

# BENCHMARKING AND VALIDATION OF HOT ROLLING SIMULATION MODELS

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## ABSTRACT

Mathematical models representing the various stages in the hot rolling process of steel production were produced. Simulated models used were validated using the EXCEL package and then subjected to a sensitivity analysis under identical conditions to predict the optimum operating conditions under which the essential process steps can be carried out.

The results showed that ferrite grain size is more sensitive to the retained strain than to the cooling rate or the austenite grain size. It was also inferred from the work that a drop in the austenite-ferrite ( $\gamma$ - $\alpha$ ) transformation temperature leads to an increase in the tensile strength of the steel. This is in agreement with industrial practices.

**Keywords:** Process modeling, hot strip rolling, mechanical properties, simulation.

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## Notation

Symbol	Meaning	Unit
D	Grain size	Microns
$D_0, d_0$	Initial grain size	Microns
T	Temperature	$^{\circ}\text{C}$
t	Time	Second
MFS, $\sigma$ , $\sigma_n$ , $\sigma_m$	Mean flow stress	MPa
$\sigma_{ss}$	Steady state stress	MPa
$\sigma_s$	Saturation stress	MPa
$\sigma_p, \sigma_{STAT}$	Peak stress	MPa
$\epsilon$	Strain	-
$\epsilon_p$	Peak strain	-
$\sigma_{DYN}$	Dynamic recrystallization stress	MPa
$\epsilon'$	Strain rate	/second
$\epsilon_{dyn}$	Dynamic recrystallization strain	-
$\epsilon_c$	Critical strain	-
$\epsilon_s$	Saturation strain	-
$\epsilon_r$	Retained strain	-
$\sigma_d$	Dislocation strengthening	MPa
$\sigma$ , YS, L.Y.S., $\sigma_y$	Lower yield stress	MPa
$\sigma_y, T_s$	Tensile strength	MPa
Z	Zener-Hollomon parameter	-
X, $X_{rec}$ , F.S., $X_{st}$ , $X_{RX}$	Fraction recrystallized	-
$t_{50}$ , $t_{0.5}$ , $t_{50\%}$ , $t_{X0.5}$	Time for 50% recrystallization	Second
Q	Activation energy	KJ/mol
R	Universal constant	KJ/mol
SRX	Static recrystallization	-

$Q_{RX}, Q_s$	Activation energy for static recrystallization	KJ/mol
$d_{NRX}, d_{RX}, D_s, D_{N1}, D_{M1}$	Static recrystallization grain size	Microns
$S_g$	Grain surface to grain volume ratio	-
$Q_d$	Deformation activation energy	KJ/mol
$D_g$	Grain size after grain growth	Microns
$t_g$	Time allowed for grain growth	Second
$Q_{gg}$	Activation energy for grain growth	KJ/mol
$t_{ip}$	Interpass time	Second
$T_{95\%}$	Time for 95% or full recrystallization	Second
$d_f, D_f$	Austenite grain size	Microns
$X_{DR}, X_{dyn}$	Dynamically recrystallized fraction	-
$D_{dyn}, d_{dyn}, D_{DRX}, D_{RX}$	Grain size after dynamic recrystallization	Microns
$D_{mdrx}, d_{md}, d_{PMRX}, D_{MD}$	Grain size after metadynamic recrystallization	Microns
$D_{mrx}, d_{mrx}$	Partially recrystallized grain size	Microns
$D_p, D_{PR}, D_c$	Partially recrystallized grain size	Microns
$C.R., \alpha, T', C_f$	Cooling rate	°C/second
$d_a, D_a$	Ferrite grain size	Microns
$C_{eq}$	Carbon equivalent	Weight %
$T_{0.05f}$	Temperature at which transformation begins	°C
$T_{mf}$	Mean transformation temperature for ferrite	°C
$V_f, X_f$	Volume fraction of ferrite	-
$S_0$	Pearlite interlamellar spacing	Microns
$H_f$	Transformed phase hardness for ferrite	Brinell
$H_p$	Transformed phase hardness for pearlite	Brinell
$V_p$	Volume fraction for pearlite	-
$H_b$	Transformed phase hardness for bainite	Brinell
$V_b$	Volume fraction for bainite	-
$Nb_{eff}$	Effective Niobium concentration	Weight %
$Q_{md}$	Activation energy for metadynamic recrystallization	KJ/mol
$T_{50MDRX}$	Time for 50% metadynamic recrystallization	Second
$[Nb]_{eq}$	Equivalent Niobium concentration	Weight %
$T_{nr}, T_{NRX}$	No recrystallization temperature	°C
$K_s$	Supersaturation ratio	-
$P_{0.05}, t_{0.05p}, t_{ps}, R_{0.05}$	Precipitation time	Second
HSLA	High strength, low alloy	-
fcc	Face centred cubic	-

**INTRODUCTION**

Historically, the development and advancement of societies have been intimately tied to members' ability to produce and manipulate materials to fill their needs (William, 1997 and Phaniraj et al, 2000). Early civilizations have been designated by the level of materials development, i.e., Stone Age, Bronze Age (Odewole, 2003). The development of many technologies that make human existence so comfortable has been intimately associated with the accessibility of suitable materials.

Steels are one of the most commonly used construction materials. A new class of steels is the low alloy, high strength variety. The versatility of steel for structural and constructional applications rests on the fact that it can be readily supplied in a

wide range of different product forms and with a useful range of materials properties, the use of method which imparts such important characteristics on them during their production should be carefully looked into, and hence, the choice of the hot rolling process for steel production.

