

# JEE I NEET I Foundation

MOTION

25000+ SELECTIONS SINCE 2007

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Javant Chittora (JC Sir) Sr. Faculty Exp. : 16 yrs



Anurag Garg (AG Sir) Sr. Faculty Exp. : 17 yrs



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## Join JEE DROPPER BATCH Kota Classroom

English & Hindi Medium

Batch Starting from: 4th August 2021

Online + Offline Mode



### SECTION -A

- Which one of the following gases is reported to retard photosynthesis? (1)CO (2) CFCs (3)  $NO_2$  (4)CO<sub>2</sub>
- **Sol.** (3) According to NCERT only NO<sub>2</sub> from the given options can retard the photosynthesis process in plants.
- **2.** Which one of the following species doesn't have a magnetic moment of 1.73 BM. (spin only value)?
  - (1)CuI
- $(2)[Cu(NH_3)_4]Cl_2$
- $(3)0_{2}^{+}$
- $(4) O_{2}^{-}$

Sol. (1)

Species must not contain single unpaired

$$O_{2}^{+} \rightarrow \sigma_{1s}^{2} < \sigma_{1s}^{*2} < \sigma_{2s}^{2} < \sigma_{2s}^{*2} < \sigma_{2pz}^{2} < \pi_{2px}^{2} = \pi_{2py}^{2} < \pi_{2px}^{*1} = \pi_{2py}^{*}$$

unpaired  $e^- = 1$   $\therefore \mu = 1.73$  BM

(1) 
$$Cu^+ I^- Cu+ \rightarrow [Ar]3d^{10}$$
... unpaired  $e^- = 0$   $I^- \rightarrow [Xe]$ ... unpaired  $e^- = 0$  therefore  $\mu = 0$ 

3.  $[Cu(NH_3)_4]Cl_2$ 

$$Cu \rightarrow [A] \ 3d^3 \therefore \ unpaired = 1 \quad \therefore \ \mu = 1.73 \ BM$$

4.  $O_2 \rightarrow 0$ 

$$\sigma_{1s}^{2} < \sigma_{1s}^{*2} < \sigma_{2s}^{*2} < \sigma_{2s}^{*2} < \sigma_{2px}^{2} < \pi_{2px}^{2} = \pi_{2py}^{2} < \pi_{2px}^{*1} = \pi_{2py}^{*}$$

(11e<sup>-</sup>)

∴ unpaired ∴ 
$$\mu = 1.73$$
 BM

0

$$NH_2$$
 KOBr A (major product)

Br

Br

0

NH<sub>2</sub> LiAlH<sub>4</sub> B 
$$H_3O^+$$
 (major product)

In the above reactions, product A and product B respectively are:

$$(1)$$
  $NH_2$   $NH_2$   $NH_2$   $NH_2$ 

$$(2) \begin{array}{c} NH_2 \\ Br \end{array}, \begin{array}{c} NH_2 \\ Br \end{array}$$

### Sol. (2)

$$NH$$
 $(KOBr)$ 
 $NH$ 
 $(Hoffmann's)$ 
 $(KOBr)$ 
 $(Hoffmann's)$ 
 $(Reduction of amide)$ 
 $(Reduction of amide)$ 

**4.** The major product (P) in the following reaction is :

CHO 
$$\stackrel{\hbox{O}}{}$$
 (i) (KOH (alc.).  $\stackrel{\hbox{P}}{}$  O  $\stackrel{\hbox{(ii)}}{}$   $\stackrel{\hbox{H}^+}{}$ ,  $\stackrel{\hbox{$\varDelta$}}{}$  (major product)

$$(1) \begin{picture}(200,0) \put(0,0){\line(1,0){100}} \put(0,0){\line(1,0$$

### Sol. (4)

$$\begin{array}{c} CH = O \\ CH_2 - CH_2 \\ O \\ \end{array} \begin{array}{c} C = O \\ (i) \text{ KOH (alc.)} \\ (i) H^+, \Delta \\ -H_2O \\ \end{array} \begin{array}{c} O \\ \end{array}$$

