

for sand casting: $P + F = C + 1$

only temp. vary

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YEAR- 1ST YEAR

CONSTITUTION OF ALLOYS

- An alloy can be defined as a substance having metallic properties, composed of two or more no. of elements, one of which atleast one is a metal.
- A phase is defined as microscopically distinct, chemically homogeneous and mechanically separable part of a system.
- No. of phases in an alloy are found from Gibbs phase rule
$$P + F = C + 2$$
where,
 P = No. of phases
 F = Degree of freedom
 C = No. of elements participating in alloy formation
 2 = stands for any two variable out of 3 namely Temperature, Pressure, concentration.

✓ BINARY ALLOYS (Made of two elements)

HOMOGENEOUS alloys

(Single phase)

→ Hume Rothery's rule

cannot be
expressed by
a formula
because solute
concentration
in solid solⁿ
does not remain
constant.

Solid solution

Compounds

Substitutional

Interstitial

Intermediate

Interstitial

Electron

Compounds

Compounds

Compounds

Mg_2Pb , Mg_3Sn

Gold, Ag, Fe, Ni

+
{ Cd, Mg, Sn }

solubility in FCC is more than BCC because
void space is greater in FCC than BCC
(void length)

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Poly phase alloys (Mixtures)

Combination of
solid solutions

Combination of
compounds

Combination of
solid solⁿ + compounds

HUME ROTHERY RULE : (for solid solⁿ)

- 1) Crystal structure factor :- crystal structure of both the elements should be same.
- 2) Relative size factor :- The diff. in atomic radii should be less than 15 %
- 3) Chemical Affinity factor :- Smaller the chemical affinity b/w the two elements, greater will be the solubility.
- 4) Relative valency factor :- Elements of lower valency can dissolve a large amount of higher valency elements of

➤ Intermetallic compounds are formed b/w chemically dissimilar metal.

➤ Interstitial compounds are formed b/w transition metal like tungsten, iron, titanium etc. with oxygen, N₂, C, B, H₂ etc.

➤ Electron compounds are formed b/w a group of alloys in whose case the ratio of valance e⁻ to no. of atoms participating in bond formation remain constant.

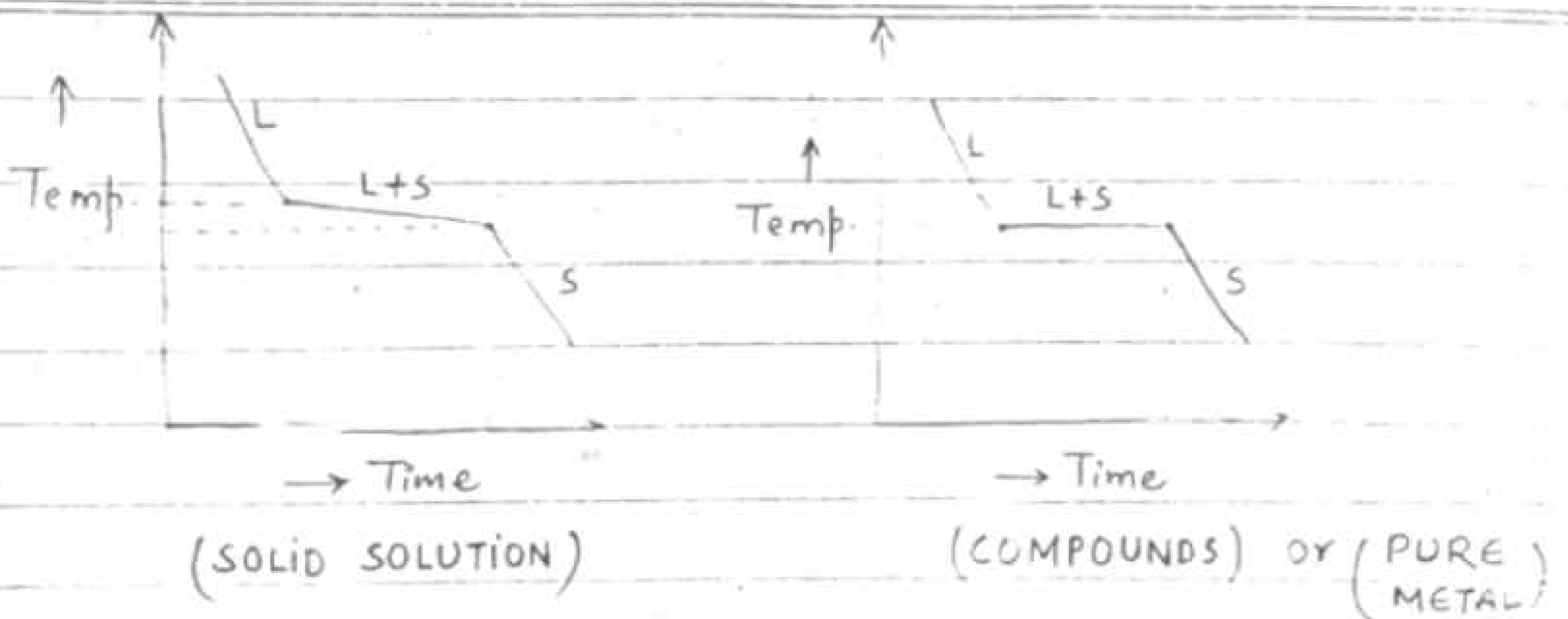
Strength $\rightarrow$ microstructure (grain structure)

