

Liquid Nitrogen Tetroxide: N_2O_4 WAHID U. MALIK
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- * Powerful oxidising agent.
- * It can oxidise reducing agent substances explosively particularly of organic nature.
- * It has low dielectric constant.
- * Poor solvent for ionic substances, but good " " organic compounds.
- * The auto ionisation of N_2O_4 is



nitrosyl nitrate ion

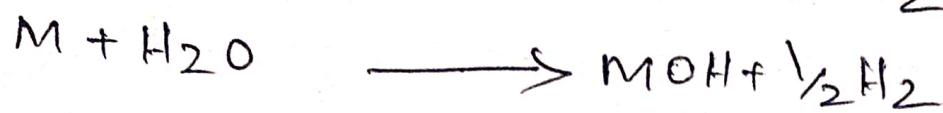
chemical rxns:(1) Acid-Base rxns:

- * formation NO^+ from auto-ionisation would behave as acids in liq. N_2O_4

- * formation of NO_3^- from auto-ionisation would behave as bases in N_2O_4

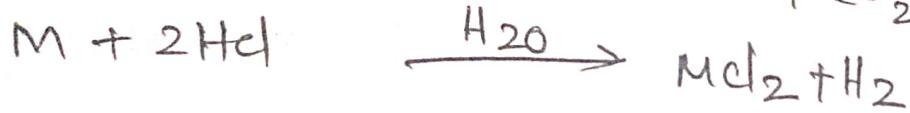
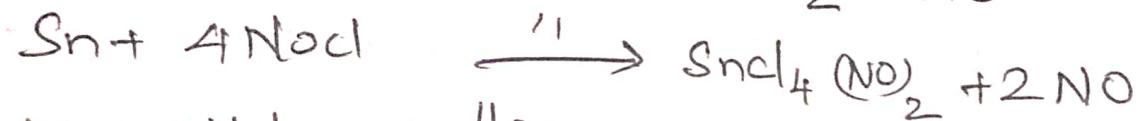
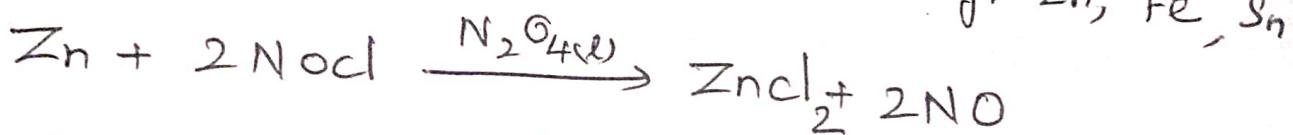
(2) Reaction with metal:

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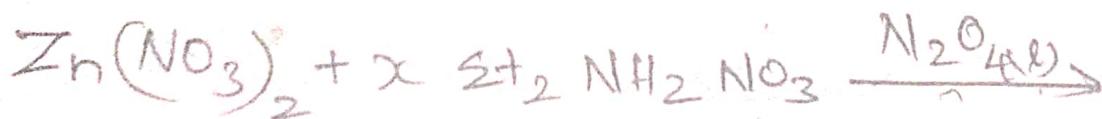
This rxn is manner as water.

- * Addition of HCl to H₂O increases the reactivity of H₂O towards metals.
- * addition of NaCl to N₂O₄ also increases its reactivity toward metals. e.g. Zn, Fe, Sn



(3) Complex formation:

- * Zn(NO₃)₂ dissolves readily in a solution of diethyl amm. nitrate Et₂NH₂NO₃ in liq. N₂O₄ to yield mta Zin Complex.



Solid



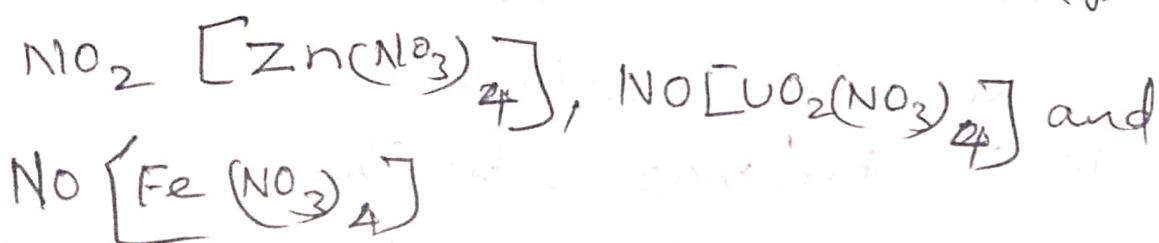
diethyl ammonium nitroato zincate

* above oxns indicates amphoteric oxns.