Research into controlled hot rolling modeling and simulation has been going on since the 1960's (Tamura et al, 1988, Laasraoui and Jonas, 1991). Many models have been suggested over the years, by researchers, to simulate the full range of the different stages in hot, controlled rolling. A detailed review of these models is available in literatures (Cser et al, 1999, Phaniraj et al, 2000). These models have saved steel industries all over the world millions of dollars; for they have considerably reduced the number of experimental test runs needed for the production of steel with improved properties needed in structural and constructional applications. Different

researchers have used different experimental methods to carry out such simulation; depending on the amount of total deformation, strain rates, temperature, and types of simulation equipment available (Tadeusz, 1992, Medina *et al.*, 1996, Bai *et al.*, 1996 and Djaka *et al.*, 1997 Phaniraj *et al.*, 2000, etc). Laboratory hot rolling, hot compression tests, hot torsion tests and tensile tests have been the most common.

Since the ultimate aim of a modeller is to predict as accurately as possible the mechanical properties of micro-alloy steels for utilization in design and production; reheating of the steel before rolling must be accurately taken care of so that the casting yield of the steel and the products quality could be effectively optimized. Other important factors include flow stress, static recrystallization, peak stress, dynamic recrystallization, metadynamic recrystallization, grain growth, non-recrystallization temperature, precipitation kinetics, partial recrystallization, accumulated strain and ferrite transformations which are thermo-mechanical properties covering the entire span of the hot rolling.

The present work aims at providing ways by which the casting yield of the different steels can be greatly improved, the development of more energy-efficient processing technology for the steels and the determination of models that will optimize products' quality and the process efficiency of the hot rolling.

## MATERIALS AND METHODS

Experiments were carried out to determine the behaviour of some steel samples to ascertain the strains, strain rates, temperatures and other parameters affecting the production and casting yield of steel. The steel samples were specially brought from Finland and some of their properties were already known and detailed in Table II. Tests on these were in uniaxial tension, torsion and compression of axially symmetrical or plane samples. Surface treatment and preparation were carried out on the specimens. These operations prepared the specimens used for a later metallographic studies conducted on them for the determination of the crystal grains, their sizes and grain growth which occurred due to the heat treatment the samples were subjected to.

The current concerns of major steel producers which involve the protection of the environment and competition from mini-mills, upgrading the quality of the product by improving the adaptive control systems, introducing tool steel rolls to reduce roll wear, reduction in the cost of production by eliminating non-essential process steps, many of which are energy-intensive and elimination of the reheating process in favour of dynamic recrystallization have become imperative. More importantly, new methods are being for direct casting and rolling of thin strips.

Mathematical modelling and simulation of the properties of hot rolled products are being used so that the models constructed could be easily manipulated to yield a motion picture of reality. For this work, these in addition to the use of the EXCEL package generated data which were later combined with the earlier obtained experimental results and adequate "adjustments" were made between these values and those available from industries and mills, with the result that better operating conditions were proposed for the hot rolling process used in production of steel.

There are several possible objectives that may justify the mathematical modelling of the hot rolling process. These include the analysis of metal flow and or the metallurgical events during and after the pass, the off-line scheduling of the draft, the on-line adaptive control of the process or the design of the mill: the rolls, the frame, the drive spindle, the bearings, or the screwdown system.

## RESULTS AND DISCUSSIONS

Appendices 1 and 2 give the values of the parameters used in the validation of the simulated models. The results obtained from the validation of the models in the process steps are as presented in Table II. A comparison of these results with those of the Carbon-Manganese (C-Mn) steels is carried out in Table III. The comparison made here are those obtained from the averaging out of the results with similar averaged values for the C-Mn steel for the hot, controlled rolling process. This reveals a possible influence of both the micro-alloys and the substitutional elements on the properties of the different steel produced.

The major microstructural changes that occurred due to the addition of both micro-alloys and substitutional elements are grain refinement due to recrystallization and strain-induced precipitation of the carbo-nitrides. The strengthening of the steel obtained are through precipitation hardening and the refinement of the ferrite grains through reduction of the austenite - ferrite ( $\gamma$ - $\alpha$ ) transformation temperature.

The percentage compositions of additives used for the validation of the simulated models are presented in Table I. From Table II, it is evident that the presence of the additives, i.e., the micro-alloys and the substitutional elements strengthens the steel, however, there is an increase in the solubility of the elements as indicated by the high temperature (1213°C) obtained when compared with that of the C-Mn steel (Table III). This rise in temperature is an indication of the increased energy used in the reheating of the slab. These strengthening factors are present from the reheat stage and up to the transformation which occurred in the ferrite stage of the hot rolling process and may therefore, be responsible for the increase in softening times for both static and metadynamic recrystallization (2 to

12.80s and 0.52 to 3.79s respectively) in comparison with the C-Mn steel.

The reduction in static and metadynamic grain sizes (from 35 to 30.5 microns, and from 32 to 20.9 microns respectively) contributed to producing steel with small, uniform ferrite grains which improves the mechanical properties of the products. Another contributing factor is reduced

grain growth after metadynamic recrystallization (from 54.60 to 43.10 microns). These occur as a result of the greatly refined microstructure formed.

The reductions and refinement obtained through the changes that occurred above led to a higher ultimate tensile strength (a change from 480.5 to 796 MPa).

