

Extraction of Metal from the concentrated ore  
Metal is extracted from the concentrated ore by the following steps

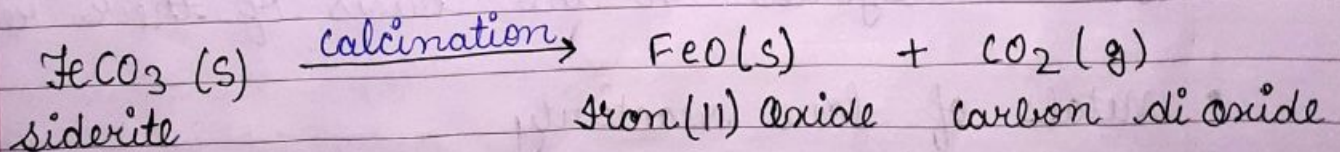
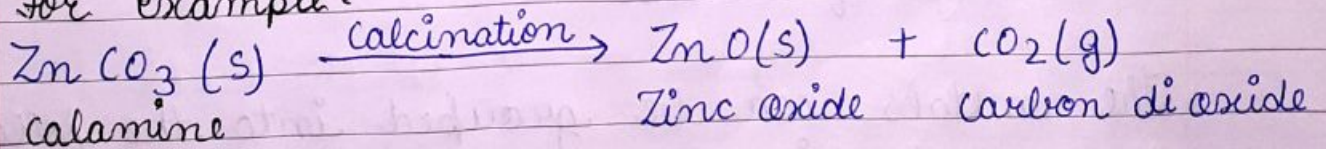
a) Conversion of the concentrated ore into its oxide :  
the production of metal from the concentrated ore mainly involves reduction process. This can be usually done by 2 processes known as calcination and roasting. The method depends upon the nature of the ore

(b) conversion of oxide to metal by reduction process

(a) Conversion of ore into Metal oxide :  
these are briefly discussed below :

- i) Calcination : it is the process of heating the concentrated ore in the absence of air  
the calcination process is used for the following changes:
  - to convert carbonate ores into metal oxide..
  - to remove water from the hydrated ores
  - to remove volatile impurities from the ore.

For example:

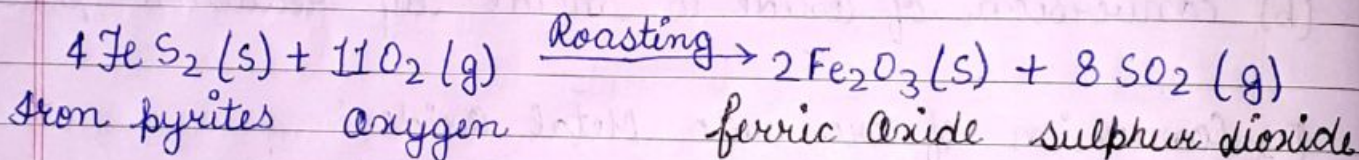
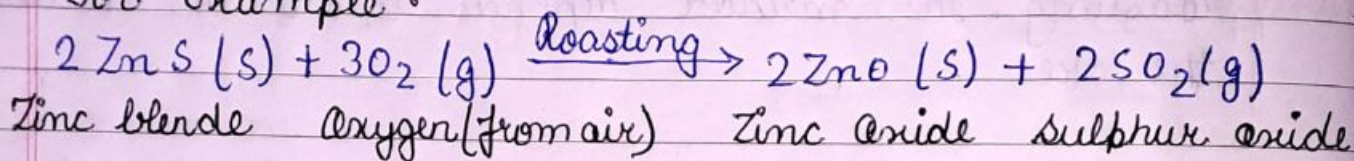


ii) Roasting : It is the process of heating the concentrated ore strongly in the presence of excess air.

This process is used for converting sulphide ores to metal oxide. In this process, the following changes take place

- the sulphide ores undergo oxidation to their oxides
- moisture is removed
- volatile impurities are removed

For example :



Note: calcination is used for hydrated and carbonate ores and roasting is used for sulphide ore.

