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Chapter-wise \& Topic-Wise A comprehensive collection of NEET \& AIPMT Questions from past 37 Years

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# Previous Solved Paper 

## Chapter-wise \& Topic-wise

## NEET CHEMISTRY

## Salient Features

- A compilation of 37 years of AIPMT/NEET questions (1988-2024) that align with the most recent syllabus
e Includes ' $1400+$ ' AIPMT/NEET MCQs
- Contains Question Papers from examination conducted twice in a year:

| - | 2015 (Re-Test) | - | 2016 (Phase II) | - | 2019 (Odisha) |
| :--- | :--- | :--- | :--- | :--- | :--- |
| - | 2020 (Phase II) | - | 2023 (Manipur) |  |  |

- Chapter-wise and Topic-wise segregation of questions
\& Year-wise flow of content concluded with the latest questions
- Solutions provided wherever required
© Graphical analysis of questions: Chapter-wise and Topic-wise


## Printed at: Print to Print, Mumbai

## PREFACE

Target's 'NEET Chemistry: PSP (Previous Solved Papers)' is a compilation of questions asked in the past 37 years (1988-2024) in the National Eligibility cum Entrance Test (NEET), formerly known as the All India Pre-Medical Test (AIPMT). The book is updated as per the latest syllabus of NEET (UG) examination.

The book consists of chapter-wise categorization of questions. Each chapter is further segregated into topics and thereafter all the questions pertaining to a topic are arranged year-wise concluding with the latest year. To aid students, we have also provided detailed solutions for questions wherever deemed necessary.

A graphical (\% wise) analysis of the topics for the past 37 years as well as 12 years (2013 onwards) has been provided at the onset of every chapter. Both the graphs will help the students to understand and analyse each topic's distribution for NEET/AIPMT (37 years) and NEET (UG) (12 Years).

We are confident that this book will comprehensively cater to needs of students and effectively assist them to achieve their goal.

The journey to create a complete book is strewn with triumphs, failures and near misses. If you think we've nearly missed something or want to applaud us for our triumphs, we'd love to hear from you.
Please write to us on: mail@targetpublications.org
A book affects eternity; one can never tell where its influence stops.
Best of luck to all the aspirants!
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## Frequently Asked Questions

## Why this book?

- This book acts as a go-to tool to find all the AIPMT/NEET questions since the past 37 years at one place.
- The topic wise arrangement of questions provides the break-down of a chapter into its important components which will enable students to design an effective learning plan.
- The graphical analysis guides students in ascertaining their own preparation of a particular topic.


## Why the need for two graphs?

Admission for undergraduate and post graduate medical courses underwent a critical change with the introduction of NEET in 2013. Although it received a huge backlash and was criticised for the following two years, NEET went on to replace AIPMT in 2016. The introduction of NEET brought in a few structural differences in terms of how the exam was conducted. Although the syllabus has majorly remained the same, the chances of asking a question from a particular topic are seen to vary slightly with the inception of NEET.

The two graphs will fundamentally help the students to understand that the (weightage) distribution of a particular chapter can vary i.e., a particular topic having the most weightage for AIPMT may not necessarily be the topic with the most weightage for NEET.

## How are the two graphs beneficial to the students?

- The two graphs provide a topic's weightage distribution over the past 37 years (for NEET/AIPMT) and over the past 12 years (for NEET-UG).
- The students can use these graphs as a self-evaluation tool by analyzing and comparing a particular topic's weightage with their preparation of the topic. This exercise would help the students to get a clear picture about their strength and weakness based on the topics.
- Students can also use the graphs as a source to know the most important as well as least important topics as per weightage of a particular chapter which will further help them in planning the study structure of a particular chapter.
(Note: The percentage-wise weightage analysis of topics is solely for the knowledge of students and does not guarantee questions from topics having the most weightage, in the future exams. Question classification of a topic is done as per the authors' discretion and may vary with respect to another individual.)


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## Chapter-wise Weightage Analysis of past 12 Years (2013 Onwards)

 ( (a)
$e^{8}$
 $\qquad$
$\qquad$

6.1 Equilibrium in physical and chemical processes and its dynamic nature
6.2 Law of chemical equilibrium and equilibrium constant
6.3 Applications of equilibrium constant
6.4 Factors affecting equilibrium: Le-Chatelier's principle
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37 Years NEET/AIPMT analysis (Percentage-wise weightage of topics)


12 Years NEET analysis (2013 onwards) (Percentage-wise weightage of topics)

6.1 Equilibrium in physical and chemical processes and its dynamic nature

1. In liquid-gas equilibrium, the pressure of vapours above the liquid is constant at $\qquad$ .
[1995]
(A) constant temperature
(B) low temperature
(C) high temperature
(D) none of these
2. Consider the following liquid-vapour equilibrium.

Liquid $\rightleftharpoons$ Vapour
Which of the following relation is CORRECT?
[Phase-I 2016]
(A) $\frac{\mathrm{dmP}}{\mathrm{dT}^{2}}=\frac{-\Delta \mathrm{H}_{\mathrm{V}}}{\mathrm{T}^{2}}$
(B) $\frac{\mathrm{d} \ln \mathrm{P}}{\mathrm{dT}}=\frac{\Delta \mathrm{H}_{\mathrm{V}}}{\mathrm{RT}^{2}}$
(C) $\frac{\mathrm{dnG}}{\mathrm{dT}^{2}}=\frac{\Delta \mathrm{H}_{\mathrm{V}}}{\mathrm{RT}^{2}}$
(D) $\frac{\mathrm{dlnP}}{\mathrm{dT}^{2}}=\frac{-\Delta \mathrm{H}_{\mathrm{V}}}{\mathrm{RT}}$

