# CHEMISTRY <br> JEE-MAIN (July-Attempt) 20 July <br> (Shift-1) Paper 

## SECTION -A

1. Compound $A$ is converted to $B$ on reaction with $\mathrm{CHCl}_{3}$ and KOH . The compound $B$ is toxic and can be decomposed by $\mathrm{C} . \mathrm{A}, \mathrm{B}$ and C respectively are :
(1) secondary amine, nitrile compound, conc. NaOH
(2) primary amine, isonitrile compound, conc. HCl
(3) secondary amine, isonitrile compound, conc. NaOH
(4) primary amine, nitrile compound, conc. HCl

## Sol. (2)


(A)
(B)
2. According to the valence bond theory the hybridization of central metal atom is $\mathrm{dsp}^{2}$ for which one of the following compounds ?
(1) $\mathrm{Na}_{2}\left[\mathrm{NiCl}_{4}\right]$
(2) $\mathrm{NiCl}_{2} \cdot 6 \mathrm{H}_{2} \mathrm{O}$
(3) $\mathrm{K}_{2}\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]$
(4) $\left[\mathrm{Ni}(\mathrm{CO})_{4}\right]$

## Sol. (3)

$\mathrm{NiCl}_{2} .6 \mathrm{H}_{2} \mathrm{O}$
$\mathrm{Ni}^{+2} \rightarrow[\mathrm{Ar}]_{18} 3 \mathrm{~d}^{8} 4 \mathrm{~s}^{0}$
CTN. $=6$ octahedral

(2) $\mathrm{K}_{2}\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]$

$$
\mathrm{Ni}^{+2} \rightarrow[\mathrm{Ar}]_{8} 3 \mathrm{~d}^{8} 4 \mathrm{~s}^{0}
$$

C. N. $=4 \quad$ Strong field ligand
C.N. $\rightarrow$ Strong field


Hydridisati
on

Square planar

3)

$$
\mathrm{d}_{\mathrm{yz}} \quad \mathrm{~d}_{\mathrm{xz}}
$$

CO- Strong field
$\mathrm{Ni} \rightarrow$



Hybridisati
4) $\mathrm{Na}_{2}\left[\mathrm{NiCl}_{4}\right]$

$$
\mathrm{Ni} \rightarrow[\mathrm{Ar}]_{18} 3 \mathrm{~d}^{8} 4 \mathrm{~s}^{0}
$$

$\mathrm{Cl}^{\ominus} \rightarrow$ weak
Field ligand

$t_{2}$ Hybridisation
C. N. $=4$
tetrahedral splitting

3. The set in which compounds have different nature is :
(1) $\mathrm{B}(\mathrm{OH})_{3}$ and $\mathrm{H}_{3} \mathrm{PO}_{3}$
(2) $\mathrm{B}(\mathrm{OH})_{3}$ and $\mathrm{Al}(\mathrm{OH})_{3}$
(3) NaOH and $\mathrm{Ca}(\mathrm{OH})_{2}$
(4) $\mathrm{Be}(\mathrm{OH})_{2}$ and $\mathrm{Al}(\mathrm{OH})_{3}$

Sol. (2)
(1) $\mathrm{B}(\mathrm{OH})_{3}$ acidic and $\mathrm{H}_{3} \mathrm{PO}_{3}$ acidic
(2) $\mathrm{B}(\mathrm{OH})_{3}$ acidic and $\mathrm{Al}(\mathrm{OH})_{3}$ amphoteric
(3) NaOH basic and $\mathrm{Ca}(\mathrm{OH})_{2}$ basic
(4) $\mathrm{Be}(\mathrm{OH})_{2}$ amphoteric and $\mathrm{Al}(\mathrm{OH})_{3}$ amphoteric
4.