Intramolecular aldol condensation

- **5.** Benzene on nitration gives nitrobenzene in presence of  $HNO_3$  and  $H_2SO_4$  mixture, where :
  - (1) HNO<sub>3</sub> acts as an acid and H<sub>2</sub>SO<sub>4</sub> acts as a base
  - (2) both H<sub>2</sub>SO<sub>4</sub> and HNO<sub>3</sub> act as a bases
  - (3) HNO<sub>3</sub> acts as a base and H<sub>2</sub>SO<sub>4</sub> acts as an acid
  - (4)both H<sub>2</sub>SO<sub>4</sub> and HNO<sub>3</sub> act as an acids
- Sol. (3)

Reagent for nitration of Benzene

$$\begin{array}{c} \textbf{H}_2\textbf{SO}_4 + \textbf{HNO}_3 \\ \text{(Acid)} & \text{(base)} \end{array} \rightleftarrows \textbf{HSO}_4^- + \textbf{H}_2 \overset{\text{\tiny o}}{\textbf{N}} \textbf{O}_2$$

$$H_2 \overset{\oplus}{N} O_3 \rightleftharpoons H_2 O + \overset{\oplus}{N} O_2$$

- **6.** Metallic sodium does not react normally with:
  - (1) tert-butyl alcohol

(2) But-2-yne

(3) Ethyne

(4) gaseous ammonia

### Sol. (2)

Metallic sodium do not react with 2-butyne because 2-butyne does not have acidic hydrogen

**7.** The hybridisations of the atomic orbitals of nitrogen in  $NO_{2}^{-}$ ,  $NO_{2}^{+}$  and  $NH_{4}^{+}$  respectively are.

(1)  $sp^2$ , sp and  $sp^3$ 

(2) sp,  $sp^2$  and  $sp^3$ 

(3) sp<sup>3</sup>, sp and sp<sup>2</sup>

(4)  $sp^3$ ,  $sp^2$  and sp

$$O = \stackrel{\oplus}{N} = O$$
  $\stackrel{2\sigma + 1}{\circ} \stackrel{lp}{sp^2}$ 

**8.** Consider two chemical reactions (A) and (B) that take place during metallurgical process:

(A) 
$$ZnCO_{3(s)} \xrightarrow{\Delta} ZnO_{(s)} + CO_{2(g)}$$

$$\text{(B)}\,2ZnS_{(s)}^{\phantom{\dagger}}+3O_{2(g)}^{\phantom{\dagger}} \xrightarrow{\phantom{\Delta}} 2ZnO_{(s)}^{\phantom{\dagger}}+2SO_{2(g)}^{\phantom{\dagger}}$$

The **correct** option of names given to them respectively is:

- (1) (A) is roasting and (B) is calcination
- (2) (A) is calcination and (B) is roasting
- (3) Both (A) and (B) are producing same product so both are calcination
- (4) Both (A) and (B) are producing same product so both are roasting
- Sol. (2)

(A) 
$$ZnCO_3$$
 (S)  $\xrightarrow{\Delta}$   $ZnO(s) + CO_2(g)$ 

Heating in absence of oxygen in calcination.

(B) 
$$2ZNs(s) + 3O_2(g) \rightarrow 2ZnO(g) + 2SO_2(g)$$

heating in presence of oxygen in roasting

Hence (A) is calcination while (B) in roasting

- **9.** The single largest industrial application of dihydrogen is :
  - (1) Rocket fuel in space research
- (2) Manufacture of metal hydrides
- (3) In the synthesis of ammonia
- (3) In the synthesis of nitric acid.

Sol. (3)

Informative, according to ncert uses of di hydrogen.

In fact NH<sub>3</sub> largest production in used to manufacture nitrogenous fertilisers.

