## **FINAL JEE-MAIN EXAMINATION – MARCH, 2021** (Held On Tuesday 16<sup>th</sup> March, 2021) TIME: 9:00 AM to 12:00 NOON





5. Match List-I with List-II :

List-I	List-II
Industrial process	Application
(a) Haber's process	(i) HNO <sub>3</sub> synthesis
(b) Ostwald's process	(ii) Aluminium
	extraction

(c) Contact process (iii) NH<sub>3</sub> synthesis

(d) Hall-Heroult process (iv)  $H_2SO_4$  synthesis Choose the correct answer from the options given below :

- (1) (a)-(ii), (b)-(iii), (c)-(iv), (d)-(i)
- (2) (a)-(iii), (b)-(iv), (c)-(i), (d)-(ii)
- (3) (a)-(iii), (b)-(i), (c)-(iv), (d)-(ii)
- (4) (a)-(iv), (b)-(i), (c)-(ii), (d)-(iii)

#### Official Ans. by NTA (3)

- **Sol.** (a) Haber's process is used for  $NH_3$  synthesis.
  - (b) Ostwald's process is used for HNO<sub>3</sub> synthesis.
  - (c) Contact process is used for  $H_2SO_4$  synthesis.
  - (d) In Hall-Heroult process, electrolytic reduction of impure alumina can be done.(Aluminium extraction)
- 6. Among the following, the aromatic compounds are :



Choose the correct answer from the following options :

- (1) (A) and (B) only
- (2) (B) and (C) only
- (3) (B), (C) and (D) only
- (4) (A), (B) and (C) only

#### Official Ans. by NTA (2)

Sol. (A) Non-Aromatic (B) Aromatic (C) Aromatic (D) Anti-Aromatic NH<sub>2</sub> <u>NaNO<sub>2</sub>, HCl</u> <u>773 - 278 K</u>"

7.

8.

In the above chemical reaction, intermediate "X" and reagent/condition "A" are :

Major Product

(1) X-
$$(1)^{N_{2}^{+}Cl^{-}}$$
; A-H<sub>2</sub>O/NaOH  
(2) X- $(1)^{NO_{2}}$ ; A-H<sub>2</sub>O/ $(1)^{N_{2}^{+}Cl^{-}}$ ;

(4) X- 
$$(4)$$
 ; A- H<sub>2</sub>O/NaOH

NO

#### Official Ans. by NTA (3)

Sol. NH<sub>2</sub>  
$$NH_2$$
  
 $NaNO_2+ HCl$   
 $Diazotisation$   
Reaction  
 $(A)$   
 $N_2^{\oplus}Cl^{\odot}$   
 $H_2O/\Delta$   
 $(B)$ 

Given below are two statements : Statement I : The E° value of Ce<sup>4+</sup> / Ce<sup>3+</sup> is + 1.74 V.

Statement II : Ce is more stable in  $Ce^{4+}$  state than  $Ce^{3+}$  state.

In the light of the above statements, choose the most appropriate answer from the options given below :

- (1) Both statement I and statement II are correct
- (2) Statement I is incorrect but statement II is correct
- (3) Both statement I and statement II are incorrect
- (4) Statement I is correct but statement II is incorrect

Official Ans. by NTA (4)



- 9. The functions of antihistamine are :
  - (1) Antiallergic and Analgesic
  - (2) Antacid and antiallergic
  - (3) Analgesic and antacid
  - (4) Antiallergic and antidepressant

#### Official Ans. by NTA (2)

- 10. Which of the following is Lindlar catalyst?
  - (1) Zinc chloride and HCl
  - (2) Cold dilute solution of  $KMnO_4$
  - (3) Sodium and Liquid  $NH_3$
  - (4) Partially deactivated palladised charcoal

#### Official Ans. by NTA (4)

Sol. Partially deactivated palladised charcoal  $(H_2/pd/CaCO_2)$  is lindlar catalyst.

H<sub>2</sub>C OH

Sol.