### 6.2 Law of chemical equilibrium and equilibrium constant

1. $K_{1}$ and $K_{2}$ are equilibrium constant for reactions (i) and (ii)
$\mathrm{N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NO}_{(\mathrm{g})}$
$\mathrm{NO}_{(\mathrm{g})} \rightleftharpoons \frac{1}{2} \mathrm{~N}_{2(\mathrm{~g})}+\frac{1}{2} \mathrm{O}_{2(\mathrm{~g})}$
[1989]
(A) $\mathrm{K}_{1}=\left(\frac{1}{\mathrm{~K}_{2}}\right)^{2}$
(B) $\mathrm{K}_{1}=\mathrm{K}_{2}^{2}$
(C) $\mathrm{K}_{1}=\frac{1}{\mathrm{~K}_{2}}$
(D) $\mathrm{K}_{1}=\left(\mathrm{K}_{2}\right)^{0}$
2. The equilibrium constant for the reaction $\mathrm{N}_{2}+3 \mathrm{H}_{2} \rightleftharpoons 2 \mathrm{NH}_{3}$ is K , then the equilibrium constant for the equilibrium $2 \mathrm{NH}_{3} \rightleftharpoons \mathrm{~N}_{2}+3 \mathrm{H}_{2}$ is $\qquad$ .
[1996]
(A) $\sqrt{\mathrm{K}}$
(B) $\sqrt{\frac{1}{\mathrm{~K}}}$
(C) $\frac{1}{\mathrm{~K}}$
(D) $\frac{1}{\mathrm{~K}^{2}}$
3. If $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ are the respective equilibrium constant for the two reactions,
$\mathrm{XeF}_{6(\mathrm{~g})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{g})} \longrightarrow \mathrm{XeOF}_{4(\mathrm{~g})}+2 \mathrm{HF}_{(\mathrm{g})}$
$\mathrm{XeO}_{4(\mathrm{~g})}+\mathrm{XeF}_{6(\mathrm{~g})} \longrightarrow \mathrm{XeOF}_{4(\mathrm{~g})}+\mathrm{XeO}_{3} \mathrm{~F}_{2(\mathrm{~g})}$, the equilibrium constant of the reaction,
$\mathrm{XeO}_{4(\mathrm{~g})}+2 \mathrm{HF}_{(\mathrm{g})} \longrightarrow \mathrm{XeO}_{3} \mathrm{~F}_{2(\mathrm{~g})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{g})}$, will be $\qquad$ .
[1998]
(A) $\mathrm{K}_{1} / \mathrm{K}_{2}$
(B) $\mathrm{K}_{1} \cdot \mathrm{~K}_{2}$
(C) $\mathrm{K}_{1} /\left(\mathrm{K}_{2}\right)^{2}$
(D) $\quad \mathrm{K}_{2} / \mathrm{K}_{1}$
4. Equilibrium constant $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ for the following equilibria:
$\mathrm{NO}_{(\mathrm{g})}+\frac{1}{2} \mathrm{O}_{2} \stackrel{\mathrm{~K}_{1}}{\rightleftharpoons} \mathrm{NO}_{2(\mathrm{~g})}$ and
$2 \mathrm{NO}_{2(\mathrm{~g})} \stackrel{\mathrm{K}_{2}}{\rightleftharpoons} 2 \mathrm{NO}_{(\mathrm{g})}+\mathrm{O}_{2(\mathrm{~g})}$ are related as
$\qquad$ [2005]
$\overline{\text { (A) }} \mathrm{K}_{2}=1 / \mathrm{K}_{1}^{2}$
(B) $\mathrm{K}_{2}=\mathrm{K}_{1}^{2}$
(C) $\quad \mathrm{K}_{2}=1 / \mathrm{K}_{1}$
(D) $\quad \mathrm{K}_{2}=\mathrm{K}_{1} / 2$
5. In which of the following equilibrium, $\mathrm{K}_{\mathrm{c}}$ and $\mathrm{K}_{\mathrm{p}}$ are NOT equal?
[2010]
(A) $2 \mathrm{C}_{(\mathrm{s})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{CO}_{2(\mathrm{~g})}$
(B) $\quad 2 \mathrm{NO}_{(\mathrm{g})} \rightleftharpoons \mathrm{N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}$
(C) $\quad \mathrm{SO}_{2(\mathrm{~g})}+\mathrm{NO}_{2(\mathrm{~g})} \rightleftharpoons \mathrm{SO}_{3(\mathrm{~g})}+\mathrm{NO}_{(\mathrm{g})}$
(D) $\quad \mathrm{H}_{2(\mathrm{~g})}+\mathrm{I}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{HI}_{(\mathrm{g})}$
6. For the reaction, $\mathrm{N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NO}_{(\mathrm{g})}$, the equilibrium constant is $\mathrm{K}_{1}$. The equilibrium constant is $\mathrm{K}_{2}$ for the reaction, $2 \mathrm{NO}_{(\mathrm{g})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NO}_{2(\mathrm{~g})}$
What is K for the reaction,
$\mathrm{NO}_{2(\mathrm{~g})} \rightleftharpoons \frac{1}{2} \mathrm{~N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}$ ?
[2011]
(A) $\frac{1}{2 \mathrm{~K}_{1} \mathrm{~K}_{2}}$
(B) $\frac{1}{4 \mathrm{~K}_{1} \mathrm{~K}_{2}}$
(C) $\left[\frac{1}{\mathrm{~K}_{1} \mathrm{~K}_{2}}\right]^{1 / 2}$
(D) $\frac{1}{\mathrm{~K}_{1} \mathrm{~K}_{2}}$
7. Given the reaction between 2 gases represented by $\mathrm{A}_{2}$ and $\mathrm{B}_{2}$ to give the compound $\mathrm{AB}_{(\mathrm{g})}$.
$\mathrm{A}_{2(\mathrm{~g})}+\mathrm{B}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{AB}_{(\mathrm{g})}$
At equilibrium, the concentration of
$\mathrm{A}_{2}=3.0 \times 10^{-3} \mathrm{M}, \mathrm{B}_{2}=4.2 \times 10^{-3} \mathrm{M}$, $\mathrm{AB}=2.8 \times 10^{-3} \mathrm{M}$
If the reaction takes place in a sealed vessel at $527^{\circ} \mathrm{C}$, then the value of $\mathrm{K}_{\mathrm{c}}$ will be $\qquad$ .
[2012]
(A)
2.0
(B)
1.9
(C) 0.62
(D) 4.5
8. Given that the equilibrium constant for the reaction, $2 \mathrm{SO}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{SO}_{3(\mathrm{~g})}$ has a value of 278 at a particular temperature. What is the value of the equilibrium constant for the following reaction at the same temperature?
$\mathrm{SO}_{3(\mathrm{~g})} \rightleftharpoons \mathrm{SO}_{2(\mathrm{~g})}+\frac{1}{2} \mathrm{O}_{2(\mathrm{~g})}$
[2012]
(A) $1.8 \times 10^{-3}$
(B) $3.6 \times 10^{-3}$
(C) $6.0 \times 10^{-2}$
(D) $1.3 \times 10^{-5}$
9. If the equilibrium constant for
$\mathrm{N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NO}_{(\mathrm{g})}$ is K ,
the equilibrium constant for
${ }_{2}^{1} \mathrm{~N}_{2(\mathrm{~g})}+\frac{1}{2} \mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons \mathrm{NO}_{(\mathrm{g})}$
will be $\qquad$ .
[Re-Test 2015]
(A) K
(B) $\mathrm{K}^{2}$
(C) $\mathrm{K}^{1 / 2}$
(D) $\frac{1}{2} \mathrm{~K}$
10. The equilibrium constants of the following are:
$\mathrm{N}_{2}+3 \mathrm{H}_{2} \rightleftharpoons 2 \mathrm{NH}_{3} \quad ; \mathrm{K}_{1}$
$\mathrm{N}_{2}+\mathrm{O}_{2} \rightleftharpoons 2 \mathrm{NO} \quad ; \mathrm{K}_{2}$
$\mathrm{H}_{2}+\frac{1}{2} \mathrm{O}_{2} \longrightarrow \mathrm{H}_{2} \mathrm{O} \quad ; \mathrm{K}_{3}$
The equilibrium constant $(\mathrm{K})$ of the reaction:
$2 \mathrm{NH}_{3}+\frac{5}{2} \mathrm{O}_{2} \stackrel{\mathrm{~K}}{\rightleftharpoons} 2 \mathrm{NO}+3 \mathrm{H}_{2} \mathrm{O}$, will be
(A) $\quad \mathrm{K}_{1} \mathrm{~K}_{3}^{3} / \mathrm{K}_{2}$
[2017, 2007, 2003]
(B) $\quad \mathrm{K}_{2} \mathrm{~K}_{3}^{3} / \mathrm{K}_{1}$
(C) $\quad \mathrm{K}_{2} \mathrm{~K}_{3} / \mathrm{K}_{1}$
(D) $\quad \mathrm{K}_{2}^{3} \mathrm{~K}_{3} / \mathrm{K}_{1}$
11. In which of the following equilibria, $\mathrm{K}_{\mathrm{p}}$ and $\mathrm{K}_{\mathrm{c}}$ are NOT equal?
[2024]
(A) $\mathrm{CO}_{(\mathrm{g})}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{g})} \rightleftharpoons \mathrm{CO}_{2(\mathrm{~g})}+\mathrm{H}_{2(\mathrm{~g})}$
(B) $2 \mathrm{BrCl}_{(\mathrm{g})} \rightleftharpoons \mathrm{Br}_{2(\mathrm{~g})}+\mathrm{Cl}_{2(\mathrm{~g})}$
(C) $\mathrm{PCl}_{5(\mathrm{~g})} \rightleftharpoons \mathrm{PCl}_{3(\mathrm{~g})}+\mathrm{Cl}_{2(\mathrm{~g})}$
(D) $\quad \mathrm{H}_{2(\mathrm{~g})}+\mathrm{I}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{HI}_{(\mathrm{g})}$

### 6.3 Applications of equilibrium constant

1. If $\alpha$ is dissociation constant, then the total number of moles for the reaction,
$2 \mathrm{HI} \longrightarrow \mathrm{H}_{2}+\mathrm{I}_{2}$ will be $\qquad$ [1996]
(A) 1
(B) $1-\alpha$
(C) 2
(D) $2-\alpha$
2. Equilibrium constant $\mathrm{K}_{\mathrm{p}}$ for following reaction $\mathrm{MgCO}_{3(\mathrm{~g})} \rightleftharpoons \mathrm{MgO}_{(\mathrm{g})}+\mathrm{CO}_{2(\mathrm{~g})}$
[2000]
(A) $\mathrm{K}_{\mathrm{p}}=\mathrm{p}_{\mathrm{CO}_{2}}$
(B) $\mathrm{K}_{\mathrm{p}}=\mathrm{p}_{\mathrm{CO}_{2}} \times \frac{\mathrm{p}_{\mathrm{CO}_{2}} \times \mathrm{p}_{\mathrm{MgO}}}{\mathrm{p}_{\mathrm{MgCO}_{3}}}$
(C) $\mathrm{K}_{\mathrm{p}}=\frac{\mathrm{p}_{\mathrm{CO}_{2}}+\mathrm{p}_{\mathrm{MgO}}}{\mathrm{p}_{\mathrm{MgCO}_{3}}}$
(D) $\mathrm{K}_{\mathrm{p}}=\frac{\mathrm{p}_{\mathrm{MgCO}_{3}}}{\mathrm{p}_{\mathrm{CO}_{2}} \times \mathrm{p}_{\mathrm{MgO}}}$
3. For any reversible reaction, if we increase concentration of the reactants, then effect on equilibrium constant $\qquad$ -
(A) depends on amount of concentration
(B) remains unchanged
(C) decreases
(D) increases
4. The reaction quotient $(\mathrm{Q})$ for the reaction $\mathrm{N}_{2(\mathrm{~g})}+3 \mathrm{H}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NH}_{3(\mathrm{~g})}$ is given by $\mathrm{Q}=\frac{\left[\mathrm{NH}_{3}\right]^{2}}{\left[\mathrm{~N}_{2}\left[\mathrm{H}_{2}\right]^{3}\right.}$. The reaction will proceed from right to left if $\qquad$ -
[2003]
(A) $\mathrm{Q}=\mathrm{K}_{\mathrm{c}}$
(B) $\mathrm{Q}<\mathrm{K}_{\mathrm{c}}$
(C) $\mathrm{Q}>\mathrm{K}_{\mathrm{c}}$
(D) $\mathrm{Q}=0$
5. In Haber's process, 30 litres of dihydrogen and 30 litres of dinitrogen were taken for reaction, which yielded only $50 \%$ of the expected product. What will be the composition of gaseous mixture under the aforesaid condition at the end?
[2003]
(A) 20 litres ammonia, 25 litres nitrogen, 15 litres hydrogen
(B) 20 litres ammonia, 20 litres nitrogen, 20 litres hydrogen
(C) 10 litres ammonia, 25 litres nitrogen, 15 litres hydrogen
(D) 20 litres ammonia, 10 litres nitrogen, 30 litres hydrogen
6. The values of $\mathrm{K}_{\mathrm{p}_{1}}$ and $\mathrm{K}_{\mathrm{p}_{2}}$ for the reactions,
$\mathrm{X} \rightleftharpoons \mathrm{Y}+\mathrm{Z}$
$A \rightleftharpoons 2 B$
are in the ratio 9:1. If degree of dissociation of X and A be equal, then total pressure at equilibrium (i) and (ii) are in the ratio $\qquad$ -
[2008]
(A) 36:1
(B) $1: 1$
(C) $3: 1$
(D) $1: 9$
7. The value of equilibrium constant of the reaction,

$$
\mathrm{HI}_{(\mathrm{g})} \rightleftharpoons{ }_{2}^{1} \mathrm{H}_{2(\mathrm{~g})}+\frac{1}{2} \mathrm{I}_{2(\mathrm{~g})} \text { is } 8.0 .
$$