$\xrightarrow[\mathrm{H}_{2} \mathrm{O}, 273 \mathrm{~K}]{\mathrm{KMnO}_{4}} \xrightarrow{\mathrm{KMajor}^{\prime}{ }^{\prime}}$ (moduct)
For above chemical reactions, identify the correct statement from the following:
(1) Compound' A 'is dicarboxylic acid and compound ' B ' is diol
(2) Compound ' A ' is diol and compound ' B ' is dicarboxylic acid
(3) Both compound ' A ' and compound ' B ' are diols
(4) Both compound 'A' and compound 'B' are dicarboxylic acids

## Sol. (1)



$\mathrm{KMnO}_{4} / \mathrm{H}_{2} \mathrm{SO}_{4} / \Delta$ act as a strong oxidising agent where as $\mathrm{KMnO} \mathrm{O}_{4} / \mathrm{H}_{2} \mathrm{O} / 273 \mathrm{~K}$ is a mild oxidising agent.
5.

(I)

(II)

(III)

(IV)

Which among the above compound/s does/do not form Silver mirror when treated with Tollen's reagent?
(1) Only (II)
(2)(I), (III) and (IV) only
(3)(III) and (IV) only
(4)Only (IV)

## Sol. (1)

Aldehydes and hemiacetal give $\oplus$ ve Tollen's Test (Silver mirror test)
(I)


CHO
(II)


Ketone
(III)

(IV)


Tollen's test Positive

Negative

Hemiacetal
6. Green chemistry in day-to-day life is in the use of:
(1)Chlorine for bleaching of paper
(2)Liquified $\mathrm{CO}_{2}$ for dry cleaning of clothes
(3)Large amount of water alone for washing clothes
(4)Tetrachloroethene for laundry

## Sol. (2)

Chlorine gas was used earlier for bleaching paper. These days, hydrogen peroxide $\left(\mathrm{H}_{2} \mathrm{O}_{2}\right)$ with suitable catalyst.
Tetra chlroroethene $\left(\mathrm{Cl}_{2} \mathrm{C}=\mathrm{CCl}_{2}\right)$ was earlier use as solvent for dry cleaning. The compound contaminates the ground water and is also asuspected carcinogen. Replacement of halogenatedsolvent by liquid $\mathrm{CO}_{2}$ will result in less harm to groundwater.
Hence given statement (2) is correct.
7.

(A)
(B)
(C)
(D)

Among the given species the Resonance stabilised carbocations are:
(1)(C) and (D) only
(2) (A), (B) and (C) only
(3)(A), (B) and (D) only
(4)(A) and (B) only

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## Sol. (4)

(A) and (B) only in Resonance
(A)


(B)

8. The metal that can be purified economically by fractional distillation method is:
(1)Ni
(2)Cu
(3)Zn
(4)Fe

Sol. (3)
Zinc can be purified economically by fractionaldistillation.
9. The conditions given below are in the context of observing Tyndall effect in colloidal solutions:
(A)The diameter of the colloidal particles is comparable to the wavelength of light used.
(B) The diameter of the colloidal particles is much smaller than the wavelength of light used.
(C)The diameter of the colloidal particles is much larger than the wavelength of light used.
(D) The refractive indices of the dispersed phase and the dispersion medium are comparable.
(E)The dispersed phase has a very different refractive index from the dispersion medium.

Choose the most appropriate conditions from the options given below:
(1) (B) and
(E) only
(2) (C) and (D) only
(3) (A) and
(E) only
(4) (A) and (D) only

## Sol. (3)

The phenomenon of scattering of light by colloidalparticles as a result of which the path of the beambecomes visible is called a tyndall effect.smaller the diameter and similar the magnitude ofrefractive indices, lesser is the scattering and hence the tyndall effect and viced-versa. The diameter of the dispersed phase particle should not be smaller than the wavelength of light usedbecause they won't be able to scatter the light so,therefore, the diameter of the dispersed particlesshould be equal or not much smaller than the wavelength of the light used.
2. The refractive indies (i.e. the ratio of the velocity of light in vacuum to the velocity of light in any medium) of the dispersed phase and thedispersion medium should differ greatly inmagnitude than only the particles will be able to scatter the light and tyndall effect will be obersved.On the other hand, if the refractive indices of the dispersed phase and dispersion medium are almostsimilar in magnitude, then there will be noscattering of light and hence, therefore, no tyndalleffect effect is observed.
Hence answer (A) and (E) are correct.
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10. Given below are two statements: One is labelled as Assertion $\mathbf{A}$ and other is labelled as

## Reason R.

Assertion A : The dihedral angles in $\mathrm{H}_{2} \mathrm{O}_{2}$ in gaseous phase is $90.2^{\circ}$ and in solid phase is $111.5^{\circ}$.