A) Solvate formation oxns:

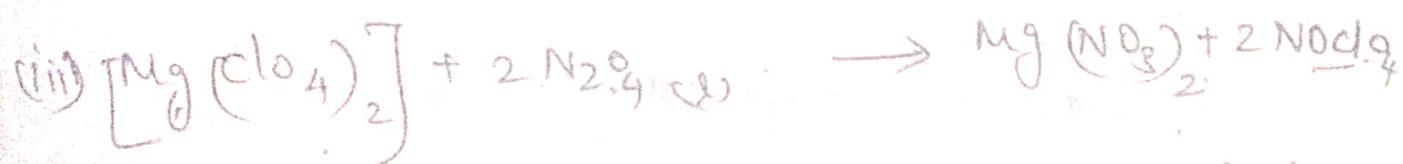
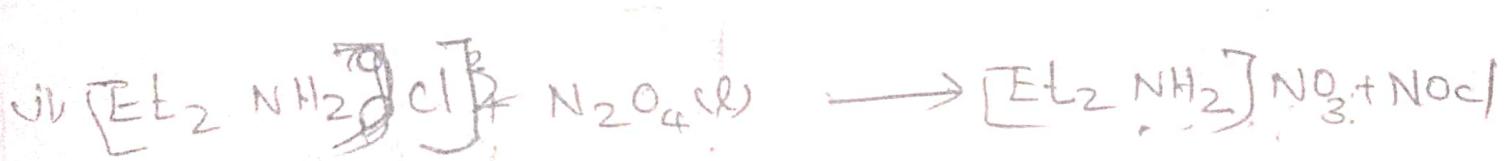
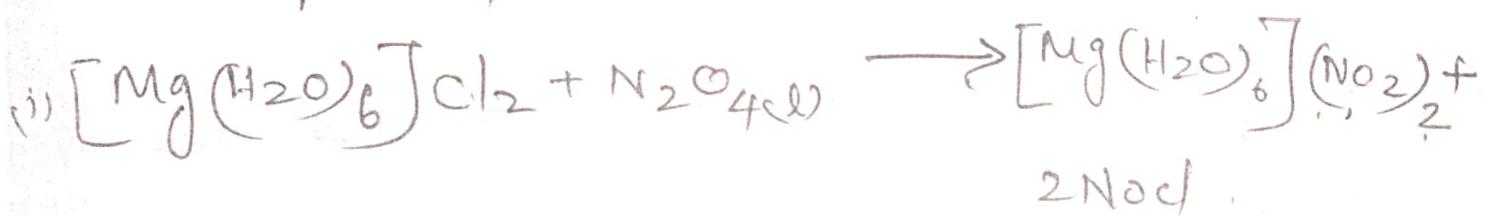
* nitrate of some metals such as Zn, U and Fe react with N_2O_4 to form the solvates. e.g $Zn(NO_3)_2 \cdot 2N_2O_4$, $UO_2(NO_3)_2 \cdot N_2O_4$ and $Fe(NO_3)_3 \cdot N_2O_4$ respectively.

* These are complex salts. e.g.

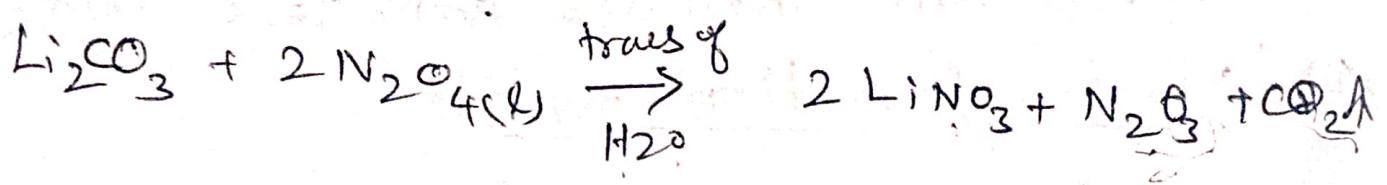


B) Solvolyisis oxns:

* no. of solvolysis rxns have been studied in liq. N_2O_4



* $LiCO_3$ gets solvated in presence of traces of H_2O .