 $\frac{20\% \text{ H}_{3}\text{PO}_{4}}{358 \text{ K}}$ 11. "A" (Major Product)



The product "A" and "B" formed in above reactions are :





- 12. Given below are two statements : Statement I : H<sub>2</sub>O<sub>2</sub> can act as both oxidising and reducing agent in basic medium. Statement II : In the hydrogen economy, the energy is transmitted in the form of dihydrogen. In the light of the above statements, choose the correct answer from the options given below : (1) Both statement I and statement II are false (2) Both statement I and statement II are true (3) Statement I is true but statement II is false (4) Statement I is false but statement II is true Official Ans. by NTA (2)
- Sol. (a)  $H_2O_2$  can acts as both oxidising and reducing agent in basic medium.

(i)  $2Fe^{2+} + H_2O_2 \rightarrow 2Fe^{3+} + 2OH^-$ In this reaction,  $H_2O_2$  acts as oxiding agent.

(ii)  $2 \stackrel{+}{M} nO_4^- + 3H_2O_2 \rightarrow 2 \stackrel{+}{M} nO_2 + 3O_2 + 2H_2O + 2OH^-$ 

- In this reaction, H<sub>2</sub>O<sub>2</sub> acts as reducing agent.
- (b) The basic principle of hydrogen economy is the transportation and storage of energy in the form of liquids or gaseous dihydrogen.

Advantage of hydrogen economy is that energy is transmitted in the form of dihydrogen and not as electric power

The type of pollution that gets increased during 13.

the day time and in the	e presence of O <sub>3</sub> is :
(1) Reducing smog	(2) Oxidising smog
(3) Global warming	(4) Acid rain
Official Ang. by NTA (2)	

**Official Ans. by NTA (2)** Sol. In presence of  $ozone(O_3)$ , oxidising smog gets increased during the day time because automobiles and factories produce main components of the photochemical smog (oxidising smog) results from the action of sunlight on unsaturated hydrocarbon and nitrogen oxide.

> Ozone is strong oxidising agent and can react with the unburnt hydrocarbons in the polluted air to produce chemicals.



+ CHBr.

CH<sub>2</sub>-CH<sub>2</sub>-NH<sub>2</sub>

List-II

(v) + 1

Oxidation state of 'P'

 $NH_3/\Delta$ 



4

- **Sol.** (a) Hypophosphorus acid :  $H_3 \underline{P}O_2$ (+1) 3 + x + (-2)2 = 0 x = +1
  - (b) Orthophosphoric acid :  $H_3\underline{P}O_4$ (+1) 3 + x + (-2)4 = 0 x = +5
  - (c) Hypophosphoric acid :  $H_4P_2O_6$ (+1) 4 + 2x + (-2)6 = 0 x = +4
  - (d) Orthophosphorous acid :  $H_3\underline{P}O_3$ (+1)3 + x + (-2)3 = 0 x = +3
- **18.** Given below are two statements : one is labelled as Assertion A and the other is labelled as Reason R :

**Assertion A :** The H–O–H bond angle in water molecule is 104.5°.

**Reason R :** The lone pair – lone pair repulsion of electrons is higher than the bond pair - bond pair repulsion.

- (1) A is false but R is true
- (2) Both A and R are true, but R is not the correct correct explanation of A
- (3) A is true but R is false
- (4) Both A and R are true, and R is the correct explanation of A

#### Official Ans. by NTA (4)

Sol. H<sub>2</sub>O

 $\theta = 104.5^{\circ}$ 

the hybridisation of oxygen is water molecule is  $sp^{3}$ .

So electron geometry of water molecule is tetrahedral and the bond angle should be 109°28" but as we know that lone pair-lone pair repulsion of electrons is higher than the bond pair-bond pair repulsion because lone pair is occupied more space areound central atom than that of bond pair.