Reason R : The change in dihedral angle in solid and gaseous phase is due to the difference in the intermolecular forces.
Choose the most appropriate answer from the options given below for $A$ and $R$.
(1)Both $A$ and $R$ are correct but $R$ is not the correct explanation of $A$.
(2)A is correct but $R$ is not correct.
(3)Both $A$ and $R$ are correct and $R$ is the correct explanation of $A$.
(4)A is not correct but $R$ is correct.

Sol. (4)

(a) Gas phase
(a) Solid phase
(a) $\mathrm{H}_{2} \mathrm{O}_{2}$ structure in gas phase, dihedral angle is $111.5^{\circ}$.
sssss(b) $\mathrm{H}_{2} \mathrm{O}_{2}$ structure in solid phase at 110 K , dihedral angle is $90.2^{\circ}$
Hence given statement (A) is not correct
But statement (B) is correct.
11. Orlon fibres are made up of:
(1) Polyacrylonitrile
(2) Cellulose
(3) Polyamide
(4) Polyesters

Sol. (1)
$\rightarrow$ orlon fibers are made up of Polyacrylonitrile

12. In the given reaction 3-Bromo-2,2-dimethyl butane


Product A is :
(1) 2-Hydroxy-3,3-dimethyl butane.
(2) 2-Ethoxy-2,3-dimethyl butane.
(3) 2-Ethoxy-3,3-dimethyl butane.
(4) 1-Ethoxy-3,3-dimethyl butane

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## Sol. (2)


13. The correct order of intensity of colors of the compounds is:
(1) $\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}>\left[\mathrm{NiCl}_{4}\right]^{2-}>\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$
(2) $\left[\mathrm{NiCl}_{4}\right]^{2-}>\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}>\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}$
(3) $\left[\mathrm{NiCl}_{4}\right]^{2-}>\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}>\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}$
(4) $\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}>\left[\mathrm{NiCl}_{4}\right]^{2-}>\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}$

## Sol. (3)

$\left[\mathrm{NiCl}_{4}\right]^{2-}>\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}>\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{-2-}$
Splitting $\Delta_{\mathrm{t}}<\Delta_{0} \quad<\Delta_{\mathrm{sq}}$
energy order
absorbed $\left[\mathrm{NiCl}_{4}\right]^{2-}<\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{2+}<\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}$
energy order
intensity of $\left[\mathrm{NiCl}_{4}\right]^{-2-}>\left[\mathrm{Ni}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6}\right]^{-2+}>\left[\mathrm{Ni}(\mathrm{CN})_{4}\right]^{2-}$
colour of
compound
14. An inorganic Compound ' X ' on treatment with concentrated $\mathrm{H}_{2} \mathrm{SO}_{4}$ produces brown fumes and gives dark brown ring with $\mathrm{FeSO}_{4}$ in presence of concentrated $\mathrm{H}_{2} \mathrm{SO}_{4}$. Also Compound 'X' gives precipitate ' Y ', when its solution in dilute HCI is treated with $\mathrm{H}_{2} \mathrm{~S}$ gas. The precipitate ' Y ' on treatment with concentrated $\mathrm{HNO}_{3}$ followed by excess of $\mathrm{NH}_{4} \mathrm{OH}$ further gives deep blue coloured solution, Compound ' X ' is :
(1) $\mathrm{Cu}\left(\mathrm{NO}_{3}\right)_{2}$
(2) $\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}$
(3) $\mathrm{Pb}\left(\mathrm{NO}_{2}\right)_{2}$
(4) $\mathrm{Co}\left(\mathrm{NO}_{3}\right)_{2}$