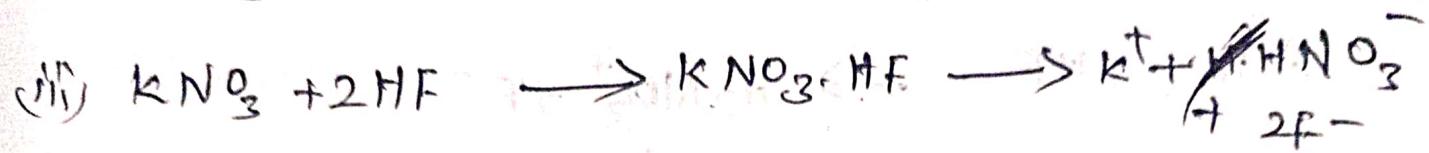
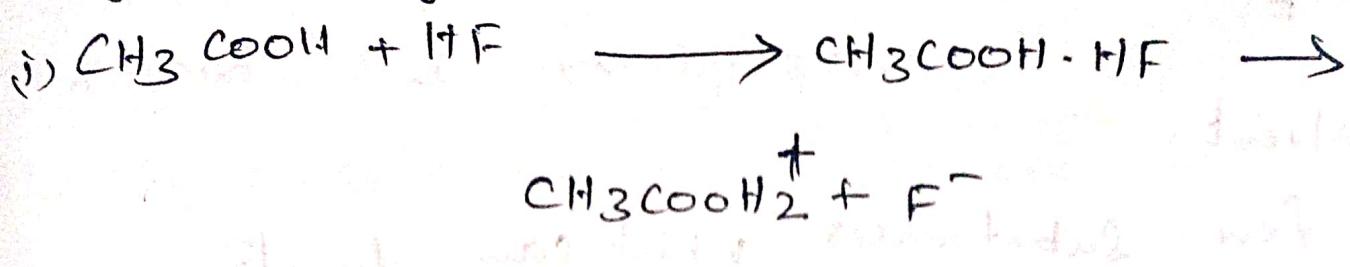
Liquid HF : (243)

- * liq. HF has a wide but convenient liquid range (-83°C to 19.4°C)
- * soluble few substances without chemical rxns and it has poisonous character.
- * High dielectric constant
- * excellent ionising solvents.
- * It dissolves many organic & inorganic compds to give highly conducting solution.
- Inorganic compds are more soluble than organic compds.
- * HF dissociates into ions ~~and~~ viz H⁺ & F⁻



Such solutions behaves as acidic solutions.

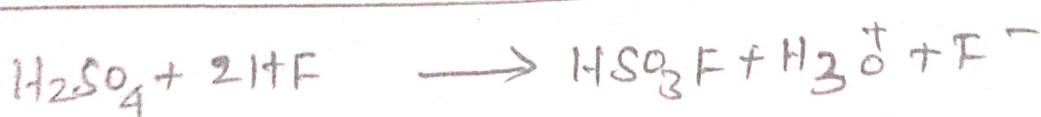
followed by dissociation to give F⁻ ion.



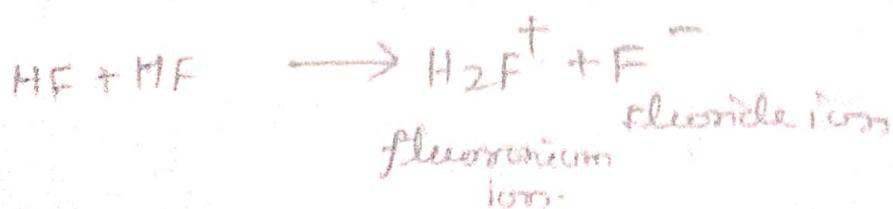
* The anion of the solute may be replaced by F⁻ ion.