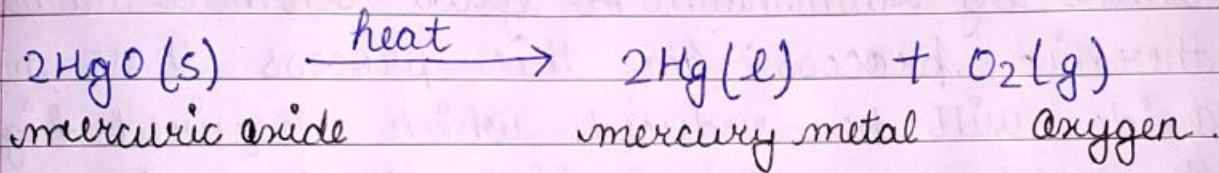
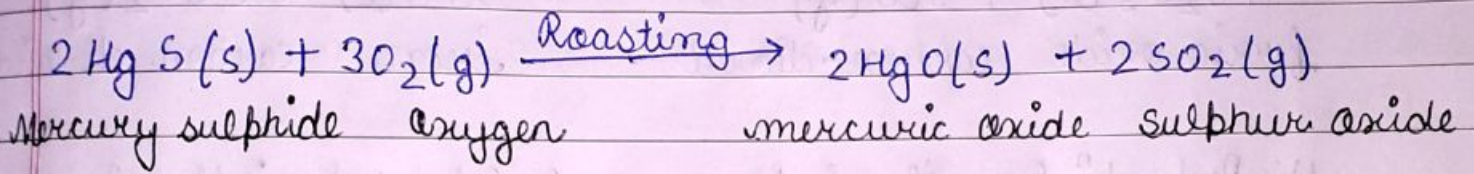
- b) conservation of metal oxide into Metal :
- the metal oxide formed after calcination or roasting is converted into metal by reduction. The method used for reduction of metal oxides depends upon the nature and chemical reactivity of metal.

The metals can be grouped into the following three categories on the basis of their reactivity.

- Metals of low reactivity
- Metals of medium reactivity
- Metals of high reactivity

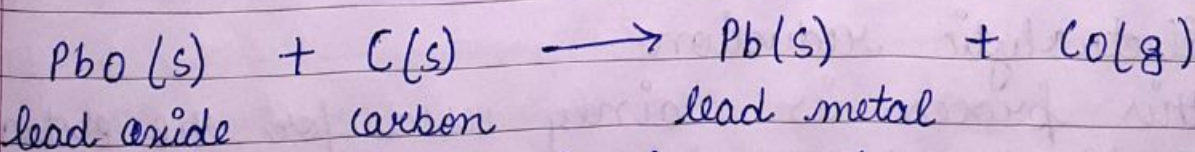
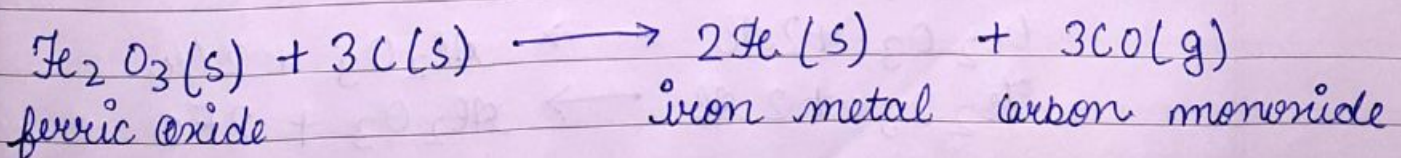
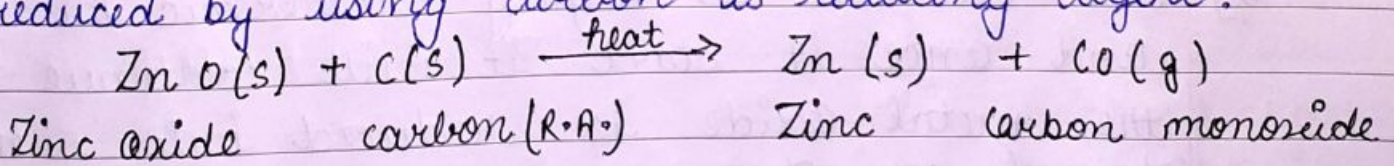
these different categories of metals are extracted by different techniques. The different steps involved in separation are as follows:

i) Reduction by heating: Metals placed low in the reactivity series are very less reactive. They can be obtained from their oxides by simply heating in air.



ii) chemical Reduction (for metals in the middle of the reactivity series):

A) Reduction with carbon: the oxides of moderately reactive metals (occurring in the middle of reactivity series) like zinc, copper, nickel, tin, lead etc. can be reduced by using carbon as reducing agent.