**Table 1: Percentage Composition of Additives Used**

*Const.	Percentage Composition															
	C	Si	Mn	Nb	S	P	V	Cr	Ni	Mo	Ti	Cu	Al	B	N	H
Mo-Ti	0.06	0.47	1.49		.005		0.08						.042		0.013	
Si-Mn	0.03	0.5	1.21	0.01	.001	.036	.031	18.1	10	0.15	.006	0.19	.055		0.03	
V-T	0.03	0-	0.24		0.02	0.00							0.02		0.003	
		0.21	-			8-										
IF-Mo	0.45		1.08			.017							.042		0.005	
A36	.155	.259	1.51		.031		0.12								0.015	
1538V	.112	1.26	1.53		.008	.013		.021	.022				.029		.0017	
Mo-VNb	0.21	0.06	0.79		.013		0.01	0.84	0.12	0.31			.046		0.02	
IF	1.1	0.23	0.63		.007	0.01	0.17	0.04	.038			.036			0.007	
Cr-Mo	0.18	0.33	1.33	.003			.003	.012			.003					
Ti-V	.310	.084	0.67					1.04					.027	.002	.0077	
Cr-B	0.18		1.30	.035												
Nb-B	.090	0.30	1.30													
Nb-Ti-V	0.14	0.47	1.60													
Nb-Ti	.150	0.35	1.39	.024	.003	.018					.018		.024	.027		3[ppm]
Nb-Ti	0.09	0.23	1.51	.033							.015					
Nb-Ti	0.08	0.39	1.65	0.04			.003			.004	.012		.026		.0039	
Cu-Nb-B	.026	0.18	1.38	.058	.006	.007					.017	2.03	.019	.003	0.006	
Mo1	0.44	0.24	0.79							0.26						
V3	.011	0.24	1.00				.093								.0144	
Mo-Ti	.063	.220	1.20		.008	.011				0.18	0.02		.029		27ppm	
Mo	.063	0.22	1.20		.008	.011				0.18	.029		.029		27 ppm	
IF	.008	.011	0.29	.055	.012	.008					0.13		.049		.0066	
310SS	0.25	1.50	0.20					25.0	22.0							
Ti1	0.15	0.24	1.12								.021				0.015	
V-T	0.12	.362	1.42		.014	.017	.042								.0095	
Mo2	0.44	0.23	0.79							0.38						
V1	0.11	0.24	1.10				.043								.0105	
Mo-NbB	.026	.120	1.70	.059						0.30	.015		.037	.004	0.003	
Mo4	.063	0.22	1.20		.008	.011				0.18	.029		.029		27 ppm	
Cr3	0.55	0.31	0.78		.002	.007		0.76	0.07	0.02		0.1	.004	.005		
Cr-Mo-V	0.47		0.66	.016			0.12	0.98	0.46	0.97					.0042	
C	.038	.009	0.30		.008	.010						.015	0.04		.0052	
Mo-B	0.08		1.5	.045			0.08				.015					

Source: Phannaj et al, 2000

\*Const.: Steel designation based on major constituent

Table II: Validated Models for Other Types of Steels

Table II A: Solubility Models

MD NO	*CONST	MODEL	Value(°C)
1	V	$T=8700[1/\{3.63-\text{Log}[V][N]\}]$	1284.58
2	V	$T=8330[1/\{3.40-\text{Log}[V][N]\}]$	1302.79
3	V	$T=9500[1/\{6.72-\text{Log}[V][C]\}]$	1213.61
4	V	$T=7840[1/\{3.02-\text{Log}[\%V][\%N]\}]$	1329.68
Grain Growth in Reheating			
NO.	CONST	MODEL	Value
Isothermal Grain Growth Model			(Microns)
5	Si-Mn	$d^n=d_0^n+k_1t^n\exp(k_2/T)$	52.85
Grain Size in Continuous Reheating Model			
6	Si-Mn	$d_g=(0.60661/l_m)^4(A_m/0.4861)^{5/2}$	46.50

Table II B: Flow Stress Models

MD. NO.	*CONST.	MODEL	Value (MPa)
7	IF	$\sigma = K(\epsilon + \epsilon_0)^n (\dot{\epsilon}' / \dot{\epsilon}_0)^m (1 + \beta(T - T_0))$	71.43
8	Mo	$\sigma_1 = \sigma_s (1 - \exp(-\epsilon - \epsilon_r)^0.5)$	380.21
9	"	$\sigma = \sigma_1 - (\sigma_s - \sigma_{ss})(1 - \exp(-\beta_{dyn}(\epsilon - \epsilon_p)^n))$	417.08
10	Cr-Mo	$MFS_{Cr-Mo} = (MFS_{misaka}(0.835 + 0.51Nb + 0.098Mn + 0.128Cr^{0.80} + 0.144Mo^{0.3} + 0.175V + 0.01Ni)) * (1 - X_{dyn}) + K\sigma_{ss}X_{dyn}$	402.11

