Martensite’s Logistic Paradigm

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Received: August 14, 2020; Revised: December 28, 2020; Accepted: April 01, 2021

This work introduces a deterministic approach to the martensite transformation curve. Martensite is a nucleation-controlled transformation that has two characteristics: autocatalysis and auto-accommodation. Only a small number of martensite units initially form owing to primary nucleation. These new units may cause the transformation of other units by autocatalysis. We call this kind of transformation chained autocatalysis. Moreover, as the transformation progresses, the auto-accommodation influences the arrangement of new units. This work assumes that the transformation-saturation relates to the exhaustion of the chained autocatalysis, which underlines the microstructure. To compare, we considered the KJMA’s extended-transformation concept that implies assuming exhaustion by impingement. Data from isothermal martensite transformations and anisothermal martensite transformations are used to validate the model. Those data comprised different grain sizes and carbon contents. The model is based upon Verhulst’s logistic concept. We propose that the model’s high fitting-capability stems from its deterministic aspect combined with martensite’s self-similarity. Additionally, we suggest that chained autocatalysis controls the rate of martensite transformation. Therefore, the relaxation of transformation strains by plasticity assisted by mutual accommodation determines the transformation’s martensite volume in the absence of post-propagation coarsening/coalescence.

Keywords: Martensitic transformations, microstructure, logistic equation, analytical methods, Avrami’s equation.

1. Introduction

The transformation curve, that is, the volume fraction transformed, \( V_f \), against time, \( t \), is a tool in research and process development and industrial operations. Modeling the transformation curve is an issue that has been studied for decades. In the late thirties and early forties of the last century, Kolmogorov, Johnson-Mehl, and Avrami1-5, KJMA, published seminal papers on this subject. KJMA used a geometrical model to obtain transformation curves. KJMA supposed that the growing regions were spherical, that their growth rate was a constant, that the nuclei were uniform randomly located in space, and that the nucleation took place in two ways: site-saturation and constant nucleation-rate. Their most important contribution was how to consider impingement6. As Liu et al.7 put it, “KJMA’s model consists of nucleation, growth, and impingement.” KJMA model was generalized in different directions. One direction was to obtain more KJMA-like expressions using mathematically exact methods when nucleation and growth took place in a way distinct from KJMA’s. Recently, Rios and Villa8 used mathematical methods for this purpose. The disadvantage of such an approach is that a limited number of situations admit an exact expression. Another possibility, suggested by Avrami herself, is to employ the well-known “Avrami’s equation,” which is an expression containing two adjustable parameters: \( k \) and \( n \)

\[
V_f = 1 - \exp\left(-k^n\right)
\]

(1)

Focusing the transformations in steels, the present authors have proposed an alternative to Avrami’s equation9-11

\[
V_f = 1 - \exp\left(-V_{fi}\left(\frac{x-x^*}{\xi - \xi^*}\right)^n\right)
\]

(2)

In the equation above, \( \xi \) is an “advance” variable, which in previous works9-11 was equal to temperature, magnetic field, mechanical deformation, and time. The \( \xi^* \) is the first datum in a dataset. \( V_{fi} \) is the integration constant resulting from the process of obtaining Equation 2. We denote this as \( V_{fi} \). This constant is a small volume fraction transformed when the martensite transformation starts. In this work, one uses \( V_{fi} \) as a fitting parameter. \( \xi^* \) and \( k^* \) are also fitting parameters. Throughout the text, one discusses the meaning of these parameters. This equation showed excellent agreement when fitted to transformations ranging from martensite to pearlite10,11.

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However, another way to approach formal kinetics is possible. Abramov’s idea\(^1\) of using Verhulst’s logistic equation\(^3\) as the basis to describe transformation kinetics represents a significant shift of paradigm. For the derivation of the transformation curve, the kinetic ideas are expressed directly by mathematics instead of mediated by the transformation’s geometry, as did KJMA. The purpose of using such an approach for modeling transformations is not new. In 1938, Austin and Rickett\(^4\) took the logistic equation as their starting point to obtain the so-called “Austin-Rickett equation”: \[ \frac{V_f}{1-V_f} = kr^n \] (3)

In 1938 not all KJMA’s papers had already been published. The so-called Austin-Rickett equation is seldom applied today, superseded by KJMA’s developments.

