

$$\Delta G = \Delta H - T \cdot \Delta S$$

if $\Delta H < 0$, $\Delta S = +ive$.

ΔG will be $-ive$ NO matter what the temp is.

- (53) X will be major if $\Delta G = +ive$ or > 0
 Y will be major if $\Delta G = -ive$ or < 0
 X & Y will be in equi. if $\Delta G = 0$
 Hit & trial

putting $T = 315K$ to $\Delta G = 120 - \frac{3}{8}T$
 $\Delta G = +ive$ or > 0 \therefore X is major.

- (54) at $100^\circ C$ & $1 atm$ $H_2O(l) \rightarrow H_2O(g)$
 $\therefore \Delta S_{system} > 0$ but surrounding $\Delta S = -\frac{q}{T}$
 $\Delta S_{surrounding} < 0$

- (55) for zero order $t_{1/2} = \frac{a}{2K}$
 $6 = \frac{12}{2K} \Rightarrow K = \frac{1}{60}$

Now $A_t = A_0 - Kt$
 $A_0 - A_t = Kt$
 $15 - 12 = \frac{1}{60} \times t$ $t = 18 \times 60 = 18h$.

~~$t = \frac{12}{2 \times 60} = 10$~~
 $\frac{600}{200}$

(56) $r = K[A]^n$ — (1)

$r8 = K[2A]^n$ — (2)

dividing (2) by (1)

$$8 = 2^n = 2^3$$

$$n = 3$$

(57) ratio expression is always defined for 1 mole

$$\therefore r = -\frac{dN_2}{dt} = -\frac{1}{3} \frac{dN_2}{dt} = +\frac{1}{2} \frac{dNN_2}{dt}$$

(58) by electrophoresis colloids can be purified & precipitated.

(59) b

(60) $Ti < Mn < Ni < Zn$. I.E $\propto Z^*$ (effective nuclear charge)

(61) In dil HNO_3 oxidation state of 'N' will be less and in conc. HNO_3 O.S of 'N' will be greater
e.o.d

(62) German silver is 56% Cu, Zn 24%, Ni 20%

(63) In transition elements O.S is less

\therefore given ligand must have lesser no. of -ive sites and in inner transition elements O.S is greater \therefore ligands must have more -ive sites \therefore C.N must be 6 & 8.

(64) given complex in H_2O gives ce^- ion and complex in option b gives NO_2^- ion.

(65) CO is a strong ligand $\therefore Cr(d^5s^1)$ will become $Cr(d^6s^0)$

1	1	1	1	1	1	1
---	---	---	---	---	---	---

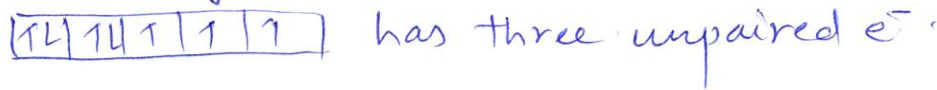
(66) O.S of Fe in $[Fe(H_2O)_5NO]SO_4$

$$x + 5 \times 0 + 1 + (-2) = 0$$

$$x = +1$$



∴ In the given complex $Fe^+(d^7)$

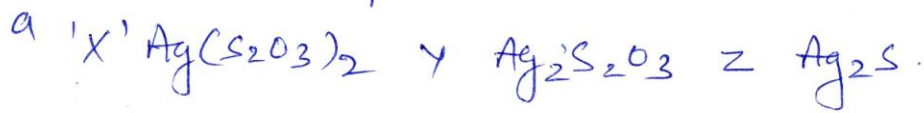


(67) Oxidation of Al is more spontaneous above $800^\circ C$ and oxidation of Zn is spontaneous below $800^\circ C$

∴ Al can reduce ZnO above $800^\circ C$

(68) Slag is always silicate or phosphate

(69) $S_2O_3^{2-}$ is fluxidant (monodentate or bidentate)