Table II C: Static Recrystallization Kinetics of Static Recrystallization Models

MD. NO.	*CONST.	MODEL	Value(s)
11	Mo-Ti-Nb	$t_{0.5} = A\epsilon^q \dot{\epsilon}^p D_s \exp(Q/RT)$	62.08
12	V	$T_{0.5} = A' \exp(Q^*/RT)$	2.55
13	"	$t_{0.5} = A_s d_0^3 \dot{\epsilon}'^{-1/3} \exp(Q_s/RT)$	0.37
14	Nb-B	$t_{0.5} = 2.86 * 10^{-19} \epsilon^{-3.80} \dot{\epsilon}^{-0.42} \exp(436000/RT)$	101.78
15	"	$t_{0.5} = 1.06 * 10^{-24} \epsilon^{-3.55} \dot{\epsilon}^{-0.33} \exp(559000/RT)$	28.20
16	Nb-Ti	$t_{0.5} = 3.677 * 10^{-20} D^2 (\epsilon - 0.08)^2 \dot{\epsilon}^{-0.176} \exp(359000/RT)$	0.01
17	"	$t_{0.5} = 1.6 * 10^{-14} \epsilon^{-1.7} \dot{\epsilon}^{-0.42} d_0^{0.7} \exp(360000/RT)$	2277.70
18	Nb-Ti-Mo	$t_{0.5} = A d_0^2 \dot{\epsilon}^{-y} \exp(Q/RT)$	81.71
19	V	$T_{0.5} = 1.033 * 10^{-10} \epsilon^{-0.44} \dot{\epsilon}^{-2.6} D \exp(158000/RT)$	1.04
20	"	$T_{0.5} = 3.7100 * 10^{-12} \epsilon^{-0.44} \dot{\epsilon}^{-2.6} D \exp(205000/RT)$	3.16
21	CrMo-VNb	$T_{0.5} = 1.57 * 10^{-14} \epsilon^{-2.9} d^2 \exp(271000/RT)$	55.44
22	Nb-Ti	$t_{0.5} = 3.677 * 10^{-20} D^2 (\epsilon - 0.08)^2 \dot{\epsilon}^{-0.176} \exp(359000/RT)$	70.01
23	Mo-Ti	$T_{0.5x} = 1.5 * 10^{-12} \epsilon^{-0.28} \dot{\epsilon}^{-2.18} D^{-0.87} \exp[232000/RT]$	28.21
24	Ti-V	$t_{0.5x} = 5 * 10^{-18} (\epsilon - 0.085)^{-3.5} D^2 \exp[280000/RT]$	7.53
25	Eutectoid	$t_{0.5x} = 2.4 * 10^{-8} \epsilon^p \dot{\epsilon}^{-0.29} D^{-0.2} \exp[160420/RT]$	1.37
26	V-T	$t_{0.5x} = 9.3 * 10^{-13} \epsilon^{-4} D^2 \exp[230000/RT]$	4.03
27	Austenitic	$T_{50} = 2.29 * 10^{-15} \epsilon^{-2} Z^{-0.32} \exp(400000/RT)$	0.17

Table II C Contd.

28	Cr-Mo	$T_{0.5}=1.57*10^{-14}d_0^2\epsilon^{-2.9}\exp(271000/RT)$	55.44
29	A36	$t_{0.5}=8.31*10^{-15}d^{1.5}\epsilon^{-1.5}\epsilon_c^{-0.33}\exp(263000/RT)$	2.05
30	4120		
30	1538V	$t_{0.5}=V2*\epsilon_c^{-0.93}d_0^{0.94}\epsilon^{-0.95}\exp(L1/RT)$	0.16
31	Cr	$t_{0.5}=2.75*10^{-11}\epsilon_c^{-0.125}\epsilon^{-2.45}d^{1.06}\exp(184000/RT)$	3.20
<b>Static Recrystallization Grain Size Models</b>			
NO.	CONST.	MODEL	Value (Microns)
32	Ti-V	$D_{rex}^y=B+CD_0^x\epsilon^{-y}[\exp(350000/RT)]^{-z}$	11.04
33	V	$d_{rex}=\Lambda d_0^{1.3}\epsilon^{-p}\exp(-Q_{gx}/RT)$	41.65
34	V-Ti-Nb	$D_{rex}^y=B+CD_0^x\epsilon^{-y}[\exp(350000/RT)]^{-z}$	11.20
35	Nb-V	$D_{rex}^y=-1.25+24.4(V+Nb)^{-0.2}N^{-0.04}D_0^{0.25}\epsilon^{-0.55}[\exp(35000/RT)]^{-0.7}$	11.80
36	Ti-V	$D_r=4.3+95.7D^{0.15}\epsilon^{-0.57}\exp[-350000/RT]$	43.01
37	Eutectoid	$D_r=9.91D^{0.54}\epsilon^{-0.65}\epsilon_c^{-0.1}\exp[-17540/RT]$	44.46
38	V-T	$D_r=4.54D^{0.67}\epsilon^{-0.53}\epsilon_c^{-0.1}\exp[-15000/RT]$	35.49
39	Austenitic	$D_r=2.24\epsilon^{-0.5}DZ^{-0.06}$	45.70
40	"	$D_r=83.9\epsilon^{-0.537}D^{0.355}Z^{-0.0655}$	124.66
41	Cr-Mo	$d_{SRX}=1.1d_0^{0.67}\epsilon^{-0.67}$	44.46
42	A36	$d_{rex}=88.96d_0^{0.36}\epsilon^{-0.368}\exp(-28060/RT)$	46.38
43	4120		
43	1538V	$d_{srxn}=V3*d_0^{0.96}\epsilon^{-0.97}\exp(-L2/RT)$	37.89

Table II C: Static Recrystallization  
Grain Growth After Static Recrystallisation

NO.	CONST.	MODEL	Value (Microns)
<b>Growth During 1st Second Model</b>			
44	Cr-Mo	$d^2=d_{SRX}^2+4.0*10^7(t_p-4.32t_{0.5})\exp(-113000/RT)$	30.12
<b>Growth After 1st Second Model</b>			
45	"	$d^7=d_{SRX}^7+1.5*10^{27}(t_p-4.32t_{0.5})\exp(-400000/RT)$	50.30