- **10.** Cu<sup>2+</sup> salt reacts with potassium iodide to give :
  - (1)  $Cu_2I_3$
- (2) CuI
- (3)  $Cu_2I_2$
- (4)  $Cu(I_3)_2$

Sol. (3)

### Motion 2 & 3

$$2Cu^{+2} + 4I^{-} \longrightarrow Cu_{2}I_{2}(s) + I_{2}$$
  
 $2Cu^{+2} + 3I^{-} \longrightarrow 2CuI + I_{2}$ 

11. Br 
$$(i) (C_6H_5CO)_2O_2$$
, HBr  $P$  (major product)

Major product P of above reaction, is:

Sol. (1) 
$$\frac{(C_6H_5CO)_2O_2, HBr}{Anti markownikoff}$$

Br 
$$\bigcirc$$
 CoF<sub>2</sub> (Swart reaction)

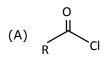
**12.** Which one of the following statements is not true about enzymes?

reaction product

- (1) Enzymes work as catalysts by lowering the activation energy of a biochemical reation.
- (2) The action of enzymes is temperature and pH specific.
- (3) Almost all enzymes of proteins.
- (4) Enzymes are non-specific for a reaction and substrate.
- Sol. (4)

Enzymes are highly specific both in the reactions that they catalyzed and in their choice of reactions, which are called substrates.

13.



$$(D) \quad R \qquad N \qquad H$$

The correct order of their reactivity towards hydrolysis at room temperature is :

Sol. (3)

Reactivity

towards A > B > C > D

Hydrolysis

**14.** Spin only magnetic moment of an octahedral complex of  $Fe^{2+}$  in the presence of a strong field ligand in B.M. is

- (1) 4.89
- (2) 0
- (3) 2.82

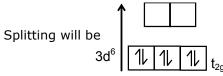
(4)3.46

Sol. (2)

In presence of SFL  $\Delta_0$ > P means pairing occurs therefore

.....

For Fe+2 3d<sup>6</sup>



 $\therefore$  No of unpaired e- (s) = 0

$$\therefore \mu = \sqrt{n(n(n+2)}BM = 0$$

[n = No of unpaired e-(s)]

In NiCl<sub>2</sub> Ni<sup>+2</sup> is having configuration 3d<sup>8</sup>

∴ Number of unpaired electron = 2

After formation of oxidised product

 $[Ni(CN)_6]^{-2} Ni^{+4}$  is obtained

 $\text{Ni}^{+4}\!\!\Rightarrow 3\text{d}^6$  and  $\text{CN}^{\scriptscriptstyle{-}}$  is strong field ligand

 $\therefore$  number of unpaired electrons = 0

 $\therefore$  The charge is 2 - 0 = 2

- **15.** In Carius method, halogen containing organic compound is heated with fuming nitric acid in the presence of :
  - (1) AgNO<sub>3</sub>
- (2) HNO<sub>3</sub>
- (3) BaSO<sub>4</sub>
- (4)CuSO<sub>4</sub>

Sol. (1)

Organic compound is heated with fuming nitric acid in the presence of silver nitrate in carius method.

Lunar caustic (AgNO $_3$ ) is used as reagent hare to distinguish Cl $^-$ , Br and I $^-$  respectively as follows.

- $Cl^{-}(aq) \xrightarrow{AgNO_3} AgCl \downarrow_{ppt} white$
- $Br^{-}(aq) \xrightarrow{AgNO_3} AgBr \downarrow_{ppt} pale yellow$
- $I^{-}(aq) \xrightarrow{AgNO_3} AgNO_3 AgI \downarrow_{ppt} Dark yellow$
- **16.** Bakelite is a cross-linked polymer of formaldehyde and :
  - (1) Buna-S
- (2) Dacron
- (3) Novolac
- (4) PHBV

Sol. (3)

Novolac (phenol formaldehyde Resin) → Bakelite

- **17.** Which one of the following pairs of isomers is an example of metamerism?
  - (1)  $CH_3CH_2CH_2CH_3$  and  $H_3C-C-CH_3$  (2)
    - (2) and
  - (3)  $H_5C_6$  OH and  $H_5C_6$
- (4)  $C_6H_5$  and  $H_5C_6$

- Sol. (2)
  - are metamers
- **18.** A solution is 0.1 M in  $\text{Cl}^-$  and 0.001 M in  $\text{CrO}_4^{2-}$ . Solid AgNO<sub>3</sub> is gradually added to it.