Ductility / Brittleness $\rightarrow$ crystal structure

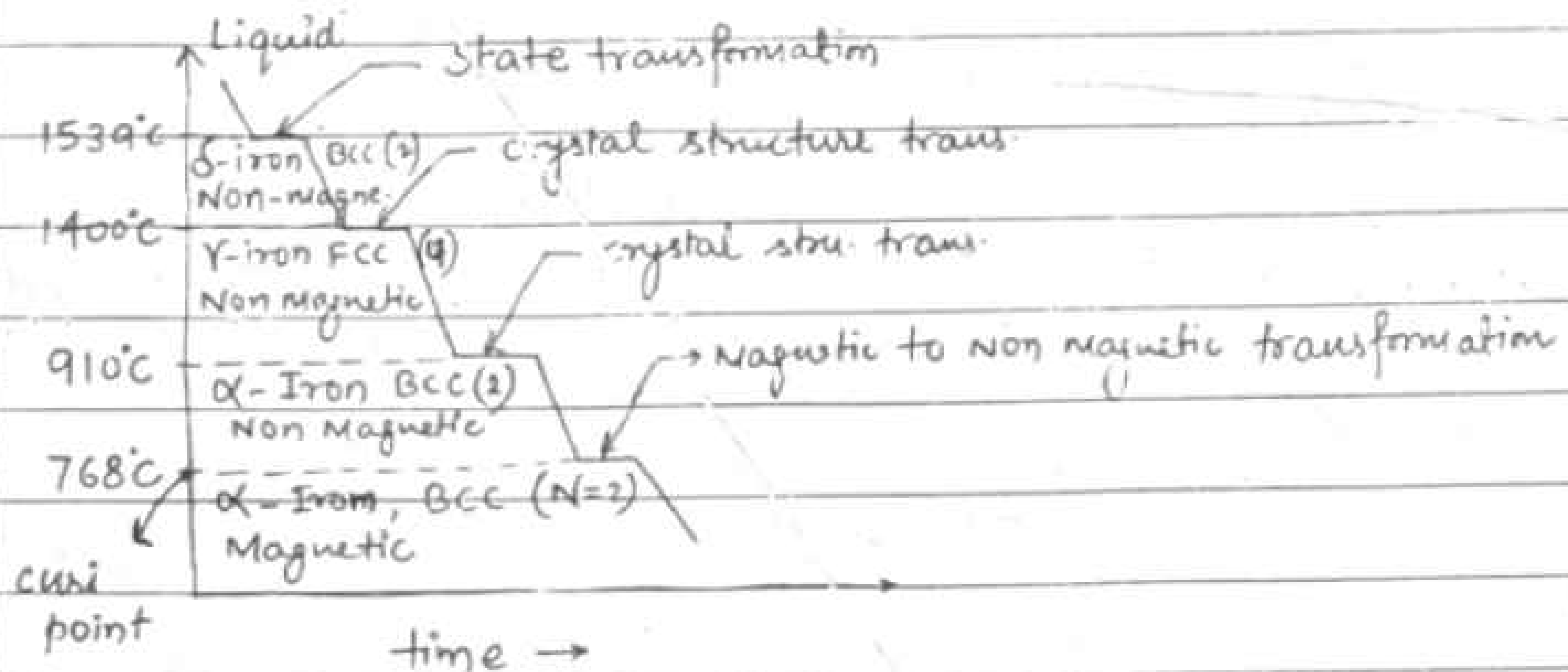
