Association of Metallurgical Engineers Serbia and Montenegro Scientific paper AME UDC:669.01-122.4-415:620.179.1.001.573=861

## MATHEMATICAL MODEL OF STRUCTURE FORMING IN HOT-ROLLED SHEETS

T. KOINOV<sup>1</sup>, D. ANGELOVA<sup>2</sup>, R. SHATALOV<sup>3</sup>

<sup>1</sup>Associated professor in the University of Chemical Technology and Metallurgy-Sofia Dept. of Physical Metallurgy and Heat Aggregates, 8 Kl. Ohridski Blvd., 1756 Sofia, Bulgaria, e-mail: info@leadert.com, <sup>2</sup>Professor in the University of Chemical Technology and Metallurgy-Sofia, Dept. of Physical Metallurgy and Heat Aggregates, 8 Kl. Ohridski Blvd., 1756 Sofia, Bulgaria, e-mail: dona\_ang@techno-link.com, <sup>3</sup>professor in the Institute "Zvetmetobrabotka" – Moscow, e-mail: lyudmila@cmet.ru

### ABSTRACT

Time necessary for 95% metal recrystallization is modelled for the process of hot-rolling of sheets on a finishing train of continuous mills. The grain size of such a sheet is calculated for the different fractions formed as a result of a gradually accumulating hardening.

A comparison is made between the results of modelling and the corresponding experimental data for some rolled low carbon steels.

**Key words:** Hot-rolling of sheets, Recrystallization, Grain size, Rolled low carbon steel, Modelling of grain size

# **INTRODUCTION**

A mathematical model is created for calculating microstructural changes in a lowcarbon steel sheets exposed to hot rolling. The model is based on torsion investigations of grain size, published by other researchers and own experiments [1] for determination of grain size, carried out on a torsion machine under controlled loading. The material is a carbon steel produced in Kremikovtzi plant.

## MATERIAL AND MECHANISM OF STRUCTURE FORMING

Material tested is a low-carbon steel Bulgarian Standard 2 K $\Pi$  (37.2 DIN) rolled on the finishing train of 1700 mill in Kremikovtzi plant, which consist of 6 continuous stands.

At hot rolling process an increase of deformation  $\varepsilon$  in some stands leads to a point of reaching its critical value,  $\varepsilon_m$ , for initiating nuclei of dynamic recrystallization [2, 3]. Since this moment the stress has become decreasing and the situation characterizes with a sufficient recrystallization. This process continues to establishment of equilibrium after deformation, described by  $\varepsilon_s$ . The equation of Zener-Hollamon presents an existing connection between the parameters of stress  $\sigma$ , strain rate v and metal temperature T by a specific characteristic z in (1), [4, 5]:

$$z = v \exp \frac{Q}{RT} = A \exp \beta \sigma \quad , \tag{1}$$

where Q is activation energy of recrystallization during deformation process, R – universal gas constant, A and  $\beta$  – coefficients characterizing steel nature; and the parameter z expresses the influence of v and T on the energy stored in metal that makes recrystallization possible.

#### DISCUSSION AND MODELLING

Before the first stand (K1) of a finishing train, the semifinished-product grain size d10 is usually accepted as equal to that before the rolling begins – d00. At the next stands it is necessary to calculate the time for metal recrystallization of 95%. So, after metal deformation in a given stand (K), that time named as t95(K) can be presented by (2):

$$t95(K) = [4.14 - 0.0035T_{exit}(K - 1, N)]ln(\sigma - \sigma_0), [s].$$
<sup>(2)</sup>

The size of already recrystallized grain is determined by (3) as a result from the deformation of a given point (N) situated along the semifinished product passing through stand (K):

$$dREX(K,N) = A(K,N) \left[ 14.9 \ln \frac{z}{8.510^9} \right]^{-\frac{2}{3}} \sqrt{\frac{d10(K-1,N)}{\epsilon}} , \ [\mu m], \tag{3}$$

where A(K,N) is a coefficient depending on steel nature at conditions (K), (N). At the same time the equation (3) represents our model of grain growth under classical recrystallization.

After the whole recrystallization of 95%, the grain size grows until the very beginning of deformation in the next stand. The time for this growth - tREX(LK), calculated by (4), is a function of parameters as: distance between stands (LK), roll velocity *V*, and time t95(K) connected with a given stand, in this case (K):

$$tREX(LK) = \frac{LK}{V} - t95(K), [s].$$
<sup>(4)</sup>

At semifinished-product rolling in the first stand (K=K1) of a finishing train with *n* stands, the time for metal recrystallization towards the back end (n >1) increases by the machine time  $\tau_m$  for rolling on the continuous finishing train, (5):

$$tREX = tREX(LK) + \frac{L_n}{\tau_m}, [s], \qquad (5)$$

where  $L_n[m]$  is metal length.