The description/rationalization of the fundamental aspects of martensite transformations\(^5\)–\(^20\) constructed the present understanding of martensite. That is, the martensite is a diffusionless and nucleation-controlled transformation. This understanding has been a particular venue to develop new. In 1938, Austin and Rickett\(^4\) took the logistic equation as their starting point to obtain the so-called “Austin-Rickett equation”:

\[
d N_f(\xi) d (\xi - \xi^*) = \varphi(\Delta) \frac{N_f(\xi)}{(\xi - \xi^*)} \left(1 - \frac{N_f(\xi)}{N_{\text{MAX}}} \right) \tag{5}
\]

where \(\xi\) is the experimental “advance” variable and the subscript “\(V\)” indicates per unit volume of material, \(\xi^*\) is the incubation delay. Since we cannot calculate \(\xi^*\) or \(\varphi(\Delta)\) from first principles, they are treated here as fitting parameters.

Then, acknowledging that transformation curves are usually described in terms of the fraction transformed, \(V_f\), we recall \(N_f(\xi) = V_f(\xi) \sqrt{N_f}(\xi)\) where \(\sqrt{N_f}(\xi)\) is the mean volume of the martensite units. We calculate \(N_{\text{MAX}} = 1/\sqrt{N_{\text{MAX}}}\). Introducing these relationships into Equation 5 includes the influence of the relaxation of the transformation strains, which affects the growth of the martensite units, into the logistic model. The \(\varphi(\Delta)\) refers to this crucial process,

\[
\frac{dV_f(\xi)}{V_f(\xi) \cdot A(V_f(\xi))} = \varphi(\Delta) \frac{d(\xi - \xi^*)}{\xi - \xi^*} \tag{6}
\]

where \(A(V_f(\xi)) = 1 - \sqrt{\frac{N_{\text{MAX}}}{\sqrt{N_f}(\xi)}}V_f(\xi)\). Equation 6 can be integrated by separation, however an exact expression for \(A(V_f(\xi))\) is not available. Thus, we considered two approximations. The invariance of the mean martensite units proposed by Magee\(^25\) and the KJMA’s approach assumes transformation in extended space\(^1\)–\(^5\). In the first case \(A(V_f(\xi)) = 1 - V_f(\xi)\), so that the integration of Equation 6 yields a formal analog of the “Austin-Rickett equation,”

\[
V_f(\xi) = V_f(\xi^*)(\frac{\xi - \xi^*}{\xi_i - \xi^*})^{\varphi(\Delta)} \tag{7}
\]

where \(\xi_i\) refers to the value of \(\xi\) at the beginning of the transformation detected in the experimental dataset. We suppose \(V_f(\xi_i) << 1\), what is reasonable in the absence of an initial transformation-burst. To use KJMA’s impingement-correction, we set \(A(V_f(\xi)) = 1\) and substitute the extended-volume fraction transformed, \(V_{\text{TE}}(\xi)\), for \(V_f(\xi)\) into Equation 6, where subscript “\(E\)” signals “extended.” The integration of Equation 6 in extended space yields

\[
V_{\text{TE}}(\xi) = V_{\text{TE}}(\xi^*)(\frac{\xi - \xi^*}{\xi_i - \xi^*})^{\varphi(\Delta)} \tag{8}
\]
Then, using KJMA’s well-known equation, 

\[ V_T = 1 - \exp\left(-\frac{V_{TE}(\xi)}{\xi_I - \xi_T}\right) \]

one also obtains

\[ V_T(\xi) = 1 - \exp\left(-\frac{V_T}{\xi_I - \xi_T}\right)^{\phi(\xi)} \]  

(9)

that is analog to Equation 2.

Summing up, we have obtained two logistic equations where autocatalytic nucleation advances the transformation, but the volume fraction transformed depends on the relaxation of the transformation strains. Noteworthy the parameter \( \phi(\xi) \) refers to the relaxation of the transformation strains which influences the growth of the martensite units, whereas the transformation exhaustion described by \( A(V_T(\xi)) \) depends on the arrangement of the martensite in the austenite grains and the spread of the transformation over the austenite grains.

