Casting Processing, MH2252, 6hp



Lecture 7a+7b
Precipitation of pores and slag inclusions at casting processes

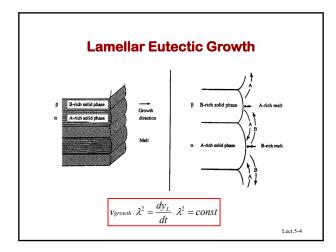
Lect 7-

Today's topics

- Repetition
 - Structure formation in cast materials
 - Microsegregation
 - Macrosegregation
- Precipitation of pores and slag inclusions at casting processes
 - Precipitation of pores
 - Gas reactions during casting
 - Pore morphology
 - Gases in steels and iron alloys/Aluminium
 - Slag inclusions
 - Cast iron

Lect.7-2

Characterisation of Dendrites - Dendrite Arm Distance and Growth Rate T_{M} $\lambda_{\text{l}} = \text{Primary Dendrite Arm Spacing}}$ $\sum_{\lambda_{\text{l}} = \text{Secondary Dendrite Arm Spacing}} T_{\text{Dendrite Arm Spacing}}$ $\sum_{\lambda_{\text{l}} = \text{Secondary Dendrite Arm Spacing}} D_{\text{rawing of a dendrite removed from a metallic melt}}$ $v_{\text{growth}} \cdot \lambda^2 = \frac{dy_L}{dt} \cdot \lambda^2 = const$ $v_{\text{growth}} \cdot \lambda^2 = \frac{dy_L}{dt} \cdot \lambda^2 = const$

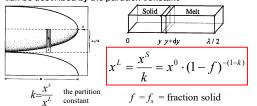


Macrostructure - Different Cooling Conditions Casting Rate Rate Coulumar Macrostructure Continuous very strong high long Figure 37 a land mould weak slow absent Figure 37 b A: Strong cooling. High casting Hedium cooling. High casting temperature temperature Lect.6-5

The Lever Rule: Equilibrium Solidification • High diffusion rates: Lever rule valid $x^L = \frac{x^S}{k} = \frac{x^0}{1 - f(1 - k)}$ $x^0 = \text{initial concentration of the alloying element}}$ f = fraction solid $k = \text{partition coefficient } (x^5/x^L) \text{ of the alloying element}}$ $D^S > 10^{-10} \text{ m}^2/\text{sek}$ See chapter 7.3.2 The Lever rule

Scheil's Model

- $\underline{f < 0.9}$ and \underline{low} diffusion rate: Scheil's model valid
- There is an even composition in the melt at every moment
- The diffusion in the solid phase is neglectable
- There exists local equilibrium between solid and liquid phase can be described by the partition constant



See chapter 7.3.1 Scheil's Model for Microsegregation

Lect.6-7

Macrosegregations in Ingots Structures Macrosegregations Macrosegregations Positive zone A-segregations V-segregations V-segregations Negative zone See chapter 11.9 Macrosegregations in Steel Ingots Lect.7-8

Sedimentati	ion Seg	grega	ition - +	/- Segregation
		H	34448	a) Equiaxed crystals in the melt before sedimentation. b) Equiaxed crystals, including their associated melt, after sedimentation
all <mark>o</mark> ying e <i>Negative</i>	lement ex segregation	ceeds th on : there	e is instead	ation of the concentration. a lack of the oncentration)

A-segregations - Ghost lines! The density of the melt increases linearly with decreasing temperature. The density of the melt within the two-phase region decreases due to microsegregation of lighter alloy elements. Convection pattern Natural convection in the melt and an **opposite convection pattern** in the two-phase region. Higher concentration of alloy elements will lower the liquidus temperature and cause **re-melting** of solid structure.

· Ghost-lines!

Distance

A-segregations - Freckles 0000 Channels (ghost lines/freckles) in castings are caused by natural convection due to concentration gradients in the two-phase region during the solidification process of an ingot. The channels are created in the partly solidified two-phase region S+L.

Lect.7-11

V-segregation Vacuum, caused by the solidification shrinkage in the lower parts of the ingot. Time after solidification Cannot be filled by melt from above because the distance is long and the 34 min resistance from the 50 min dendrite arms is great. The pressure difference, which arises between the 10 min different parts of the ingot results in a general settling of the crystals in the middle

Porosity in castings

- Deteriorate quality of a casting
- Lower mechanical properties
 - Reduced elasticity
 - Decreased impact resistance
- · Risk for crack formation
- · Reduced surface quality during machining
- · Gas Porosity
 - Gas entrapped during casting
 - Precipitation of dissolved gas
 - Smooth surface of porosity
- Shrinkage Porosity
 - Solidification shrinkage
 - Uneven surface of porosity

Lect.7-13

Shrinkage porosity - gas porosity

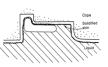




- Pores and porosity (micro / macro) cause surface defects and poor mechanical properties.
- Shrinkage porosity, arises because insufficient feeding of melt to the last solidified areas.
- Gas porosity, resulting from a high content of gas-forming substances in the melt.

