AgNO<sub>3</sub> solution
- C oscopic constant is the freezing point 21. pression when the concentration of solution is

(A)	1 M	(B)	1 m
(C)	1 ppm	(D)	1% (W/V)

22. When an electrolyte is dissociated in solution, the van't Hoff factor (i) is (A) = 0 $(\mathbf{R})$ = 1

$$(C) > 1$$
  $(D) < 1$ 

- Which of the following is an example of a 23. solution of gas in liquid?
  - Hydrogen in palladium (A)
  - Carbonated water (B)
  - (C) Chloroform in nitrogen
  - (D) Gasoline

 $\pi =$ 

24. The CORRECT equation for osmotic press of dilute solution is given as

(A) 
$$\pi V = K$$
 (B)  $\pi V =$ 

$$\frac{W_2 RT}{M}$$
 (D)  $\pi V$ 

w'

 $M_2$ 

25. The mass of ascorbic ac  $(C A_8C)$  to be dissolved in 100 g cetic old to ower its freezing point by 1.5 c g is  $(K_f \text{ of acetic acid} K_k \text{ kg} n^{-1})$ 0.015 (A) (B) 0.6 (C) 0.30 ר) 0.66

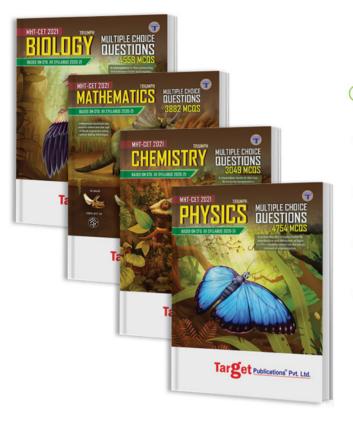
Ar

(C)

#### wers 1, Eva. Jation Test

3. (D) 4. (C) (C, **(B)** 1. 5. (C)6. (C) 7. (D) 8. **(B)** 9. D) 10. (A) 11. (B) 12. (B) 12 (A) 14. (D) 15. **(B)** 16. **(B)** (A) 11. (B) 18. 19. (C) 20. (C) 22. 23. 24. 21. **(B)** (C) **(B) (B)** 25. (B)

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