3. Experimental Data

As in the previous work, we imported databases from papers found in peer-reviewed scientific journals to validate the proposed equations. To fit the analytical expression to the experimental values, one calculated the sum-of-squares, \( \Sigma SQ \), between experimental and calculated values of \( V_T(\xi) \). The \( \Sigma SQ \) gives a “global” idea of the fitting quality. One may also define the relative distance, \( \delta \), between the experimental data and the analytical solution predictions

\[ \delta(\%) = \frac{\left| V_{T\exp}(x) - V_{T\delta}(x) \right|}{V_{T\exp}(x)} \times 100 \]  

(10)

where \( V_{T\exp}(x) \) means volume fraction imported from experimental data and the \( V_{T\delta}(x) \) means the volume fraction predicted by the analytical equations. As already established, experimental procedures may be subject to errors. One can consider a reasonable error of 5% for metallurgical experiments. Regarding the error of 5%, Tables 1-5 show the percentage of the number of points below the error of 5%. This number can help to give a quantitative basis for the fitting besides \( \Sigma SQ \) and visual inspection.

4. Isothermal Transformation

Table 1. Fitting Parameters - FeNiMn - Isothermal Martensite.

| \( T \), K | \( \phi(T) \) | \( V_{T1} \) | \( \tau, \) s | \( \Sigma SQ \) | \( \delta < 5\% \) | \( \phi(T) \) | \( V_{T2} \) | \( \tau, \) s | \( \Sigma SQ \) | \( \delta < 5\% \) |
|----------|-------------|-----------|------------|-------------|----------------|-------------|-----------|------------|-------------|----------------|
| 77       | 2.8         | 2.3x10^4  | 2.8x10^2  | 6.6x10^5   | 79.8\%         | 1.7         | 8.1x10^4  | 6.9x10^2  | 4.6x10^5   | 58.3\%        |
| 133      | 1.7         | 6.4x10^4  | 2.4x10^2  | 1.5x10^5   | 71.4\%         | 0.9         | 6.8x10^3  | 2.6x10^2  | 4.1x10^5   | 53.6\%        |
| 143      | 1.5         | 5.5x10^4  | 1.7x10^2  | 1.5x10^5   | 75.0\%         | 0.7         | 2.7x10^4  | 1.7x10^2  | 6.8x10^5   | 61.9\%        |
| 163      | 1.3         | 1.1x10^4  | 7.5x10^2  | 4.9x10^5   | 72.6\%         | 0.5         | 5.8x10^3  | 7.6x10^2  | 1.0x10^5   | 65.5\%        |
| 173      | 1.5         | 3.3x10^4  | 7.5x10^2  | 4.7x10^5   | 72.8\%         | 0.7         | 3.6x10^3  | 7.7x10^2  | 1.0x10^5   | 66.7\%        |
| 193      | 2.5         | 2.1x10^4  | 1.9x10^2  | 8.5x10^5   | 35.7\%         | 2.6         | 1.5x10^3  | 1.9x10^2  | 2.6x10^5   | 19.6\%        |
| 203      | 2.6         | 4.9x10^4  | 3.4x10^2  | 3.3x10^5   | 54.1\%         | 2.8         | 2.5x10^3  | 3.4x10^2  | 3.2x10^5   | 54.1\%        |

Table 2. Fitting Parameters - FeCrNi maraging steel - Isothermal Martensite.