Table II D: Dynamic Recrystallization

Critical and Peak Strains Models

MD. NO.	*CONST.	MODEL	Value
46	CrMoV	$\epsilon_p=0.0040d^{0.50}[\epsilon'\exp(241000/RT)]^{0.17}$	1.36
47	Nb	$\epsilon_c=0.84\epsilon_p$	1.14
48	V	$\epsilon_p=4.3*10^{-3}D^{0.20}Z^{0.174}$	1.31
49	"	$\epsilon_p=5.9*10^{-3}D^{0.20}Z^{0.174}$	1.79
50	Mo	$\epsilon_p=5.24*10^{-4}D_0^{0.5}Z^{0.18}$	0.61
51	Mo-Ti	$\epsilon_p=7.07*10^{-4}dZ^{0.15}$	2.49
52	4120		
52	1538V	$\epsilon_p=A1*d_0^{b1}\epsilon^{b2}\exp(P1/RT)$	0.50
53	"	$\epsilon_c=0.83\epsilon_p$	0.42
54	A36	$\epsilon_p=0.01318d_0^{0.174}\epsilon^{0.165}\exp(2926/T)$	0.26

Table II D Contd.

55	"	$\epsilon_c=(5/6)\epsilon_p$	0.22
56	4120 1538V	$\epsilon_c=C(V1*d_o^{g1}*Z^{g2})$	0.33
57	Cr	$\epsilon_p=4.2*10^{-4}d^{1/2}Z^{0.15}$	0.21
<b>Kinetics of Dynamic Recrystallization Models- (Volume Fraction X)</b>			
NO.	CONST.	MODEL	Value
58	Cr-B	$X_{DR}(\epsilon)=1-\exp\{-2.996[(\epsilon-\epsilon_c)/(\epsilon_s-\epsilon_c)]^n\}$	0.35
59	"	$X_{DR}(\epsilon)=1-\exp\{-2.996(\epsilon/\epsilon_s)^n\}$	0.27
60	"	$X_{DS}(\epsilon)=1-\exp\{-3[\epsilon/\epsilon_s(\epsilon',T)]^{n(\epsilon)}\}$	0.27
61	Mo	$X_{dyn}=(\sigma_s^*-\sigma)/(\sigma_s^*-\sigma_{ss})=1-\exp(-\beta_{dyn}(\epsilon-u\epsilon_p)^n)$	0.13

Table II E: Metadynamic Recrystallization Kinetics of Metadynamic Recrystallization Models

MD. NO.	*CONST.	MODEL	Value(s)
62	Mo-Ti	$t_{0.5}=6.66*10^{-6}\epsilon^{-0.51}\exp(136000/RT)$	2.54
63	"	$t_{0.5}=6.66*10^{-6}\epsilon^{-0.71}\exp(110000/RT)$	2.20
64	Mo	$t_{50\%}=6.66*10^{-6}\epsilon^{-0.61}\exp(123000/RT)$	7.50
65	V	$t_{0.5}=A_{md}\epsilon^{-2/3}\exp(Q_{md}/RT)$	2.50
66	V-Ti	$t_{50\%}=1.8*10^{-6}d_o^{0.24}\epsilon^{-0.75}\exp(13974/T)$	2.70
67	V	$t_{0.5}=(1.5([Si]+[V]+2.8)*10^{-8}Z^{-0.6}\exp(380/RT)$	3.20
68	"	$t_{0.5}=(1.5([Si]+[V]+2.8)*10^{-8}\epsilon^{-0.6}\exp(170/RT)$	4.28
69	A36	$t_{0.5}=2.13*10^{-6}\epsilon^{-0.68}\exp(132800/RT)$	6.00
70	4120 1538V	$t_{0.5}=V4*Z^{-98}\exp(L3/RT)$	2.53
71	"	$t_{0.5}=V4*\epsilon^{-n1}d_o^{n2}\epsilon^{-n3}\exp(L3/RT)$	2.91
72	Cr-Mo	$t_{0.5}=1.84*[\exp(330000/RT)]^{-0.86}\exp(271000/RT)$	5.50
73	Cr	$t_{0.5}=3.5*10^{-2}\epsilon^{-0.785}\exp(24800/RT)$	3.60
<b>Metadynamic Recrystallization Grain Size Models</b>			
NO.	CONST.	MODEL	Value (Microns)
74	Mo	$D_{mdrx}=1.66*10^3Z^{-0.16}$	17.78
75	Mo-Ti	$D_{MDRX}=1.19*10^3Z^{-0.16}$	12.75
76	Cr-Mo	$d_{MDRX}=1370*\epsilon^{-0.13}\exp(-45000/RT)$	20.41
77	4120 1538V	$d_{Drxn}=V5*Z^{-99}$	31.07
<b>Grain Growth After Metadynamic Recrystallization</b>			
NO.	CONST.	MODEL	Value (Microns)
<b>Growth During 1st Second Model</b>			
78	Cr-Mo	$d^2=d_{MDRX}^2+1.2*10^7(t_p-2.65t_{0.5})\exp(-113000/RT)$	58.79
<b>Growth After 1st Second Models</b>			
79	Cr-Mo	$d^7=d_{MDRX}^7+8.2*10^{25}(t_p-2.65t_{0.5})\exp(-400000/RT)$	32.50
80	"	$d^{4.5}=d_o^{4.5}+4.1*10^{23}*t_p*\exp(-435000/RT)$	46.09

**Table II F: Non-Recrystallization Temperature Models**

MD. NO.	*CONST.	MODEL	Value(°C)
81	Mo-B	$T_{nr}=A\epsilon^{-0.12}\epsilon_r^{-0.01}t^{-0.1}$	815.20
82	Nb-B	$T_{nr}=(A'\text{Log}[C][\text{Nb}]+B')\epsilon^{-0.12}\epsilon_r^{-0.01}t^{0.04}$	945.63

