

## LECTURE 27 (16/08/2023)

### 125] Steel Making

became  
totally  
obsolete  
75 years ago  
(but it's  
imp. bcz  
it is the  
mother of  
all other  
processes)

1. Bessemer Process → Acid Bessemer  
Basic Bessemer
2. Open Hearth Process → Acid O. H.  
Basic O. H.
3. Oxygen Steel making
  - (a) Top Blown Process → LD, LD-AC
  - (b) Bottom Blown Process → Q-BOP, OBM
  - (c) Combined Blowing
  - (d) Kaldo Process
  - (e) Rotor Process
4. Electric Steel Process
 

have  
become  
obsolete

↓  
electric  
furnace,  
induction furnace

{ Basic oxygen furnace }  
or even more  
commonly as  
BOF ↑  
Now more  
commonly known  
as BOP  
↑ (Basic oxygen  
process)  
(Linz-Donawitz)

{ "quiet" }  
(Oxygen Bottom  
Maxhuette)

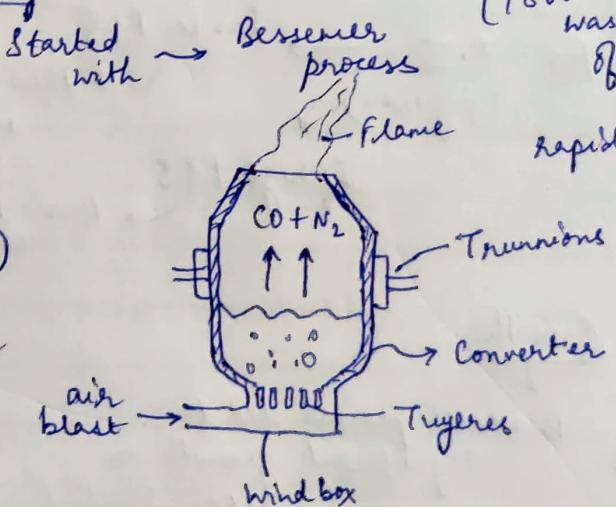
name of  
the German  
company  
that developed it

### World Steel Production

BOF - 55%      Electric Steel - 45%

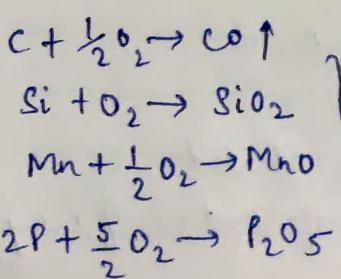
### 126] Changing Pattern of Steel Making

#### Growth & Decay:



(blowing time is ~25 mins.)  
this entire refining  
process takes place  
↑ in 1 hour

(1860-1910  
was 2<sup>nd</sup> phase  
of Industrialization)  
↓  
rapid growth occurred  
during this period



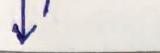
\* (when room temp. air is passed through molten metal ore ↓ instead: the impurities in it get oxidized which is an exothermic process )

it does NOT freeze the molten metal

exothermic

127] When Bessemer

Came up with his process



He used  $\text{SiO}_2$  as flux

( $\because$  it came from Sweden)

& he wasn't aware of other kinds of flux)

this is not that good as it can't remove P from ore

lead to Acid slag & acid lining } }

that is why it was called

### Acid Bessemer Process

Later on a scientist

S. G. Thomas (in 1878) came up with

$\text{CaO}$  as the basic flux to be used in this process

Can remove Phosphorus  
(as:  $\text{CaO} + \text{P}_2\text{O}_5$ )

\* leading to Basic Bessemer process

(basic slag & basic lining)

NOTE: For P. I. → has low P&S  
(pig iron) → thus Acid Bessemer is okay to use

If high P&S

→ Basic Bessemer used

The Basic lining consists of:

$\text{MgO}$  or  
 $\text{MgO}, \text{CaO}$   
(Dolomite)

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### Growth

(i) Bessemer process led to significant growth in

converter production &

steel production

(ii) But after about 1910

decay of converters started

(i.e. generation of scrap)

However this couldn't be  
recycled ( bcz Bessemer converters  
couldn't generate enough  
heat )  
it only had enough heat  
to melt original metal  
& the cold scrap couldn't  
be melted

(iii) No composition control (leading to bad quality of steel)

(iv) High nitrogen content

dissolved from air

about 0.018%

which was enough to produce Brittleness in steel

( Thus it couldn't be used for wire drawing )

{ LECTURE 28 attached  
on next page }

{ From Manish Verma's Notes }

Lecture 27

Charging pattern of Steel Making process →

Boilerie process → Growth and decay.

Disadvantages →

- (i) Scrap could not be processed from cold.
- (ii) Control of composition of steel was poor due to process being very fast.
- (iii) High nitrogen content - 0.013% N making it stiffer and harder to process.
- (iv) Some heat was lost in the flame / Burning of CO outside the converter.

