

Kinetics Model of Isothermal Pearlite Formation in a 0.4C-1.6Mn Steel

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Abstract

The present article is concerned with a theoretical and experimental study of the growth kinetics of pearlite in a 0.4C-1.6Mn medium carbon steels. Factors controlling the isothermal formation of this microconstituent are explored in this work. In this sense, the transition temperature between local equilibrium (LE) and no partition local equilibrium (NPLE) growth mechanisms is theoretically determined. Moreover, the nucleation of pearlite has been considered as a cementite precipitation process on a moving austenite- ferrite interface. Finally, a theoretical model is presented in this work to calculate the evolution of austenite-to-pearlite transformation with time at a very wide temperature range.

Keywords phase transformations, kinetics, steels, forging, structural

1- Introduction

Pearlite is probably the most familiar microstructural feature in the whole science of metallography. It was discovered by Sorby over 100 years ago, who assumed it to be a lamellar mixture of ferrite and cementite. Pearlite is a very common constituent of a wide variety of steels, where it provides a substantial contribution to strength, so it is not surprising that this phase has received intense study [1].

It is now generally agreed that during pearlite growth the alloying element distributes between the ferrite and cementite at low supersaturations (LE mechanism), and the growth is controlled by alloying element boundary diffusion. At higher supersaturations pearlite growth occurs without any partitioning of the alloying element (NPLE mechanism) and it is controlled by carbon volume diffusion [2]. The partitioning of alloying elements has been experimentally observed by Razik et al. [3], Al-Salman et al. [4], and Chance and Ridley [5] in a number of Fe-C-Mn, Fe-C-Si, and Fe-C-Cr alloys. In all these studies, the partitioning was observed at low supersaturations, whereas below a characteristic temperature of the steel no partition was found.

Recent works have demonstrated that medium carbon forging steels with acicular ferrite microstructure can be manufactured at industrial scale [6-8]. The main interest of this microstructure lies in the good combination of mechanical properties that presents as compared with bainite and especially with ferritic-pearlitic microstructures. In those steels, acicular ferrite is always formed after the growth of allotriomorphic ferrite and pearlite. As a consequence, acicular ferrite transformation is inevitably influenced by previous allotriomorphic ferrite and pearlite formation. The role of the allotriomorphic ferrite to promote the formation of acicular ferrite to the detriment of bainite in a two stages heat treatment has been reported in previous works [9-10]. Thus, the amount of acicular ferrite increases as allotriomorphic ferrite is formed along the austenite grain boundaries because saturation of nucleation sites occurs. However, if pearlite transformation follows the allotriomorphic ferrite one, there will not be untransformed austenite to obtain a massive acicular ferrite transformation as temperature decrease. Therefore, a deep understanding of the decomposition of austenite in allotriomorphic ferrite and pearlite is needed in order to control the total amount of acicular ferrite present in the microstructure in medium carbon forging steels.

Recently, the austenite-to-allotriomorphic ferrite transformation has been reported by the authors [11-13]. In this work, a mathematical transformation model is presented for the purpose of simulating the austenite-to-pearlite transformation which occur during the isothermal decomposition of austenite. The model is quite general but was targeted at modern forging steels

compositions, which typically contain 0.4 wt.% C, <0.6 wt.% Si and 1.6 wt.% Mn.

2- Experimental Procedure

Cylindrical dilatometric test pieces of 2 mm in diameter and 12 mm in length were machined parallel to the rolling direction of the bar. Experimental validation of the pearlite transformation kinetics model developed in this work was carried out using the heating and cooling devices of an Adamel Lhomargy DT1000 high-resolution dilatometer described elsewhere [14]. The heating device consists of a very low thermal inertia radiation furnace. The power radiated by two tungsten filament lamps is focussed on the specimen by means of a bi-elliptical reflector. The temperature is measured with a 0.1 mm diameter Chromel – Alumel (Type K) thermocouple welded to the specimen. Cooling is carried out by blowing a jet of helium gas directly onto the specimen surface. These devices ensure an excellent efficiency in controlling the temperature and holding time of isothermal treatments and as well as fast cooling in quenching processes.

Austenitisation conditions were fixed to avoid the influence of the austenite grain size on the kinetics of pearlite formation. Since the transformation rate of pearlite is higher the finer the prior austenite grain size (*PAGS*), a coarse *PAGS* of 76 μm was selected to make easier the experimental study of the transformation kinetic of pearlite. Thus, specimens were austenitised at 1523 K for 1 min and subsequently isothermally transformed at temperatures ranging from 943 to 873 K during different times. In order to freeze the microstructure at those temperatures, specimens were quenched to room temperature by helium gas flow at a cooling rate of 200 K/s.

Specimens were polished in the usual way for metallographic examination. Nital - 2pct etching solution was used to reveal the ferrite+pearlite microstructure by optical microscopy. The volume fraction of pearlite (V_p) was statistically estimated by a systematic manual point counting procedure [15]. The maximum volume fraction of pearlite (V_p^{EQ}) formed during the isothermal decomposition of austenite at 943, 933, 913 and 873 K was again determined by a combination of dilatometric and metallographic analysis.

Special metallographic preparation was required to reveal the interlamellar spacing of pearlite. Deep primary etching with a solution of picric acid in isopropyl alcohol with several drops of Vilella's reagent was used to ensure that any deformed layer introduced by polishing was removed. This etching was eliminated using the 1 and 0.25 μm diamond pads with almost no pressure being exerted on the sample for no longer than 3 to 4 min. The sample was then etched again, this time lightly, and polished carefully on the 1 and 0.25 μm diamond pads. Finally, a light etch was given to the sample. This preparation procedure was carefully detailed in Ref. [16]. The values of the mean true interlamellar spacing (S_o) were derived from electron micrographs according to Underwood's intersection procedure described in Refs. [16] and [17].

The austenite-to-allotriomorphic ferrite (Ae_3) and austenite-to-pearlite (Ae_1) critical temperatures were experimentally determined by dilatometric and metallographic analysis. Initially, both temperatures were estimated from a dilatometric curve obtained by continuous cooling at a rate of 0.05 K/s. This is the rate normally used for considering quasi-equilibrium conditions [18]. Figure 1 shows the dilatometric curve obtained during austenite transformation by continuous cooling processes. Since the difficulty for monitoring separately the pearlite and proeutectoid ferrite transformation, the Ae_1 temperature was more accurately determined after several isothermal heat treatments at temperatures ranging from 953 to 933 K. Thus, a temperature of 948 K could be defined as the Ae_1 temperature for the studied steel.

