Numerical Model for the Minimization of Intermixed Round Bars in a Four Line Continuous Caster

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The grade transition process in a four line round billets continuous caster was analyzed. A numerical model was developed to determine, quickly and accurately, the length and location of intermixed steel in the bar. The model is based on "tank in series" or "volumes" mathematical models and was calibrated with physical water model measurements and full three-dimensional (3-D) turbulent numerical calculations. Comparison of the model with measurements performed on intermixed steel bars is also presented. The good agreement between these plant measurements and the numerical results validated the model. The model was implemented on PCs with a visual interface that allowed the easy input and output of data, and the program is operated in a steel shop. Finally, the code was employed to evaluate different strategies to fill the tundish during grade transitions. The "double dilution" process, which consists of filling the tundish in two steps, proved to be useful to reduce the amount of downgraded steel.

SEQUENTIAL casting of different steel grades during

speed and time-dependent inlet flow rate. This model could

the continuous casting process is often employed in steel

change, to find the best sequence of grades to

I. INTRODUCTION of casting conditions, including time-dependent casting

Several publications^[1,8,9] on grade transition models have
focused on mixing in the tundish. Diener *et al.*^[10] incorpo-
rated the mixing in the mold into the transition models,
while Huang and Thomas^[2] included

models" or "volume models." All these models, however,
consider a one-step filling of the tundish, with the tundish
volume increasing monotonically until its operating value.
The aim of this work is to develop a numerical to caster with any number of lines are discussed. The last Section is devoted to the conclusions.

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An important point to keep in mind is that, as the flow

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An important point to

Table I. Description of the Basic Volumes

Equation	Mixing Volume (V_m)	Dead Volume (V_d)	Plug Volume (V_p)
	Q^{out} m $_{OIII}$		Q ^{out} ´out
Mass conservation	$\frac{dV_m}{dt} = Q_{\rm out} - Q_{\rm in}$	dV_d $= \pm Q$	$\frac{dV_{p}}{dt} = Q_{\rm out}$ $-Q_{\rm in}$
Mass conservation of a chemical elements	$\frac{dC_{\text{out}}}{dt} = \frac{Q_{\text{in}}}{V_{\text{in}}} (C_{\text{in}} - C_{\text{out}})$	$C \neq f(Q,V)$	$C_{\text{out}} = C_{\text{in}} t$ $Q_{\rm in}$

measurements described in Section V. Thus, concentration calculations were performed for a single dimensionless function $(\tilde{C}(t))$ defined by

$$
\tilde{C}(t) = \frac{C(t) - C_{\text{old}}}{C_{\text{new}} - C_{\text{old}}}
$$
 [1]

where C_{old} and C_{new} represent the concentration of a given element in the old and new steel grade, respectively. This dimensionless concentration can take any value between 0 (old steel grade) and 1 (new steel grade). Once the dimensionless concentration is calculated, the concentration of a given chemical element $(C(t))$ may be obtained from Eq. [1].

The model is composed of three types of volumes: mixing volume, dead volume, and plug volume. The basic equations of the model may be derived by imposing mass conservation for the fluid and for the chemical elements in each volume. Fig. 1—Tundish mixing model. Mixing volumes are assumed to be well mixed, having a uniform concentration equal to the outgoing one for each time interval. Plug flow volumes introduce a delay (t_p) in $the outgoing concentration. Table I provides the descriptions$ and the equations for the three volumes.

The time $t = 0$ is considered when the ladle with the new steel grade is opened.

The model consists of three submodels: tundish, mold, and $final$ composition.

III. TUNDISH MIXING MODEL

The flow inside the tundish was modeled by a tank in series mathematical model. Several "tanks" or "volumes" were considered to model a four line symmetric tundish with isothermal flow. In Figure 1, we show the mixing volume model of half the tundish with the internal and external continuous casting lines.

There are two well-defined groups of volumes: one of them associated with the internal lines (superscript int) and the other associated with the external lines (superscript ext), with an extra recirculating mixing volume (V_m^R) , which dC_{m2}^{ext} allows the mass flow between both groups of volumes.