The equilibrium constant of the reaction $\mathrm{H}_{2(\mathrm{~g})}+\mathrm{I}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{HI}_{(\mathrm{g})}$ will be $\qquad$ .
[2008]
(A) 16
(B) $1 / 8$
(C) $1 / 16$
(D) $1 / 64$
8. If the concentration of $\mathrm{OH}^{-}$ions in the reaction $\mathrm{Fe}(\mathrm{OH})_{3(\mathrm{~s})} \rightleftharpoons \mathrm{Fe}_{(\mathrm{aq})}^{3+}+3 \mathrm{OH}_{(\mathrm{aq})}^{-}$is decreased by $1 / 4$ times, then equilibrium concentration of $\mathrm{Fe}^{3+}$ will increase by $\qquad$ .
[2008]
(A) 64 times
(B) 4 times
(C) 8 times
(D) 16 times
9. The dissociation equilibrium of a gas $\mathrm{AB}_{2}$ can be represented as:

$$
2 \mathrm{AB}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{AB}_{(\mathrm{g})}+\mathrm{B}_{2(\mathrm{~g})}
$$

The degree of dissociation is x and is small compared to 1 . The expression relating the degree of dissociation ( x ) with equilibrium constant $K_{p}$ and total pressure $P$ is
.
[2008]
(A) $\left(2 \mathrm{~K}_{\mathrm{p}} / \mathrm{P}\right)^{1 / 2}$
(B) $\left(\mathrm{K}_{\mathrm{p}} / \mathrm{P}\right)$
(C) $\quad\left(2 \mathrm{~K}_{\mathrm{p}} / \mathrm{P}\right)$
(D) $\left(2 \mathrm{~K}_{\mathrm{p}} / \mathrm{P}\right)^{1 / 3}$
10. The reaction $2 \mathrm{~A}_{(\mathrm{g})}+\mathrm{B}_{(\mathrm{g})} \rightleftharpoons 3 \mathrm{C}_{(\mathrm{g})}+\mathrm{D}_{(\mathrm{g})}$ is begun with the concentration of A and B both at an initial value of 1.00 M . When equilibrium is reached, the concentration of $D$ is measured and found to be 0.25 M . The value for the equilibrium constant for this reaction is given by the expression $\qquad$ .
[2010]
(A) $\left[(0.75)^{3}(0.25)\right] \div\left[(1.00)^{2}(1.00)\right]$
(B) $\left[(0.75)^{3}(0.25)\right] \div\left[(0.50)^{2}(0.75)\right]$
(C) $\left[(0.75)^{3}(0.25)\right] \div\left[(0.50)^{2}(0.25)\right]$
(D) $\quad\left[(0.75)^{3}(0.25)\right] \div\left[(0.75)^{2}(0.25)\right]$
11. If the value of an equilibrium constant for a particular reaction is $1.6 \times 10^{12}$, then at equilibrium the system will contain
[2015]
(A) all reactants
(B) mostly reactants
(C) mostly products
(D) similar amounts of reactants and products
12. $3 \mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{O}_{3(\mathrm{~g})}$
for the above reaction at $298 \mathrm{~K}, \mathrm{~K}_{\mathrm{c}}$ is found to be $3.0 \times 10^{-59}$. If the concentration of $\mathrm{O}_{2}$ at equilibrium is 0.040 M , then concentration of $\mathrm{O}_{3}$ in M is $\qquad$ .
[2022]
(A) $2.4 \times 10^{31}$
(B) $1.2 \times 10^{21}$
(C) $4.38 \times 10^{-32}$
(D) $1.9 \times 10^{-63}$
13. The equilibrium concentrations of the species in the reaction $\mathrm{A}+\mathrm{B} \rightleftharpoons \mathrm{C}+\mathrm{D}$ are $2,3,10$ and $6 \mathrm{~mol} \mathrm{~L}^{-1}$, respectively at $300 \mathrm{~K} . \Delta \mathrm{G}^{\circ}$ for the reaction is $(\mathrm{R}=2 \mathrm{cal} / \mathrm{mol} \mathrm{K}) \quad$ [2023]
(A) -137.26 cal
(B) -1381.80 cal
(C) -13.73 cal
(D) $\quad 1372.60 \mathrm{cal}$
14. For the reaction $2 \mathrm{~A} \rightleftharpoons \mathrm{~B}+\mathrm{C}, \mathrm{K}_{\mathrm{c}}=4 \times 10^{-3}$. At a given time, the composition of reaction mixture is: $[\mathrm{A}]=[\mathrm{B}]=[\mathrm{C}]=2 \times 10^{-3} \mathrm{M}$.
Then, which of the following is correct?
[2024]
(A) Reaction has a tendency to go in backward direction.
(B) Reaction has gone to completion in forward direction.
(C) Reaction is at equilibrium.
(D) Reaction has a tendency to go in forward direction.

### 6.4 Factors affecting equilibrium: <br> Le-Chatelier's principle

1. Which one of the following information can be obtained on the basis of Le Chatelier's principle?
[1992]
(A) Dissociation constant of a weak acid.
(B) Entropy change in a reaction.
(C) Equilibrium constant of a chemical reaction.
(D) Shift in equilibrium position on changing value of a constraint.
2. According to Le Chatelier's principle, adding heat to a solid and liquid in equilibrium will cause the $\qquad$ .
[1993]
(A) temperature to increase
(B) temperature to decrease
(C) amount of liquid to decrease
(D) amount of solid to decrease
3. Standard state Gibbs free energy change for isomerisation reaction,
cis-2-Pentene $\rightleftharpoons$ trans-2-Pentene
is $-3.67 \mathrm{~kJ} / \mathrm{mol}$ at 400 K . If more trans-2-pentene is added to the reaction vessel then $\qquad$ .
[1995]
(A) equilibrium remains unaffected
(B) equilibrium is shifted in the forward direction
(C) more cis-2-pentene is formed
(D) additional trans-2-pentene is formed
4. Reaction $\mathrm{BaO}_{2(\mathrm{~s})} \rightleftharpoons \mathrm{BaO}_{(\mathrm{s})}+\mathrm{O}_{2(\mathrm{~g})}$;
$\Delta \mathrm{H}=+\mathrm{ve}$. In equilibrium condition, pressure of $\mathrm{O}_{2}$ depends on $\qquad$ _.
[2002]
(A) increase mass of $\mathrm{BaO}_{2}$
(B) increase mass of BaO
(C) increase temperature on equilibrium
(D) increase mass of $\mathrm{BaO}_{2}$ and BaO both
5. For the reaction:
$\mathrm{CH}_{4(\mathrm{~g})}+2 \mathrm{O}_{2(\mathrm{~g})} \rightleftharpoons \mathrm{CO}_{2(\mathrm{~g})}+2 \mathrm{H}_{2} \mathrm{O}_{(l)}$,
$\Delta \mathrm{H}_{\mathrm{r}}=-170.8 \mathrm{~kJ} \mathrm{~mol}^{-1}$. Which of the following statements is INCORRECT?
[2006]
(A) The reaction is exothermic.
(B) At equilibrium the concentrations of $\mathrm{CO}_{2(\mathrm{~g})}$ and $\mathrm{H}_{2} \mathrm{O}_{(l)}$ are not equal.
(C) The equilibrium constant for the reaction is given by $\mathrm{K}_{\mathrm{p}}=\frac{\left[\mathrm{CO}_{2}\right]}{\left[\mathrm{CH}_{4}\right] \cdot\left[\mathrm{O}_{2}\right]}$
(D) Addition of $\mathrm{CH}_{4(\mathrm{~g})}$ or $\mathrm{O}_{2(\mathrm{~g})}$ at equilibrium will cause a shift to the right.
6. The value of $\Delta \mathrm{H}$ for the reaction
$\mathrm{X}_{2(\mathrm{~g})}+4 \mathrm{Y}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{XY}_{4(\mathrm{~g})}$ is less than zero. Formation of $\mathrm{XY}_{4(\mathrm{~g})}$ will be favoured at
[2011]
(A) high temperature and high pressure
(B) low pressure and low temperature
(C) high temperature and low pressure
(D) high pressure and low temperature
7. $\mathrm{KMnO}_{4}$ can be prepared from $\mathrm{K}_{2} \mathrm{MnO}_{4}$ as per the reaction,
$3 \mathrm{MnO}_{4}^{2-}+2 \mathrm{H}_{2} \mathrm{O} \rightleftharpoons 2 \mathrm{MnO}_{4}^{-}+\mathrm{MnO}_{2}+4 \mathrm{OH}^{-}$
The reaction can go to completion by removing $\mathrm{OH}^{-}$ions by adding $\qquad$ .
[2013]
(A) $\quad \mathrm{CO}_{2}$
(B) $\mathrm{SO}_{2}$
(C) HCl
(D) KOH
8. For a given exothermic reaction, $\mathrm{K}_{\mathrm{p}}$ and $\mathrm{K}_{\mathrm{p}}^{\prime}$ are the equilibrium constants at temperatures $\mathrm{T}_{1}$ and $\mathrm{T}_{2}$, respectively. Assuming that heat of reaction is constant in temperature range between $\mathrm{T}_{1}$ and $\mathrm{T}_{2}$, it is readily observed that $\qquad$ .
(A) $\quad K_{p}>K_{p}^{\prime}$
(B) $\mathrm{K}_{\mathrm{p}}<\mathrm{K}_{\mathrm{p}}^{\prime}$
(C) $\mathrm{K}_{\mathrm{p}}=\mathrm{K}_{\mathrm{p}}^{\prime}$
(D) $\mathrm{K}_{\mathrm{p}}=\frac{1}{\mathrm{~K}_{\mathrm{p}}^{\prime}}$
9. For the reversible reaction,
$\mathrm{N}_{2(\mathrm{~g})}+3 \mathrm{H}_{2(\mathrm{~g})} \rightleftharpoons 2 \mathrm{NH}_{3(\mathrm{~g})}+$ Heat
The equilibrium shifts in forward direction
$\qquad$ [2014]
(A) by increasing the concentration of $\mathrm{NH}_{3(\mathrm{~g})}$
(B) by decreasing the pressure
(C) by decreasing the concentration of $\mathrm{N}_{2(\mathrm{~g})}$ and $\mathrm{H}_{2(\mathrm{~g})}$
(D) by increasing pressure and decreasing temperature
10. A 20 litre container at 400 K contains $\mathrm{CO}_{2(\mathrm{~g})}$ at pressure 0.4 atm and an excess of SrO (neglect the volume of solid SrO ). The volume of the container is now decreased by moving the movable piston fitted in the container. The maximum volume of the container, when pressure of $\mathrm{CO}_{2}$ attains its maximum value, will be $\qquad$ .
(Given that: $\mathrm{SrCO}_{3(\mathrm{~s})} \rightleftharpoons \mathrm{SrO}_{(\mathrm{s})}+\mathrm{CO}_{2(\mathrm{~g})}$, $\left.\mathrm{K}_{\mathrm{p}}=1.6 \mathrm{~atm}\right)$
[2017]
(A) 5 litre
(B) 10 litre
(C) 4 litre
(D) 2 litre
11. Which one of the following conditions will favour maximum formation of the product in the reaction?
$\mathrm{A}_{2(\mathrm{~g})}+\mathrm{B}_{2(\mathrm{~g})} \rightleftharpoons \mathrm{X}_{2(\mathrm{~g})} ; \Delta_{\mathrm{r}} \mathrm{H}=-x \mathrm{~kJ}$
[2018]
(A) Low temperature and high pressure
(B) Low temperature and low pressure
(C) High temperature and high pressure
(D) High temperature and low pressure