- 19. In chromotography technique, the purification of compound is independent of : (1) Mobility or flow of solvent system (2) Solubility of the compound (3) Length of the column or TLC Plate (4) Physical state of the pure compound Official Ans. by NTA (4) Sol. In chromotography technique, the purification of a compound is independent of the physical state of the pure compound. 20. A group 15 element, which is a metal and forms a hydride with strongest reducing power among group 15 hydrides. The element is : (1) Sb (2) P(3) As (4) Bi Official Ans. by NTA (4) Sol. In group 15  $\begin{bmatrix} N \\ P \end{bmatrix} \rightarrow Non metal$  $As_{Sb}$   $\rightarrow$  Metalloid Bi  $\rightarrow$  Metal Hydrides of group 15 elements are NH, PH, AsH, SbH, BiH<sub>3</sub> In NH<sub>2</sub>, hydrogen atom gets partial positive charge due to less electronegativity. But in BiH<sub>3</sub>, hydrogen atom gets partial negative charge because hydrogen is more electronegative than bismuth. i.e. BiH<sub>3</sub> is a strong reducing agent than others because we know that H<sup>-</sup> is a strong reducing agent. **SECTION-B** 1.
  - For the reaction A(g)  $\rightleftharpoons$  B(g) at 495 K,  $\Delta_r G^o = -9.478 \text{ kJ mol}^{-1.}$

If we start the reaction in a closed container at 495 K with 22 millimoles of A, the amount of B is the equilibrium mixture is \_\_\_\_\_ millimoles. (Round off to the Nearest Integer).

 $[R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}; \ \ln 10 = 2.303]$ 

Official Ans. by NTA (20)

3.

**Sol.**  $\Delta G^{\circ} = -RT \ \ell n \ K_{ac}$ Given  $\Delta G^{\circ} = -9.478$  KJ/mole T = 495K R = 8.314 J mol<sup>-1</sup> So  $-9.478 \times 10^3 = -495 \times 8.314 \times \ln K_{eq}$  $ln K_{eq} = 2.303$  $= \ell n 10$ So  $K_{eq} = 10$  $A(g) \rightleftharpoons B(g)$ Now t = 022 0 22-x t = tx  $K_{eq} = \frac{[B]}{[C]} = \frac{x}{22 - x} = 10$ or x = 20So millmoles of B = 202. Complete combustion of 750 g of an organic

compound provides 420 g of CO<sub>2</sub> and 210 g of H<sub>2</sub>O. The percentage composition of carbon and hydrogen in organic compound is 15.3 and respectively. (Round off to the Nearest Integer)

Official Ans. by NTA (3)

Sol. 44 gm CO<sub>2</sub> have 12 gm carbon

So, 420 gm 
$$\text{CO}_2 \Rightarrow \frac{12}{44} \times 420$$

 $\Rightarrow \frac{1260}{11}$  gm carbon

 $\Rightarrow$  114.545 gram carbon

18 gm H<sub>2</sub>O  $\Rightarrow$  2 gm H<sub>2</sub>

So, % of carbon = 
$$\frac{114.545}{750} \times 100$$

 $\simeq 15.3\%$ 

$$210 \text{ gm} \Rightarrow \frac{2}{18} \times 210$$
  
= 23.33 gm H<sub>2</sub>  
So, % H<sub>2</sub>  $\Rightarrow \frac{23.33}{750} \times 100 = 3.11\%$   
 $\approx 3\%$ 

 $2 \operatorname{Mn} O_4^- + b C_2 O_4^{2-} + c \operatorname{H}^+ \rightarrow x \operatorname{Mn}^{2+} + y \operatorname{CO}_2$  $+ z H_2O$ If the above equation is balanced with integer coefficients, the value of c is (Round off to the Nearest Integer). Official Ans. by NTA (16) Sol. Writting the half reaction oxidation half reaction  $MnO_4^- \rightarrow Mn^{2+}$ balancing oxygen  $MnO_4^- \rightarrow Mn^{2+} + 4H_2O$ balancing Hydrogen  $8H^+ + MnO_4^- \rightarrow Mn^{2+} + 4H_2O$ balancing charge  $5e^{-} + 8H^{+} + MnO_4^{-} \rightarrow Mn^{2+} + 4H_2O$ Reduction half  $C_2O_4^{2-} \rightarrow CO_2$ Balancing carbon  $C_2 O_4^{2-} \rightarrow 2CO_2$ Balancing charge  $C_2O_4^{2-} \rightarrow 2CO_2 + 2e^{-1}$ Net equation