**Table II G: Kinetics of Precipitation Models**

MD. NO.	*CONST.	MODEL	Value(s)
83	Nb-Ti	$R_{0.05}=0.33*10^{-19}d_0^2\epsilon^{-4}\exp(360000/RT)$	2.89
84	"	$R_{0.05}=2.89*10^{-38}d_0^2\epsilon^{-4}\exp(780000/RT)$	8.81

**Table II H: Ferrite Grain Size Models**

MD. NO.	*CONST.	MODEL	Value (Microns)
85	V-Ti	$D_\alpha = 3.75+0.18D+1.4Cr^{-0.5}$	13.27
86	"	$D_\alpha = (1-0.45\epsilon_r^{1/2})\{3+1.4Cr^{-1/2}+17[1-\exp(-1.5*10^2D)]\}$	19.08
87	V-T	$D_\alpha = (D/(1+(0.036+0.0233Cr^{0.5})D))$	12.09
88	Nb-V-Ti	$D_V/D_\alpha = 1+(0.0026+0.053\%C+0.006\%Mn+0.009\%Nb+4.23\%V+N-0.081\%Ti)*(1.5+\alpha^{1/2})*D_V$	10.12

**Table II I: Lower Yield Stress Models**

MD. NO.	*CONST.	MODEL	Value (MPa)
89	V	$LYS=62.6+26.1[Mn]+60.2[Si]+759[P]+212.9[Cu]+3286.0[N]+\sigma_{ppn}+19.7d\alpha^{-0.5}$	562.05
<b>Models Relate Lower Yield Stress to Mean Free Path in Ferrite</b>			
90	Eutectoid	$\sigma_y=308+0.07(1/M)$	308.23
91	"	$\sigma_y=259+0.087(1/M)$	259.44
92	V	$LYS=70+84Si+32Mn+680P+33Ni+38Cu-30Cr+1.81d^{-1/2}$	163.45

**Table II J: Models Describing Contribution of Precipitation to Strengthening(L.Y.S)**

MD. NO.	*CONST.	MODEL	Value (MPa)
93	V	$\sigma_p=57\text{LogCr}+700[V]+7800[N]+19$	16.20
94	"	$\sigma_p=19.9[C]+552.8[C]^2+590[V]+8650[N]+19.9\ln(\text{Cr})$	15.82
95	Nb-Ti	$\sigma_p=2500([\text{Nb}]^{-0.5}[\text{Ti}])$	50.00

**Table II K: Tensile Strength Models**

MD. NO.	*CONST.	MODEL	Value (MPa)
96	V	$TS=164.9+634.7[C]+53.6[Mn]+99.7[Si]+651.9[P]+472.6[Ni]+3339.4[N]+\sigma_{ppn}+11d\alpha^{-0.5}$	796.49
97	Eutectoid	$\sigma_u=706+0.072(1/M)+122[Si]$	763.58
98	"	$\sigma_u=773+0.058(1/M)+122[Si]$	830.63



Table III: Comparisons Between C-Mn and Other Types of Steels

Models	Steels	*C-Mn	Other Types
Solubility (°C)		1203	1213
Grain Growth in Reheating (Microns)		40.7	50
Flow Stress (MPa)		208	392
Static Recrystallization, t <sub>50</sub> (Second)		2	12.8
Static Recrystallization Grain Size (Microns)		35	30.5
Static Recrystallization Grain Growth (Microns)		45.5	43.8
Metadynamic Recrystallization, t <sub>50</sub> (Second)		0.52	3.79
Metadynamic Recrystallization Grain Size (Microns)		32	20.9
Metadynamic Recrystallization Grain Growth (Microns)		54.6	43.1
Ferrite Grain Size (Microns)		10.8	13.5
Lower Yield Stress, L.Y.S (MPa)		331.2	323
Ultimate Tensile Strength, U.T.S (MPa)		480.5	796

\*: Phaniraj et al, 2000.

**CONCLUSION**

The following conclusions can be drawn:

1. Ferrite grain size is more sensitive to the retained strain than to the cooling rate or the austenite grain size. However, there is a limit to the size of the ferrite grain that can be obtained by increasing the retained strain or the cooling rate.
2. A drop in the austenite-ferrite ( $\gamma - \alpha$ ) transformation temperature leads to an increase in the tensile strength of the steel.
3. The use of the computer in the validation of the simulated models reduces the number of experimental test runs needed for the formulation and determination of process parameters required in steel production.

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**Appendix 1**

**Values Used in the Validation of the Simulated Models**

Unless otherwise stated, the general values of the parameters utilized in validating the simulated models are as follows:

**1. Reheating Before Rolling**

*Solubility Models*

C = 0.1 wt %, N = 0.02 wt %, V = 0.09 wt %, Al = 0.025 wt %, D<sub>0</sub> = 40 microns, T = 1473K.

*Grain Growth in Reheating*

D<sub>0</sub> = 25 microns, T = 1473 K, t = 30 mins, R = 8.312 KJ/mol.

**2. Flow Stress**

Strain Rate = 1/s, R = 8.312 KJ/mol, T = 1273 K, [C] = 0.03 wt %, [Mn] = 0.27 wt %, X<sub>d<sub>0</sub></sub> = 0.8, Strain = 0.2, Q<sub>rx</sub> = 300000 J/mol, Q<sub>d</sub> = 330000 J/mol, Peak Strain, c<sub>p</sub> = 0.35.