3- Calculation of the interface compositions

The interface compositions at the a/g and a/q boundaries under LE and NPLE growth mechanisms have been calculated according to the method firstly reported by Kirkaldy and co-workers [19-20]. Thermodynamic data for the calculation of the compositions in ferrite (a), austenite (g) and cementite (q) phases in a Fe-C-Mn system are listed in the Appendix. The equilibrium condition can be expressed by the equality of the chemical potentials of each element in both phases at the interface. The chemical potentials of carbon, Mn and Fe in ferrite, austenite and cementite are

calculated assuming Hillert - Staffanson regular solution model, and are expressed as follows. The numbers 0, 1 and 2 denote, respectively, Fe, C and Mn. Likewise, the average mole fractions are designated as \bar{x}_i ($i=1$ to 2), and the mol fractions in each phase as x_i ($i=0$ to 2).

In austenite and ferrite;

$$m_0 = G_0 + RT \ln x_0 - \frac{RT}{2} (e_{11}x_1^2 + e_{22}x_2^2) - RTe_{12}x_1x_2 \quad (1)$$

$$m_1 = G_1 + RT \ln x_1 + RT(e_{11}x_1 + e_{12}x_2) \quad (2)$$

$$m_2 = G_2 + RT \ln x_2 + RT(e_{12}x_1 + e_{22}x_2) \quad (3)$$

and in cementite;

$$m_{Fe_3C} = G_{Fe_3C} + \frac{3}{4} RT \ln y_0 + \frac{3}{4} (1 - y_0) w_{02} y_2 \quad (4)$$

$$m_{M_3C} = G_{M_3C} + \frac{3}{4} RT \ln y_2 + \frac{3}{4} (1 - y_2) w_{02} y_0 \quad (5)$$

where w_{02} is the Fe-Mn interaction coefficient, e 's are the Wagner's interaction coefficients, and $y_i = 4x_i^{ag}/3$. In the case of LE mechanism the equilibrium conditions can be expressed as follows.

$$m_i^a = m_i^f \quad (i=0 \text{ to } 2) \quad (6)$$

$$4m_{Fe_3C} = 3m_0^f + m_1^f \quad (7)$$

$$4m_{M_3C} = 3m_2^f + m_1^f \quad (8)$$

Equations (6) for carbon ($i=1$) and manganese ($i=2$), and considering that $e_{12} = e_{12}^a = e_{12}^g$, are rewritten in terms of the partition coefficients A_1 and A_2 as

$$x_1^{ag} = A_1 x_1^{ga} = \frac{\exp(DG_1^{ag}/RT + e_{11}^g x_1^{ga})}{1 + e_{11}^a x_1^{ga} \exp(DG_1^{ag}/RT)} x_1^{ga} \quad (9)$$

$$x_2^{ag} = A_2 x_2^{ga} = \frac{\exp(DG_2^{ag}/RT + e_{12}^g x_1^{ga})}{1 + e_{12}^a x_1^{ga} \exp(DG_1^{ag}/RT)} x_2^{ga} \quad (10)$$

where DG_1^{ag} and DG_2^{ag} are the free energy change between austenite and ferrite for carbon and manganese, respectively. According to Hashiguchi et al. [20] the tie-line for the manganese concentration in austenite in the a/g boundary may be expressed by,

$$x_2^{gf} = \frac{\bar{x}_2(1 - A_1)x_1^{gf}}{(1 - A_2)\bar{x}_1 + (A_2 - A_1)x_1^{gf}} \quad (11)$$

On the other hand, equation (6) for iron ($i=0$) can be rewritten as follows:

$$DG_0^{ag} + RT \ln \frac{\bar{e}_1 - x_1^{gf} - x_2^{gf}}{\bar{e}_1 - x_1^{ag} - x_2^{ag}} \frac{\bar{u}}{\bar{u}} - \frac{RT}{2} \left[\left(e_{11}^g x_1^{gf^2} - e_{11}^a x_1^{ag^2} \right) + \left(e_{22}^g x_2^{gf^2} - e_{22}^a x_2^{ag^2} \right) \right] - RT e_{12} \left(x_1^{gf} x_2^{gf} - x_1^{ag} x_2^{ag} \right) = 0 \quad (12)$$

where DG_0^{ag} is the free energy change between austenite and ferrite in iron. This equation (12) contains only one unknown variable, x_1^{gf} , because the x_1^{ag} and x_2^{ag} , as well as x_2^{gf} , are functions of x_1^{gf} as expressed in equations (9), (10) and (11), respectively. Therefore, it is possible to numerically calculate the value of x_1^{gf} for each temperature.

An expression for the equilibrium between cementite and austenite may be obtained from the subtraction of equation (7) from (8). Expanding the equations for the chemical potentials according to the expressions from (1) to (3), and restricting attention to low alloy steels where $x_2 \ll 1$, $y_2 \ll 1$, $x_0 \approx 1$ and $y_0 \approx 1$, equation (7) is approximated by

$$y_2 = B_2 x_2^{gf} = x_2^{gf} \exp \left\{ \left[DG_0^{gf} - DG_2^{gf} - w_{02} \right] / RT + e_{12} x_1^{gf} \right\} \quad (13)$$

According to Sharma et al. [21], the tie line for the Mn concentration is expressed by the following equation

$$x_2^{gf} = \frac{\bar{x}_2(4x_1^{gf} - 1)}{3B_2(x_1^{gf} - \bar{x}_1) + 4x_1 - 1} \quad (14)$$

Expanding equation (7) according to equations (1) to (3) the following equation is yield

$$DG_0^{gf} = RT \ln \frac{\bar{e}_1 - x_1^{gf} - x_2^{gf}}{\bar{e}_1 - y_2} \frac{\bar{u}}{\bar{u}} + \frac{RT}{3} \ln x_1^{gf} - \frac{RT}{2} \left(e_{11} x_1^{gf^2} + e_{12} x_2^{gf^2} \right) - \frac{RT}{3} \left(3e_{12} x_1^{gf} x_2^{gf} - e_{11} x_1^{gf} - e_{12} x_2^{gf} \right) - y_2^2 w_{02} \quad (15)$$

where DG_0^{ag} is the free energy change between austenite and cementite phases under equilibrium conditions for iron. Since x_2^{ag} and y_2 are expressed as function of x_1^{ag} , this equation is only function of x_1^{ag} . This parameter can be solved in the same way that x_1^{gp} in equation (12).