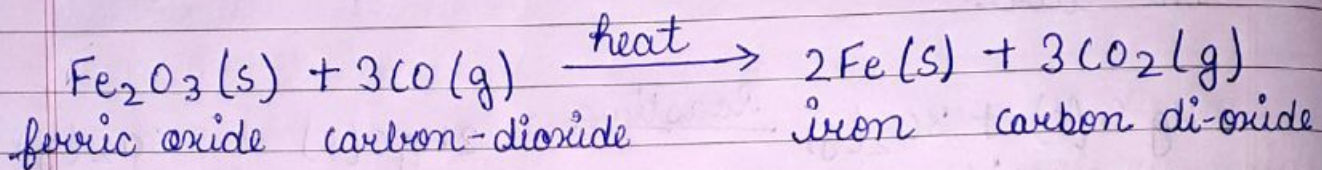


One disadvantage of using carbon as reducing agent is that small traces of carbon are added to metal as impurity. Therefore, it contaminates the metals

\* Note:

coke is very commonly used as reducing agent because it is cheap

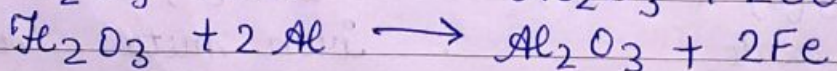
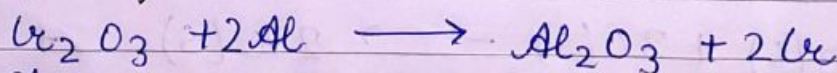
B) Reduction with carbon monoxide: Metals can be obtained from oxides by reduction with carbon monoxide in the furnace



b) Reduct<sup>n</sup> by Aluminium or Gold schmidt aluminothermic process: In this process those metal oxide will be reduced which required high temperature and at high temp carbon react with metal to form metal carbide. In this process we use aluminium as a reducing agent due to

i) Al has greater affinity towards oxygen as it forms most stable oxide ( $\text{Al}_2\text{O}_3$ )

ii) This reaction is highly exothermic in nature and once it start it will continue till at the metal oxide is reduced into metal  
For Cr, Mn, Fe



iv) Electrolytic reduction

this process is mainly used for the extraction of highly electropositive metals. IA, IIA, & Al  
Electrolysis is carried out in a large cell and a small amount of another suitable

electrolyte is added which :

- lowers the melting point of the main electrolyte
- enhances its conductivity
- reduces corrosion troubles

eg:- manufacture of metallic sodium (Down's process)

Molten NaCl containing a little  $\text{CaCl}_2$  is electrolysed between graphite anode and iron cathode. The various reactions that take place are:

On fusion:  $\text{NaCl} \rightleftharpoons \text{Na}^+ + \text{Cl}^-$  (Ions become mobile)

On electrolysis: At cathode:  $\text{Na}^+ + e^- \rightarrow \text{Na}(\text{red}^n)$

At anode:  $2\text{Cl}^- \rightarrow \text{Cl}_2(\text{g}) + 2e^-$

Hall heroult process:

This process is used for extraction of Al from alumina. The extraction of Al from  $\text{Al}_2\text{O}_3$  is quite difficult because

- Fusion temp. of Alumina is quite high ( $2050^\circ\text{C}$ ) even more than boiling point of Al ( $1150^\circ\text{C}$ )
- It is bad conductor of electricity. To overcome these difficulties we mix some amount of neutral flux [ $\text{Na}_3\text{AlF}_6 + \text{CaF}_2$ ]. Neutral flux provides free ion to the sol<sup>n</sup> which decreases the fusion temp. of Alumina from  $2050^\circ\text{C}$  to  $950^\circ\text{C}$ .