Assuming that the addition does not change in volume and  $K_{sp}(AgCl) = 1.7 \times 10^{-10} \text{ M}^2$  and  $K_{sp}(Ag_2CrO_4) = 1.9 \times 10^{-12} \text{ M}^3$ .

Select correct statement from the following:

- (1) AgCl will precipitate first as the amount of Ag<sup>+</sup> needed to precipitate is low.
- (2) AgCl precipitates first because its  $K_{sp}$  is high.
- (3) Ag<sub>2</sub>CrO<sub>4</sub> precipitates first because the amount of Ag<sup>+</sup> needed is low.
- (4) Ag<sub>2</sub>CrO<sub>4</sub> precipitates first as its K<sub>sp</sub> is low.
- Sol. (1)
  - (i) [Ag<sup>+</sup>] required to ppt AgCl(s)

$$Ksp = IP = [Ag^+] [CI^-] = 1.7 \times 10^{-10}$$

 $[Ag^+] = 1.7 \times 10^{-9}$ 

(ii) [Ag<sup>+</sup>] required to ppt Ag<sub>2</sub>CrO<sub>4</sub>(s)

$$Ksp = IP = [Ag^{+}]^{2} [CrO_{4}^{-2}] = 1.9 \times 10^{-12}$$

 $[Aq^+] = 4.3 \times 10^{-5}$ 

[Ag<sup>+</sup>] required to ppt AgCl is low so AgCl will ppt 1<sup>st</sup>.

**19.** Outermost electronic configuration of a group 13 element, E, is 4s², 4p¹. The electronic configuration of an element of p-block period-five placed diagonally to element, E is :

(1) [Xe]  $5d^{10} 6s^2 6p^2$ 

(2) [Kr]  $4d^{10} 5s^2 5p^2$ 

(3) [Kr]  $3d^{10} 4s^2 4p^2$ 

(4) [Ar]  $3d^{10} 4s^2 4p^2$ 

Sol. (2)

The element E is Ga and the diagonal element of  $5^{th}$  period is  $_{50}$ Sn having outer electronic configuration will be [Kr]  $5s^2$   $4d^{10}$   $5p^2$ .

20. 
$$NH_2$$

$$273 - 278K$$

$$(major product)$$

$$NH_2$$

$$CH_3$$

$$CH_3$$

$$CH_3$$

$$CH_3$$

$$(major product)$$

Consider the above reaction, compound B is :

(1) 
$$HO_3S$$
  $\longrightarrow$   $N = N$   $\longrightarrow$   $N$   $CH_3$   $CH_3$ 

(2) 
$$N = N - N - N$$

$$CH_3$$

$$CH_3$$

(3) 
$$HO_3S$$
  $N = N$   $CH_3$ 

Sol. (1)

$$NH_2$$
 $NH_2$ 
 $NH_2$ 

### SECTION -B

- For a given chemical A  $\rightarrow$  B at 300 K the free energy change is -49.4 kJ mol<sup>-1</sup> and the enthalpy of reaction is 51.4 kJ mol<sup>-1</sup>. The entropy change of the reaction is \_\_\_\_\_\_ J K<sup>-1</sup> mol<sup>-1</sup>.
- Sol. (360) NTA, (336) Motion

Given chemical reaction:

A 
$$\longrightarrow$$
 B [ $\Delta$ G]<sub>P, T</sub> = -49.4 kJ/mol 
$$\Delta$$
H<sub>rxn</sub> = 51.4 kJ/mol 
$$\Delta$$
S<sub>rxn</sub> = ?

 $\Rightarrow$  From the relation  $[\Delta G]_{P, T} = \Delta H - T\Delta S$ 

$$\Rightarrow \Delta S_{rxn} = \frac{\Delta H_{rxn} - [\Delta G]_{P,T}}{T}$$

$$= \frac{[51.4 - (-49.4)] \times 100}{300} \frac{J}{\text{mol K}}$$

$$\Rightarrow \Delta S_{rxn} = 336 \frac{J}{\text{mol K}}$$

- 2. 100 ml of 0.0018% (w/v) solution of Cl<sup>-</sup> ion was the minimum concentration of Cl<sup>-</sup>reaquired to precipitate a negative sol in one h. The coagulating value of Cl<sup>-</sup> ion is \_\_\_\_\_\_. (Nearest integer)
- Sol. (1) NTA, (Bonus) Motion,
- An aqueous solution of  $NiCl_2$  was heated with excess sodium cyanide in presence of strong oxidizing agent to form  $[Ni(CN)_6]^{2-}$ . The total change in number of unpaired electrons on metal centre is \_\_\_\_\_\_.
- Sol. (2)