Sources for Common Gases in Metals

- Entrapped Air
- From atmosphere
 - Oxygen, Nitrogen
 - Hydrogen from moisture
 H₂0+M=2<u>H</u>+MO and 2<u>H</u>=H₂(g)
- · In Steels and Iron
 - Carbon monoxide, <u>C</u>+O₂=2CO(g)
- Reaction with mould materials
 - H_2 (moisture in sand), O_2 , CO, CO_2 , SO_2



Entrapped Air



Precipitated gas porosity

Lect.8-15

Condition for pore precipitation (Pressure in a Gas Bubble)

$$p_{gas} = p_{alm} + p_h + \frac{2\sigma}{r}$$

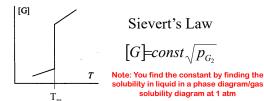
$$p_h = \rho g h \qquad p_{gas} > p_{alm}$$

$$p_{gas} > p_{alm}$$
Condition for pore precipitation (simplified)

Note: the lower surface tension - the easier it is to form bubbles (gas pores)

See chapter 9.3.1 Gas Reactions and Gas Precipitation in Metal Melts

Gas Solubility in Metal Melts

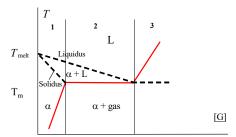


The concentration of the dissolved gas atoms in the melt or in the solid phase is proportional to the square root of the partial pressure of the gas in the surrounding atmosphere

See chapter 9.7.1 Hydrogen in Steel and Iron Alloys

Lect.7-17

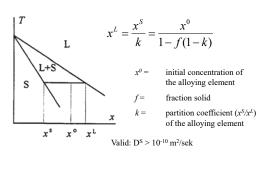
Gas Solubility and the Phase diagram



Concentration of solved gas atoms

Region 1: Gas content totally dissolved in solid.
Region 2: Precipitation of gas porosity during solidification
Region 3: Precipitation of gas porosity during melt cooling
followed by gas porosity during solidification

The Lever Rule



Lect.7-19

Pore precipitation in solidifying melts

1.
$$x^L = (\frac{x^S}{k}) = \frac{x^0}{1 - f(1 - k)}$$
 The Lever Rule.

- Always valid for gaseous elements
- f^S in relation to x^L

 $[G]=const\sqrt{p_{G_2}}$

- Sievert's Law Valid for two atomic gas molecules ${\bf x}^{\rm L}$ in relation to ${\bf P_G}^2$

3. $p_{gas} > p_{atm}$

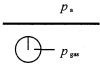
- $\begin{array}{l} \textbf{Condition for pore precipitation} \\ \textbf{-} \textbf{Simplified, close to the melt surface} \\ \textbf{-} \textbf{P}_{\text{G}}^2 \textbf{in relation to P}_{\text{atm}} \end{array}$

The partition coefficient

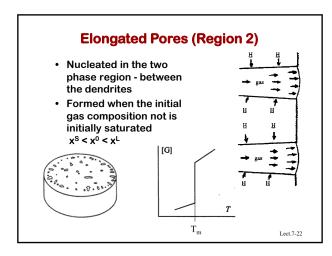
- close to 1, low enrichment, << 1 large enrichment to the liquid

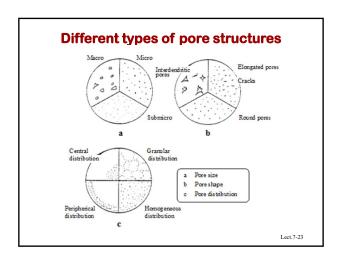
Lect.8-20

Rounded Pores (Region 3)



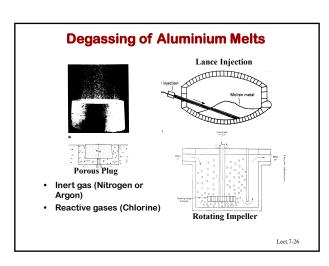
Precipitated in melt Composition in melt $x^0 > x^L$