The condition tREX > 0 means that between passes metal is wholly (95%) recrystallised. In such a case the size of grain already grown at a static recrystallization

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between passes d10(K,N) can be calculated. However the investigations show that the grain growth has different rate at metal temperatures over and below  $900^{\circ}C$ .

At  $T > 900^{\circ}C$ , the grain size d10(K,N) is represented by (6):

$$d10(K,N) = dREX(K,N) + 3.8710^{14} tREX \exp \frac{-400}{RT + 273}.$$
 (6)

At  $T \le 900^{\circ}C$ , a condition characterizing continuous mills, the grain size d10(K,N) adopt the form of expression (7):

$$d10(K,N) = dREX(K,N) + 6.0210^{39} tREX \exp \frac{-914}{RT + 273}.$$
(7)

The condition tREX < 0 means that recrystallization between passes does not established wholly. During whichever next pass, a structure is formed that consists of recrystallized grains with sizes of dREX(K,N), and non-recrystallised ones - dREX(K-I,N) still with unchanged sizes equal to those before the previous pass.

This analysis shows that it can be accepted the following statement: the grains dREX(K,N) do not have residual stresses being wholly recrystallized, while the grains dREX(K-1,N) have residual stresses arisen from the previous pass. Then for the latter grains the stress in the rolled metal obtained during the pass (K+1) summarizes with that of the pass (K). Here we adopt the idea that each grain size has been studied independently by itself and not been influenced by the other grains. This approach is simplified and schematic, but there still exist many difficulties for describing structure forming at tREX < 0 including simultaneously both, recrystallized and non-recrystallized grains. It is important to note that at every next pass (under tREX < 0) the number of grains with different sizes doubles, a fact showing formation of complex structures corresponding to different residual stresses. This makes the algorithm for calculating different grain sizes very complicated.

For simplifying calculations it is necessary to make readings of sizes and relative parts taken of four grain-groups at the exit of the finishing train. Consequently, at tREX < 0 the readings made for the different (by size) grain-groups begin two passes before the exit of the train (K=Kn-1). Such a precision is enough for research and practice. Even if there we can find more than four grain-groups under tREX < 0 and before the pass (Kn-1), their corresponding relative parts will be negligible [1, 6].

We mark the size of recrystallized grain at the entrance of (Kn-1) stand as dREXI and the size of non-recrystallized grain after stand (Kn-2) as dREX2. Then we insert the following symbols: X1 for the corresponding relative part of recrystallized fraction dREX1; and X2 for the corresponding relative part of non-recrystallized fraction dREX2. All this results in (8, 9, 10, 11):

$$dREX1 = dREX(Kn - 1, n);$$
(8)

$$dREX2 = dREX(Kn - 2, n);$$
<sup>(9)</sup>

$$X1 = 1 - exp\left(-2.996\frac{LK}{V}t95\right);$$
 (10)

$$X2 = 1 - X1. (11)$$

At the entrance of the last stand of train (Kn) four different grain sizes are calculated. The sizes of recrystallized grains are the already mentioned dREX1, and dREX3 and dREX4 with a common (summarized) corresponding relative part X3; dREX3 forms from dREX1 and dREX4 - from dREX2. Then dREX5 and dREX6 are symbols for the grains formed from dREX1 and dREX2, which are non-recrystallized until the metal enters the last stand. Their corresponding relative part is X4. The corresponding relative parts of the fractions dREX3 - dREX6 are X5-X8 respectively or:

$$X4 = 1 - X3;$$
  
 $X5 = X1X3;$   
 $X6 = X2X3;$   
 $X7 = X1X4;$   
 $X8 = X2X4.$ 

The histograms of grain sizes obtained experimentally and by the model are shown on Fig. 1. The experimental readings are done by automatic quantitative metallographic analysis.



*Fig. 1. Histogram of grain-size distribution for*  $4.0 \times 1250$  *mm sheet of steel*  $2K\Pi$ *: the model data are shown by thick line and the experimental ones – by dash line* 

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### SUMMARY

The proposed model gives us possibilities for investigation of influence of different technological parameters and regimes on metal structure forming, then for control of structure forming during deformation process, and for design of new regimes for controlled sheet rolling.

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