| Field (kOe) | \( \phi(T) \) | \( V_{T1} \) | \( \tau, \) s | \( \Sigma SQ \) | \( \delta < 5\% \) | \( \phi(T) \) | \( V_{T2} \) | \( \tau, \) s | \( \Sigma SQ \) | \( \delta < 5\% \) |
|-------------|-------------|-----------|------------|-------------|----------------|-------------|-----------|------------|-------------|----------------|
| 0           | 3.5         | 2.0x10^4  | 1.2x10^2  | 9.5x10^5   | 75.0\%         | 2.8         | 4.0x10^3  | 1.2x10^2  | 5.5x10^5   | 100.0         |
| 20          | 2.7         | 3.2x10^4  | 9.3x10^2  | 5.7x10^4   | 95.5\%         | 1.9         | 1.6x10^3  | 1.3x10^2  | 2.7x10^4   | 72.7          |
| 40          | 2.1         | 1.3x10^4  | 7.6x10^2  | 1.0x10^4   | 100.0\%        | 1.4         | 4.7x10^3  | 1.2x10^2  | 2.2x10^4   | 72.2          |
| 60          | 2.1         | 2.5x10^4  | 2.8x10^2  | 2.4x10^4   | 75.0\%         | 1.6         | 1.1x10^4  | 2.7x10^2  | 5.2x10^5   | 100.0         |
| 90          | 1.7         | 2.5x10^4  | 3.3x10^3  | 6.7x10^3   | 60.9\%         | 1.4         | 1.3x10^3  | 3.2x10^2  | 7.2x10^3   | 95.7          |

Table 3. Fitting Parameters - FeMnSiMo - Athermal Martensite.

| \( D, \) mm | \( T, \) K | \( \tilde{\phi}_G \) | \( V_{T1} \) | \( \tau^*, \) K | \( \Sigma SQ \) | \( \delta < 5\% \) | \( \tilde{\phi}_G \) | \( V_{T1} \) | \( \tau^*, \) K | \( \Sigma SQ \) | \( \delta < 5\% \) |
|-------------|--------|----------------|-----------|---------------|-------------|----------------|----------------|-----------|---------------|-------------|----------------|
| 0.185       | 628    | 4.0           | 2.3x10^4  | 646.6         | 3.5x10^5   | 83.3\%         | 1.4         | 2.8x10^3  | 628.1        | 4.9x10^5   | 69.7\%        |
| 0.067       | 628    | 3.1           | 4.5x10^4  | 643.5         | 8.9x10^4   | 87.7\%         | 1.3         | 7.1x10^4  | 623.1        | 8.1x10^5   | 58.5%         |
| 0.025       | 638    | 3.6           | 4.5x10^4  | 643.5         | 1.8x10^4   | 79.4\%         | 1.3         | 5.3x10^4  | 613.1        | 1.1x10^5   | 39.7%         |
| 0.015       | 608    | 2.7           | 2.2x10^4  | 613.2         | 2.4x10^4   | 88.7\%         | 1.0         | 7.0x10^4  | 607.8        | 1.2x10^5   | 59.7%         |
| 0.006       | 608    | 3.3           | 6.3x10^4  | 613.2         | 1.2x10^4   | 83.6\%         | 1.4         | 4.5x10^4  | 607.8        | 1.7x10^4   | 47.5%         |
For isothermal martensite transformation, one replaces time for the advancing variable in the equations of the previous section, so Equation 7 becomes

$$V_T(t) = V_{VI}(\tau - T_{t})^\delta G$$

where $\phi$ is the proportionality factor, $E_a$, and $\Delta Ga$ are apparent energies, $T$ is the reaction temperature and $k_B$ is the Boltzmann constant. The charts in Figure 3 yield $E_a \approx 5$ kJ/mol - 13 kJ/mol, which is compatible with dislocation processes, and $\Delta Ga \approx 0.9 - 1.3$ kJ/mol, which has the same magnitude as the elastic free-energy (0.9 kJ/mol) of an oblate spheroid with a typical 0.05 aspect-ratio in a constrained matrix. The FeNiMn isothermal martensite undergoes a substructure change at low transformation temperatures. Thus, we propose that the variation in $\phi(T)$ refers to changes in the relaxation of transformation strains. The variation in the $V_{VI}$ corroborate the variation in $\phi(T)$. However, the variations in $1/\tau$ show the opposite trends, see Figure 4. Such specific behavior point to differences in the martensite propagation. Martensite propagation at incubation depends on the probability that austenite defects sustain coordinated atomic groups to cross the nucleation path. By contrast, the nucleation’s post-incubation is determined by a previously formed martensite barrier for converting such groups into nuclei. Hence, the chemical driving force controls the incubation. Instead, at low temperatures (higher driving forces), a thermal barrier controls the martensite incubation/nucleation. In this regard, it is noteworthy that the apparent activation energy obtained from the incubation time, ~ 6 kJ/mol, compare with the ~ 5 kJ/mol obtained from the parameter $\phi(T)$, which refers to the accommodation of the shape strain at high transformation temperatures. This comparison says that dislocation processes are present in both processes (relaxations of lattice-misfit and the shape strain). At this time, the analysis of the temperature variation in the parameter $V_{VI}$ was not conclusive.