The transition from the LE to the NPLe conditions occurs when $x_2^{ag} = \bar{x}_2$ and $x_2^{gp} = \bar{x}_2$ for the a/g and g/q interface, respectively. Therefore, considering that the value of x_2^{gp} may be obtained from equation (10), i.e. $x_2^g = \bar{x}_2/A_2$, the corresponding value of x_1^{gp} at the a/g interface under NPLe consideration is derived from equation (12) assuming these values for x_2^{ag} and x_2^{gp} . In the same way, the value of x_1^{gp} under NPLe mechanism at the g/q interface is calculated from equation (15) but considering $x_2^{ag} = \bar{x}_2$ and $x_2^{gp} = 4\bar{x}_2/3B_2$ (equation (13)).

4-Theoretical determination of the maximum volume fraction of pearlite (V_p^{EQ})

The V_p^{EQ} formed after the complete isothermal decomposition of austenite can be determined by applying the lever rule at the phase diagram schematically presented in Fig. 2 known as Hultgren's extrapolation. As it was pointed out by Christian [22], pearlite starts to form at temperature below Ae_1 , and there is a temperature (T^* in Fig. 2) at which pearlite is the only decomposition product of austenite. In this sense, the maximum amount of pearlite obtained at temperatures ranging from Ae_1 and T^* could be expressed as,

$$V_p^{EQ} = \frac{x_1^{gp} - x_1^{gq}}{x_1^{gp} - x_1} \quad (16)$$

where x_1^{gp} and x_1^{gq} are the carbon concentration in austenite which are under either LE or NPLe conditions with ferrite and cementite, respectively.

Figure 3(a) shows the section of the Fe-Mn-C system for a manganese concentration of 1.6 wt.-%. From this figure the carbon content at the a/g and g/q interface can be easily obtained, and thus V_p^{EQ} at different temperatures can be derived using equation (16). Figure 3(b) shows experimental

and calculated values of V_p^{EQ} as a function of T considering either LE or NPLe conditions. According to Fig. 3(b), the formation of pearlite proceeds under LE consideration at temperatures of 943 and 933 K, whereas no partitioning seems to be the dominant mechanism at temperature of 873 K. This behaviour is consistent with the experimental observations carried out by several authors [1,3,23] in Fe-C-Mn eutectoid steels, which reported partitioning temperature of 933 K for manganese concentration of 1.08 wt.-% [1], and partitioning temperature of 895 K for manganese concentration of 1.8 wt.-% [3,23].

5- Evaluation of the transition between LE and NPLe mechanisms

The partitioning coefficient of manganese, K^p , between ferrite and cementite is calculated using the following expression

$$K^p = \frac{x_2^{gq}}{x_2^{ag}} \quad (17)$$

At temperatures closer than the theoretical eutectoid temperatures, K^p is expected to have a larger value since the supersaturation of austenite is small, and the mobility of Mn atom is high at those temperatures. Thus, the value of K^p decreases with temperature and reaches the unity when no partitioning condition occurs at both ga and gq interfaces. Figure 4(a) shows a comparison between the measured K^p values reported by Ridley [1] in a Fe-0.7C-1.08Mn steel, and the calculated values according to the model presented in this work. This figure suggests that the change with temperature in K^p is successfully predicted for the model in a Fe-0.7C-1.08Mn steel. Once the reliability of the model for K^p calculation has been proved, the variation of K^p with isothermal temperature for the Fe-0.4C-1.6Mn steel studied in this work has been calculated. The effect of previous proeutectoid ferrite transformation has been taken into account assuming carbon enrichment in the residual austenite. Therefore, the change in the bulk carbon concentration is expressed in the following equation:

$$\bar{x}' = \frac{\bar{x}}{1 - F} \quad (18)$$

where F is the volume fraction of proeutectoid ferrite transformed during the isothermal decomposition of austenite. Thus, the change in bulk carbon concentration at each isothermal temperature should be taken into account in the calculations of the interface concentrations of equation (17). Figure 4(b) shows the evolution of K^p with the isothermal temperature in the studied steel considering the corresponding carbon enrichment in austenite due to proeutectoid ferrite formation. This figure suggests that the isothermal formation of pearlite occurs with partitioning of manganese at a temperature of 893 K, which is consistent with the value of ~895 K reported by Tewari and Sharma in a 0.69C-1.8Mn eutectoid steel [23].

6- The onset of pearlite transformation

Since proeutectoid ferrite is usually the first phase to develop on isothermal heat treatment, pearlite nodules nucleate on the austenite- proeutectoid ferrite (a_p) interface. It has been observed that the formation of pearlite requires the establishment of cooperative growth of ferrite and cementite [24]. The previous formation of proeutectoid ferrite enriches in carbon the surrounding austenite promoting the formation of cementite nucleus at the g/a_p interface and the local reduction of carbon content in the austenite that surrounds the cementite nucleus leads to the ferrite formation of pearlite aggregate. The simultaneous ferrite and cementite formation process yields to the characteristic lamellar structure of pearlite.

Aaronson *et al.* [25] analysed the conditions under which nucleation is feasible at moving disordered interface boundary. The restriction that the migration rate of the g/a_p boundary, G_{ag}^* , at which nucleation may take place must not exceed that which displaces this boundary a distance equal to the austenite lattice parameter in the time required for an embryo to develop to the critical nucleus size should be satisfied. He concluded that G_{ag}^* at which a crystal of cementite can nucleate is

$$G_{ag}^* = \frac{-a_g^3 D_C^g x_1^{g_a} D G_V^3}{16(1 - \cos \gamma) \sqrt{\rho k_B T S_{ag}^5 K}} \quad (19)$$

where a_g is the austenite lattice parameter; x_1^{gs} is the carbon concentration in austenite; k_B is the Boltzmann constant; T is the isothermal temperature; D_C^g is the carbon diffusion coefficient in austenite at the isothermal temperature; K is the ratio between the volume of the double spherical cap critical nucleus and that of a sphere of the same radius; DG_V is the volume free energy change; $\cos\gamma$ is the ratio between the interfacial energies of disordered $a/g(s_{ag})$ and $a/q(s_{aq})$ boundaries, and is defined as $\cos\gamma = s_{ag}/2s_{aq}$.

In this work, a value of $K = 0.0001$ has been considered [25]. Likewise, a_g has been calculated as reported by Dyson and Holmes considering the dependence of alloying elements on the lattice parameter of austenite [26]. Likewise, the value of DG_V for cementite nucleation at 913 and 873 K has been calculated as reported by Zener [27]. Values of a_g and DG_V at 913 and 873 K are listed in Table 1.