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## Sol. (1)



$$
\begin{array}{cc}
\mathrm{FeSO}_{4}+ & \mathrm{H}_{2} \mathrm{SO}_{4}+\mathrm{NO}_{3}^{-} \\
\mathrm{Sol}^{n} & \text { conc. } \\
& \downarrow \\
{\left[\mathrm{Fe}\left(\mathrm{H}_{2} \mathrm{O}\right)_{5}\right.} & (\mathrm{NO})] \mathrm{SO}_{4}
\end{array}
$$

(Dark brown ring)
$\mathrm{Cu}^{2+}+\left(\mathrm{dil} \mathrm{HCl}+\mathrm{H}_{2} \mathrm{~S}\right)$
$\underset{\text { (cation) }}{\mathrm{X}} \quad$ (Group - II reagent)
$\downarrow$
CuS $\downarrow$
(Black ppt)
(Y)

$\therefore \mathrm{X} \rightarrow \mathrm{Cu}\left(\mathrm{NO}_{3}\right)_{2}$
15. Identify the incorrect statement from the following:
(1) Glycogen is called as animal starch
(2) $\beta$-Glycosidic linkage makes cellulose polymer
(3) Amylose is a branched chain polymer of glucose
(4) Starch is a polymer of $\alpha-D$ glucose

## Sol. (3)

Amylose is a linear chain polymer of $\alpha-D$-glucosewhile amylopectine is branched chain polymer of $\alpha$-D-glucose.
16. The correct structure of Rhumann's Purple, the compound formed in the reaction of ninhydrin with proteins is:
(1)

(2)

(3)

(4)


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## Sol. (2)




Ninhydrin Test
17. Given below are two statements. One is labelled as Assertion A and the other is labelled as

Reason R.
Assertion A: Sharp glass edge becomes smooth on heating it upto its melting point.
Reason $\mathbf{R}$ : The viscosity of glass decreases on melting.
Choose the most appropriate answer from the options given below.
(1) Both $\mathbf{A}$ and $\mathbf{R}$ are true and $\mathbf{R}$ is the correct explanation of $\mathbf{A}$
(2) $\mathbf{A}$ is true but $\mathbf{R}$ is false.
(3) $\mathbf{A}$ is false but $\mathbf{R}$ is true.
(4) Both $\mathbf{A}$ and $\mathbf{R}$ are true but $\mathbf{R}$ is NOT the correct explanation of $\mathbf{A}$.

## Sol. (4)

Hence given assertion (A) is correct
reason (R) is correct
but reason is not correct explanation
18. Chemical nature of the nitrogen oxide compound obtained from a reaction of concentrated nitric acid and $\mathrm{P}_{4} \mathrm{O}_{10}$ (in 4:1 ratio) is:
(1) acidic
(2) amphoteric
(3) neutral
(4) basic

Sol. (1)
$4 \mathrm{HNO}_{3}+\mathrm{P}_{4} \mathrm{O}_{10}$
$\downarrow$
$2 \mathrm{~N}_{2} \mathrm{O}_{5}+\left(\mathrm{HPO}_{3}\right)_{4}$
Ans. $\mathrm{N}_{2} \mathrm{O}_{5}$ is acidic in nature.
19. A s-block element (M) reacts with oxygen to form an oxide of the formula $\mathrm{MO}_{2}$. The oxide is pale yellow in colour and paramagnetic. The element (M) is :
(1) Na
(2) K
(3) Ca
(4) Mg

Sol. (2)
(A) $2 \mathrm{Mg}+\mathrm{O}_{2} \rightarrow 2 \mathrm{MgO}$ (Diamagnetic)
(B) $2 \mathrm{Na}+\mathrm{O}_{2} \rightarrow \mathrm{Na}_{2} \mathrm{O}$ (Diamagnetic)
$\underset{\text { (excess) }}{2 \mathrm{O}} \rightarrow \mathrm{Na}_{2} \mathrm{O}_{2}$ (Diamagnetic) (excess)
(C) $2 \mathrm{Ca}+\mathrm{O}_{2} \rightarrow \mathrm{CaO}_{2}$ (Diamagnetic)
$\mathrm{Ca}+\mathrm{O}_{2} \rightarrow \mathrm{CaO}_{2}$ (Diamagnetic)
(D) $\mathrm{K} \underset{\text { (excess) }}{+\mathrm{O}_{2} \rightarrow \mathrm{KO}_{2}}$ (Paramagnetic)
20. The species given below that does NOT show disproportionation reaction is :
(1) $\mathrm{BrO}_{2}^{-}$
(2) $\mathrm{BrO}_{4}^{-}$
(3) $\mathrm{BrO}^{-}$
(4) $\mathrm{BrO}_{3}^{-}$