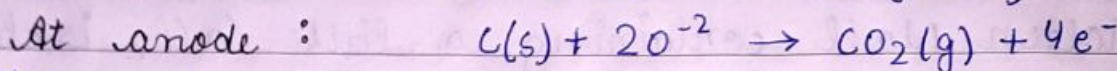
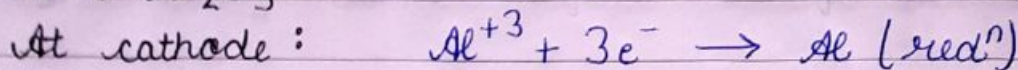
i) From cryolite  $\rightarrow \text{AlF}_3 \rightarrow \text{Al}^{+3} + 3\text{F}^-$   
At cathode:  $\text{Al}^{+3} + 3e^- \rightarrow \text{Al}(\text{red}^n)$   
At anode:  $2\text{F}^- - 3e^- \rightarrow \frac{3}{2}\text{F}_2$

$\text{F}_2$  reacts with  $\text{Al}_2\text{O}_3$   $3\text{F}_2 + \text{Al}_2\text{O}_3 \rightarrow 2\text{AlF}_3 + \frac{3}{2}\text{O}_2$

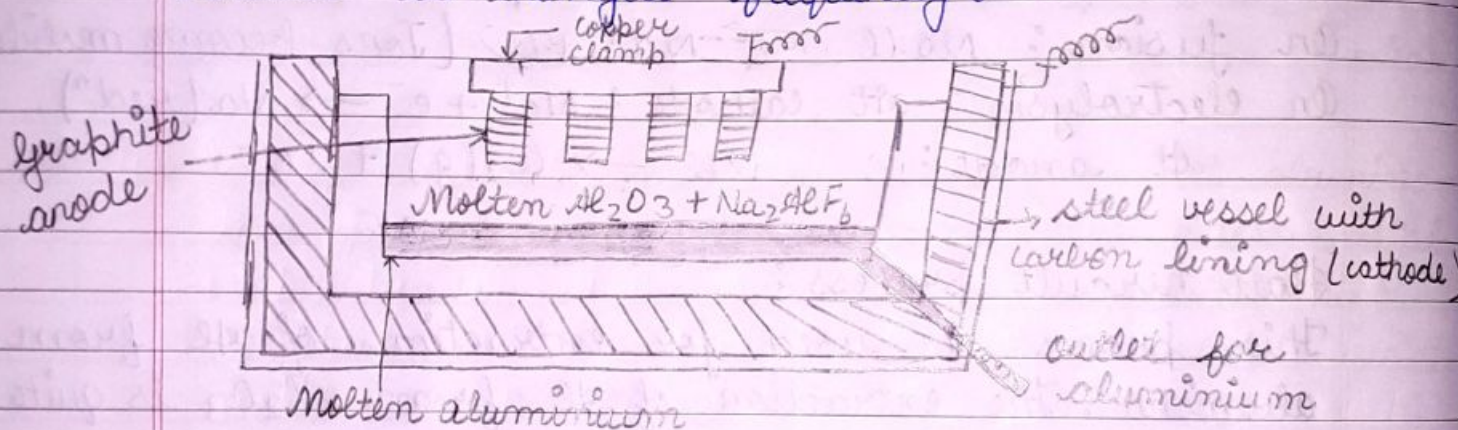
so at anode  $O_2$  gas is liberated which graphite anode convert into  $CO_2$  + destroy the anode



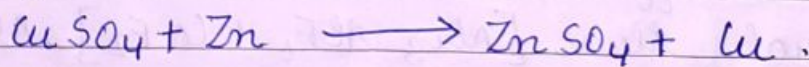
ii) From  $Al_2O_3$



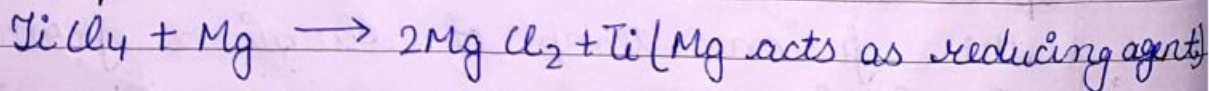
The main drawback of this process is that anode should be changed frequently.



(III) Metal displacement method: In this method, compound is reacted with a more electropositive & more reactive metal which displaces, the metal from the sol<sup>n</sup>.