### 6.5 Ionic equilibrium

1. Aqueous solution of which of the following compounds is the best conductor of electric current?
[2015]
(A) Hydrochloric acid, HCl
(B) Ammonia, $\mathrm{NH}_{3}$
(C) Fructose, $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$
(D) Acetic acid, $\mathrm{C}_{2} \mathrm{H}_{4} \mathrm{O}_{2}$
6.6 Acids and bases: Various concepts
2. Which of the following is NOT a Lewis acid?
[1996]
(A) $\mathrm{SiF}_{4}$
(B) $\mathrm{C}_{2} \mathrm{H}_{4}$
(C) $\mathrm{BF}_{3}$
(D) $\mathrm{FeCl}_{3}$
3. The strongest conjugate base is $\qquad$ .
[1999]
(A) $\mathrm{SO}_{4}^{2-}$
(B) $\mathrm{Cl}^{-}$
(C) $\mathrm{NO}_{3}^{-}$
(D) $\mathrm{CH}_{3} \mathrm{COO}^{-}$
4. Conjugate acid of $\mathrm{NH}_{2}^{-}$is $\qquad$ .
[2000]
(A) NH
(B) $\mathrm{NH}_{4}^{+}$
(C) $\mathrm{NH}_{2}$
(D) $\mathrm{NH}_{3}$
5. Which of the following molecules acts as a Lewis acid?
[2009]
(A) $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{~B}$
(B) $\left(\mathrm{CH}_{3}\right)_{2} \mathrm{O}$
(C) $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{P}$
(D) $\left(\mathrm{CH}_{3}\right)_{3} \mathrm{~N}$
6. Which of the following is least likely to behave as Lewis base?
[2011]
(A) $\mathrm{OH}^{-}$
(B) $\mathrm{H}_{2} \mathrm{O}$
(C) $\mathrm{NH}_{3}$
(D) $\mathrm{BF}_{3}$
7. Which of these is least likely to act as a Lewis base?
[2013]
(A) $\mathrm{BF}_{3}$
(B) $\quad \mathrm{PF}_{3}$
(C) CO
(D) $\mathrm{F}^{-}$
8. Which of the following fluoro compounds is most likely to behave as a Lewis base?
[Phase-II 2016]
(A) $\quad \mathrm{SiF}_{4}$
(B) $\mathrm{BF}_{3}$
(C) $\mathrm{PF}_{3}$
(D) $\mathrm{CF}_{4}$
9. Conjugate bases for Bronsted acids $\mathrm{H}_{2} \mathrm{O}$ and HF are $\qquad$ [2019]
(A) $\mathrm{H}_{3} \mathrm{O}^{+}$and $\mathrm{F}^{-}$, respectively
(B) $\mathrm{OH}^{-}$and $\mathrm{F}^{-}$, respectively
(C) $\mathrm{H}_{3} \mathrm{O}^{+}$and $\mathrm{H}_{2} \mathrm{~F}^{+}$, respectively
(D) $\mathrm{OH}^{-}$and $\mathrm{H}_{2} \mathrm{~F}^{+}$, respectively
10. Which of the following CANNOT act both as Bronsted acid and as Bronsted base?
[Odisha 2019]
(A) $\mathrm{HSO}_{4}^{-}$
(B) $\mathrm{HCO}_{3}^{-}$
(C) $\mathrm{NH}_{3}$
(D) HCl
11. Amongst the given options which of the following molecules/ ion acts as a Lewis acid?
[2023]
(A) $\mathrm{H}_{2} \mathrm{O}$
(B) $\mathrm{BF}_{3}$
(C) $\mathrm{OH}^{-}$
(D) $\mathrm{NH}_{3}$