## Sol. (2)

In $\mathrm{BrO}_{4}^{\ominus}, \mathrm{Br}$ is in highest oxidation state (+7), Soit cannot oxidise further it only reduced hence it cannot show disproportionation reaction
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## SECTION -B

1. The number of lone pairs of electrons on the central I atom in $\mathrm{I}_{3}^{-}$is $\qquad$
Sol. 3
$\mathrm{I}_{3}^{-}$:


The number of lone pairs of electron on the central atom is 3 .
2. An average person needs about 10000 kJ energy per day. The amount of glucose (molar mass $=180.0 \mathrm{~g} \mathrm{~mol}^{-1}$ ) needed to meet this energy requirement is $\qquad$ g.
(Nearest integer)
(Use : $\Delta_{\mathrm{C}} \mathrm{H}$ (glucose) $=-2700 \mathrm{~kJ} \mathrm{~mol}^{-1}$ )
Sol. 667
2700 kJ energy requires $=180 \mathrm{gm}$
10000 kg energy requires $=\frac{180 \times 10000}{2700} \mathrm{gm}$
mass of glucose $=667 \mathrm{gm}$
3. To synthesise 1.0 mole of 2-methylpropan-2-ol from Ethylethanoate $\qquad$ equivalents of $\mathrm{CH}_{3} \mathrm{MgBr}$ reagent will be required. (Integer value)

## Sol. 2



2 - Methylpropan - 2 ol
4. The Azimuthal quantum number for the valence electrons of $\mathrm{Ga}^{+}$ion is $\qquad$ .
(Atomic number of $\mathrm{Ga}=31$ )

## Sol. 0

$\mathrm{Ga}^{+}$: $\mathrm{Is}^{2} 2 \mathrm{~s}^{2} 2 \mathrm{P}^{6} 3 \mathrm{~s}^{2} 3 \mathrm{p}^{6} 3 \mathrm{~d}^{10} 4 \mathrm{~s}^{2}$
last orbital $=\mathrm{s}$
The azimuthal quantum number for the valenceelectrons (4s-subshell) of Ga+ ion is zero(0).
5. At $20^{\circ} \mathrm{C}$, the vapour pressure of benzene is 70 torr and that of methyl benzene is 20 torr. The mole fraction of benzene in the vapor phase at $20^{\circ} \mathrm{C}$ above an equimolar mixture of benzene and methyl benzene is $\qquad$ $\times 10^{-2}$ (Nearest integer)

Sol. 78
$P_{B}^{\circ}=70 \quad P_{T}^{\circ}=20 \quad X_{B}=0.5=X_{M}$
Now. $\mathrm{Y}_{\mathrm{B}}=\frac{\mathrm{X}_{B} \mathrm{P}_{B}^{\circ}}{\mathrm{X}_{\mathrm{B}} \mathrm{P}_{B}^{\circ}+\mathrm{X}_{M} \mathrm{P}_{M}^{\circ}}$
$=\frac{70 \times 0.5}{70 \times 0.5+20 \times 0.5}$
$=0.777 \Rightarrow 77.7 \times 10^{-2} \Rightarrow 78 \times 10^{-2}$
6. 250 mL of 0.5 M NaOH was added to 500 mL of 1 M HCl The number of unreacted HCl molecules in the solution after complete reaction is $\qquad$ $\times 10^{21}$. (Nearest integer) $\left(N_{A}=6.022 \times 10^{23}\right)$