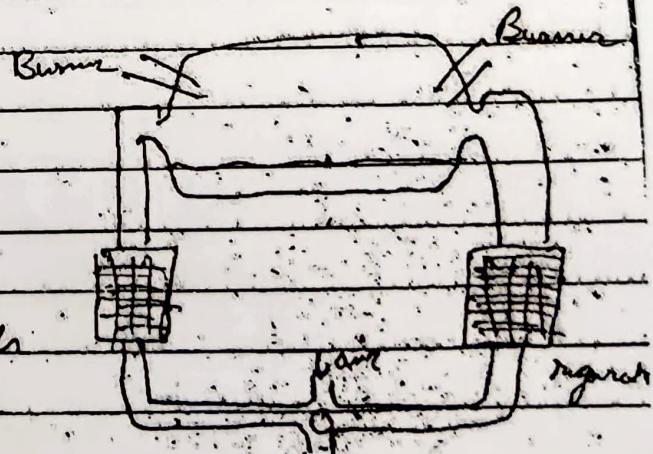
Despite it all, it remained for 100 years (1860-1960)

(utilised in 1861 by Martin brothers and Siemens Brothers).

→ Open Hearth Process → In 1857 John Stearling invented recuperative system. That means the heat going out ~~was~~ in puddling or glass making furnace, ~~was~~ ~~the~~ was not utilised. The maximum temperature that could be attained was 1400°C and the slag and metal became

a paste due to metal solidifying which was separated by heating as much as possible and wrought iron was produced. The outgoing gases were allowed to pass through the chickwork and the bricks are heated.

At one time, brick of one side is used while the other side heats from the exhaust. Then the process is reversed. This allowed the temperature to reach 1600°C. Now, it was possible to melt the metal and slag.



Initially its use was to melt the steel scrap.  
Later on the pig iron refining could also be done.

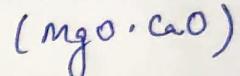
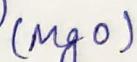
[Written from  
Manish  
Verma's  
Notes]

Similar to other methods

→ Acid open hearth  
→ Basic open hearth

Initially it was Acidic, but later modified to Basic by K.G. Thomas in 1884. Removal of phosphorous and buffer by using Calcium oxide was possible.

Basic refractories were used; — Magnesite and Dolomite



Magnesite  $\Rightarrow$   $MgCO_3$  is calcinated at  $1000^{\circ}C$  to produce  $MgO$ , but  $MgO$  is not Magnesite still it is called this; why?

\* Magnesite is a misnomer  
(misnomer: "false name")

It is called open hearth because the hearth is wide and shallow compared to converter with long pear shape.

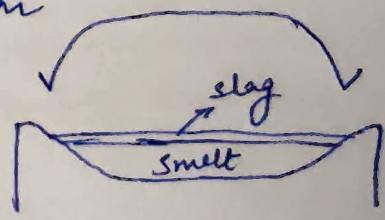
It does not depend on the oxidation of impurities to produce heat. Therefore it can take any type of charge:

- (i) 100% scrap
- (ii) 50% scrap + 50% P.I.
- (iii) 100% P.I.

Here the heating is done immediately.

A layer of slag was allowed to form and the oxygen has to ...

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on next  
page}



diffuse through the core layer of slag. Due to slow rate of oxidation, the process is controllable. It takes 6-10 hours to complete one cycle and produces one batch of steel.

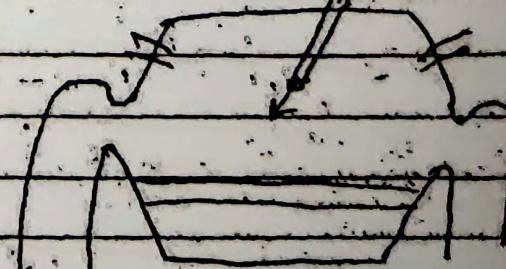
\* Advantages →

- 1) tiny amount of scrap can be added
- 2) control of composition. A very good quality of steel could be made.
- 3) Here, the nitrogen content is confined to 0.003%. So the steel is very ductile.

\* Drawbacks (Limitations) →

- 1) It has a high capital investment.
- 2) Low productivity due to the process being slow and produces very small quantity.
- 3) High fuel consumption.
- 4) Long time is required (6-10 hrs) to do one batch.

This technology's limitations were reduced by using oxygen lance which reduces the impurity even further and oxygen in the air was combined and used to burn the fuel.

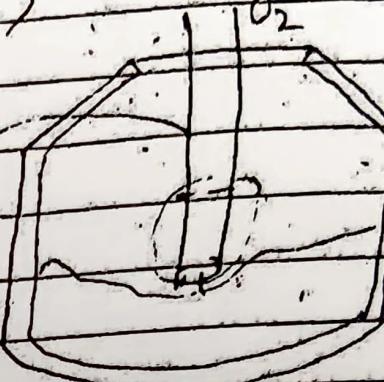


Later open hearth became obsolete and oxygen & steel making furnaces were adopted.

\* Oxygen Steel Making → It started in 1950's

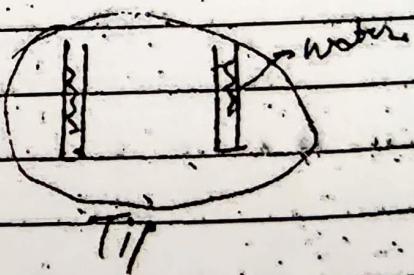
Oxygen before 1942 could only be produced in labs and after 1942, oxygen could be produced in commercial amount. Oxygen was introduced in steel making in 1951. The first experiment was done in Europe in Austria in Linz' and Donawitz (L D process).

Solid bottom like Bessemer converter, there are just two holes.



In Bessemer, the oxygen was used instead of air, oxygen would have melted due to heat and it was not available commercially. So people realising this started using oxygen through water cooling lance. The tip was made of copper which was double layered so water could be passed.

The oxygen was injected at jet speed to make the oxygen go super.



1959, 1954 Kaldo Rotor