Kroll's process



### REFINING OF METALS

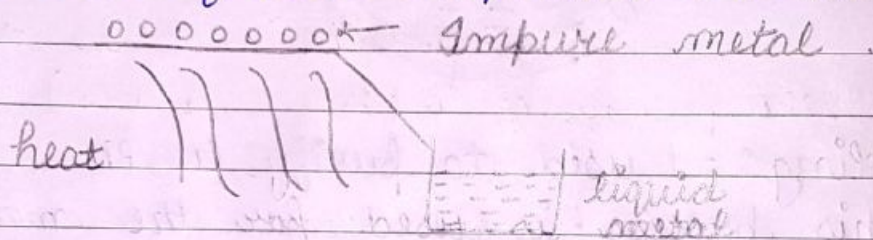
Metals obtained by the reduction of its compound still contains some objectionable substance and have to be refined. Depending upon the nature of the metal and impurities, the following methods are used for purification of the metals.

## (I) Physical Process

i) **Liquation**: this method is used for the refining of metals having low melting point and are associated with high melting impurities

Ex: Pb, Sn, Sb and Bi

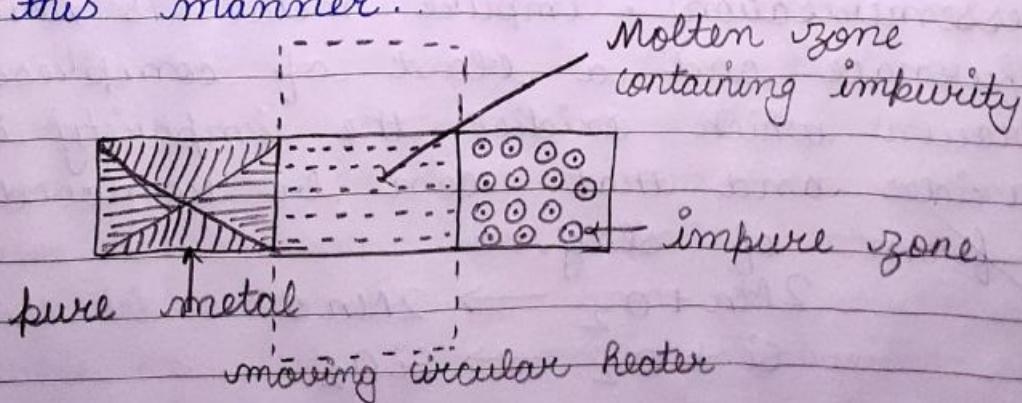
the impure metal is heated on the sloping hearth of a furnace. The pure metal flows down leaving behind the non-fusible impurities on the hearth.



ii) **Distillation**: Metals having low boiling point are refined by this method, for example, Zinc, cadmium and mercury

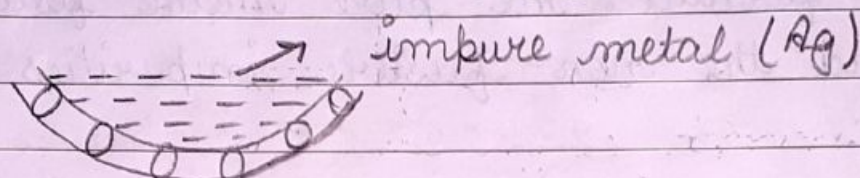
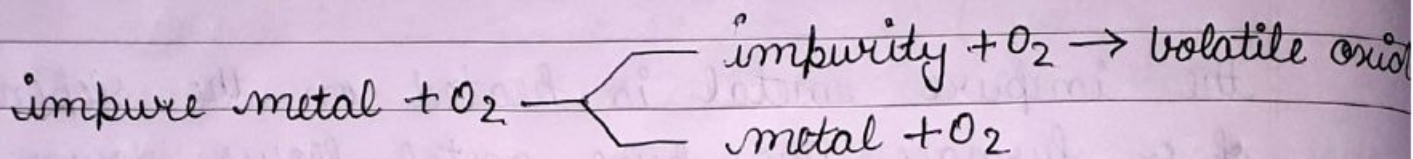
iii) **Zone refining**: Metals of very high purity are obtained by zone refining this refining method is based on the fact that impurities tend to remain dissolved in molten metal.