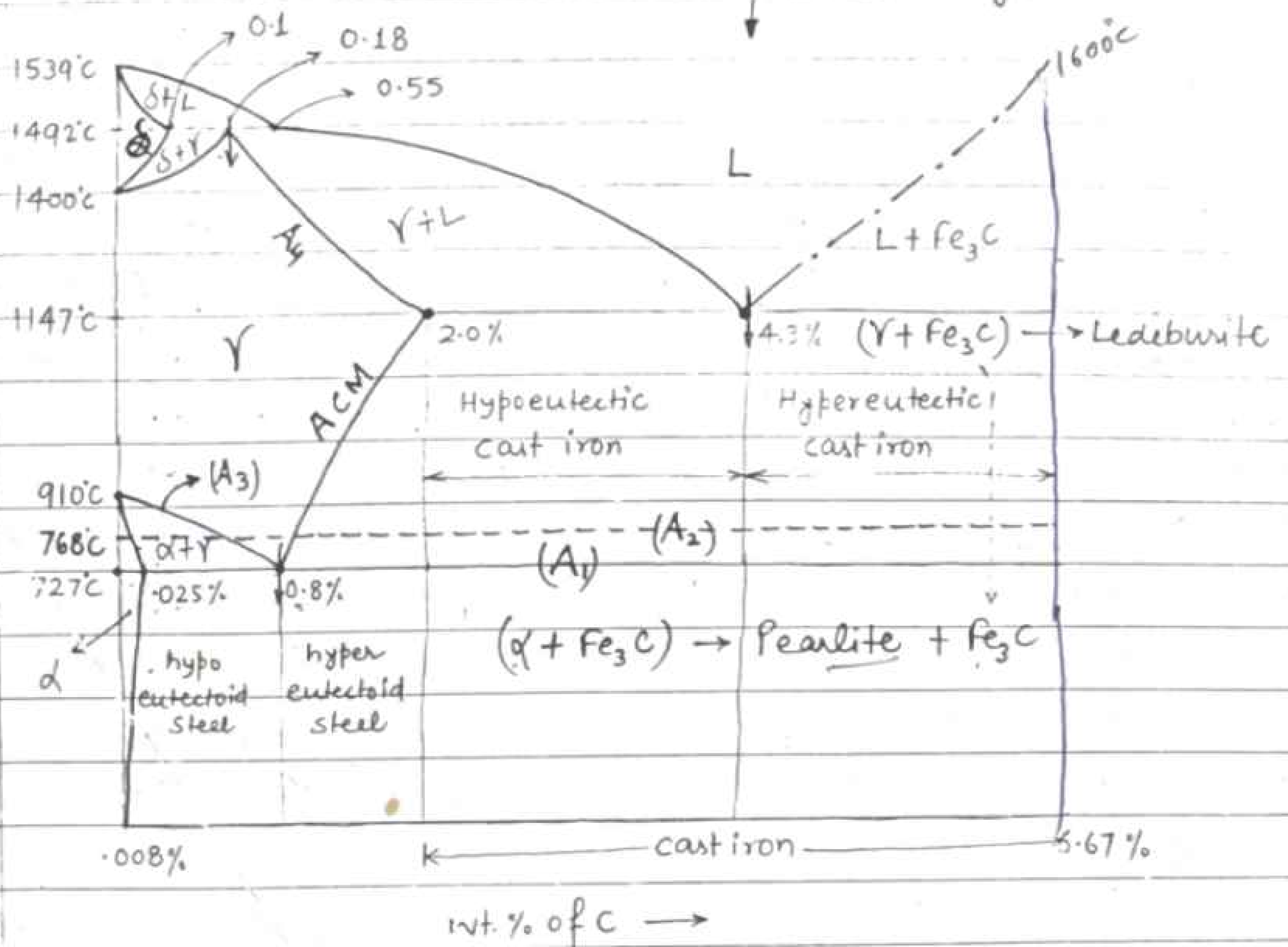
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Iron - Iron Carbide Diagram



