burn body.



(4) pig iron, scrap iron and steel

Official Ans. by NTA (2)

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Two compounds A and B with same molecular 6. The shape/structure of $[XeF_5]^-$ and XeO_3F_2 , formula (C_3H_6O) undergo Grignard's reaction respectively, are : with methylmagnesium bromide to give (1) pentagonal planar and trigonal bipyramidal products C and D. Products C and D show following chemical tests. Test С D Ceric Positive Positive Official Ans. by NTA (1) ammonium nitrate Test Lucas Test Turbidity Turbidity obtained after obtained five minutes immediately Iodoform Test Positive Negative C and D respectively are : XeF_5^- (1) $C=H_3C-C-OH$; sp³d³ Pentagonal planar D=H₃C-CH₂-CH-CH₃ I OH 7. major product The (2) $C = H_3C - CH_2 - CH_2 - CH_2 - OH$; is: $D = H_{3}C - C - OH$ Rr OH (3) $C = H_3C - CH_2 - CH_2 - CH_3$; СН₃ D=H₃C-С-ОН СН₃ (4) $C = H_3C - CH_2 - CH_2 - CH_2 - OH$; D=H₃C-CH₂-CH-CH₃ OH Official Ans. by NTA (3) Official Ans. by NTA (4) Sol. $\begin{array}{c} & & OH \\ H \\ CH_3 - CH_2 - C - H \xrightarrow{CH_3MgBr} CH_3 - CH_2 - CH - CH_3 \\ (A) & 2^{\circ} Alcohol \end{array}$ Sol. more acidic H (C) CAN test for alcohol : \checkmark Iodoform test : \checkmark Better leaving group $CH_{3}-C-CH_{3}\xrightarrow{CH_{3}MgBr}CH_{3}-C-CH_{3}\xrightarrow{CH_{3}MgBr}CH_{3}-C-CH_{3}\xrightarrow{CH_{3}MgBr}CH_{3}$ 3° Alcohol CAN test for alcohol : \checkmark

(2) trigonal bipyramidal and pentagonal planar (3) octahedral and square pyramidal (4) trigonal bipyramidal and trigonal bipyramidal Sol. $\begin{bmatrix} F & & F \\ F & & F \\ F & & F \end{bmatrix}^{\circ} & \begin{bmatrix} O & & O \\ F & & F \\ F & & O \end{bmatrix}^{\circ} = O$ XeO_3F_2 sp³d Trigonal bipyramidal obtained from E₂-elimination of 3-bromo-2-fluoropentane

(1)
$$CH_{3}CH_{2}$$
- CH - CH = CH_{2}
Br
(2) CH_{3} - CH_{2} - C = CH - CH_{3}
(3) CH_{3} - CH = CH - CH - $CH_{-}CH_{3}$
(4) $CH_{3}CH_{2}CH$ = C - F
 CH_{3}



Stable alkene having 5 αH

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Three elements X, Y and Z are in the 3rd period Strength of ligand $F^- < NCS^- < NH_3$ Sol. of the periodic table. The oxides of X, Y and $[MF_{6}]^{(-6+n)} [M(NCS)]^{(-6+n)} [M(NH_{3})_{6}]^{n+1}$ Z, respectively, are basic, amphoteric and acidic. The correct order of the atomic numbers (ii) of X, Y and Z is : (1) Z < Y < X(2) X < Z < Y(3) X < Y < Z(4) Y < X < ZAs given in graph : A < B < C (λ_{max}) Official Ans. by NTA (3) \therefore Correct matching is A-(iii), B-(i), C-(ii) Sol. When we are moving from left to right in a periodic 11. Consider the reaction sequence given below : table acidic character of oxides increases (as well as atomic number of atom increases) Br $\frac{OH^{e}}{H_{0}O}$ X < Y < Z*.*.. (acidic character) X < Y < Z(atomic number) 9. The number of subshells associated with n = 4and m = -2 quantum numbers is : (1) 4(2) 8(3) 16 (4) 2 Official Ans. by NTA (4) **Sol.** For n = 4 $\ell = 0, 1, 2, 3$ Which of the following statements is true : (1) Changing the concentration of base will m $-1 \ 0 \ +1$ -2 -1 0 +1 +2 -3 -2 -1 0 +1 +2 +3 have no effect on reaction (1) \therefore 4d & 4f subshell associated with n = 4, m = -2 (2) Changing the concentration of base will Simplified absorption spectra of three complexes 10. have no effect on reaction (2) $((i), (ii) \text{ and } (iii)) \text{ of } M^{n+} \text{ ion are provided below;}$ (3) Changing the base from OH^{\ominus} to $^{\Theta}OR$ will their λ_{max} values are marked as A, B and C have no effect on reaction (2) respectively. The correct match between the (4) Doubling the concentration of base will complexes and their λ_{max} values is : double the rate of both the reactions. Official Ans. by NTA (1) Sol. Reaction $1 : SN_1$ Reaction $2 : E_2$ Absorption SN₁ is independent of concentration of nucleophile/base The results given in the below table were obtained 12. during kinetic studies of the following reaction: λ_{max} λ_{max} λ_{max}