Ge, Si, and Ga used as semiconductors are refined in this manner.



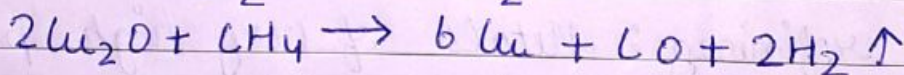
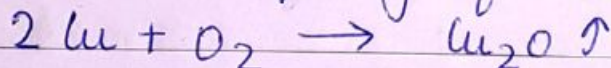
chemical process :-

Cupellation: this process is used to purify silver containing the impurities of Pb. This process is used when impurity have greater affinity towards  $O_2$  while metal does not have.

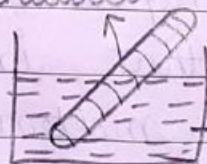


Poling: - used to purify Cu, Pb

this process is used for the metal having the impurity of their own oxide. In this process a green wooden pole is heated with molten metal, which provide C and H to metal oxide which reduce impurity of metal oxide to metal

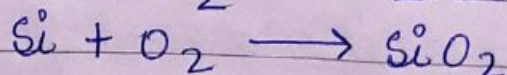
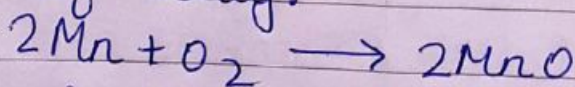


greenwood

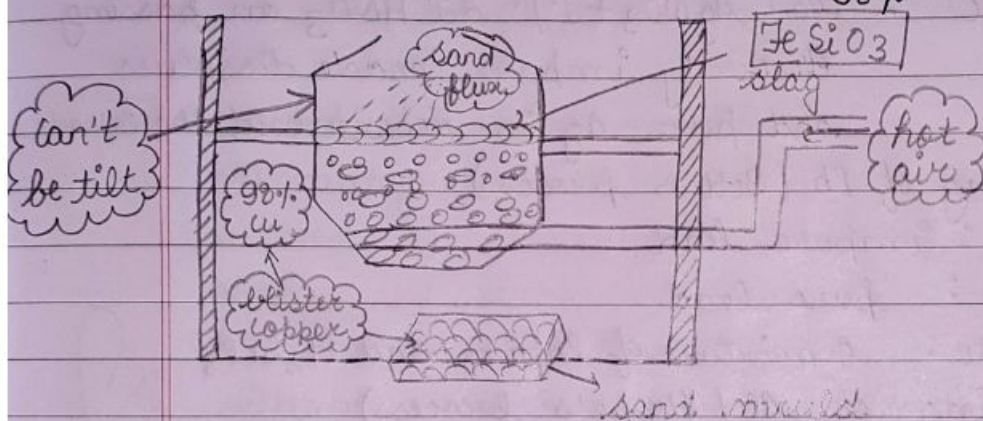
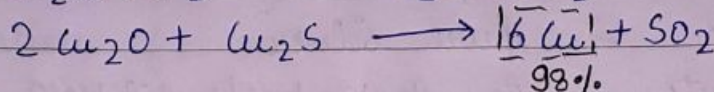
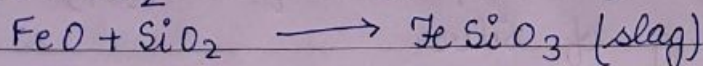
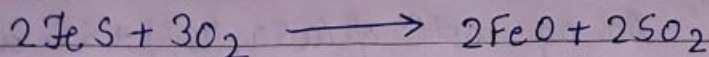


impure metal

Bessemerisation: impure metal is heated in a furnace and a blast of compressed air is blown which oxidise the impurity into their oxides and that can be removed in the form of slag.