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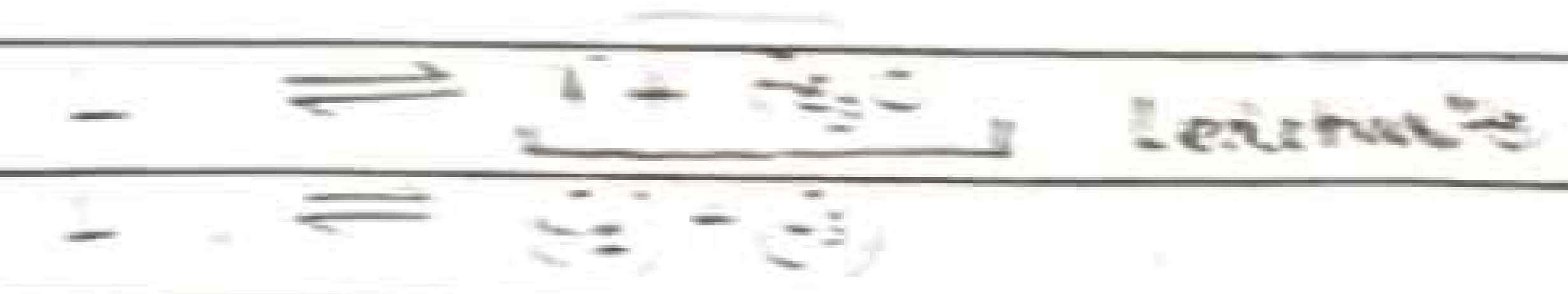
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Important Reaction:

Exothermic reaction (Heat is released)



Exothermic reaction (Heat is released)



As Temp $\uparrow$, magnetic dipoles of iron get randomized, so they will cancel out their effect and become non magnetic.

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$\delta \rightarrow$ Ferrite

$\gamma \rightarrow$ Austenite

$\alpha \rightarrow$ cementite ferrite

1.) δ -ferrite: It is an interstitial solid solⁿ of carbon in δ iron which is of BCC structure with $N_{av}=2$ and non magnetic in character.

2.) γ -Austenite: It is defined as an interstitial solid solⁿ of C in γ iron which is of FCC structure with avg. no. of atoms 4 and non magnetic in char.

3.) α -ferrite: It is an interstitial solid solⁿ of C in α iron which is of BCC structure with $N_{av}=2$ and magnetic in character only up to 768°C .

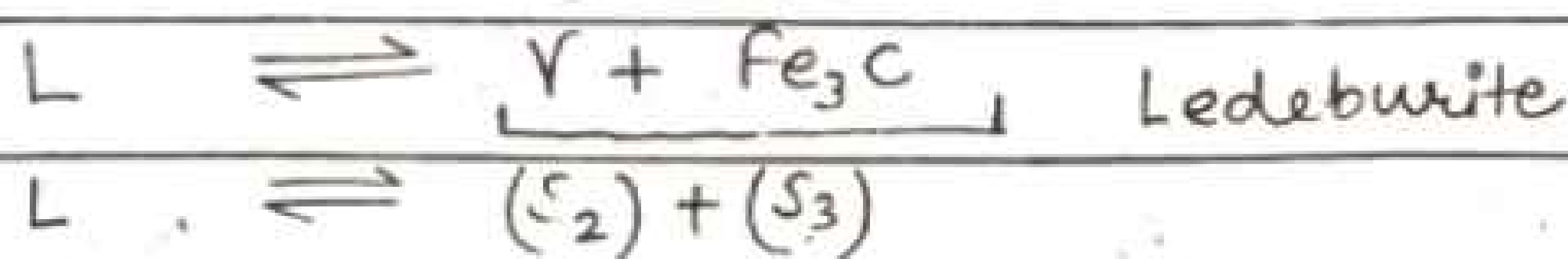
4.) Cementite (Fe_3C) It is an intermetallic compound of iron and carbon with a fixed carbon content of 6.67% in it.