### 6.7 Ionization of acids and bases

1. Aqueous solution of acetic acid contains
$\qquad$ .
[1991]
(A) $\mathrm{CH}_{3} \mathrm{COO}^{-}$and $\mathrm{H}^{+}$
(B) $\mathrm{CH}_{3} \mathrm{COO}^{-}, \mathrm{H}_{3} \mathrm{O}^{+}$and $\mathrm{CH}_{3} \mathrm{COOH}$
(C) $\mathrm{CH}_{3} \mathrm{COO}^{-}, \mathrm{H}_{3} \mathrm{O}^{+}$and $\mathrm{H}^{+}$
(D) $\mathrm{CH}_{3} \mathrm{COOH}, \mathrm{CH}_{3} \mathrm{COO}^{-}$and $\mathrm{H}^{+}$
2. At $80^{\circ} \mathrm{C}$, distilled water has $\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$ concentration equal to $1 \times 10^{-6}$ mole/litre. The value of $\mathrm{K}_{\mathrm{w}}$ at this temperature will be $\qquad$ .
[1994]
(A) $1 \times 10^{-12}$
(B) $1 \times 10^{-15}$
(C) $1 \times 10^{-6}$
(D) $1 \times 10^{-9}$
3. The ionic product of water at $25^{\circ} \mathrm{C}$ is $10^{-14}$. Its ionic product at $90^{\circ} \mathrm{C}$ will be, $\qquad$ -.
[1996]
(A) $1 \times 10^{-14}$
(B) $1 \times 10^{-16}$
(C) $1 \times 10^{-20}$
(D) $1 \times 10^{-12}$
4. Correct relation between dissociation constants of a dibasic acid is $\qquad$ .
[2000]
(A) $\mathrm{K}_{\mathrm{a}_{1}}=\mathrm{K}_{\mathrm{a}_{2}}$
(B) $\mathrm{K}_{\mathrm{a}_{1}}>\mathrm{K}_{\mathrm{a}_{2}}$
(C) $\mathrm{K}_{\mathrm{a}_{1}}<\mathrm{K}_{\mathrm{a}_{2}}$
(D) $\mathrm{K}_{\mathrm{a}_{1}}=\frac{1}{\mathrm{~K}_{\mathrm{a}_{2}}}$
5. Ionisation constant of $\mathrm{CH}_{3} \mathrm{COOH}$ is $1.7 \times 10^{-5}$ and concentration of $\mathrm{H}^{+}$ions is $3.4 \times 10^{-4}$. Then find out initial concentration of $\mathrm{CH}_{3} \mathrm{COOH}$ molecules.
[2001]
(A) $3.4 \times 10^{-4}$
(B) $3.4 \times 10^{-3}$
(C) $6.8 \times 10^{-4}$
(D) $6.8 \times 10^{-3}$
6. At $25^{\circ} \mathrm{C}$, the dissociation constant of a base, BOH , is $1.0 \times 10^{-12}$. The concentration of hydroxyl ions in 0.01 M aqueous solution of the base would be $\qquad$ $\mathrm{L}^{-1}$
[2005]
(A) $1.0 \times 10^{-5} \mathrm{~mol} \mathrm{~L}^{-1}$
(B) $1.0 \times 10^{-6} \mathrm{~mol} \mathrm{~L}^{-1}$
(C) $2.0 \times 10^{-6} \mathrm{~mol} \mathrm{~L}^{-1}$
(D) $1.0 \times 10^{-7} \mathrm{~mol} \mathrm{~L}^{-1}$
7. A weak acid, HA, has a $\mathrm{K}_{\mathrm{a}}$ of $1.00 \times 10^{-5}$. If 0.100 mol of this acid is dissolved in one litre of water, the percentage of acid dissociated at equilibrium is closet to $\qquad$ .
[2007]
(A) $1.00 \%$
(B) $99.9 \%$
(C) $0.100 \%$
(D) $\quad 99.0 \%$
8. The dissociation constant for acetic acid and HCN at $25^{\circ} \mathrm{C}$ are $1.5 \times 10^{-5}$ and $4.5 \times 10^{-10}$ respectively. The equilibrium constant for the equilibrium,
$\mathrm{CN}^{-}+\mathrm{CH}_{3} \mathrm{COOH} \rightleftharpoons \mathrm{HCN}+\mathrm{CH}_{3} \mathrm{COO}^{-}$
would be $\qquad$ .
[2009]
(A) $3.0 \times 10^{-5}$
(B) $3.0 \times 10^{-4}$
(C) $3.0 \times 10^{4}$
(D) $3.0 \times 10^{5}$
9. What is $\left[\mathrm{H}^{+}\right]$in $\mathrm{mol} / \mathrm{L}$ of a solution that is 0.20 M in $\mathrm{CH}_{3} \mathrm{COONa}$ and 0.10 M in $\mathrm{CH}_{3} \mathrm{COOH}$ ? $\left(\mathrm{K}_{\mathrm{a}}\right.$ for $\left.\mathrm{CH}_{3} \mathrm{COOH}=1.8 \times 10^{-5}\right)$
[2010]
(A) $3.5 \times 10^{-4}$
(B) $1.1 \times 10^{-5}$
(C) $1.8 \times 10^{-5}$
(D) $9.0 \times 10^{-6}$
10. The percentage of pyridine $\left(\mathrm{C}_{5} \mathrm{H}_{5} \mathrm{~N}\right)$ that forms pyridinium ion $\mathrm{C}_{5} \mathrm{H}_{5} \stackrel{+}{\mathrm{N}} \mathrm{H}$ in a 0.10 M aqueous pyridine solution is $\qquad$ ( $\mathrm{K}_{\mathrm{b}}$ for $\mathrm{C}_{5} \mathrm{H}_{5} \mathrm{~N}=1.7 \times 10^{-9}$ ) [Phase-II 2016]
(A) $1.6 \%$
(B) $0.0060 \%$
(C) $0.013 \%$
(D) $0.77 \%$
11. For a weak acid HA, the percentage of dissociation is nearly $1 \%$ at equilibrium. If the concentration of acid is $0.1 \mathrm{~mol} \mathrm{~L}{ }^{-1}$, then the CORRECT option for its $K_{a}$ at the same temperature is:
[Manipur 2023]
(A) $1 \times 10^{-5}$
(B) $1 \times 10^{-3}$
(C) $1 \times 10^{-4}$
(D) $1 \times 10^{-6}$
12. Consider the following reaction in a sealed vessel at equilibrium with concentrations of $\mathrm{N}_{2}=3.0 \times 10^{-3} \mathrm{M}, \mathrm{O}_{2}=4.2 \times 10^{-3} \mathrm{M}$ and $\mathrm{NO}=2.8 \times 10^{-3} \mathrm{M}$.
$2 \mathrm{NO}_{(\mathrm{g})} \rightleftharpoons \mathrm{N}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}$
If 0.1 mol L - ${ }^{-1}$ of $\mathrm{NO}_{(\mathrm{g})}$ is taken in a closed vessel, what will be degree of dissociation $(\alpha)$ of $\mathrm{NO}_{(\mathrm{g})}$ at equilibrium?
[2024]
(A) 0.8889
(B) 0.717
(C) 0.00889
(D) 0.0889

### 6.8 Relative strength of acids and bases

1. The hydride ion $\mathrm{H}^{-}$is a stronger base than its hydroxide ion $\mathrm{OH}^{-}$. Which of the following reactions will occur if sodium hydride $(\mathrm{NaH})$ is dissolved in water?
[1997]
(A) $\mathrm{H}_{(\text {aq) }}^{-}+\mathrm{H}_{2} \mathrm{O} \longrightarrow \mathrm{H}_{3} \mathrm{O}_{\text {(aq) }}^{-}$
(B) $\mathrm{H}_{(\mathrm{aq})}^{-}+\mathrm{H}_{2} \mathrm{O}_{(\mathrm{l})} \longrightarrow \mathrm{OH}_{(\mathrm{aq})}^{-}+\mathrm{H}_{2(\mathrm{~g})}$
(C) $\mathrm{H}_{(\mathrm{aq})}^{-}+\mathrm{H}_{2} \mathrm{O}_{(I)} \longrightarrow$ No reaction
(D) $\mathrm{H}_{(\mathrm{aq})}^{-}+\mathrm{H}_{2} \mathrm{O}_{(l)} \longrightarrow \mathrm{H}^{+}+\mathrm{H}_{3} \mathrm{O}^{+}$
2. In $\mathrm{HS}^{-}, \mathrm{I}^{-}, \mathrm{R}-\mathrm{NH}_{2}, \mathrm{NH}_{3}$ order of proton accepting tendency will be $\qquad$ .
[2001]
(A) $\mathrm{I}^{-}>\mathrm{NH}_{3}>\mathrm{R}-\mathrm{NH}_{2}>\mathrm{HS}$
(B) $\mathrm{NH}_{3}>\mathrm{R}-\mathrm{NH}_{2}>\mathrm{HS}^{-}>\mathrm{I}^{-}$
(C) $\mathrm{R}-\mathrm{NH}_{2}>\mathrm{NH}_{3}>\mathrm{HS}^{-}>\mathrm{I}^{-}$
(D) $\mathrm{HS}^{-}>\mathrm{R}-\mathrm{NH}_{2}>\mathrm{NH}_{3}>\mathrm{I}^{-}$
3. Which one of the following statements is not true?
[2003]
(A) Among halide ions, iodide is the most powerful reducing agent.
(B) Fluorine is the only halogen that does not show a variable oxidation state.
(C) HOCl is a stronger acid than HOBr .
(D) HF is a stronger acid than HCl .

### 6.9 Hydrogen ion concentration: Concept of pH

1. The compound whose water solution has the highest pH is $\qquad$ .
[1988]
(A) NaCl
(B) $\mathrm{NaHCO}_{3}$
(C) $\mathrm{Na}_{2} \mathrm{CO}_{3}$
(D) $\quad \mathrm{NH}_{4} \mathrm{Cl}$
2. The pH value of a 10 M solution of HCl is
[1995]
(A) equal to 1
(B) equal to 2
(C) less than 0
(D) equal to 0
3. The pH value of $\mathrm{N} / 10 \mathrm{NaOH}$ solution is
[1996]
(A) 12
(B) 13
(C) 10
(D) 11
4. The concentration of $\mathrm{H}^{+}$and concentration of $\mathrm{OH}^{-}$of a 0.1 M aqueous solution of $2 \%$ ionized weak acid is $\qquad$
[Ionic product of water $=\overline{1 \times 10^{-1} 4}$ ]
[1999]
(A) $2 \times 10^{-3} \mathrm{M}$ and $5 \times 10^{-12} \mathrm{M}$
(B) $1 \times 10^{3} \mathrm{M}$ and $3 \times 10^{-11} \mathrm{M}$
(C) $0.02 \times 10^{-3} \mathrm{M}$ and $5 \times 10^{-11} \mathrm{M}$
(D) $3 \times 10^{-2} \mathrm{M}$ and $4 \times 10^{-13} \mathrm{M}$
5. Which statement is INCORRECT about pH and $\mathrm{H}^{+}$?
[2000]
(A) pH of neutral water is not zero.
(B) Adding 1 N solution of $\mathrm{CH}_{3} \mathrm{COOH}$ and 1 N solution of NaOH , pH will be seven.
(C) $\left[\mathrm{H}^{+}\right]$of dilute and hot $\mathrm{H}_{2} \mathrm{SO}_{4}$ is more than concentrated and cold $\mathrm{H}_{2} \mathrm{SO}_{4}$.
(D) Mixing solution of $\mathrm{CH}_{3} \mathrm{COOH}$ and HCl , pH will be less than 7 .
6. The hydrogen ion concentration of a $10^{-8} \mathrm{M}$ HCl aqueous solution at $298 \mathrm{~K}\left(\mathrm{~K}_{\mathrm{w}}=10^{-14}\right)$ is
(A) $1.0 \times 10^{-8} \mathrm{M}$
(B) $1.0 \times 10^{-6} \mathrm{M}$
(C) $1.0525 \times 10^{-7} \mathrm{M}$
(D) $9.525 \times 10^{-8} \mathrm{M}$
7. Calculate the pOH of a solution at $25^{\circ} \mathrm{C}$ that contains $1 \times 10^{-10} \mathrm{M}$ of hydronium ions, i.e., $\mathrm{H}_{3} \mathrm{O}^{+}$.
[2007]
(A) 4.000
(B) 9.000
(C) 1.000
(D) 7.000
8. Equal volumes of three acid solutions of $\mathrm{pH} 3,4$ and 5 are mixed in a vessel. What will be the $\mathrm{H}^{+}$ion concentration in the mixture?
[2008]
(A) $3.7 \times 10^{-3} \mathrm{M}$
(B) $1.11 \times 10^{-3} \mathrm{M}$
(C) $1.11 \times 10^{-4} \mathrm{M}$
(D) $3.7 \times 10^{-4} \mathrm{M}$
9. Equimolar solutions of the following substances were prepared separately. Which one of these will record the highest pH value?
[2012]
(A) $\mathrm{BaCl}_{2}$
(B) $\mathrm{AlCl}_{3}$
(C) LiCl
(D) $\mathrm{BeCl}_{2}$
10. What is the pH of the resulting solution when equal volumes of 0.1 M NaOH and 0.01 M HCl are mixed?
[Re-Test 2015]
(A)
(B) 1.04
(C) 12.65
(D)
2.0
11. Following solutions were prepared by mixing different volumes of NaOH and HCl of different concentrations:
i. $\quad 60 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{HCl}+40 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{NaOH}$
ii. $\quad 55 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{HCl}+45 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{NaOH}$
iii. $\quad 75 \mathrm{~mL} \frac{\mathrm{M}}{5} \mathrm{HCl}+25 \mathrm{~mL} \frac{\mathrm{M}}{5} \mathrm{NaOH}$
iv. $\quad 100 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{HCl}+100 \mathrm{~mL} \frac{\mathrm{M}}{10} \mathrm{NaOH}$
pH of which one of them will be equal to 1 ?
[2018]
(A) ii.
(B) $i$.
(C) iv.
(D) iii.
12. The pH of $0.01 \mathrm{M} \mathrm{NaOH}_{(\mathrm{aq})}$ solution will be
$\qquad$ [Odisha 2019]
9 .
(B) 7.01
(C) 2
(A) 9
(D) 12