Each of these groups consists of two mixing volumes $(V_{m1}$ and V_{m2}), one dead volume (V_d) , and one plug flow volume (V_p) . In this system of equations, the volume fractions $f_i = V_i/$

I and assembling them according to Figure 1, we reach the also a known function of time. following system of equations, There are three flow rate fractions to be calibrated:

$$
dC_{m2}^{\text{ext}}/dt = (Q_f C_{ml}^{\text{ext}} + Q_h C^R - (Q_f + Q_h) C_{m2}^{\text{int}})/V_{m2}^{\text{ext}}
$$

$$
dC^R/dt = (Q_d C_{m2}^{\text{ext}} + Q_c C_{m2}^{\text{int}} - (Q_d + Q_e) C^R)/V_m^{\text{ext}}
$$

 V_T are parameters to be calibrated with the water model. The total flow rate entering (Q^{en}) and leaving $(Q^{int} + Q^{ext})$
the tundish are known functions of time, which the user Using the basic equations previously described in Table \qquad must specify. Consequently, the half-tundish volume (V^T) is

Fig. 2—Calibration of the tundish mixing model.

$$
x_a = \frac{Q_a}{Q_{en}}
$$
 (fraction of entering flow rate directed to the

 $x_d = \frac{Q_d}{Q_j}$

$$
x_c = \frac{Q_c}{Q_i}
$$
 (fraction between the two flow rates coming out of V_{m2}^{int}).

The new grade composition prescribes the concentration of the chemical elements at the tundish entrance, C^{en} . The time delay introduced by the plug flow volumes is given

 $P_P^{\text{int}}/Q_k^{\text{int}}$ and $t_P^{\text{ext}} = V_P^{\text{ext}}/Q_l^{\text{ext}}$. solved with a fourth-order Runge Kutta algorithm, with the old grade composition as initial composition.

sented in Figure 2. The time is represented in terms of the theoretical mean residence time $\hat{t} = tQ^e/V_T$, $t = 0$ being the moment when the salt solution was injected. The nondimen- to fill the tundish. sional concentration in Figure 2 was defined in such a way (2) *Double dilution procedure*: The filling of the tundish is

Fig. 3—Scheme of a simple dilution procedure. Full lines indicate the weight of the steel in tundish and dotted lines the flow rate of the steel entering the tundish.

that the area beneath the innerline curve is unity, $\hat{C} =$ ∞

$$
C/\int\limits_0^\cdot C(\hat{t})d\hat{t}.
$$

To determine the group of parameters that best fits the internal line); **RTD** curves obtained by the water model experiment, the function $\sum_{k} (C_{\text{int}}(t_k) - C_{\text{int}}^{vm}(t_k))^2 + (C_{\text{ext}}(t_k) - C_{\text{ext}}^{vm}(t_k))^2$ was considered. In this expression, the superscript *wm* indicates concentrations obtained by the water model experience and out of V_{m2}^{ext} ; and t_k represents the time intervals when concentrations were evaluated. The parameter space was explored in order to

find the set of parameters that minimizes this function.

The resulting parameters are as follows:

out of
$$
V_{m2}^{\text{int}}
$$
).
\n $f_m^{\text{out}} = f_{ml}^{\text{int}} = 0.0004$ $f_{m2}^{\text{int}} = 0.33$ $f_{m2}^{\text{ext}} = 0.44$
\nnew grade composition prescribes the concentration
\nhemical elements at the tundish entrance, C^{en} . The
\nlaw introduced by the plua flow volumes is given
\n $f_d^{\text{ext}} = f_{d2}^{\text{int}} = 0.0371$ $x_a = 0.53$ $x_d = x_c = 0.36$
\n $x_d = x_c = 0.36$

The nondimensional concentrations obtained by setting The system of coupled ordinary differential equations was these parameters in the numerical model are plotted in Fig-
Ived with a fourth-order Runge Kutta algorithm, with the ure 2.

D. *Simple and Double Dilution*

C. *Water Model Calibration* During normal operation, the steel weight in the tundish The tundish model just described was calibrated with
experimental measurements performed by $[AS^{[11]}]$ on a physical
cal water model. The water model is a 1:3 scale reproduction
of the tundish at Siderca S.A.I.C. (Campana

- The tracking of concentration was accomplished by (1) *Simple dilution procedure*: When the new ladle opens, injecting a salt solution into the tundish entrance and analyz-
the flow rate entering the tundish is set to a value Q_{sd} ing the flow coming out of each line for different time above Q_{steady} until the operating weight W_{tundish} is instants. **reached.** Then the caster is in normal operation again, The residence time distribution (RTD) curves are pre-

the *W*_{tundish} and Q_{steady} . Figure 3 shows the time evolu-

tion of the weight of steel in the tundish and the inlet flow rate when a simple dilution procedure is adopted
	-