INVARIANT REACTION :

1.) Peritectic reaction (1492°C , 0.18% C)



2.) Eutectic reaction (1147°C , 4.3% C)



- 1) When liquid is cooled down, it formed two solids one is austenite (solid soln) and second cementite (a compound).
- 2) Ledeburite is defined as the eutectic mixture of ^{austenite} ferrite and cementite.
Ledeburite is defined as eutectic reaction product of liquid

- 3) Eutectoid reaction (727°C , $0.8\% \text{C}$):



Pearlite is defined as eutectoid mixture of ferrite and cementite. 'or' It is eutectoid decomposition product of austenite.

→ SOLUBILITY OF Carbon in various phases :

- 1) Max^m C solubility in δ -ferrite phase is 0.01% .
- 2) " " " " γ -austenite " " 0.2% .
- 3) " " " " α -ferrite " " 0.025% .
- 4) Room temp. solubility of C in α -ferrite is 0.008% .

→ Significance of some important line in diagrams :

$A_1 \rightarrow \text{Pearlite} \rightarrow \text{Austenite}$

$A_2 \rightarrow \text{ferrite} \rightarrow \text{Austenite}$

$A_{cm} \rightarrow \text{Cementite} \rightarrow \text{Austenite}$

$A_4 \rightarrow$

→ A_1 line is called as lower critical temp. line. When

Eutectoid steels are heated to a temp. above A_1 line, pearlite transforms into austenite.

➤ A_2 line occurs at 768°C . It signifies the magnetic to non magnetic transformation on heating.

➤ A_3 line is known as upper critical temp. line for hypo eutectoid steel. It signifies the transformation of ferrite into austenite upon heating to a temp. above A_3 line.

➤ ACM line is known as upper critical temp. line for hyper eutectoid steel. This line signifies the transformation of cementite into austenite upon heating.

➤ Mild steel gives a very mild steel response to heat treatment process. (low % of C) ✓

✓ b/s cementite is in needle form shape but thermal cycle around A_1 of pearlite gives rise to spheroidal shape cementite (softest)

→ HEAT TREATMENT OF STEELS ←

- 1) Annealing
 - 2) Normalising
 - 3) Hardening
 - 4) Surface hardening
 - carburizing
 - Nitriding
 - Cyaniding
 - Induction hardening
 - Flame hardening
- } low 'C' steels
- } high 'C' steels

1) ANNEALING :

Steel sample is heated to completely austenitic range, held at the particular temp. for sufficiently long time for homogeneous austenitization to take place. It is then allowed to cool in the furnace itself. gradually.

This process results in development of uniformly coarse grain size on all surfaces and the resulting phase structure is pearlite (coarse).

Annealing result in relieving of internal stresses, recovering of ductility, increase of machinability, softening of material etc.

When it is applied to cold worked metal, the process is known as recrystallization annealing or stress relief annealing. The operating temp. in cold worked material case is less than A_1 .

The oxidation of annealed surface can be prevented by circulating the inert gases in the furnace

Box $\rightarrow$ CI $\rightarrow$ CI chips has high affinity towards air

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(bright annealing) or by enclosing the steel sample in a box (box annealing)

2) NORMALISING :

Steel is heated to austenitic range and taken out of the furnace and allowed to cool in the open air. This process results in non uniform grain structure and fine pearlitic phase structure in steel due to fast rate of cooling.

In normalising residual stress are not completely relieved (due to fast cooling), hence strength and hardness of normalised steel is always more than that of annealed steel.

Machinability of normalised steel is inferior to annealed steel.