Lastly, mind that impingement of martensite on the austenite grain boundary generates a stress-field capable of fostering martensite propagation into the next grain. However, such an “intergrain-spread” is hindered if the austenite plasticity hails the radial propagation of a martensite unit. Such a possibility is comparable to “soft-impingement.”

### Table 4. Fitting Parameters - FeCrNi - Athermal Martensite.

| wt%C | $T_i$, K | $\bar{\phi}_G$ | $V_{VI}$ | $\tau^*$, K | $\Sigma SQ$ | $\delta < 5\%$ | $\bar{\phi}_G$ | $V_{VI}$ | $\tau^*$, K | $\Sigma SQ$ | $\delta < 5\%$ |
|-------|----------|----------------|----------|------------|-------------|----------------|----------------|----------|------------|-------------|----------------|
| 0.002 | 290      | 3.7            | $1.2\times10^4$ | 339.7      | 7.0x10^4     | 88.9           | 1.3            | 8.6x10^4 | 299.2      | 8.1x10^4     | 93.3          |
| 0.020 | 265      | 4.0            | $3.7\times10^2$ | 304.7      | 3.6x10^4     | 96.9           | 1.5            | 6.4x10^4 | 267.8      | 8.8x10^4     | 96.9          |
| 0.050 | 254      | 5.5            | $1.2\times10^2$ | 304.1      | 2.3x10^4     | 88.9           | 2.4            | 8.0x10^4 | 265.4      | 4.1x10^4     | 77.8          |
| 0.100 | 193      | 6.1            | $4.6\times10^2$ | 242.6      | 1.1x10^4     | 90.0           | 2.8            | 6.6x10^4 | 218.4      | 3.2x10^4     | 90.0          |

### Table 5. Fitting Parameters - Carbon Steels - Athermal Martensite.

| wt%C | $T_i$, K | $\bar{\phi}_G$ | $V_{VI}$ | $\tau^*$, K | $\Sigma SQ$ | $\delta < 5\%$ | $\bar{\phi}_G$ | $V_{VI}$ | $\tau^*$, K | $\Sigma SQ$ | $\delta < 5\%$ |
|-------|----------|----------------|----------|------------|-------------|----------------|----------------|----------|------------|-------------|----------------|
| 0.46  | 586      | 1.8            | $1.7\times10^4$ | 587.5      | 3.0x10^2     | 65.1%          | 1.0            | 8.7x10^4 | 586.3      | 8.5x10^2     | 49.2%         |
| 0.66  | 535      | 2.3            | $1.6\times10^2$ | 546.0      | 1.6x10^2     | 69.7%          | 1.2            | 2.9x10^3 | 535.9      | 4.0x10^2     | 53.0%         |
| 0.80  | 502      | 2.2            | $1.6\times10^2$ | 513.9      | 6.3x10^2     | 84.2%          | 1.2            | 7.9x10^3 | 503.8      | 2.3x10^2     | 52.6%         |
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5. Martensite “Athermal” Transformation

To describe the transformation curve of time-independent, driving-force induced (“athermal”) martensite, one replaces temperature for the advancing variable in Equations 11 and 12 that gives:

\[
\frac{V_f(T)}{1-V_f(T)} = V_{f_1} \left( \frac{T^* - T}{T^* - T_f} \right)^{\Phi_G} \tag{14}
\]

and

\[
V_f(T) = 1 - \exp \left( -V_{f_1} \frac{T^* - T}{T^* - T_f}^{\Phi_G} \right) \tag{15}
\]

where \( T^* \) is the upper temperature for martensite nucleation, and \( T_f \) is the highest experimental temperature in a data set. The variables, \( V_{f_1} \), \( T^* \) and \( \Phi_G \) are fitting parameters. We
quantities seem to be inverted when compared with \(\frac{t - \tau}{t_i - \tau}\). The reason for this is that time increases and temperature decreases. Thus, the terms are inverted so that the subtractions remain positive.