- $16H^{+} + 2MnO_{4}^{-} + 5C_{2}O_{4}^{2-} \rightarrow 10CO_{2} + 2Mn^{2+} + 8H_{2}O_{2}$ So c = 16
- 4.  $AB_2$  is 10% dissociated in water to  $A^{2+}$  and B<sup>-</sup>. The boiling point of a 10.0 molal aqueous solution of AB<sub>2</sub> is \_\_\_\_\_oC. (Round off to the Nearest Integer).

[Given : Molal elevation constant of water  $K_{\rm b} = 0.5 \text{ K kg mol}^{-1}$  boiling point of pure water  $= 100 \circ C$ 

Official Ans. by NTA (106)

**Sol.**  $AB_2 \rightarrow A^{2+} + 2B^$ t = 0 a 0 0  $t = t \quad a - a\alpha \quad a\alpha$ 2aα  $n_{T} = a - a\alpha + a\alpha + 2a\alpha$  $= a (1 + 2\alpha)$ so  $i = 1 + 2\alpha$ Now  $\Delta T_{h} = i \times m \times K_{h}$  $\Delta T_{b} = (1 + 2\alpha) \times m \times K_{b}$  $\alpha = 0.1$  m = 10 K<sub>b</sub> = 0.5  $\Delta T_{\rm h} = 1.2 \times 10 \times 0.5$ = 6 So boiling point = 106

5. The equivalents of ethylene diamine required to replace the neutral ligands from the coordination sphere of the trans-complex of CoCl<sub>3</sub>.4NH<sub>3</sub> is \_\_\_\_\_. (Round off to the Nearest Integer).

### Official Ans. by NTA (2)

**Sol.** trans -  $CoCl_3.4NH_3$ 

ALLEN

8

or

trans- $[Co(NH_3)_4Cl_2]C\ell$ 





As we know that ethylene diamine is a bidentate ligand and ammonia is a mono dentate ligand.

It means overall two ethylene diamine is required to replace the all neutral ligands (four ammonia) from the coordination sphere of this complex.

A 6.50 molal solution of KOH (aq.) has a density of 1.89 g cm<sup>-3</sup>. The molarity of the solution is \_\_\_\_\_ mol dm<sup>-3</sup>. (Round off to the Nearest Integer).

[Atomic masses: K :39.0 u; O :16.0 u; H :1.0 u] Official Ans. by NTA (9)

Sol. 6.5 molal KOH = 1000gm solvent has  
6.5 moles KOH  
so wt of solute = 
$$6.5 \times 56$$
  
=  $364$  gm  
wt of solution =  $1000 + 364 = 1364$ 

Volume of solution = 
$$\frac{1364}{1.89}$$
m $\ell$   
Molarity =  $\frac{\text{mole of solute}}{V_{\text{solution}} \text{ in Litre}}$   
=  $\frac{6.5 \times 1.89 \times 1000}{1364}$   
= 9.00

When light of wavelength 248 nm falls on a metal of threshold energy 3.0 eV, the de-Broglie wavelength of emitted electrons is \_\_\_\_\_\_ Å. (Round off to the Nearest Integer).

[Use : 
$$\sqrt{3} = 1.73$$
, h = 6.63 × 10<sup>-34</sup> Js  
m<sub>e</sub> = 9.1 × 10<sup>-31</sup> kg ; c = 3.0 × 10<sup>8</sup> ms<sup>-1</sup> ;  
1eV = 1.6 × 10<sup>-19</sup>J]  
Official Ans. by NTA (9)