**Bassmerization** :- the process of further heating of molten matte in the presence of flux is called



Bassimer

↓  
Shape

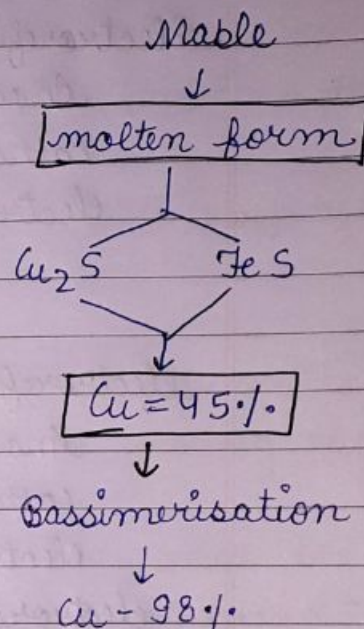
↓  
pear like

$\text{Cu}_2\text{S} \rightarrow$  cuprous sulphide

$\text{FeS} \rightarrow$  ferrous sulphide

$\text{FeO} \rightarrow$  " oxide

$\text{Cu}_2\text{O} \rightarrow$  cuprous oxide



### (III) Electro refining of metals:

metals such as Cu, Ag, Zn, Sn, Pb, Al, Ni, Cr are refined by this method

The impure metal is made the anode of an electrolytic cell, while cathode is thin plate of pure metal. Electrolyte is the sol<sup>n</sup> of a salt of the metal.

On passing the electric current pure metal from the anode dissolves and gets deposited at the cathode.

The soluble impurities go into the solution while insoluble or less electropositive impurities settle down below the anode as anode mud or sludge.

For example :-

## → Electrorefining of copper

Anode : blister copper (98%)

Cathode : pure copper

Electrolyte : An aq. sol<sup>n</sup> of  $\text{CuSO}_4$  (15%) + 5% dil  $\text{H}_2\text{SO}_4$

## Electrorefining of silver

Anode : impure silver

Cathode : pure silver

Electrolyte : Aq.  $\text{AgNO}_3$  + 1% dil  $\text{HNO}_3$  on passing electricity impure anode dissolves and pure Ag is deposited at the cathode.

## Electrorefining of Pb (Bett's process)

Anode : impure lead

Cathode : pure lead

Electrolyte : A mixture of  $\text{PbSiF}_6$  and  $\text{H}_2\text{SiF}_6$

## Electrorefining of Al (Hoopes's process)

Anode : impure Al

Cathode : pure Al

Electrolyte : a mixture of  $\text{Na}_3\text{AlF}_6$  +  $\text{CaF}_2$

## (IV) Vapour phase refining (thermal decomposition)

### i) Van-Arkel process :

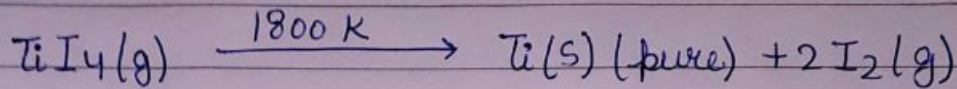
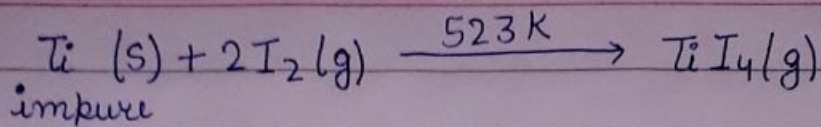
This method is very useful for removing all the oxygen & nitrogen present in the form of impurity

i) employed to get metal in very pure form of small quantities

ii) In this method, the metal is converted into a volatile unstable compound (eg. iodide) and impurities are not affected during compound formation.

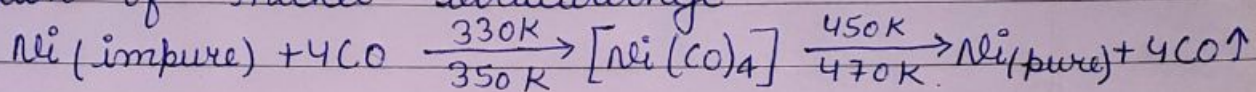
iii) The compound thus obtained is decomposed to get the pure metal.

iv) Employed for purification of metals like titanium and zirconium



ii) Mond's process :-

i) Nickel is purified by using CO gas. This involves the formation of nickel tetracarbonyl



iii) chromatographic methods :-

chromatography is the most modern and versatile method of separation, purification and testing the purity of elements and their compounds.

The method is based upon the principle that the different components of a mixture are absorbed to different extents of an adsorbent.