 $[Ni(CN)_6]^{2-}$ 

 $Ni^{+4} \rightarrow d^6$  strong field ligand

1	1	1	

Pairing will be there zero unpaired electron

$$NiCl_2 \rightarrow Ni^{2+} \rightarrow d^8$$

Change = 2

When 0.15 g of an organic compound was analyzed using Carius method for estimation of bromine, 0.2397 g of AgBr was obtained. The percentage of bromine in the organic compound is \_\_\_\_\_. (Nearest integer)

[Atomic mass : Silver = 108, Bromine = 80]

Sol. (68)

Moles of Br = Moles of AgBr obtained

$$\Rightarrow \qquad \text{Mass of Br} = \frac{0.2397}{188} \times 80g$$

therefore % Br in the organic compound

$$= \frac{W_{Br}}{W_{T}} \times 100$$

$$= \frac{0.2397 \times 80}{188 \times 0.15} \times 100 = 0.85 \times 80$$

$$= 68$$

- ⇒ Nearest integer is '68'
- The vapour pressures of A and B at 25°C are 90 mm Hg and 15 mm Hg respectively. If A and B are mixed such that the mole fraction of A in the mixture is 0.6, then the mole fraction of B in the vapour phase is  $x \times 10^{-1}$ . The value of x is \_\_\_\_\_. (Nearest integer)
- Sol. (1)

Given  $P_A^{\circ} = 90 \text{ mm Hg}$ , at 25°C

 $P_{B}^{\circ} = 15 \text{ mm Hg}$ 

and 
$$X_A = 0.6$$
  
 $X_B = 0.4$   $P_T = X_A T_A^o + X_B P_B^o$   
 $= (0.6 \times 90) + (0.4 \times 15)$   
 $= 54 + 6 = 60 \text{ mm}$ 

Now mol fraction of B in the vapour phase

i.e. 
$$Y_B = \frac{P_B}{P_T} = \frac{X_B P_B^o}{60} = 0.1 = 1 \times 10^{-1}$$

Therefore: x = 1

- 6. Dimond has a three dimensional structure of C atoms formed by covalent bonds. The structure of diamond has face centred cubic lattice where 50% of the tetrahedral voids are also occupied by carbon atoms. The number of carbon atoms present per unit cell of diamond is \_\_\_\_\_\_.
- Sol. (8)

Carbon atoms occupy FCC lattice points as well as half of the tetrahedral voids

From face z = 4

From 50 % o.v. z = 4

Total = 8

7. 
$$PCl_{5}(g) \rightarrow PCl_{3}(g) + Cl_{2}(g)$$

In the above first order reaction the concentration of  $PCl_5$  reduces from initial concentration 50 mol  $L^{-1}$  to 10 mol  $L^{-1}$  in 120 minutes at 300 K. The rate constant for the reaction at 300 K is X  $\times$  10<sup>-2</sup> min<sup>-1</sup>. The value of x is \_\_\_\_\_. [Given log 5 = 0.6989]

$$\begin{array}{l} \text{PCI}_{5}(g) \xrightarrow{\frac{1 \text{ order}}{300 \text{K}}} \text{PCI}_{3(g)} + \text{CI}_{2(g)} \\ & t = 0 \qquad 50 \text{ M} \\ & t = 120 \text{ min } 10 \text{ M} \\ \\ \Rightarrow & K = \frac{2.303}{t} log \frac{\left[A_{0}\right]}{\left[A_{t}\right]} \\ \\ \Rightarrow & K = \frac{2.303}{t} log \frac{50}{10} \\ \\ \Rightarrow & K = \frac{2.303}{120} \times 0.6989 = 0.013413 \text{ min}^{-1} \\ \\ = & 1.3413 \times 10^{-2} \text{ min}^{-1} \\ 1.34 \Rightarrow \text{Nearest integer} = 1 \end{array}$$