Wavelength (nm)

(1) A-(ii), B-(i), C-(iii) (2) A-(iii), B-(i), C-(ii)

(3) A-(ii), B-(iii), C-(i) (4) A-(i), B-(ii), C-(iii)

(ii) $[MF_6]^{(-6+n)}$

(i) $[M(NCS)_6]^{(-6+n)}$

Official Ans. by NTA (2)

(iii) $[M(NH_3)_6]^{n+1}$

 $2A + B \longrightarrow C + D$

Initial Experiment [A]/molL⁻¹ $[B]/molL^{-1}$ rate/molL⁻¹ \min^{-1} 6.00×10^{-3} Ι 0.1 0.1 Π 0.1 0.2 2.40×10^{-2} 1.20×10^{-2} III 0.2 0.1 7.20×10^{-2} IV Х 0.2 V 2.88×10^{-1} 0.3 Y

(iii)

 \rightarrow OH + Br^{Θ}(1)

 $+ HOH + Br^{\Theta} \dots (2)$

rate = k[t-BuBr] $[OH^{\Theta}]$

rate = k[t-BuBr]

 $\rightarrow \Delta_{Oh}$ 1

(i)

CH₃

CH.

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X and Y in the given table are respectively : (1) 0.3, 0.4(2) 0.4, 0.3(3) 0.4, 0.4(4) 0.3, 0.3Official Ans. by NTA (1) **Sol.** From rate law $r = -\frac{1}{2}\frac{d[A]}{dt} = \frac{-d[B]}{dt}$ $= K[A]^{x} [B]^{y}$ $6 \times 10^{-3} = K(0.1)^x (0.1)^y$(1) $2.4 \times 10^{-2} = K(0.1)^{x} (0.2)^{y} \dots (2)$ $1.2 \times 10^{-2} = K(0.2)^{x} (0.1)^{y} \dots (3)$ $(3) \div (1) \implies x = 1$ $(2) \div (3) \implies x = 2$ So, order with respect to A = 1Order with respect to B = 2 $(4) \div (3)$ $\left(\frac{x}{0.2}\right) \times \left(\frac{0.2}{0.1}\right)^2 = \frac{7.2 \times 10^{-2}}{1.2 \times 10^{-2}}$ $x = \frac{6 \times 0.2}{4}$ x = 0.3 M $(5) \div (4)$ $\left(\frac{y}{0.2}\right)^2 = \frac{2.88 \times 10^{-1}}{7.2 \times 10^{-2}}$ $y^2 = 4 \times 0.2^2$ y = 0.4 M13. An organic compound 'A' $(C_0H_{10}O)$ when treated with conc. HI undergoes cleavage to yield compounds 'B' and 'C'. 'B' gives yellow precipitate with AgNO₃ where as 'C' tautomerizes to 'D'. 'D' gives positive idoform

(1)
$$\bigcirc$$
 -O-CH=CH-CH₃
(2) \bigcirc -CH₂-O-CH=CH₂
(3) \bigcirc -O-CH₂-CH=CH₂
(4) H₃C \bigcirc -O-CH=CH₂

test. 'A' could be :



Sol. Both Li and Mg form nitride when reacts directly with nitrogen.