### 6.10 Hydrolysis of salts

1. 0.1 M solution of which one of these substances will act basic?
[1992]
(A) Sodium borate
(B) Ammonium chloride
(C) Calcium nitrate
(D) Sodium sulphate
2. Which has the highest pH ?
[2002]
(A) $\mathrm{CH}_{3} \mathrm{COOK}$
(B) $\mathrm{Na}_{2} \mathrm{CO}_{3}$
(D) $\mathrm{NH}_{4} \mathrm{Cl}$
(D) $\quad \mathrm{NaNO}_{3}$
3. The rapid change of pH near the stoichiometric point of an acid-base titration is the basis of indicator detection. pH of the solution is related to ratio of the concentrations of the conjugate acid (HIn) and base ( $\mathrm{In}^{-}$) forms of the indicator by the expression $\qquad$ -
[2004]
(A) $\quad \log \frac{\left[\mathrm{In}^{-}\right]}{[\mathrm{HIn}]}=\mathrm{pK}_{\mathrm{In}}-\mathrm{pH}$
(B) $\quad \log \frac{[\mathrm{HIn}]}{\left[\mathrm{In}^{-}\right]}=\mathrm{pH}_{\mathrm{In}}-\mathrm{pH}$
(C) $\log \frac{[\mathrm{HIn}]}{\left[\mathrm{In}^{-}\right]}=\mathrm{pH}-\mathrm{pK}_{\text {In }}$
(D) $\quad \log \frac{\left[\mathrm{In}^{-}\right]}{[\mathrm{HIn}]}=\mathrm{pH}-\mathrm{pK}_{\text {In }}$
4. The ionization constant of ammonium hydroxide is $1.77 \times 10^{-5}$ at 298 K . Hydrolysis constant of ammonium chloride is $\qquad$ .
[2009]
(A) $6.50 \times 10^{-12}$
(B) $5.65 \times 10^{-13}$
(C) $5.65 \times 10^{-12}$
(D) $5.65 \times 10^{-10}$
5. Which of the following salts will give the highest pH in water?
[2014]
(A) KCl
(B) NaCl
(C) $\quad \mathrm{Na}_{2} \mathrm{CO}_{3}$
(D) $\mathrm{CuSO}_{4}$
6. Which among the following salt solutions is basic in nature?
[Phase-II 2020]
(A) Sodium acetate
(B) Ammonium chloride
(C) Ammonium sulphate
(D) Ammonium nitrate
7. The $\mathrm{pK}_{\mathrm{b}}$ of dimethylamine and $\mathrm{pK}_{\mathrm{a}}$ of acetic acid are 3.27 and 4.77 respectively at $\mathrm{T}(\mathrm{K})$. The CORRECT option for the pH of dimethylammonium acetate solution is $\qquad$ .
[2021]
(A) 5.50
(B) 7.75
(C) 6.25
(D) 8.50

### 6.11 Buffer solutions

1. The pH value of blood does not appreciably change by a small addition of an acid or a base, because the blood $\qquad$ [1995]
(A) can be easily coagulated
(B) contains iron as a part of the molecule
(C) is a body fluid
(D) contains serum protein which acts as buffer
2. Solution of $0.1 \mathrm{~N} \mathrm{NH}_{4} \mathrm{OH}$ and $0.1 \mathrm{~N} \mathrm{NH}_{4} \mathrm{Cl}$ has pH 9.25 . Then find out $\mathrm{pK}_{\mathrm{b}}$ of $\mathrm{NH}_{4} \mathrm{OH}$.
[2002]
(A) 9.25
(B) 4.75
(C) 3.75
(D) 8.25
3. Which of the following pairs constitutes a buffer?
[2006]
(A) HCl and KCl
(B) $\mathrm{HNO}_{2}$ and $\mathrm{NaNO}_{2}$
(C) NaOH and NaCl
(D) $\mathrm{HNO}_{3}$ and $\mathrm{NH}_{4} \mathrm{NO}_{3}$
4. What is the $\left[\mathrm{OH}^{-}\right]$in the final solution prepared by mixing 20.0 mL of 0.050 M HCl with 30.0 mL of $0.10 \mathrm{M} \mathrm{Ba}(\mathrm{OH})_{2}$ ?
[2009]
(A) $\quad 0.40 \mathrm{M}$
(B) 0.0050 M
(C) $\quad 0.12 \mathrm{M}$
(D) $\quad 0.10 \mathrm{M}$
5. In a buffer solution containing equal concentration of $\mathrm{B}^{-}$and HB , the $\mathrm{K}_{\mathrm{b}}$ for $\mathrm{B}^{-}$is $10^{-10}$. The pH of buffer solution is $\qquad$ .
[2010]
(A) 10
(B) 7
(C) 6
(D) 4
6. A buffer solution is prepared in which the concentration of $\mathrm{NH}_{3}$ is 0.30 M and the concentration of $\mathrm{NH}_{4}^{+}$is 0.20 M . If the equilibrium constant, $\mathrm{K}_{\mathrm{b}}$ for $\mathrm{NH}_{3}$ equals $1.8 \times 10^{-5}$, what is the pH of this solution?
[2011]
(A) 8.73
(B) 9.08
(C) 9.43
(D) 11.72
7. Buffer solutions have constant acidity and alkalinity because $\qquad$ .
[2012]
(A) these give unionized acid or base on reaction with added acid or alkali
(B) acids and alkalies in these solutions are shielded from attack by other ions
(C) they have large excess of $\mathrm{H}^{+}$or $\mathrm{OH}^{-}$ions
(D) they have fixed value of pH
8. Which one of the following pairs of solution is NOT an acidic buffer?
[Re-Test 2015]
(A) $\mathrm{H}_{2} \mathrm{CO}_{3}$ and $\mathrm{Na}_{2} \mathrm{CO}_{3}$
(B) $\mathrm{H}_{3} \mathrm{PO}_{4}$ and $\mathrm{Na}_{3} \mathrm{PO}_{4}$
(C) $\quad \mathrm{HClO}_{4}$ and $\mathrm{NaClO}_{4}$
(D) $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COONa}$
9. Which will make basic buffer?
[2019]
(A) 100 mL of $0.1 \mathrm{M} \mathrm{CH}_{3} \mathrm{COOH}+100 \mathrm{~mL}$ of 0.1 M NaOH
(B) 100 mL of $0.1 \mathrm{M} \mathrm{HCl}+200 \mathrm{~mL}$ of $0.1 \mathrm{M} \mathrm{NH}_{4} \mathrm{OH}$
(C) 100 mL of $0.1 \mathrm{M} \mathrm{HCl}+100 \mathrm{~mL}$ of 0.1 M NaOH
(D) 50 mL of $0.1 \mathrm{M} \mathrm{NaOH}+25 \mathrm{~mL}$ of $0.1 \mathrm{M} \mathrm{CH}_{3} \mathrm{COOH}$
10. The pH of the solution containing 50 mL each of 0.10 M sodium acetate and 0.01 M acetic acid is $\qquad$ .
[Given $\mathrm{pK}_{\mathrm{a}}$ of $\mathrm{CH}_{3} \mathrm{COOH}=4.57$ ]
[2022]
(A) 4.57
(B) 2.57
(C) 5.57
(D) 3.57
11. An acidic buffer is prepared by mixing:
[Manipur 2023]
(A) strong acid and its salt with strong base.
(B) strong acid and its salt with weak base.
(C) weak acid and its salt with strong base.
(D) equal volumes of equimolar solutions of weak acid and weak base. (The $\mathrm{pK}_{\mathrm{a}}$ of $\operatorname{acid}=\mathrm{pK}_{\mathrm{b}}$ of the base)