**8.** Potassium chlorate is prepared by electrolysis of KCl in basic solution as shown by following equation.

$$6OH^{-} + Cl^{-} \rightarrow ClO_{3}^{-} + 3H_{2}O + 6e^{-}$$

A current of xA has to be passed for 10h to produce 10.0 g of potassium chlorate. The value of x is \_\_\_\_\_. (Nearest integer)

(Molar mass of KClO<sub>3</sub> = 122.6 g mol<sup>-1</sup>, F = 96500 C)

### Sol. (1)

Given balanced equation is

$$60^{\circ}H + CI \rightarrow CIO_3^- + 3H_2O + 6e^-$$

$$\rightarrow$$
 10g KClO<sub>3</sub> $\Rightarrow \frac{10}{122.6}$  mol KCO3 in obtained

 $\rightarrow$  From the above reaction, it is concluded that by 6F charge 1 mol KClO<sub>3</sub> is obtained.

 $\rightarrow$  By the passage of 6F charge = 1 mol KClO<sub>3</sub>

.. By the passage of 
$$\frac{x \times 10 \times 60 \times 60}{96500}$$
 F charge

$$= \frac{1}{6} \times \frac{\mathbf{x} \times 10 \times 60 \times 60}{96500}$$

Now 
$$\frac{x \times 10 \times 60 \times 60}{6 \times 96500} = \frac{10}{122.6}$$

$$\Rightarrow$$
 x =  $\frac{10 \times 965}{60 \times 122.6} = \frac{965}{735.6} = 1.311 \simeq 1$ 

OR

$$W = \frac{E}{F} \times I \times t$$

$$10 = \frac{122.6}{96500 \times 6} \times x \times 10 \times 3600$$

$$X = 1.311$$

**9.** The wavelength of electrons accelerated from rest through a potential difference of 40 kV is  $X \times 10^{-12}$  m. The value of x is \_\_\_\_\_. (Nearest integer)

Given: Mass of electrons =  $9.1 \times 10^{-31}$  kg

Charge on an electron =  $1.6 \times 10^{-19}$  C

Planck's constant =  $6.63 \times 10^{-34}$  Js

Sol. (6)

De-broglie-wave length of electron:

$$\lambda_{c} = \frac{h}{\sqrt{2m(KE)}} \begin{cases} \because e^{-i}s \text{ accelerated} \\ \text{from rest} \\ \Rightarrow KE = q \times V \end{cases}$$

$$\lambda = \frac{h}{\sqrt{2mqv}}$$

$$= \frac{6.63 \times 10^{-34}}{\sqrt{2 \times 1.6 \times 10^{-19} \times 9.1 \times 10^{-31} \times 40 \times 10^{3}}}$$

$$= 0.614 \times 10^{-11} \text{m}$$

$$= 6.16 \times 10^{-12} \text{m}$$

Nearest integer = 6

OR

$$\lambda = \frac{12.3}{\sqrt{V}}\,\text{\AA}$$

$$=\frac{12.3}{200}=6.15\times10^{-12}m$$

Ans is 6

- 4g equimolar mixture of NaOH and Na<sub>2</sub>CO<sub>3</sub> contains x g of NaOH and y g of Na<sub>2</sub>CO<sub>3</sub>. The value of x is \_\_\_\_\_ g. (Nearest integer)
- Sol. (1)

Total mass = 4g

Now

$$W_{NaOH} + W_{Na_2CO_3} = 4$$

$$\Rightarrow$$
 40a + 106 a = 4

$$\Rightarrow$$
 a =  $\frac{4}{146}$  mol

$$\Rightarrow$$
 therefore mass of NaOH is :  $\frac{4}{146} \times 40g$ 

$$= 1.095 \approx 1$$

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Akhilesh Kanthe (AKK Sir)



/ishal Joshi (VJ Sir)



Surendra K. Mishra (SKM Sir)



Gavesh Bhardwa

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