Ref\(^{36}\) describes a database typical of Fe-0.2wt%C, 3.5wt%Mn, 1.5wt%Si, 0.5wt%Mo with different grains sizes (0.006 mm - 0.185 mm). Ref\(^{37}\) provides data on Fe-18wt%Cr, 8wt%Ni, with varying contents of carbon.

In FeMnSiMo, the transformation took place in a dilatometer. In addition to allowing the models’ validation, the database permits to characterize the austenite grain size’s influence on the transformation curve. Bearing scatter in \(T_i\), we expressed \(\tilde{\varphi}_{G} = \lambda T_i\), and fitted \(\lambda\) until the sum-of-squares, \(\Sigma SQ\), between experimental and calculated values of \(V\) became invariant, see Figure 5. Table 3 lists the values of the obtained parameters of Equations 14 and 15. Inspection of the values of \(\Sigma SQ\) indicates that Equation 14 provides the best fittings with a minor variation in \(\Sigma SQ\). By contrast, the values of \(\Sigma SQ\), which characterize the fittings with Equation 15, increase with increasing the austenite grain-size.

A similar procedure was used to fit the Fe-18wt%Cr, 8wt%Ni data. The results are shown in Figure 6 and Table 4. The fit is excellent.

Concentrating on the FeMnSiMo, Figures 7 and 8 show the values of \(\tilde{\varphi}_{G}\) and \(T^*\) graphed after Hall-Petch, complying with the experimental variable (austenite grain size). The \(T_i\) estimates the martensite start temperature, \(M_S\). Observe that both fittings yielded decreasing values of \(\tilde{\varphi}_{G}\) and \(T^*\), with decreasing the austenite grain size down to 0.015 mm, then upshifts. The decreasing in \(\tilde{\varphi}_{G}\) suggests the influence of the austenite strength on the martensite propagation, possibly related to the mobility of the martensite-austenite interfaces or dislocation processes\(^{38}\). The reported\(^{36}\) coarser aspect ratio of the martensite units formed in the material with 0.006 mm grain size is coherent with a heightened variant-selection (auto-accommodation), also acknowledged in ref\(^{39}\). The decreasing values of \(T^*\) imply higher stability in the fine-grained austenite. Thence, the variations in \(\tilde{\varphi}_{G}\) and \(T^*\) are in qualitative agreement with the conclusions in ref\(^{36}\). The high values of \(\Sigma SQ\), typical of the fittings of the dataset with Equation 15, are due to the severe effect of crystallographic-variance in fine-grain austenite. This crystallographic-variance affects the microstructure’s local-randomicity, which is a requirement for utilizing the KJMA’s methodology\(^{1-5}\). Again, the behavior of the parameter \(\delta\) indicated a better agreement between Equations 14 and 15, which assumes exhaustion by nucleation and by impingement, respectively.

Lastly, we consider the influence of the carbon in the martensite, transformed by continuous cooling. Typical plain carbon-steels with similar austenite grain-sizes were considered: Fe46C(0.46wt%C, 0.71wt%Mn, 0.26wt%Si, 0.1wt%Ni, 0.2wt%Cr), Fe66C(0.66wt%C, 0.69wt%Mn, 0.30wt%Si, 0.1wt%Ni, 0.2wt%Cr), and Fe80C(0.80wt%C, 0.61wt%Mn, 0.41wt%Si, 0.2wt%Ni, 0.3wt%Cr). These databases were imported from ref\(^{40}\). The fittings with Equations 14 and 15 are
Figure 5. Comparison of fittings for the FeMnSiMo with different grains sizes, transformed by continuous cooling. (a) 0.006 mm; (b) 0.015 mm; (c) 0.025 mm; (d) 0.067 mm; (e) 0.185 mm. Data extracted from ref. 36. Graphs were constructed and fitted with Equations 14 and 15.

Figure 6. FeCrNi stainless steels athermal transformation during continuous cooling. Data extracted from Ref. 37. Graphs were constructed and fitted with Equations 14 and 15.