Calculations of D_C^g have been carried out according to Bhadeshia [28]. The author considers both the kinetic and equilibrium thermodynamic behaviour of carbon in austenite. These calculations takes also into account the concentration dependence of the activity of carbon in austenite, and the repulsive interactions between the nearest neighbouring carbon atoms located in octahedral interstitial sites. Thus, D_C^g is calculated by two factors: one of them is a concentration dependent factor and the other one is independent

$$D_C^g = \chi(q) \frac{k_B T}{h} \frac{\exp\left(-\frac{DG^*}{k_B T}\right)}{g_n} \exp\left(-\frac{DG^*}{k_B T}\right) \quad (20)$$

where $\chi(q)$ is the carbon concentration dependent factor obtained according to Bhadeshia's calculations [28] and takes values listed in Table 1; DG^* is the activation energy for diffusion; g_n is an activity coefficient assumed constant; l is the distance between the {002} austenite planes and h is the Planck's constant. Bhadeshia [28] found that $DG^*/k_B = 21230$ K and $\ln(g_n/l^2) = 31.84$. The values of D_C^g for temperatures of 913 K and 873 K are also listed in Table 1.

Finally, assuming that the g_{a_p} interface is a planar disordered boundary and of infinite extent, the velocity of the moving interface (G_{ag}) diminishes with time as follows,

$$G_{ag} = \frac{a_1}{2t^{1/2}} \quad (21)$$

where a_1 is the one-dimensional parabolic growth rate constant and t represents the growth time. Bradley and Aaronson [29] reported a comparison between measured allotriomorphic ferrite growth kinetics data in a Fe-C-Mn with the predictions of three different models. They concluded that paraequilibrium model is the most satisfactory of those available. The value of a_1 under paraequilibrium conditions can be obtained by numerical solution from the equation [11,22]

$$\frac{p}{4D_c} \frac{\partial}{\partial t} a_1 \exp\left(\frac{a_1^2}{4D_c} \operatorname{erfc}\left(\frac{a_1}{2\sqrt{D_c t}}\right)\right) = \frac{x_1^{gs} - \bar{x}_1}{x_1^{gs} - x_1^{ag}} = W \quad (22)$$

The values of W and a_1 for temperatures of 913 and 873 K are listed in Table 1.

A comparison between equations (19) and (21) allows us to evaluate the time required to start to form pearlite during the isothermal decomposition of austenite in the steel studied (t^*). However, it is necessary to evaluate the value of s_{aq} and s_{ag} in the studied steel in order to determine G_{ag}^* . An estimation of the value of this interface energy is yield by measuring the interlamellar spacing of pearlite as shown below.

7- Calculation of ferrite-cementite interfacial energy (s_{aq})

When the growth rate of pearlite is controlled by the bulk diffusion of carbon in austenite, Zener [27] proposed the following relationship for the interlamellar spacing, S_o , and the theoretical critical spacing at zero growth rate, S_c , based in the maximum growth rate criterion:

$$S_o = 2S_c = \frac{4Ae_1 s_{aq}}{DH_v (Ae_1 - T)} \quad (23)$$

where T is the formation temperature; Ae_1 is the eutectoid temperature; s_{aq} is the interfacial energy per unit area of the ferrite-cementite lamellar boundary in pearlite, and DH_v is the change in enthalpy of transformation per unit volume.

However, when the partitioning of the substitutional alloying elements is substantial during the growth of pearlite, boundary diffusion of the alloying elements may control the growth rate of pearlite. In that case, the maximum growth rate criterion of Zener gives an expression for S_o as a function of the pearlite formation temperature as follows:

$$S_o = \frac{3Ae_1 s_{aq}}{DH_v (Ae_1 - T)} \quad (24)$$

Figure 5 shows scanning electron micrographs of pearlite obtained after full decomposition of austenite at 943, 933, 913 and 873 K. The S_o value at these temperatures has been measured from micrographs in Fig. 5. Figure 6 represent the variation of the interlamellar spacing as a function of the temperature formation. The s_{aq} value is derived from equation (23) or (24) depending on the rate controlling mechanism that occurs at each temperature. A value of $DH_v = 6.7 \cdot 10^8 \text{ J m}^{-3}$ [27] has been considered in the determination of s_{aq} . In this sense, a value of $s_{aq} = 0.68 \pm 0.06 \text{ J m}^{-2}$ has been achieved. This value is consistent with that attained by Kirchner et al. [30] ($s_{aq}=0.6 \text{ J m}^{-2}$) for Fe-C-Mn steels.

Finally, in order to determine G_{aq}^* , a value of $\cos\gamma=0.08$ in equation (19) has been obtained from the interfacial energies reported by Reed and Bhadeshia [31] ($s_{aq}=0.1 \text{ J m}^{-2}$) and the above derived s_{aq} ($s_{aq}=0.68 \text{ J m}^{-2}$). Figure 7 shows a comparison between calculated and experimental t^* values for the studied steel. From this figure it can be concluded that a good agreement between experimental and predicted values of t^* exists.

8- Modelling of austenite-to-pearlite transformation

Puls and Kirkaldy [32], in their review on the pearlite reaction showed the following expression for the pearlite growth velocity based on Hillert's theory when the growth rate of pearlite is controlled by the bulk diffusion of carbon in austenite ahead of the interface or NPLE condition [33]:

$$G_{NPLE} = \frac{D_C^g (x_1^{ga} - x_1^{gg})}{g (x_1^{ga} - x_1^{ag})} \frac{S_o}{S_a S_q} \frac{\dot{\epsilon}}{\dot{\epsilon}} - \frac{S_C}{S_o} \frac{\dot{u}}{\dot{u}} \quad (25)$$

where g is a geometric factor equal to 0.72; D_C^g is the carbon diffusion coefficient in austenite; S_c is the theoretical critical spacing at zero growth rate; S_q and S_a are the thickness of cementite and ferrite lamellae, respectively. The ratio between S_q and S_a was assumed to be 7.

The pearlite growth in the partitioned reaction may be controlled by the alloying element interface diffusion (boundary diffusion model) [34]. The growth rate, in that case is expressed as follows:

$$G_{LE} = 12K^P D_B^g d \frac{(x_2^{ga} - x_2^{gg})}{x_2} \frac{1}{S_a S_q} \frac{\dot{\epsilon}}{\dot{\epsilon}} - \frac{S_C}{S_0} \frac{\dot{u}}{\dot{u}} \quad (26)$$

where K^P is the boundary segregation coefficient calculated according to equation (17); D_B^g is the boundary diffusion coefficient of substitutional alloying element, i.e. Mn; and d is the thickness of the boundary. Assuming that the activation energy for boundary diffusion of Mn is the half of that for self-diffusion and the boundary thickness is 0.25 nm, $D_B^g d$ can be expressed as $12.5 \cdot 10^{-14} \exp(-139108/RT)$ in $m^3 s^{-1}$ [35].