### 6.12 Solubility product

1. Which one of the following is most soluble?
[1994]
(A) $\mathrm{Bi}_{2} \mathrm{~S}_{3}\left(\mathrm{~K}_{\text {sp }}=1 \times 10^{-70}\right)$
(B) $\quad \mathrm{Ag}_{2} \mathrm{~S}\left(\mathrm{~K}_{\text {sp }}=6 \times 10^{-51}\right)$
(C) $\quad \mathrm{CuS}\left(\mathrm{K}_{\text {sp }}=8 \times 10^{-37}\right)$
(D) $\operatorname{MnS}\left(\mathrm{K}_{\mathrm{sp}}=7 \times 10^{-16}\right)$
2. The solubility product of $\mathrm{CuS}, \mathrm{Ag}_{2} \mathrm{~S}$ and HgS are $10^{-31}, 10^{-44}$ and $10^{-54}$ respectively. The solubilities of these sulphides are in the order .
[1997]
(A) $\quad \mathrm{HgS}>\mathrm{Ag}_{2} \mathrm{~S}>\mathrm{CuS}$
(B) $\mathrm{CuS}>\mathrm{Ag}_{2} \mathrm{~S}>\mathrm{HgS}$
(C) $\mathrm{Ag}_{2} \mathrm{~S}>\mathrm{CuS}>\mathrm{HgS}$
(D) $\mathrm{AgS}>\mathrm{HgS}>\mathrm{CuS}$
3. The solubility of a saturated solution of calcium fluoride is $2 \times 10^{-4}$ moles per litre. Its solubility product is $\qquad$ -
[1999]
(A) $22 \times 10^{-1}$
(B) $14 \times 10^{-4}$
(C) $2 \times 10^{-2}$
(D) $32 \times 10^{-12}$
4. Solubility of $\mathrm{M}_{2} \mathrm{~S}$ salt is $3.5 \times 10^{-6}$ then find out solubility product.
[2001]
(A) $1.7 \times 10^{-6}$
(B) $1.7 \times 10^{-16}$
(C) $1.7 \times 10^{-18}$
(D) $1.7 \times 10^{-12}$
5. Solubility of $\mathrm{MX}_{2}$ type electrolytes is $0.5 \times 10^{-4}$ mole/lit., then find out $\mathrm{K}_{\mathrm{sp}}$ of electrolytes.
[2002]
(A) $5 \times 10^{-12}$
(B) $25 \times 10^{-10}$
(C) $1 \times 10^{-13}$
(D) $5 \times 10^{-13}$
6. The solubility product of AgI at $25^{\circ} \mathrm{C}$ is $1.0 \times 10^{-16} \mathrm{~mol}^{2} \mathrm{~L}^{-2}$. The solubility of AgI in $10^{-4} \mathrm{~N}$ solution of KI at $25^{\circ} \mathrm{C}$ is approximately
$\qquad$ . (in $\mathrm{mol} \mathrm{L}^{-1}$ )
[2003]
(A) $\quad 1.0 \times 10^{-16}$
(B) $1.0 \times 10^{-12}$
(C) $1.0 \times 10^{-10}$
(D) $1.0 \times 10^{-8}$
7. The solubility product of a sparingly soluble salt $\mathrm{AX}_{2}$ is $3.2 \times 10^{-11}$. Its solubility
(in moles/litres) is $\qquad$ -
[2004]
(A) $2 \times 10^{-4}$
(B) $4 \times 10^{-4}$
(C) $5.6 \times 10^{-6}$
(D) $3.1 \times 10^{-4}$
8. In qualitative analysis, the metals of group I can be separated from other ions by precipitating them as chloride salts. A solution initially contains $\mathrm{Ag}^{+}$and $\mathrm{Pb}^{2+}$ at a concentration of 0.10 M . Aqueous HCl is added to this solution until the $\mathrm{Cl}^{-}$concentration is 0.10 M . What will the concentrations of $\mathrm{Ag}^{+}$and $\mathrm{Pb}^{2+}$ be at equilibrium?
$\left(\mathrm{K}_{\text {sp }}\right.$ for $\mathrm{AgCl}=1.8 \times 10^{-10}, \mathrm{~K}_{\text {sp }}$ for $\mathrm{PbCl}_{2}=1.7 \times 10^{-5}$ )
[2011]
(A) $\left[\mathrm{Ag}^{+}\right]=1.8 \times 10^{-7} \mathrm{M}$, $\left[\mathrm{Pb}^{2+}\right]=1.7 \times 10^{-6} \mathrm{M}$
(B) $\left[\mathrm{Ag}^{+}\right]=1.8 \times 10^{-11} \mathrm{M}$, $\left[\mathrm{Pb}^{2+}\right]=8.5 \times 10^{-5} \mathrm{M}$
(C) $\left[\mathrm{Ag}^{+}\right]=1.8 \times 10^{-9} \mathrm{M}$, $\left[\mathrm{Pb}^{2+}\right]=1.7 \times 10^{-3} \mathrm{M}$
(D) $\left[\mathrm{Ag}^{+}\right]=1.8 \times 10^{-11} \mathrm{M}$, $\left[\mathrm{Pb}^{2+}\right]=1.7 \times 10^{-4} \mathrm{M}$
9. pH of a saturated solution of $\mathrm{Ba}(\mathrm{OH})_{2}$ is 12 . The value of solubility product $\left(\mathrm{K}_{\text {sp }}\right)$ of $\mathrm{Ba}(\mathrm{OH})_{2}$ is
$\qquad$ .
[2012, 2010]
(A) $3.3 \times 10^{-7}$
(B) $5.0 \times 10^{-7}$
(C) $4.0 \times 10^{-6}$
(D) $5.0 \times 10^{-6}$
10. Identify the CORRECT order of solubility in aqueous solution.
[2013]
(A) $\mathrm{Na}_{2} \mathrm{~S}>\mathrm{ZnS}>\mathrm{CuS}$
(B) $\mathrm{CuS}>\mathrm{ZnS}>\mathrm{Na}_{2} \mathrm{~S}$
(C) $\mathrm{ZnS}>\mathrm{Na}_{2} \mathrm{~S}>\mathrm{CuS}$
(D) $\mathrm{Na}_{2} \mathrm{~S}>\mathrm{CuS}>\mathrm{ZnS}$
11. Using the Gibb's energy change $\Delta \mathrm{G}^{\circ}=+63.3 \mathrm{~kJ}$, for the following reaction,
$\mathrm{Ag}_{2} \mathrm{CO}_{3(\mathrm{~s})} \rightleftharpoons 2 \mathrm{Ag}_{\text {(aq) }}^{+}+\mathrm{CO}_{3(\text { aq) }}^{2-}$
The $\mathrm{K}_{\text {sp }}$ of $\mathrm{Ag}_{2} \mathrm{CO}_{3(\mathrm{~s})}$ in water at $25^{\circ} \mathrm{C}$ is .$\left(\mathrm{R}=8.314 \mathrm{JK}^{-1} \mathrm{~mol}^{-1}\right)$
[2014]
$\overline{\text { (A) } \quad 3.2} \times 10^{-26}$
(B) $8.0 \times 10^{-12}$
(C) $2.9 \times 10^{-3}$
(D) $7.9 \times 10^{-2}$
12. The $\mathrm{K}_{\text {sp }}$ of $\mathrm{Ag}_{2} \mathrm{CrO}_{4}, \mathrm{AgCl}, \mathrm{AgBr}$ and AgI are respectively, $1.1 \times 10^{-12}, 1.8 \times 10^{-10}, 5.0 \times 10^{-13}$ and $8.3 \times 10^{-17}$. Which one of the following salts will precipitate last if $\mathrm{AgNO}_{3}$ solution is added to the solution containing equal moles of $\mathrm{NaCl}, \mathrm{NaBr}, \mathrm{NaI}$ and $\mathrm{Na}_{2} \mathrm{CrO}_{4}$ ?
[2015]
(A) AgI
(B) AgCl
(C) AgBr
(D) $\quad \mathrm{Ag}_{2} \mathrm{CrO}_{4}$
13. MY and $\mathrm{NY}_{3}$, two nearly insoluble salts, have the same $\mathrm{K}_{\text {sp }}$ values of $6.2 \times 10^{-13}$ at room temperature. Which statement would be TRUE in regard to MY and $\mathrm{NY}_{3}$ ?
[Phase-I 2016]
(A) The salts MY and $\mathrm{NY}_{3}$ are more soluble in 0.5 M KY than in pure water.
(B) The addition of the salt of KY to solution of MY and $\mathrm{NY}_{3}$ will have no effect on their solubilities.
(C) The molar solubilities of MY and $\mathrm{NY}_{3}$ in water are identical.
(D) The molar solubility of MY in water is less than that of $\mathrm{NY}_{3}$.
14. The solubility of $\mathrm{AgCl}_{(\mathrm{s})}$ with solubility product $1.6 \times 10^{-10}$ in 0.1 M NaCl solution would be
$\qquad$ -
[Phase-II 2016]
(A) zero
(B) $1.26 \times 10^{-5} \mathrm{M}$
(C) $1.6 \times 10^{-9} \mathrm{M}$
(D) $1.6 \times 10^{-11} \mathrm{M}$
15. Concentration of the $\mathrm{Ag}^{+}$ions in a saturated solution of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ is $2.2 \times 10^{-4} \mathrm{~mol} \mathrm{~L} \mathrm{~L}^{-1}$. Solubility product of $\mathrm{Ag}_{2} \mathrm{C}_{2} \mathrm{O}_{4}$ is $\qquad$ -
[2017]
(A) $2.42 \times 10^{-8}$
(B) $2.66 \times 10^{-12}$
(C) $4.25 \times 10^{-11}$
(D) $5.3 \times 10^{-12}$
16. The solubility of $\mathrm{BaSO}_{4}$ in water is $2.42 \times 10^{-3} \mathrm{~g} \mathrm{~L}^{-1}$ at 298 K . The value of its solubility product $\left(\mathrm{K}_{\mathrm{sp}}\right)$ will be $\qquad$ -.
(Given : molar mass of $\mathrm{BaSO}_{4}=233 \mathrm{~g} \mathrm{~mol}^{-1}$ )
[2018]
(A) $1.08 \times 10^{-10} \mathrm{~mol}^{2} \mathrm{~L}^{-2}$
(B) $1.08 \times 10^{-12} \mathrm{~mol}^{2} \mathrm{~L}^{-2}$
(C) $1.08 \times 10^{-14} \mathrm{~mol}^{2} \mathrm{~L}^{-2}$
(D) $1.08 \times 10^{-8} \mathrm{~mol}^{2} \mathrm{~L}^{-2}$
17. pH of a saturated solution of $\mathrm{Ca}(\mathrm{OH})_{2}$ is 9 . The solubility product $\left(\mathrm{K}_{\text {sp }}\right)$ of $\mathrm{Ca}(\mathrm{OH})_{2}$ is $\qquad$ .
(A) $0.25 \times 10^{-10}$
(B) $0.125 \times 10^{-15}$
(C) $0.5 \times 10^{-10}$
(D) $0.5 \times 10^{-15}$
18. Find out the solubility of $\mathrm{Ni}(\mathrm{OH})_{2}$ in 0.1 M NaOH . Given that the ionic product of $\mathrm{Ni}(\mathrm{OH})_{2}$ is $2 \times 10^{-15}$.
[Phase-I 2020]
(A) $2 \times 10^{-8} \mathrm{M}$
(B) $1 \times 10^{-13} \mathrm{M}$
(C) $1 \times 10^{8} \mathrm{M}$
(D) $2 \times 10^{-13} \mathrm{M}$
19. The solubility product for a salt of the type $A B$ is $4 \times 10^{-8}$. What is the molarity of its saturated solution?
[Phase-II 2020]
(A) $4 \times 10^{-4} \mathrm{~mol} / \mathrm{L}$
(B) $2 \times 10^{-4} \mathrm{~mol} / \mathrm{L}$
(C) $16 \times 10^{-16} \mathrm{~mol} / \mathrm{L}$
(D) $2 \times 10^{-16} \mathrm{~mol} / \mathrm{L}$