Figure 7. Values of $\Theta_G$ graphed after Hall-Petch for the FeMnSiMo.
shown in Figure 9, and the values of the respective model-
parameters are listed in Table 5. Again, visual inspections
of the charts and the variations in $\Sigma SQ$ indicate that the
Equation 14 provided better fittings, especially concerning
the transformation-charts’ progressive induction. These
fittings were consistent with the behavior of the parameter $\delta$.
We ascribe the variation in $T^*$ to the influence of the carbon
on the austenite stability. The variation in $\Phi_C$ is related to the
influence of carbon content on the transformation microstructure
since increasing carbon enhances the partitioning of the
austenite grains into finer packets and blocks$^{41}$.

Like the isothermal transformation curves above analyzed,
the different modes of considering the transformation-
saturation provided proper fittings of the data. Nonetheless,
the values of the fitting parameters are model-dependent,
as might be expected.

6. Discussion

The classical Verhulst’s logistic equation, Equation (4),
proposed to describe constrained population growth, provides
a venue to express transformation curves$^{12}$. Specifically, note
that Equation 6 has two independent terms, $\varphi(\Delta) \frac{d(\xi - \xi^*)}{\xi - \xi^*}$
that refers to the advance of the microstructure, and
$\frac{dV_T(\xi)}{V_T(\xi) \cdot d(V_T(\xi))}$ that refers to the accommodation of
the transformation in the austenite grains. The integrated
Equations 11 and 14 and Equations 12 and 15 only differ
in the specification of the exhaustion factor, $A(V_T(\xi))$. By
adjusting their fitting parameters, suitable descriptions of
the transformation curves are possible.

The similar variations in the parameters $\varphi(T)$ and $\tau$
obtained with Equations 11 and 12 are coherent with the
pertinence of self-similarity in the martensite transformation.
Notably, the variations in the parameters obtained from the
FeMnSiMo database typical of martensite transformation by
cooling (“athermal”) exhibit similarity and are in qualitative
agreement with the results reported in the referenced
paper$^{36}$. Thence, the experimentalists may choose the more
appropriate expression to analyze their data and describe the transformation under consideration. Nonetheless, the meaning of the physical parameters obtained from formal models depends on the models’ premises. We assert that autocatalysis and transformation-saturation by nucleation-exhaustion are realistic premises to model martensite transformation curves as provided by Equations 11 and 14.

It is worth discussing the fitting parameters displayed in Tables 1-5.

First, we would like to offer some background on the use of phenomenological equations and fitting parameters to describe a specific kinetic curve. In the present case, to fit a $V_f(t)$ curve. The first possible approach to describe experimental measurements by an analytical expression is to employ an arbitrary function to fit the experimental curve. This fit may be useful if one has an analytical theory that takes a continuous function as its input, for example, Ref. On another extreme, one may fit an expression derived from fundamental theories. Generally, these are not easy to come by. An intermediary approach is to use the so-called formal kinetics. These provide exact expressions when one specifies the nucleation and growth rates. The pioneering work is, of course, KJMA theory. More recently, Rios and Villa derived several new expressions. Still, the number of exact solutions is limited.

Yet another possibility is to use functions that have some physical or mathematical basis. Such as Avrami’s, see Equation 1. Or Austin-Rickett, see Equation 3. Here we used a generalization of both Avrami and Austin-Rickett equations, containing an extra parameter related to the beginning of the transformation.

From the equation employed here, one expects: I) that they give a good fit; II) that we can extract some information from the fitting parameters. Notice that the functional form is different for Equations 14 and 15. Therefore it comes as no surprise that the absolute value of the fitting parameters differs. Nonetheless, Tables 6, 7, and 8 show that they do not differ by the same magnitude. In the case of Table 6, the differences between the fitting parameters were calculated as follows: (Equation 12 parameter – Equation 11 parameter)/ (mean value of Equation 11 and Equation 12 parameters). The same reasoning was adopted for Tables 7 and 8, but with Equations 11 and 12 replaced by Equations 14 and 15.