> The hydrogen carbonate of both Li and Mg does not exist in solid state.

> All alkali metal hydrogen carbonate exist in solid state except LiHCO₃.

16. The one that is not expected to show isomerism Match the type of interaction in Column A with 18. the distance dependence of their interaction is : energy in Column B: (1) $[Ni(NH_3)_4(H_2O)_2]^{2+}$ (2) $[Ni(NH_3)_2Cl_2]$ A B (4) $[Ni(en)_3]^{2+}$ (3) $[Pt(NH_3)_2Cl_2]$ Official Ans. by NTA (2) (a) $\frac{1}{r}$ (I) ion - ion Sol. $[Ni(NH_3)_2Cl_2]$ is tetrahedral complex, therefore does not show geometrical and optical isomerism. (b) $\frac{1}{r^2}$ (II) dipole - dipole [Ni(NH₃)₂Cl₂] does not show structural isomerism (c) $\frac{1}{r^3}$ (III) London dispersion $[Ni(NH_3)_4(H_2O)_2]^{2+}$ & $[Pt(NH_3)_2Cl_2]$ show geometrical isomerism (d) $\frac{1}{r^6}$ $[Ni(en)_3]^{2+}$ show optical isomerism Amongst the following statements regarding 17. (1) (I)-(a), (II)-(b), (III)-(c) adsorption, those that are valid are : (2) (I)-(a), (II)-(c), (III)-(d) (a) ΔH becomes less negative as adsorption (3) (I)-(a), (II)-(b), (III)-(d) proceeds. (4) (I)-(b), (II)-(d), (III)-(c) (b) On a given adsorbent, ammonia is adsorbed Official Ans. by NTA (3) more than nitrogen gas. Official Ans. by (2)(c) On adsorption, the residual force acting along **Sol.** Type of interaction Interaction Energy(E) the surface of the adsorbent increases. $E \propto \frac{1}{r}$ (d) With increase in temperature, the equilibrium ion - ion concentration of adsorbate increases. (1) (b) and (c) (2) (a) and (b) $E \propto \frac{1}{r^3}$ dipole - dipole (4) (c) and (d) (3) (d) and (a) Official Ans. by NTA (2) $E \propto \frac{1}{r^6}$ London dispersion **Sol.**(a) Since adsorption is exothermic process, as adsorption proceeds number of active sites 19. The correct observation in the following present over adsorbent decreases, so less heat reactions is : is evolved. Sucrose $\xrightarrow{Glycosidic bond} A + B \xrightarrow{Seliwanoff's} ?$ (b) Since NH₃ has higher force of attraction on adsorbent due to its polar nature (high value of (Hydrolysis) 'a'). (1) Formation of blue colour (c) As the adsorption increases, residual forces (2) Formation of violet colour over surface decreases. (3) Formation of red colour (d) Since process is exothermic, on increasing temperature it shift to backward direction, so (4) Gives no colour concentration of adsorbate particle decreases.



Sol. Seliwanoff's test is used to distinguish between aldose and ketose sugars; when added to a solution containing ketose, red colour is formed rapidly.

Sucrose $\xrightarrow{\text{Hydrolysis}}$ Glucose $\xrightarrow{\text{Seliwanoff's}}$ Red + $\xrightarrow{\text{reagent}}$ Red Fructose

- **20.** The molecular geometry of SF_6 is octahedral. What is the geometry of SF_4 (including lone pair(s) of electrons, if any) ?
 - (1) Trigonal bipyramidal
 - (2) Square planar
 - (3) Tetrahedral
 - (4) Pyramidal

Official Ans. by NTA (1)



 4σ bonds +1 lone pair

∴ Shape (including lone pair of electrons) is Trigonal bipyramidal

21. The heat of combustion of ethanol into carbon dioxide and water is -327 kcal at constant pressure. The heat evolved (in cal) at constant volume and 27°C (if all gases behave ideally) is (R = 2 cal mol⁻¹ K⁻¹)

Official Ans. by NTA (-326400.00)

Official Ans. by (326400.00)
Sol.
$$C_2H_5OH_{(\ell)} + 3O_{2(g)} \longrightarrow 2CO_{2(g)} + 3H_2O_{(\ell)}$$

 $\Delta n_g = 2 - 3 = -1$
 $\Delta_c H = \Delta_c U + (\Delta n_g) RT$
 $\Delta_c H = \Delta_c U - RT$
 $\Delta_c U = \Delta_c H + RT$
 $= -327 \times 10^3 + 2 \times 300$
 $= -326400$ cal.
 \therefore Heat evolved

= 326400 cal.