3) HARDENING :

Steel is heated in the temp. range of $1100-1300^{\circ}\text{C}$ and quenched in a quenching medium. This sudden cooling results in the transformation of austenite into martensite with the following characteristics -

- $\rightarrow$ Martensitic transformation occurs at the speed of sound in the quenching medium.
- $\rightarrow$ It occurs by a process of shear. (diffusionless process)
- $\rightarrow$ Martensitic transformation does not depend upon time
- $\rightarrow$ Martensitic transformation never goes to 100% completion at room temp. (ie retained austenite will be formed)

The following three methods can be applied for the elimination of retained austenite -

- i) Subzero cooling (This process transforms retained austenite into martensite)
- ii) Plastic deformation of previously hardened component :
It changes the austenite into martensite by shear distortion of lattice.
- iii) High temp. tempering : transforms retained austenite into bainite and also relieves residual stresses caused by quenching.
- High temp. tempering is the only technique which ↑ the toughness of previously hardened components.
- A hardening process which employs interrupted quenching is known as martempering. (2-3 stages). The objective of martempering is to prevent the occurrence of quenched cracks and distortion of physical geometry.
- In austempering, the steel sample is heated to hardening temp. and quenched in a molten salt bath maintained at 550°C . Then it is allowed to cool gradually along with the salt bath. This process results in the development of tougher forms of steels than the commercial steel available.

• SURFACE HARDENING :

- CARBURISING : It denotes the process of ↑ carbon concentration on the surface, by heating steel in a carbonaceous medium (pack carburising, liquid

Charcoal

Na_2CO_3

Ethene

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carburising, gas carburising)

➤ Diffusion of C atoms into the steel surface makes the surface rich in carbon.

➤ Carburising is always followed by hardening and quenching, in order to convert surface austenite into martensite. This process developed a hardness of 800 vhn but it takes a very long time.

b) NITRIDING: Steel is heated in the presence of ammonia gas at a temp of $550-580^\circ\text{C}$. The ammonia gas dissociates into nitrogen and hydrogen atoms. These nitrogen atoms diffused into surface of steel and results in iron nitride formation.

➤ Nitriding process can be completed quickly and hardness $\uparrow$ to 1000-1100 vhn. The nitrated surface will acquire very high brittleness and hence the life is less.

c) CYANIDING: Steel is heated to a temp. of 800°C in molten cyanide salt bath. This process must be followed by quenching in water or oil to take advantage of martensitic transformation.

Cyanided cases will have higher hardness than carburised cases but higher life than nitrated cases.

d) INDUCTION HARDENING Applicable to relatively small sized components made of high C steel.

One surface of steel is brought to red hot condition by

Plain Carbon Steel

Plain Carbon Steel

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0-10% Low alloy steel

10% or more high alloy steel

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Alloying Element	Effects
Sulphur	→ formation of iron sulphide which is hard & brittle → Addⁿ of S results in hot cracking tendency in steel (HOT SHORTNESS) → S ↑ Machinability
Phosphorous	→ Powerful solid solⁿ strengthener. It ↑ strength and hardness → P ↑ Machinability → P ↑ brittleness in steel. → It induces (COLD SHORTNESS) in steel.
Silicon	→ It dissolves in ferrite. → It is a strong deoxidiser. → It is used to make magnetically soft material for use in transformers.
Manganese	→ It ↑ strength, hardness, toughness. → It ↑ machinability of steel. → It is used to produce extremely tough, wear resistant and non magnetic steel called as HADFIELD Mn Steel. → It is an austenite stabilizer. → It improves abrasion resistance
Nickel	→ It ↑ toughness without ↓ ductility. → It is an austenite stabilizer → It ↑ corrosion resistance. → It is used to produce alloys with zero coefficient of thermal expansion. (INVAR steel)

Chromium	It is a carbide former. It ↑ wear resistance. It ↑ corrosion resistance.
W, Mo, V	They are carbide forming element. They ↑ wear resistance They refine grain size.
Titanium	It is a strong carbide former. It ↑ strength and hardness. Refine grain size.
Aluminium	It is a powerful deoxidizer. It refines grain size. It's a graphite forming element. ✓

→ Ceramic - features & classification ←