The overall transformation kinetics of pearlite was described by Johnson-Mehl-Avrami theory, and here is adapted for hemispherical particles of radius r_p nucleating at the ga_p boundaries at a rate I after incubation time t . The particles grow with a constant rate G_{LE} or G_{NPLE} depending on the temperature range studied. In order to determine the volume fraction of pearlite formed at a given temperature, a series of planes parallel to the boundary and spaced a distance dy apart are considered. If the radius of a particle exceeds the distance y of a plane from the boundary, then the area of intersection of that particle with the plane is determined as Fig. 8 illustrates. The total of such areas of intersection on one plane for all particles growing from the boundary is the *extended area* of transformation on that plane.

If the area of intersection is $\rho r_p^2 = \rho(G^2(t - t)^2 - y^2)$, the change in extended area of pearlite on one plane due to particles emanating from one boundary in the time interval from t to $t+dt$ can be expressed as:

$$dO_e = \rho O_b I [G^2(t - t)^2 - y^2] dt \quad (27)$$

for $r_p > y$, otherwise $dO_e = 0$. The O_b is the total area of the plane. Assuming that I is constant, substituting $f = y/Gt$, and integrating over all incubation times, from $t=0$ to $t=t-(y/G)$, the total extended area of pearlite on one plane is obtained:

$$O_e = \frac{\rho}{3} O_b I G^2 t^3 (1 - 3f^2 - 2f^3) \quad (28)$$

The actual area O_p which intersects the plane O_b will be smaller than the extended area, since the extended area includes a fraction $[1 - (O_p/O_b)]$ of 'phantom' area which has already transformed to pearlite. The relationship between the extended area and the actual area O_p is then given by [36]

$$\frac{O_e}{O_b} = \ln \frac{O_b}{O_p} - \frac{O_p}{O_b} \quad (29)$$

If it is assumed that there is no interference from particles emanating from other g/a_p boundaries, then the total volume of pearlite originating from one boundary, V_b , can be calculated by integrating the actual area over all the planes y . Thus,

$$V_b = O_b G t \int_0^1 \exp\left\{-\frac{\rho}{3} I G^2 t^3 (1 - 3f^2 - 2f^3)\right\} df = O_b G t f(G, I, t) \quad (30)$$

If V is the total volume of the assembly and S_V the g/a_p boundary surface area per unit volume, the total *extended* volume transformed from all boundaries (V_e) can be calculated from the following equation not considering the hard impingement or overlapping of regions emanating from different boundaries:

$$\frac{V_e}{V} = S_V G t f(G, I, t) \quad (31)$$

This can be converted into the actual volume of pearlite, V_p , using the following equation (related to equation (31))

$$\frac{V_p}{V_p^{EQ}} = 1 - \exp\left\{-\frac{S_v}{V_p} G t f(G, I, t)\right\} \quad (32)$$

where V_p^{EQ} is the maximum amount of pearlite obtained after the complete isothermal decomposition of austenite calculated according to equation (16). It follows that the extent of the pearlite reaction, x , is therefore:

$$x = \frac{V_p}{V_p^{EQ}} = 1 - \exp\left\{-\frac{S_v}{V_p} G t f(G, I, t)\right\} \quad (33)$$

The $g a_p$ boundary area per unit volume, S_v , can be estimated assuming that proeutectoid ferrite nucleates at the austenite grain surface under the condition of site saturation and grow within the austenite grains. Such assumption gives [37]

$$S_v = S_g (1 - F)^{2/3} \quad (34)$$

where S_g is the austenite grain boundary per unit volume; F is the volume fraction of proeutectoid ferrite transformed before pearlite reaction starts. Assuming that austenite grains have a tetrakaidehedra geometry [36], $S_g = 3.35/d_g$ where d_g is the austenite grain diameter.

The nucleation rate of pearlite on $g a_p$ boundary per unit area (I) has been calculated as reported by Reed and Bhadeshia [31]. The values of the parameters used in the calculation of pearlite volume fraction as a function of time at 913 and 873 K are listed in Table 2.

Figure 9 shows the experimental and predicted evolution of V_p during the isothermal decomposition of austenite at 913 and 873 K, assuming either full partition of alloying elements (boundary diffusion) or carbon partition (bulk diffusion) between austenite and pearlite. It could be concluded from this figure that partitioning of manganese is the dominant mechanism controlling pearlite growth during isothermal decomposition of austenite at 913 K, whereas partitioning of carbon occurs at 873 K.

9- Conclusions

1. The kinetics of austenite-to-pearlite transformation has been described in a wide temperature range for a 0.4C-1.6Mn forging steel. The proposed kinetic model successfully considers the change in pearlite growth mechanisms at low and high supersaturation.
2. The transition temperature between LE and NPLE growth mechanisms has been theoretically determined at 893 K for a 0.4C-1.6Mn. This result is consistent with experimental measurements of this temperature carried out by Tewari and Sharma [23] in a similar steel.
3. The onset of pearlite transformation occurs when carbon concentration of saturated austenite falls into the denominated Hultgren's extrapolation of the phase diagram, and simultaneously, the g_{a_p} interface progresses at a rate lower than G_{ag}^* to allow cementite precipitation on a moving interface. The mathematical procedure presented in this work successfully predicts the incubation time for pearlite formation.
4. The overall transformation kinetics of pearlite described by Johnson-Mehl-Avrami theory has been successfully adapted for hemispherical particles nucleating at the g_{a_p} boundaries. The assumption of pearlite growth under LE at temperatures above 898 K and under NPLE at temperatures below has been confirmed by means of the good agreement between experimental and calculated results in the studied steel.
5. The model presented in this work is based on physical and metallurgical principles of phase transformations. Although the proposed model has only been validated for a 0.4C-1.6Mn steel, in principle this model is able to predict the isothermal decomposition of austenite in pearlite for a wide range of steels. This model is different to those empirical and semiempirical models created by fitting equations to experimental data.