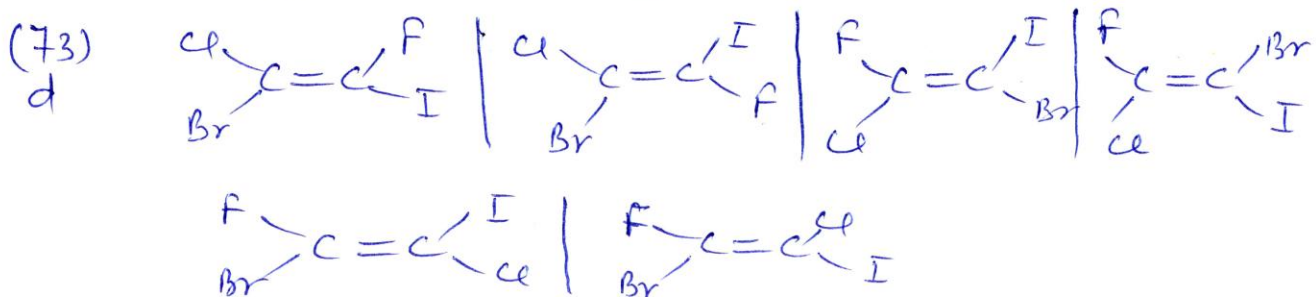
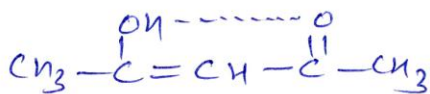
(70) c Variation in oxidation state is measured by magnetic moment

(71) Sulphate is estimated as $BaSO_4$ (ppt)

b Ionic radii of $Mg^{2+} < Ba^{2+}$ but this is not the correct explanation.

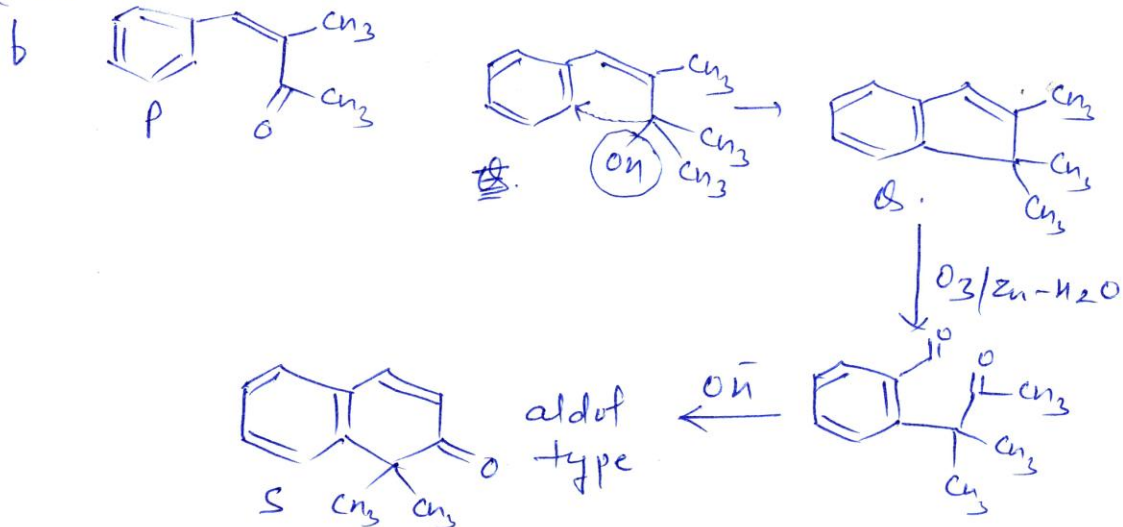
(72) maximum enol content, stronger acidic H

b due to more withdrawing nature of ketone and chelation



(74) a

(75)