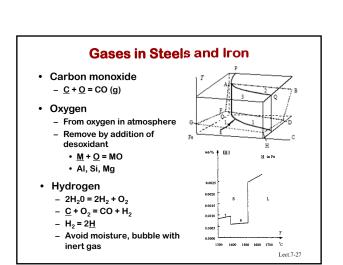




Methods of lowering the dissolved gas content in metal melts · Solidification - re-melting Stirring - Subsonic stirring Lowering the partial pressure Rotating Impeller - Vacuum treatment <u>Common gases</u> H₂: Hydrogen - Inert gas bubbling - Active gas bubbling N_2 : Nitrogen • Desoxidation - precipitation O_2 : Oxygen of secondary phases CO: Carbon monoxid

Pick-Up in Aluminium Temperature. °F 22 1000 1400 1600 10 1400 1600 10 14





Hydrogen concentration in a steel melt 0.05 0.02 0.01 0.0015 0.00 0.0010 0.0005 1300 1400 1500 · Hydrogen pick up

$-2H_20 = 2H_2 + O_2$

 $[\underline{H}]^2 \cdot [\underline{O}] = const$

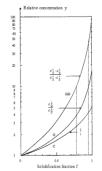
Note: a decrease of the oxygen concentration leads to an increased hydrogen concentration as a consequence of the solubility product of water

Oxygen and Carbon Monoxide in a Steel melt

$$\begin{split} &c_{\underline{C}}^{\ L} \cdot c_{\underline{Q}}^{\ L} = 0.0019 \cdot p_{\ CO} \qquad \text{(w/o and atm)} \\ &y_{\underline{O}} = c_{\underline{D}}^{\ L} / c_{\underline{O}}^{\ o} = \frac{1}{1 - f \cdot \left(1 - 0.054\right)} \\ &y_{\underline{C}} = c_{\underline{D}}^{\ L} / c_{\underline{O}}^{\ o} = \frac{1}{1 - f \cdot \left(1 - 0.020\right)} \end{split}$$

$$\mathbf{y}_{\underline{\mathbf{co}}} = \mathbf{y}_{\underline{\mathbf{c}}} \cdot \mathbf{y}_{\underline{\mathbf{o}}} = \frac{c_{\underline{\mathbf{c}}}^{\mathrm{L}} \cdot c_{\underline{\mathbf{o}}}^{\mathrm{L}}}{c_{\underline{\mathbf{c}}}^{\mathrm{o}} \cdot c_{\underline{\mathbf{o}}}^{\mathrm{o}}}$$

Both C and O are present in all steel and iron melts. When the steel solidifies both C and O concentrate in the melt and the saturation limit may be exceeded



See chapter 9.7.3 Oxygen and Carbon Monoxide in Steel and Iron Alloys

Slag Inclusions (Cinders)

- Slag:
 - Non-Metallic Inclusions in Metals
- · Microslags:
 - Not detectable by naked eye (<0.1 mm)
 - Most often formed interdendritic during solidification
 - M+O = MO where M = desoxidation element (Si, Al etc.) Solubility Product [M][O]= K_{MO}
- · Macroslags:
 - Visible by eye (>0.1 mm)
 - Formed in the melt before solidification
 - Reaction products from desoxidation
 - Refractory material from mould walls



oxide skin (Al₂O₃)



Cast Iron - Fe-C alloys with carbon content > 2.1 w/o C 1800 1400 1200 1000 800 **Promotes Grey Iron** Si - Important Ni, Cu, Co Steel < 2% C Slow Cooling Rate Cast Iron > 2% C (2.5-4.3 %C) - Inoculation Promotes White Iron T_E= Stable (1153 °C) Cr, V, TiHigh Cooling Rate T_E = Metastable (1147 °C)

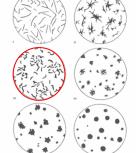
White Cast Iron



- Mixture of cementite and austenite
- Eutectic is called ledeburite
- · Very hard
- Very low ductility = fragile
- High wear resistance
- Friction material

Lect.5-32

Different types of grey cast iron



I, II : Flake graphite

III : Compacted graphite, CGI

IV,V: Vermicular graphite

VI : Nodular graphite

Lect.5-33

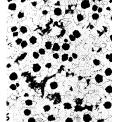
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A. Flake Graphite

- Spherical cells
- Very good castability
- Good damping properties

Lect.5-34

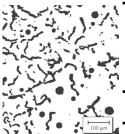
B. Nodular Graphite (Ductile Iron)



- Spheroidal graphite morphology
- By adding Cr or Mg
- More difficult to cast
- Good ductility
- High strenght

Lect.5-35

C. Compact Graphite Iron (CGI)



- Intermediate between flake and spheriodal graphite - with respect to structure castability and strength
- By adding small amounts of Cr or Mg
- Difficult to control microstructure

Lect.5-36

Recommended reading in "Materials Processing during Casting", by Hasse Fredriksson and Ulla Åkerlind Chapter: 9.1 – 9.9