Acknowledgements

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Appendix

The thermodynamic parameters for this work are taken from Uhrenius' tabulation [38] which has been adapted to a sub-regular solution model.

$$DG_0^{gr} = 8933 - 14.41T + 12.08 \cdot 10^{-3}T^2 - 11.51 \cdot 10^{-6}T^3 + 5.23 \cdot 10^{-9}T^4 \quad (T < 1000 \text{ K})$$

$$DG_0^{gr} = 71659 - 216.8T + 24.77 \cdot 10^{-2}T^2 - 12.66 \cdot 10^{-5}T^3 + 24.4 \cdot 10^{-9}T^4 \quad (T \geq 1000 \text{ K})$$

$$DG_2^{gr} = -20520 + 4.088T + 1500S_{mag}^a$$

$$DG_1^{gr} = -65562 + 32.949T$$

$$DG_0^{gr} = 1.332 \cdot 10^4 - 64.718T + 7.481T \ln T - G^{gr}$$

$$DG_2^{gr} = -14263 + 10T - G^{gr}$$

The values of S_{mag}^a and G^{gr} are, respectively,

$$S_{mag}^a = -11.91 \cdot 10^{-4}T + 8.27 \cdot 10^{-6}T^2 - 15.1 \cdot 10^{-9}T^3 + 12.86 \cdot 10^{-12}T^4 \quad (T < 1075 \text{ K})$$

$$S_{mag}^a = 208.24 - 36710/T - 23.973 \ln T \quad (1075 \text{ K} \leq T \leq 1500 \text{ K})$$

$$S_{mag}^a = 7.87 - 4.18 \cdot 10^{-4}T \quad (1500 \text{ K} > T)$$

$$G^{gr} = 15383.3 - 6.402T$$

The Wagner interaction coefficients for ferrite and austenite phases are listed as follow

$$e_{11}^g = 4.786 + 5066/T$$

$$e_{11}^a = 1.3$$

$$e_{22}^g = 2.406 - 175.6/T$$

$$e_{22}^a = 3.082 - 4679/T + 1509.8 S_{mag}^a / T$$

$$e_{12} = - 4811/T$$

Finally, the Fe-Mn interaction coefficient is $w_{02} = 8351 - 15.19T$

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Figure captions

Fig. 1- A_{e3} and A_{e1} critical temperatures marked on the cooling segment of a dilatometric curve obtained by continuous cooling at 0.05 K/s.

Fig. 2- Schematic equilibrium diagram for pearlite transformation showing the extrapolated phase boundaries (Hultgren's extrapolation).

Fig. 3- Calculated (a) isopleth of the Fe-C-Mn system (Mn content of 1.6 wt.-%), and (b) evolution of pearlite volume fraction under LE and NPLe.

Fig. 4- Calculated K^P coefficient (a) as compared with experimentally obtained in a 0.7C-1.08Mn eutectoid steel, and (b) in the studied steel.

Fig. 5- SEM images obtained after full decomposition of austenite at (a) 943 K, (b) 933 K, (c) 913 K, and (d) 873 K.

Fig. 6- Evolution of S_0 as a function of the undercooling below A_{e1} (DT).

Fig. 7- Comparison between calculated (open circles) and experimental (black filled squares) t^* values.

Fig. 8- Schematic illustration of semi-spherical particles growing from a a/g boundary and intersecting a parallel plane a distance y form the boundary.

Fig. 9- Experimental and predicted evolution of V_P during isothermal decomposition of austenite at (a) 913 K and (b) 873 K.

Tables

Table 1. Calculated values of a_g , DG_v , $\chi(q)$, D_C^g , W and a_1 parameters .

T , (K)	a_g mm	DG_v , J m ⁻³	$\chi(q)$	D_C^g , m ² s ⁻¹	W	a_1 , m s ^{-1/2}
913	3.626	$2.5 \cdot 10^7$	0.016	$12.0 \cdot 10^{-14}$	0.60	$0.4 \cdot 10^{-7}$
873	3.622	$5.8 \cdot 10^7$	0.023	$5.8 \cdot 10^{-14}$	0.70	$1.2 \cdot 10^{-6}$

Table 2. Values of I_b , S_V , G_{NPLE} and G_{LE} at the two tested temperatures.

T , K	G_{NPLE} , m s ⁻¹	G_{LE} , m s ⁻¹	I , m ⁻² s ⁻¹	S_V , m ⁻¹
913	$11.1 \cdot 10^{-6}$	$1.2 \cdot 10^{-6}$	$1.93 \cdot 10^7$	32564
873	$13.3 \cdot 10^{-6}$	$1.5 \cdot 10^{-6}$	$5.8 \cdot 10^6$	35602

Figures

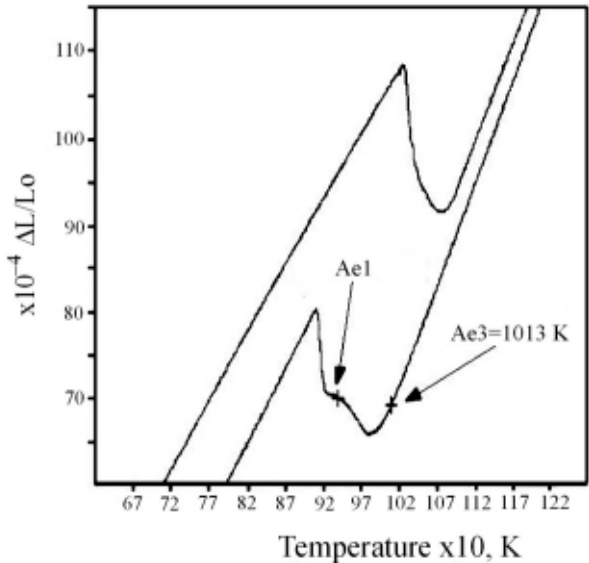


Figure 1

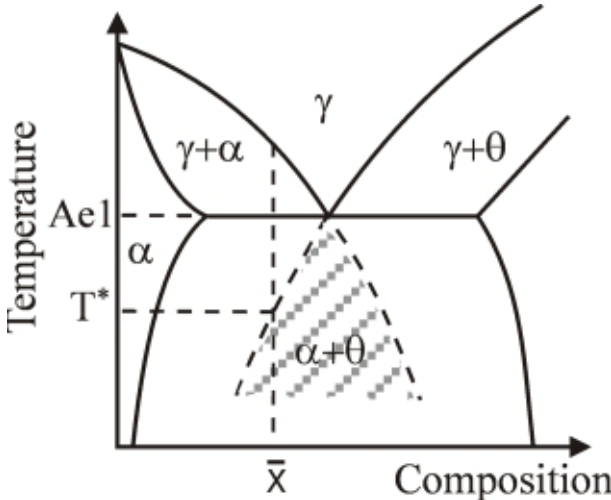


Figure 2

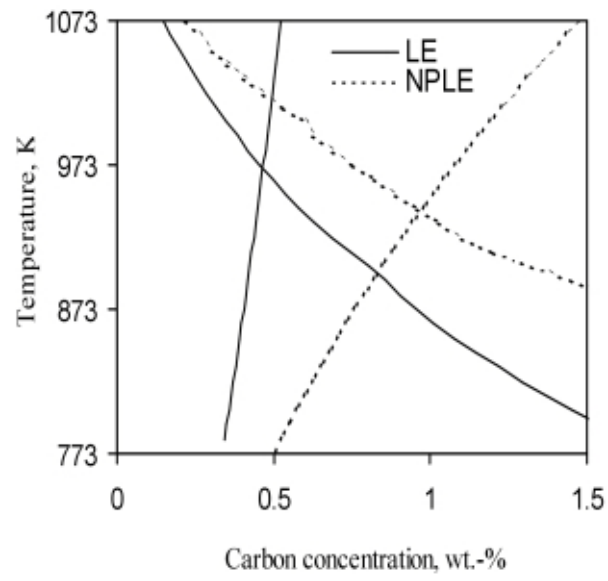


Figure 3(a)