For the disproportionation reaction 22. $2Cu^+$ (aq) $\implies Cu(s) + Cu^{2+}(aq)$ at 298 K, ln K (where K is the equilibrium constant) is _____ × 10⁻¹. Given $(E_{Cu^{2+}/Cu^{+}}^{0} = 0.16V$ $E^0_{Cu^+/Cu} = 0.52V$ $\frac{\mathrm{RT}}{\mathrm{F}} = 0.025)$ Official Ans. by NTA (144.00) Sol. $Cu^+ \longrightarrow Cu^{2+} + e^ Cu^+ + e^- \longrightarrow Cu(s)$ $2Cu^+ \longrightarrow Cu^{2+} + Cu$ $E_{cell}^{o} = E_{Cu^{+}/Cu}^{o} - E_{Cu^{2+}/Cu^{+}}^{o}$ = 0.52 - 0.16= 0.36 VAt equilibrium $\rightarrow E_{cell} = 0$

$$E_{cell}^{o} = \frac{RT}{nF} ln K$$

$$ln K = \frac{E_{cell}^{o} \times nF}{RT}$$

$$ln K = \frac{0.36 \times 1}{0.025}$$

$$= 14.4 = 144 \times 10^{-1}$$

- 23. The oxidation states of transition metal atoms in K₂Cr₂O₇, KMnO₄ and K₂FeO₄, respectively, are x, y and z. The sum of x, y and z is _____.
 Official Ans. by NTA (19.00)
- Sol. $K_2Cr_2O_7$ 2 (+1) + 2x + 7(-2) = 0 x = +6In $K_2Cr_2O_7$, Transition metal (Cr) present in +6 oxidation state. KMnO₄ (+1) + y + 4(-2) = 0x = +7

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> In $KMnO_4$, transition metal (Mn) present in +7 25. oxidation state K₂FeO₄ 2(+1) + z + 4(-2) = 0x = +6In K_2 FeO₄, transition metal (Fe) present in +6 oxidation state So, x = +6y = +7z = +6x + y + z = 1924. The ratio of the mass percentages of 'C & H' and 'C & O' of a saturated acyclic organic compound 'X' are 4 : 1 and 3 : 4 respectively. Then, the moles of oxygen gas required for complete combustion of two moles of organic compound 'X' is _ Official Ans. by NTA (5.00) **Sol.** C : H = 4 : 1C : O = 3 : 4Mass ratio C : H : O = 12 : 3 : 16Mole ratio C : H : O = 1 : 3 : 1Empirical formula = CH_3O Molecular formula = $C_2H_6O_2$ (saturated acyclic organic compound) $C_2H_6O_2 + \frac{5}{2}O_2 \longrightarrow 2CO_2 + 3H_2O$

> > 2 mole 5 mol Moles of O_2 required = 5 moles

25. The work function of sodium metal is 4.41 × 10⁻¹⁹ J. If the photons of wavelength 300 nm are incident on the metal, the kinetic energy of the ejected electrons will be $\frac{10^{-21} \text{ J.}}{(\text{h} = 6.63 \times 10^{-34} \text{ Js}; \text{ c} = 3 \times 10^8 \text{ m/s})}$ Official Ans. by NTA (222.00) Sol. E = W + K·E_{max} K·E_{max} = E - W $= \frac{\text{hc}}{\lambda} - 4.41 \times 10^{-19}$ $= \frac{6.63 \times 10^{-34} \times 3 \times 10^8}{300 \times 10^{-9}} - 4.41 \times 10^{-19}$ $= 2.22 \times 10^{-19} \text{ J}$ $= 222 \times 10^{-21} \text{ J}$