* Chemical reactions involving more than simple replacement may take place



* Auto ionisation

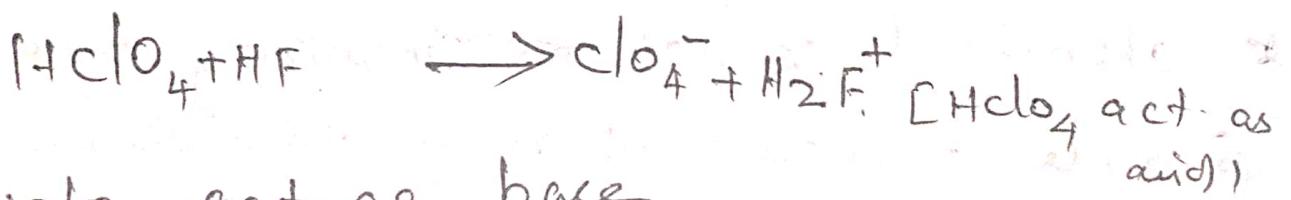
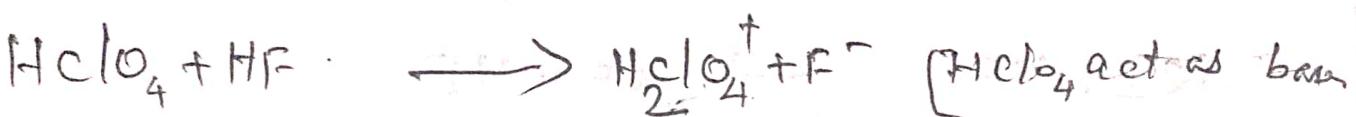


* H_2F^+ ions will behave as an acid
 F^- ions will behave as a base in solvent.

* few substances which can donate protons readily to HF.



* HClO_4 strongest acid in aq. sol. & is amphoteric in HF.

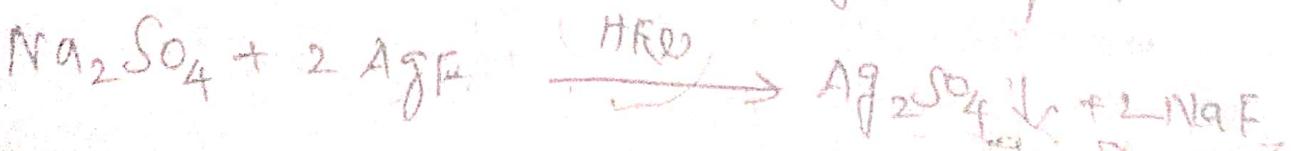


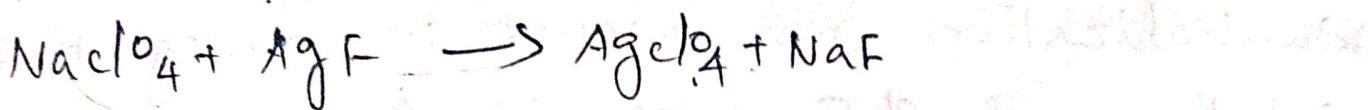
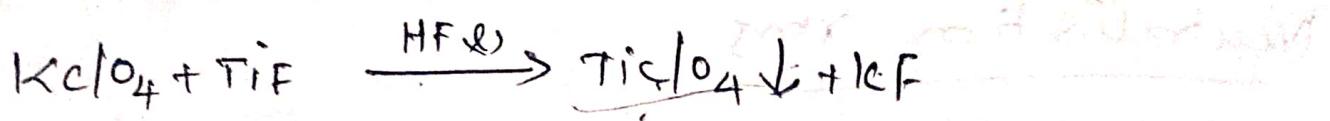
* LiClO_4 acts as base

* Compounds acting as acids in liq. HF



* Precipitation reactions occurring in liq. HF





* The precipitation can are



* HF also form a no. of addition compounds with metallic fluoride, eg. $KF \cdot HF$, $KF \cdot 2HF$, $KF \cdot 3HF$, $NH_4F \cdot HF$ etc.

Acoti Liquid bromine Trifluoride

* $liq \cdot BrF_3$ indicates ioniser.



* SbF_5 act as an acid in $liqBrF_3$



SbF_5 act as an acid in BrF_3 .

* KF act as an base in HF



Neutralisation rxns

* Neutralisation rxns occurs on mixing the solutions of SbF_5 + KF in liq. BrF_3 .



* Neutralisation rxns help us to synthesise many compounds.