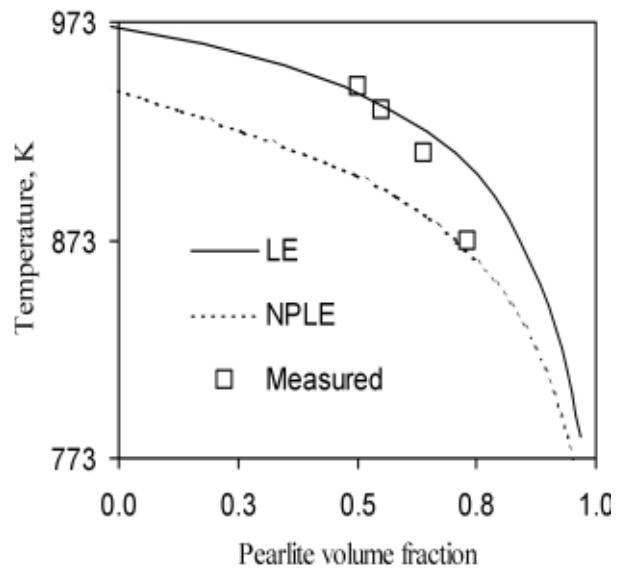


Figure 3(b)

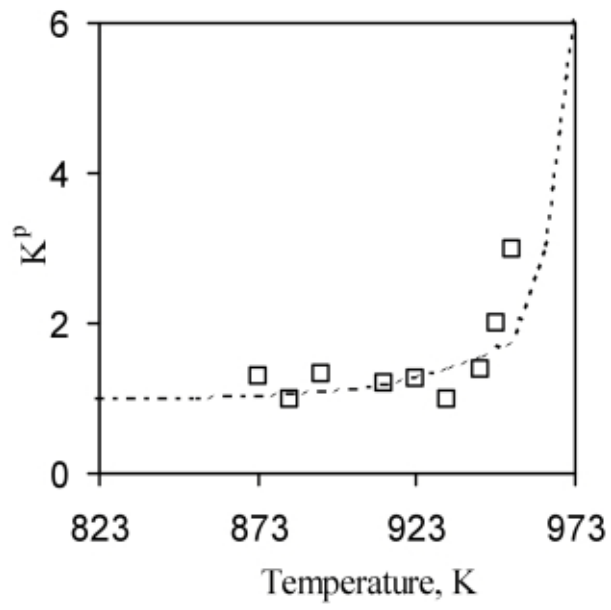


Figure 4(a)

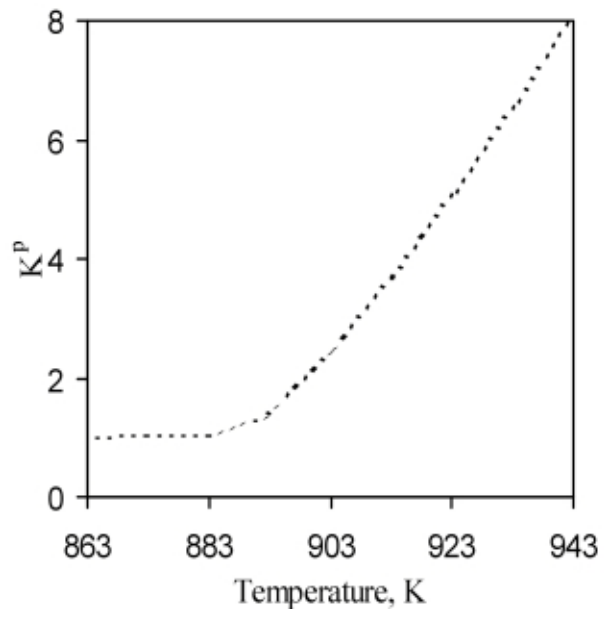


Figure 4(b)

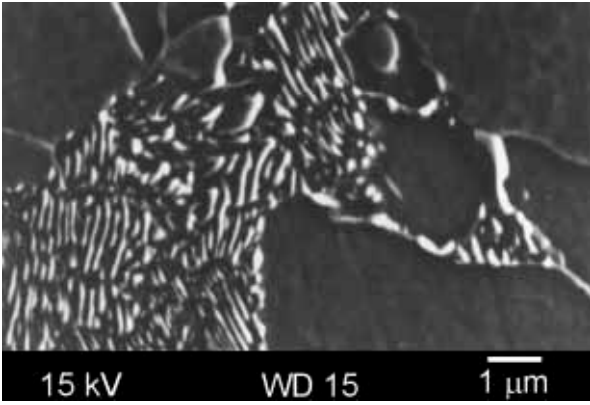


Figure 5(a)

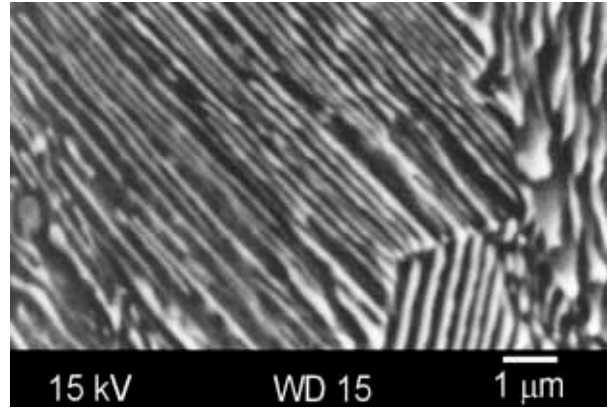


Figure 5(b)