### 6.13 Common ion effect

1. The solubility of AgCl will be minimum in
$\qquad$ .
[1995]
(A) $\quad 0.01 \mathrm{M} \mathrm{CaCl}_{2}$
(B) pure water
(C) $\quad 0.001 \mathrm{M} \mathrm{AgNO}_{3}$
(D) 0.01 M NaCl
2. $\mathrm{H}_{2} \mathrm{~S}$ gas when passed through a solution of cations containing HCl precipitates the cations of second group of qualitative analysis but not those belonging to the fourth group. It is because $\qquad$ -.
[2005]
(A) presence of HCl decreases the sulphide ion concentration
(B) solubility product of group II sulphides is more than that of group IV sulphides
(C) presence of HCl increases the sulphide ion concentration
(D) sulphides of group IV cations are unstable in HCl .
3. Consider the nitration of benzene using mixed conc. $\mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{HNO}_{3}$. If a large amount of $\mathrm{KHSO}_{4}$ is added to the mixture, the rate of nitration will be $\qquad$ - [Phase-I 2016]
(A) unchanged
(B) doubled
(C) faster
(D) slower
4. The molar solubility of $\mathrm{CaF}_{2}$
$\left(\mathrm{K}_{\mathrm{sp}}=5.3 \times 10^{-11}\right)$ in 0.1 M solution of NaF will be $\qquad$ .
[Odisha 2019]
(A) $5.3 \times 10^{-10} \mathrm{~mol} \mathrm{~L}^{-1}$
(B) $5.3 \times 10^{-11} \mathrm{~mol} \mathrm{~L}^{-1}$
(C) $5.3 \times 10^{-8} \mathrm{~mol} \mathrm{~L}^{-1}$
(D) $5.3 \times 10^{-9} \mathrm{~mol} \mathrm{~L}^{-1}$
5. HCl was passed through a solution of $\mathrm{CaCl}_{2}$, $\mathrm{MgCl}_{2}$ and NaCl . Which of the following compound(s) crystallise(s)? [Phase-I 2020]
(A) Only NaCl
(B) Only $\mathrm{MgCl}_{2}$
(C) $\mathrm{NaCl}, \mathrm{MgCl}_{2}$ and $\mathrm{CaCl}_{2}$
(D) Both $\mathrm{MgCl}_{2}$ and $\mathrm{CaCl}_{2}$

## Std. XI

## 1. Some Basic Concepts of Chemistry

### 1.1 Units of measurement

1. (B)

| Quantity | Dimensions |
| :---: | :---: |
| Pressure | $\left[\mathrm{M} \mathrm{L}^{-1} \mathrm{~T}^{-2}\right]$ |
| Force per unit volume | $\left[\mathrm{M} \mathrm{L}^{-2} \mathrm{~T}^{-2}\right]$ |
| Energy per unit volume | $\left[\mathrm{M} \mathrm{L}^{-1} \mathrm{~T}^{-2}\right]$ |
| Force | $\left[\mathrm{M} \mathrm{L} \mathrm{T}^{-2}\right]$ |
| Energy | $\left[\mathrm{M} \mathrm{L}^{2} \mathrm{~T}^{-2}\right]$ |

### 1.2 Uncertainty in measurement

1. (D)

161 has three significant figures as all are non-zero digits.
0.161 has three significant figures as zero on the left of the first non-zero digit is not significant.
0.0161 also has three significant figures as zeros on the left of the first non-zero digit are not significant.

### 1.3 Atomic and molecular masses

1. (A)

Average atomic mass
$=\frac{\text { Sum of (Isotopic mass } \times \text { its abundance) }}{100}$
Average atomic mass $=\frac{(19 \times 10)+(81 \times 11)}{100}$

$$
=10.81 \approx 10.8
$$

2. (D)

Average atomic mass
$=\frac{\text { Sum of (Isotopic mass } \times \text { its abundance) }}{100}$
Average isotopic mass of X
$=\frac{(200 \times 90)+(199 \times 8)+(202 \times 2)}{100}$
$=200$ a.m.u.

### 1.4 Mole concept and molar mass

1. (D)

## At NTP,

$1 \mathrm{~mol} \mathrm{~N}_{2} \mathrm{O}=22400$ cc $\mathrm{N}_{2} \mathrm{O}=6.02 \times 10^{23} \mathrm{~N}_{2} \mathrm{O}$ molecules
$\therefore \quad 1 \mathrm{cc}$ of $\mathrm{N}_{2} \mathrm{O}=\frac{6.02 \times 10^{23}}{22400}$ molecules
Each $\mathrm{N}_{2} \mathrm{O}$ molecule contains 3 atoms, Hence,
$\therefore \quad 1 \mathrm{cc} \mathrm{N}_{2} \mathrm{O}=\frac{3 \times 6.02 \times 10^{23}}{22400}=\frac{1.8 \times 10^{22}}{22400}$
Nitrogen contains 7 electrons while $O$ contains 8 electrons. Hence, the number of electrons in one molecule of $\mathrm{N}_{2} \mathrm{O}$ is 22 .
Hence,
Number of electrons in 1 cc $\mathrm{N}_{2} \mathrm{O}$

$$
=\frac{6.02 \times 10^{23}}{22400} \times 22=\frac{1.32}{224} \times 10^{23} \text { electrons }
$$

2. (A)

Number of moles in 4.4 g of $\mathrm{CO}_{2}$
$=\frac{4.4}{44}=0.1$
Number of oxygen atoms in 1 mole of $\mathrm{CO}_{2}$ $=2 \times \mathrm{N}_{\mathrm{A}}$
$\therefore \quad$ Number of oxygen atoms in 0.1 mole of $\mathrm{CO}_{2}$
$=0.1 \times 2 \times \mathrm{N}_{\mathrm{A}}$
$=0.2 \times 6.022 \times 10^{23}$
$=1.20 \times 10^{23}$
3. (C)

One litre of $\mathrm{O}_{2}$ contains N molecules at $15{ }^{\circ} \mathrm{C}$ and 150 mmHg pressure. If 1 L of one gas contains N molecules then 2 L of any gas under the same conditions will contain 2 N molecules.
4. (D)

1 L of air $=1000 / 0.21=210 \mathrm{~mL}$ of $\mathrm{O}_{2}$
$\because \quad 22400 \mathrm{~mL}=1 \mathrm{~mole}$
$\therefore \quad 210 \mathrm{~mL}=\frac{1}{22400} \times 210=0.0093 \mathrm{~mol}$
5. (B)

Weight of volatile gas $=0.24 \mathrm{~g}$
Volume of gas $=45 \mathrm{~mL}=0.045 \mathrm{~L}$
Density $=\frac{\text { Mass }}{\text { Volume }}$
Mass of 45 mL of $\mathrm{H}_{2}=0.089 \times 0.045$

$$
=4.005 \times 10^{-3} \mathrm{~g}
$$

Vapour density
$=\frac{\text { Mass of certain volume of vapour }}{\text { Mass of same volume of hydrogen }}$
$=\frac{0.24}{4.005 \times 10^{-3}}=59.93$

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