**3. Kinetics of Static Recrystallization**

R = 8.312 KJ/mol, T = 1273 K, D<sub>0</sub> = 50 microns, Q<sub>d</sub> = 330000 J/mol, Q<sub>rx</sub> = 480000 J/mol, X\* = 0.5, Strain rate = 1/s, Strain = 0.2, [Mn] = 1.4 wt %, Static Recrystallization Grain Size

D<sub>0</sub> = 50 microns, Strain = 0.2, R = 8.312 KJ/mol, T = 1273 K, Q<sub>d</sub> = 330000 J/mol, Grain Growth after Static Recrystallization

t<sub>0.5</sub> = 10s, t<sub>1/2</sub> = t<sub>tp</sub> = 50s, d<sub>rx</sub> = 30 microns, t<sub>0.9</sub> = 20s, R = 8.312 KJ/mol, T = 1273 K.

**4. Dynamic Recrystallization**

Critical and Peak Strain

Q<sub>rx</sub> = 312000 KJ/mol, Q<sub>d</sub> = 330000 J/mol, D<sub>0</sub> = 50 microns, T = 1273 K.

**5. Kinetics of Metadynamic Recrystallization**

D<sub>0</sub> = 40 microns, Strain Rate = 1/s, K = 1.7, R = 8.312 KJ/mol, T = 1273 K, Metadynamic Recrystallization Grain Size

Q<sub>d</sub> = 330000KJ/mol, Strain Rate = 1/s, T = 1273 K, R = 8.312 KJ/mol, Grain Growth after Metadynamic Recrystallization

t<sub>tp</sub> = 20s, T = 1273 K, D<sub>mdrx</sub> = 22 microns, R = 8.312KJ/mol, t<sub>0.5</sub> = 1s.

**6. No Recrystallisation Temperature, T<sub>nr</sub>**

t = 15s, T = 1123 K.

**7. Ferrite Grain Size**

C<sub>eq</sub> = 0.14 wt %, D = 22 microns, α = 3°C/s, T = 1273K, Retained Strain = 0.1, X<sub>f</sub> = 0.9, D = 40 microns

**8. Lower Yield Stress**

Si = 0.33 wt %, N = 0.02 wt %, Mn = 1.3 wt %, X<sub>f</sub> = 0.9, D = 10 microns, C = 0.18 wt %, α = 3°C/s, D = 40microns, T = 1273 K, Cu = 0.009 wt %, [P] = 0.009 wt %,

**9. Contribution of Precipitation to Lower Yield Stress (L.Y.S)**

[V] = 0.09, [N] = 0.02, [C] = 0.1, C.R. = 3°C/s, [Ti] = 0.003.

**10. Tensile Strength,**

[C] = 0.1 wt %, [Mn] = 1.33 wt %, [Si] = 0.25 wt %, [P] = 0.009 wt %, [Ni] = 0.009 wt %, D = 10 microns, X<sub>f</sub> = 0.9.

**Appendix 2**

The parameters listed below are specific and have been used to validate the simulated models listed above either wholly by themselves or in conjunction with the general parameters listed in appendix 1.

**SPECIFIC PARAMETERS, VALUES AND REMARKS**

When the soaking temperature is lower than 1150°C logn = 0.531 + 307.404/T, K <sub>1</sub> = 8.74*10 <sup>4</sup> , K <sub>2</sub> = 4.264*10 <sup>3</sup>
When the soaking temperature is in the range 1150 – 1250°C logn = 5.214 + 8406/T, K <sub>1</sub> = 1.29*10 <sup>-54</sup> , K <sub>2</sub> = 1.94*10 <sup>5</sup> . t = soaking time.
l <sub>m</sub> = 0.60661d <sub>g</sub> exp(5(lns) <sup>2</sup> /2), A <sub>m</sub> = 0.4861d <sup>2</sup> exp(4(lns) <sup>2</sup> ) s = exp(ln(1.321l <sub>m</sub> <sup>2</sup> A <sub>m</sub> )) <sup>1/2</sup> , d <sub>m</sub> = d <sub>g</sub> exp((lns) <sup>5</sup> /2), d <sub>g</sub> – Median grain diameter d <sub>m</sub> – Mean grain diameter, l <sub>m</sub> = 10 microns, A <sub>m</sub> = 200 μm <sup>2</sup>

Appendix 2 Contd.