The parameters that mark the beginning of the transformation, such as initial transformation temperature and incubation time, are physical parameters. Tables 7 and 8 demonstrates that the values of $\tau$ lie quite close when Equations 14 and 15 determine them. Table 6 shows the values of $\tau$, obtained from Equations 11 and 12, behave similarly but with an apparent discrepancy at the highest and lowest temperatures. The absolute values of the other parameters have a significantly higher difference. This behavior is unavoidable as the proper functions are different. This result suggests that the function

| Table 6. Difference between the fitting parameters shown in Table 1 obtained by Equation 11 and Equation 12 for the FeNiMn - Isothermal Martensite. |
|---|---|---|---|
| $T_i$, K | $\varphi(T)$ Difference | $V_{fi}$ Difference | $\tau$ Difference |
| 77 | 48.9% | 200.0% | -85.6% |
| 133 | 61.5% | 161.3% | -5.0% |
| 143 | 72.7% | 181.4% | -2.1% |
| 163 | 88.9% | -133.9% | -2.0% |
| 173 | 72.7% | -9.0% | -2.0% |
| 193 | -3.9% | 200.0% | -163.6% |
| 203 | -7.4% | 200.0% | -163.7% |

| Table 7. Difference between the fitting parameters shown in Table 3 obtained by Equation 14 and Equation 15 for the FeMnSiMo - Athermal Martensite. |
|---|---|---|---|---|
| $D$, mm | $T_i$, K | $\bar{\varphi}(T)$ Difference | $V_{fi}$ Difference | $T^*$ Difference |
| 0.185 | 628 | 96.3% | 156.6% | 2.9% |
| 0.067 | 628 | 81.8% | -44.8% | 3.2% |
| 0.025 | 638 | 93.9% | -168.7% | 4.8% |
| 0.015 | 608 | 91.9% | 103.4% | 0.9% |
| 0.006 | 608 | 80.8% | 173.3% | 0.9% |

| Table 8. Difference between the fitting parameters shown in Table 5 obtained by Equation 14 and Equation 15 for the Carbon Steels - Athermal Martensite. |
|---|---|---|---|
| wt% C | $T_i$, K | $\bar{\varphi}(T)$ Difference | $V_{fi}$ Difference | $T^*$ Difference |
| 0.46 | 586 | 57.1% | 64.6% | 0.2% |
| 0.66 | 535 | 62.9% | 138.6% | 1.9% |
| 0.80 | 502 | 58.8% | 67.8% | 2.0% |
form of Equations 11, 12, 14 and 15 strongly influence parameters, such as, $V_f$ and $\varphi$
.

One cannot expect Equations 11, 12, 14 and 15 to be more than they are. They are equations with a physical or mathematical background, but they are still approximations. And it is well-known that fitting parameters carry the error made by assuming a certain approximation. But, as shown above, the parameters are not influenced in the same way. Here, parameters that have a direct physical interpretation tend to be almost independent of the fitting expression. By contrast, parameters that are more directly related to the functional form of the fitting expressions tend to have more considerable differences.

7. Conclusions

1. The utilization of the logistic formalism to describe isothermal and continuous cooling martensite transformations yielded quality-fittings of experimental data. These quality-fittings are consistent with current views regarding martensite’s nucleation-controlled, autocatalytic kinetics, and self-similarity.

2. The apparent activation energies obtained from Equation 13, 5 kJ/mol - 13 kJ/mol, compares with the activation energies for martensite nucleation reported in refs[26-27]. Therefore, one may suggest that there are two kinds of active dislocation processes. One dislocation process acts in the conversion of coordinated atomic groups into nuclei. The other, intrinsically different dislocation process relates to the relaxation of the martensite shape-strain[26].

3. The incorporation of self-similarity into Verhulst’s logistic formalism allowed good descriptions of the martensite transformation curves as well as characterizations of kinetic aspects of isothermal or “athermal” transformations.

Acknowledgments

This study was financed in part by the Coordenação de Aperfeiçoamento de Pessoal de Nível Superior - Brasil (CAPES) - Finance Code 001. P. R. Rios, and A. L. M. Alves are also grateful to Conselho Nacional de Desenvolvimento Científico e Tecnológico, CNPQ, and Fundação de Amparo à Pesquisa do Estado do Rio de Janeiro, FAPERJ, for the financial support.

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