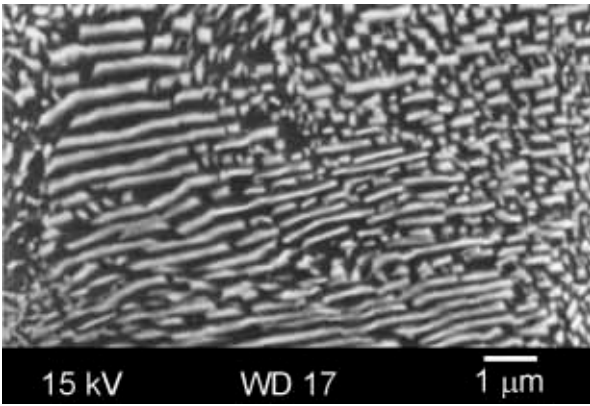


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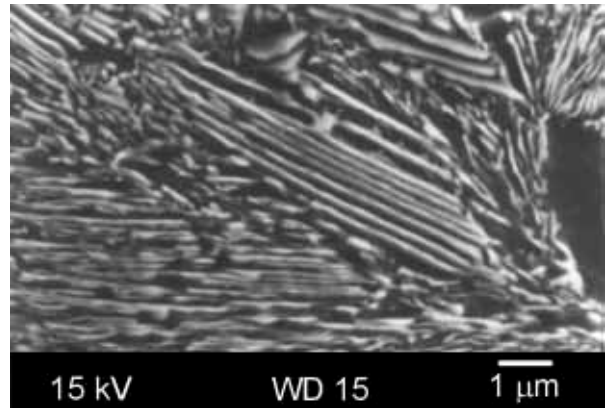


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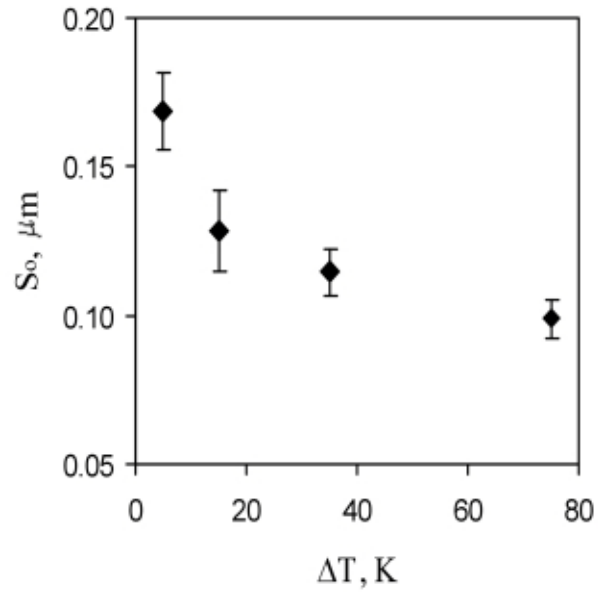


Figure 6

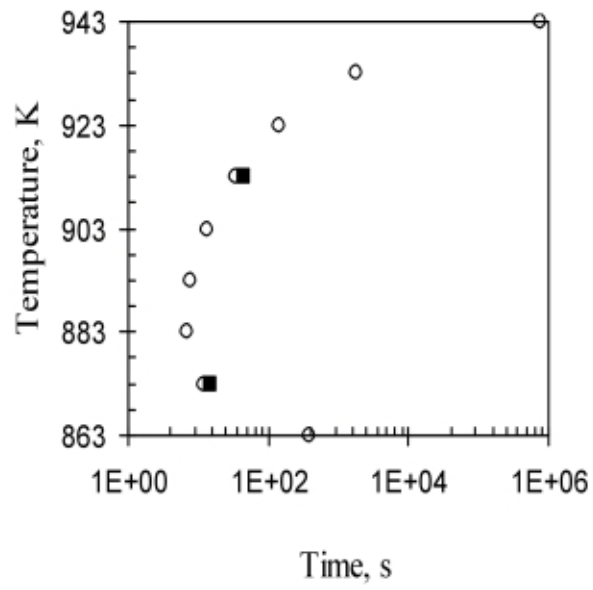


Figure 7

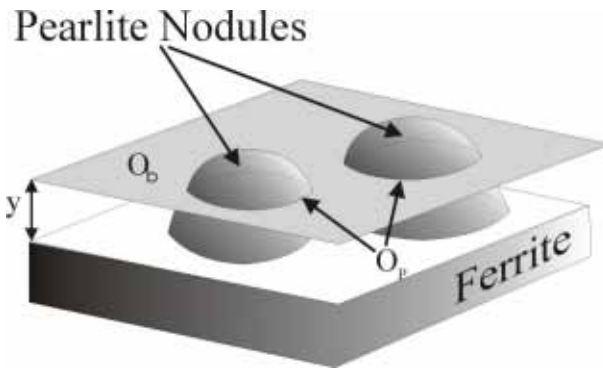


Figure 8

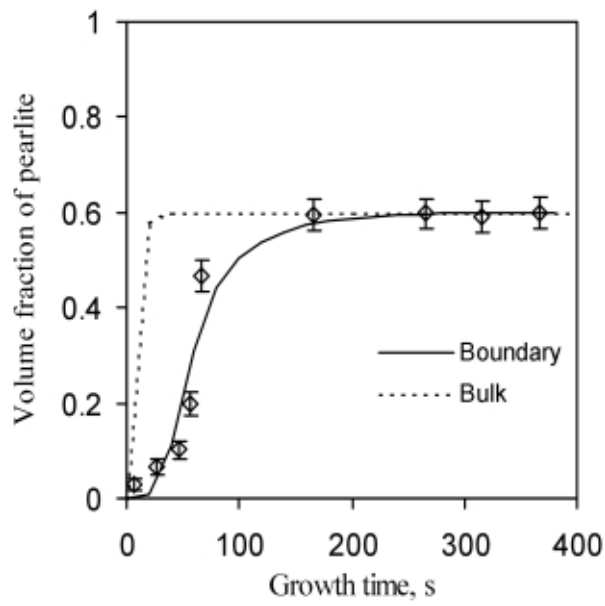


Figure 9(a)

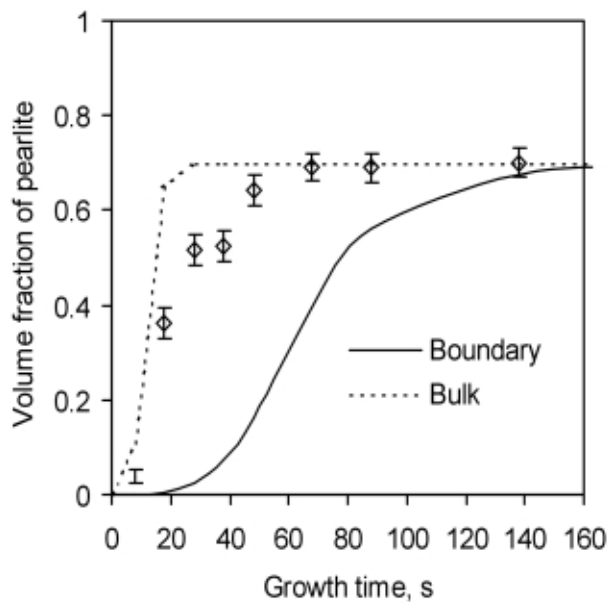


Figure 9(b)