**Sol.** Energy incident = 
$$\frac{nc}{2}$$

$$= \frac{6.63 \times 10^{-34} \times 3.0 \times 10^8}{248 \times 10^{-9} \times 1.6 \times 10^{-19}} \text{ eV}$$
$$= \frac{6.63 \times 3 \times 100}{248 \times 1.6}$$
$$= 0.05 \text{ eV} \times 100 = 5 \text{ eV}$$
Now using  
E =  $\phi$  + K.E.  
5 = 3 + K.E.  
K.E. = 2eV =  $3.2 \times 10^{-19}$  J

for debroglie wavelength  $\lambda = \frac{h}{mv}$ 

K.E = 
$$\frac{1}{2}$$
mv<sup>2</sup>  
so  $v = \sqrt{\frac{2KE}{m}}$   
hence  $\lambda = \frac{h}{\sqrt{2KE \times m}}$   
=  $\frac{6.63 \times 10^{-34}}{\sqrt{2 \times 3.2 \times 10^{-19} \times 9.1 \times 10^{-31}}}$   
=  $\frac{6.63}{7.6} \times \frac{10^{-34}}{10^{-25}} = \frac{66.3 \times 10^{-10}}{7.6}$   
=  $8.72 \times 10^{-10}$  m  
 $\approx 9 \times 10^{-10}$  m  
=  $9$ Å

8. Two salts  $A_2X$  and MX have the same value of solubility product of  $4.0 \times 10^{-12}$ . The ratio of their molar solubilities i.e.  $\frac{S(A_2X)}{S(MX)} =$ \_\_\_\_\_.

(Round off to the Nearest Integer).

Official Ans. by NTA (50)

**Sol.** For  $A_2X$ 

$$A_{2}X \rightarrow 2A^{+} + X^{2-}$$

$$2S_{1} \qquad S_{1}$$

$$K_{sp} = 4S_{1}^{3} = 4 \times 10^{-12}$$

$$S_{1} = 10^{-4}$$
for MX
$$MX \rightarrow M^{+} + X^{-}$$

$$S_{2} \qquad S_{2}$$

$$K_{sp} = S_{2}^{2} = 4 \times 10^{-12}$$

$$S_{2} = 2 \times 10^{-6}$$
so 
$$\frac{S_{A_{2}X}}{S_{MX}} = \frac{10^{-4}}{2 \times 10^{-6}} = 50$$

9. A certain element crystallises in a bcc lattice of unit cell edge length 27 Å. If the same element under the same conditions crystallises in the fcc lattice, the edge length of the unit cell in Å will be \_\_\_\_\_. (Round off to the Nearest Integer).

[Assume each lattice point has a single atom]

[Assume  $\sqrt{3} = 1.73$ ,  $\sqrt{2} = 1.41$ ]

Official Ans. by NTA (33)

**Sol.** For BCC  $\sqrt{3} a = 4r$ 

so 
$$r = \frac{\sqrt{3}}{4} \times 27$$
  
for FCC  $a = 2\sqrt{2}r$   
 $= 2 \times \sqrt{2} \times \frac{\sqrt{3}}{4} \times 27$   
 $= \frac{\sqrt{3}}{\sqrt{2}} \times 27$   
 $= 33$ 

10. The decomposition of formic acid on gold surface follows first order kinetics. If the rate constant at 300 K is  $1.0 \times 10^{-3}$  s<sup>-1</sup> and the activation energy  $E_a = 11.488 \text{ kJ mol}^{-1}$ , the rate constant at 200 K is  $\_$  × 10<sup>-5</sup> s<sup>-1</sup>. (Round of to the Nearest Integer). (Given :  $R = 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$ ) Official Ans. by NTA (10) **Sol.**  $K_{300} = 10^{-4}$  $K_{200} = ?$  $E_a = 11.488 \text{ KJ/mole}$  R = 8.314 J/mole-K so  $ln\left(\frac{K_{300}}{K_{200}}\right) = \frac{E_a}{R}\left(\frac{1}{200} - \frac{1}{300}\right)$  $ln\left(\frac{K_{300}}{K_{200}}\right) = \frac{11.488 \times 1000 \times 100}{8.314 \times 200 \times 300}$ = 2.303 $= \ell n 10$ so  $\frac{K_{300}}{K_{200}} = 10$  $K_{200} = \frac{1}{10} \times K_{300} = 10^{-4}$ 

$$= 10 \times 10^{-5} \text{ sec}^{-1}$$