$K=566\text{MPa}$ , $\epsilon_0=-0.014$ , $n=0.219$ , $\beta=-0.0012$ , $m=0.018$ , $\epsilon'_0=0.02/\text{s}$ , $T_0=298\text{K}$
$Q_d=296\text{kJ/mol}$ , $\epsilon_p=2.84 \cdot 10^{-4} D_0^{0.5} Z^{0.17}$ , $\beta_{d_{\text{dyn}}}=48\epsilon'^{-0.22} \exp(-41060 \text{ RT})$ $n=0.23\epsilon'^{0.11} \exp(18000/\text{RT})$ , $\sigma_{\text{ss}}=7.4Z^{0.096}$
Valid for 0.52 – 0.66%Mn, 0-0.8% Nb, 0.83 – 1.38% Cr, 0-0.46%Ni, 0.15-0.97% Mo, and 0-0.27% V $\sigma_{\text{ss}}=1.18[\epsilon' \exp(330000/\text{RT})]^{0.15}$
$Q=124714+28285.68[\text{Mn}]+64716.68[\text{Si}]+72775.40[\text{Mo}]+76830.32[\text{Ti}]^{0.215}+121100.37[\text{Nb}]^{0.1}$ , $A=3.869 \cdot 10^{-4} \exp(-7.921 \cdot 10^{-5} Q)$ , $s=1$ , $q=-0.53$ , $p=3.7d^{-0.137}$
$A=25.3318 \exp(-1.0147 \cdot 10^{-4} Q^*)$ , $Q^*=Q \exp[H R(1/T-1/\text{SRCT})]$ $H=400000 \text{ Jmol}^{-1}$ , $\text{SRCT}(\text{K})_V=T_s-1.05 \cdot 10^{-3} D^{-0.35} \epsilon^{0.50}$
$A_s=5.15 \cdot 10^{-15} \text{ s}$ , $\beta=2.0$ , $Q_s=262 \text{ kJ/mol}$
$A=1.8 \cdot 10^{-28}$ , $y=-4.6$ , $\text{Strain}=0.59$
$T > 915^\circ\text{C}$ , $Z=\epsilon' \exp(248000/\text{RT})$ $T > 915^\circ\text{C}$
Steel 0.06% C, 1.2% Mn, 0.22% Si, 0.18% Mo, 0.02% Ti, 0.029% Al, 0.0027% N
Steel 0.13% C, 1.4 Mn, 0.01% Ti, 0.04% V, 0.01% N
Steel 0.72% C, 1.2% Mn, 0.28% Si, 0.002% Al, 0.21% Cr.
Steel 0.2% C, 1.36% Mn, 0.26% Si, 0.09% V, 0.001% Al, 0.006% N
$B=1.1$ , $C=170$ , $x=0.15$ , $y=0.57$ , $z=0.11$ for 0.01 Ti – (0.08-0.13V)-0.023N
$\Lambda=8.9 \cdot 10^1 \mu\text{m}^{-2.3}$ , $p=0.368$ , $Q_{\text{ex}}=28.1 \text{ kJ/mol}$
$B=1.08$ , $C=176.2$ , $x=0.02$ , $y=0.66$ , $z=0.09$ for 0.01 Ti-0.04V-0.025Nb-0.008N
0.04-0.13%V, 0-0.025%Nb, 0.008-0.023%N
Steel 0.13% C, 1.4%Mn, 0.01%Ti, 0.04% V, 0.01%N
Steel 0.72% C, 1.2% Mn, 0.28% Si, 0.002% Al, 0.21% Cr
Steel 0.2% C, 1.36% Mn, 0.26% Si, 0.09% V, 0.001% Al, 0.006% N
If $d_{\text{rex}} > d_0$ , then $d_{\text{rex}} = d_0$
$T > 915^\circ\text{C}$
$T > 915^\circ\text{C}$
$n=0.27\epsilon'^{0.08} \exp(15.8 \text{ kJ/mol RT})$ , $\beta_{d_{\text{dyn}}}=3512-\epsilon'^{0.3} \exp(-84.7 \text{ kJ/mol RT})$
$950 < T < 1000^\circ\text{C}$
$A_{\text{nd}}=1.54 \cdot 10^{-6} \text{ s}$ , $Q_{\text{nd}}=127 \text{ kJ/mol}$
$A=88.1^\circ\text{C per wt}\%$ , $B=1156^\circ\text{C}$ , $t \leq 12.5 \text{ s}$ , $Nb_{\text{eq}}=\text{Nb}+0.31\text{Ti}+0.15\text{Al}$
$A'=63.5^\circ\text{C per wt}\%$ , $12.5 \text{ s} \leq t \leq 30 \text{ s}$ , $B'=885^\circ\text{C}$
$T > 1000^\circ\text{C}$
$1000 > T > 890^\circ\text{C}$
Steel 0.2% C, 1.36% Mn, 0.26% Si, 0.09% V, 0.001% Al, 0.006% N
$D_v/D_a$ - Grain refinement ratio
$\sigma_{\text{ppt}}=57 \log T' + 700[\text{V}]+7800[\text{N}]+19$
$M$ - Mean free path for slip, $M=2S_0(1-0.15[\text{C}])$
$S_0 \geq 0.15 \mu\text{m}$
$S_0 < 0.15 \mu\text{m}$
$\sigma_p$ - Precipitation stress
$\sigma_{\text{ppn}}=57 \log T' + 700[\text{V}]+7800[\text{N}]+19$
Steel 0.72% C, 1.2% Mn, 0.28% Si, 0.002% Al, 0.21% Cr in wt. %
$S_0 \geq 0.15 \mu\text{m}$
$S_0 < 0.15 \mu\text{m}$ , Where $M=2S_0(1-0.15[\text{C}])$
$S_0 \{129.3-54.4[\text{Mn}]-4.4[\text{Cr}]-17.5[\text{Si}]-0.18-0.07[\text{Mn}]-0.012[\text{Cr}]-0.027[\text{Si}]\Gamma_p\}^{-1}$

When the soaking temperature is lower than 1150°C  
 $\log n = 0.241 + 307.404/T$ ,  $K_1 = 8.74 \cdot 10^{-4}$ ,  $K_2 = 4.594 \cdot 10^{-4}$   
 When the soaking temperature is in the range 1150 – 1250°C  
 $\log n = 2.14 + 8400/T$ ,  $K_1 = 1.29 \cdot 10^{-2}$ ,  $K_2 = 1.94 \cdot 10^{-2}$ ,  $t = \text{soaking time}$   
 $\Gamma_p = 0.00014 \exp(4 \ln \Gamma_p)$ ,  $A_p = 0.48619 \exp(4 \ln \Gamma_p)$   
 $\Gamma_p = \exp(\ln(1/25))^{1/2}$ ,  $d_p = d_0 \exp(\ln \Gamma_p)$ ,  $d_p$  - Median grain diameter  
 $d_0$  - Mean grain diameter,  $d_0 = 10 \mu\text{m}$ ,  $A_p = 200 \mu\text{m}$