(76) all have CHO group with α -H

(77) all are stronger acid than H_2CO_3 except phenol

(78) $SOCl_2$ changes $-OH$ into $-Cl$

(79) order of acidic strength \propto E.W strength $\propto \frac{1}{E.R.}$

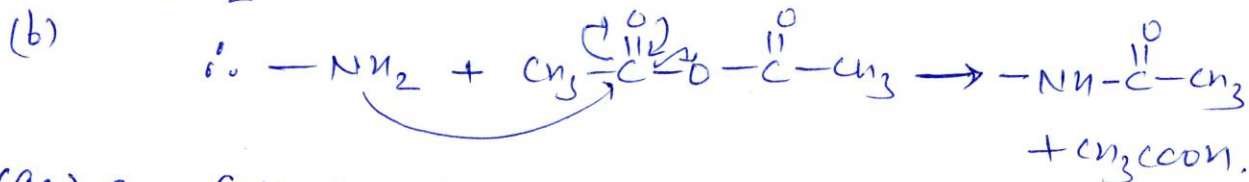
$NO_2 > Cl$



(80)

a

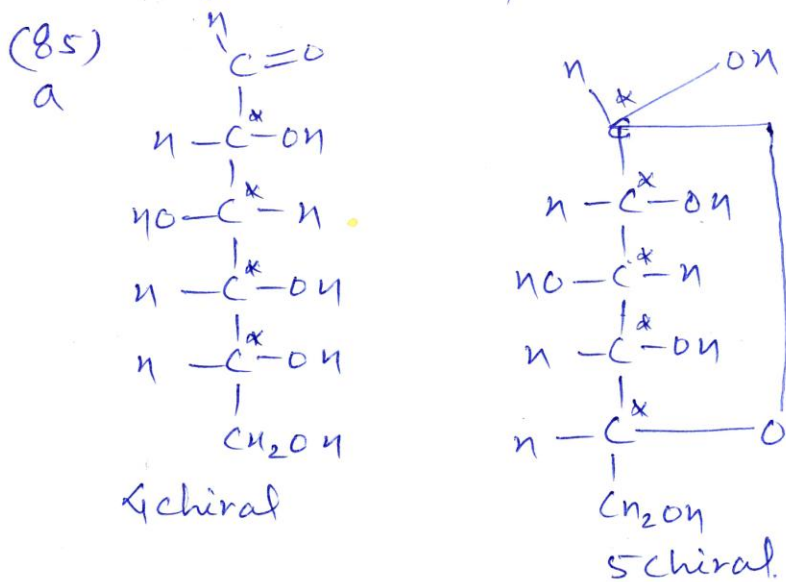
(81) $-NH_2$ is more donor



(82) c $C_6H_5SO_2Cl$

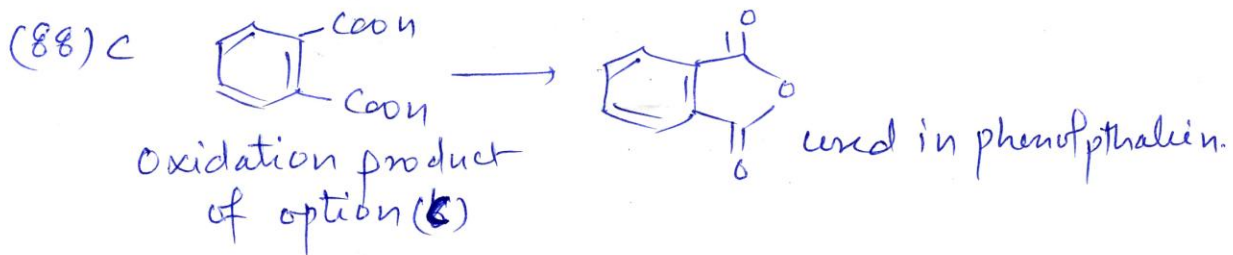
(83) c for SN_1 stability of carbocation & steric hindrance.

(84) b hydride transfer is slowest step.



(86) d

(87) b



(89) DiBAL-H is weak reducing agent and source of H^-