## Sol. 226

We known that no. of moles $=$ Vlitre $\times$ Molarity\& No. of millimoles $=\mathrm{V}_{\mathrm{ml}} \times$ Molarity so millimoles of $\mathrm{NaOH}=250 \times 0.5=125$
Millimoles of $\mathrm{HCl}=500 \times 1=500$
Now reaction is
$\mathrm{NaOH}+\mathrm{HCL} \rightarrow \mathrm{NaCl}+\mathrm{H}_{2} \mathrm{O}$
$\mathrm{t}=0 \quad 125 \quad 500 \quad 0 \quad 0$
$\begin{array}{lllll}\mathrm{t}=0 & 0 & 375 & 125 & 125\end{array}$
so millimoles of HCl left $=375$
Moles of $\mathrm{HCl}=375 \times 10^{-3}$
No. of HCl molecules $=6.022 \times 10^{23} \times 375 \times 10^{-3}$
$=225.8 \times 10^{21}$
$\approx 226 \times 10^{21}=226$
7. The number of nitrogen atoms in a semicarbazone molecule of acetone is $\qquad$
Sol. 3

8. The spin-only magnetic moment value for the complex $\left[\mathrm{Co}\left(\mathrm{CN}_{6}\right)\right]^{4-}$ is $\qquad$ BM [At. no. of $\mathrm{Co}=27$ ]
Sol. 2
$\left[\mathrm{CO}(\mathrm{CN})_{6}\right]^{4-}$
$x+6 \times(-1)=-4$
$x=+2$
$\mathrm{Co}^{2+}$ : [Ar] 3d ${ }^{7}$
and $\mathrm{CN}^{-}$is a strong field ligand which can pair electron of central atom.

Transference of electrons


It has one unpaired electron ( $n$ ) in 4d-subshell.So spin only magnetic moment $(\mu)=\sqrt{n(n+2)}$ B.M where $\mathrm{n}=$ number of unpaired electrons.
$\mu=\sqrt{3}$ B.M $\quad \mu=1.73 \mathrm{BM}$
9. $2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{SO}_{3}(\mathrm{~g})$

In an equilibrium mixture, the partial pressures are
$\mathrm{P}_{\mathrm{SO}_{3}}=43 \mathrm{kPa} ; \mathrm{P}_{\mathrm{O}_{2}}=530 \mathrm{~Pa}$ and
$\mathrm{P}_{\mathrm{SO}_{2}}=45 \mathrm{kPa}$ The equilibrium constant $\mathrm{K}_{\mathrm{p}}=$ $\qquad$ $\times 10^{-2}$. (Nearest integer)
Sol. 172 (BY NTA) Motion (17228)
$2 \mathrm{SO}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})}=2 \mathrm{SO}_{3(9)}$
$\mathrm{K}_{\mathrm{p}}=\frac{\left(\mathrm{pSO}_{3(9)}\right)^{2}}{\mathrm{pSO}_{2(9)}} \times \mathrm{pO}_{2(9)}$

$$
=\frac{43 \times 43}{45 \times 45} \times 530 \mathrm{~Pa}^{-1}=172.28 \times 10^{-5} \mathrm{~Pa}^{-1}
$$

$=174.498 \mathrm{~atm}^{-1}$
$=17449.8 \times 10^{-2} \mathrm{~atm}^{-1}$

## Ans. is 17228

10. The inactivation rate of a viral preparation is proportional to the amount of virus. In the first minute after preparation, $10 \%$ of the virus is inactivated. The rate constant for viral inactivation is $\qquad$ $\times 10^{-3} \mathrm{~min}^{-1}$. (Nearest integer)
[Use: $\ln 10=2.303 ; \log _{10} 3=0.477 ;$ property of $\log$ arithm $: \log x^{y}=y \log x$ ]

## Sol. 106

As the unit of rate constant is min-1 so it must be afirst order reaction
$K \times t=2.303 \log A_{0} / A_{t}$
in $1 \mathrm{~min} 10 \%$ is in activated so tabing
$A_{0}=100 A_{t}=90$ in 1 min
So $K \times 1=2.303 \times \log \frac{100}{90}$
$=2.303 \times(\log 10-2 \log 3)$
$=2.303 \times(1-2 \times 0.477)=0.10593$
$=105.93 \times 10^{-3}$
$\approx 106$