Fig. 4—Scheme of a double dilution procedure. Full lines indicate the weight of the steel in tundish and dotted lines the flow rate of the steel entering the tundish.

performed in two steps, as depicted in Figure 4. When the ladle with the new grade opens, it fills the tundish only up to a certain weight W_1 with a flow rate Q_{dd}^1 . Then it closes for a while (in fact, it is not completely closed, a little flow rate Q_{dd}^2 still entering the tundish), allowing for a second draining of the tundish. When a second minimum steel weight W_{drain2} is attained, the ladle opens with Q_{dd}^3 flow rate, for the tundish to reach the operating weight *W*_{tundish}. Then the flow rate is reduced to the steady-state value *Q*steady , in order to keep a constant steel weight in the tundish.

Fig. 5—Mold mixing model. The time duration of each step is determined by the weight and the flow rate entering and leaving the tundish.

and consists of a plug flow volume and a mixing volume, followed by a region where convective and diffusive effects were supposed to take place (Figure 5). In this latter region, B. *Validation with a 3-D Finite Element Model* a one-dimensional convection (1-D) convection-diffusion The mold mixing model was calibrated in the following equation was solved with the coordinate *z* (taken along the way.

$$
\frac{dC_m}{dt} = \frac{Q}{V_m}(C_T - C_m)
$$

$$
C_p = C_m \left(t - \frac{V_p}{Q} \right)
$$

$$
\frac{\partial C_{Zf}}{\partial t} + v_z \frac{\partial C_{Zf}}{\partial z} = \frac{\partial}{\partial z} \left(D_{\text{eff}} \frac{\partial C_{zf}}{\partial z} \right); \quad z_i < z < z_f
$$

where V_m is the mold mixing volume; V_p is the mold plug

flow volume; C_m , C_p , and C_{Zf} are the concentrations exiting each region; v_z is the casting speed; D_{eff} is the turbulent **IV. MOLD MIXING MODEL AND FINAL** each region, v_z is the casting speed, D_{eff} is the unbulent diffusivity; $z_i = (V_m + V_p)/\pi\phi^2$ and z_f are the initial and final positions of the convection-diffusion region; and ϕ

A. *Volumes Model* the bar radius.
The values of Q and C_T are taken from the tundish mixing The model in the mold and upper part of the bars considers
the first three meters of fluid below the meniscus. It was
validated with a numerical full 3-D turbulence model^[12-16] while the values of V_m , V_p , and D_{eff}

- bar) being the only relevant variable.
The following set of equations for the concentration is
obtained.
 (1) The steel flow inside the system nozzle-mold-bar was
numerically calculated using a 3-D code.^[17] This code
 Mixing volume: $[4]$ predictor/ (ε) -corrector iterative algorithm. This was addressed in our previous publications.[12–16] These calculations were performed considering different casting speeds and mold diameters. The modeled domain Plug flow volume: includes the nozzle, the mold, and part of the bar to 3 m below the meniscus.
- (2) Once the velocity field was obtained, the transport equation for a chemical element in a turbulent stream was tion for a chemical element in a turbulent stream was Convection-diffusion region: calculated by solving a transient turbulent convection diffusion equation:

$$
\frac{\partial C}{\partial t} + \underline{v} \cdot \underline{\nabla} C = \underline{\nabla} \cdot [(D + D') \underline{\nabla} C] \tag{5}
$$

species; *y* is the time-averaged velocity; *D* is the molecular surface.
diffusivity for the species; and *D'* is the turbulent diffusivity. The complete solution of Eq. [5] proved that diffusive diffusivity for the species; and D^t is the turbulent diffusivity. The value of D^t was considered to be equal to the turbulent effects can also be neglected in the liquid pool. This is due $\sqrt{\rho}$ (μ^t : turbulent viscosity given by the *k*- ε model, and ρ : density) and $D \ll D^t$. Therefore, part of the line and was verified by direct measurements of all species have the same concentration distribution behavior. steel composition performed in the plant (Figure 8).

Petrov Galerkin technique,^[18,19] and a standard eight-node lated by convection of the elements concentration due to (3-D) isoparametric finite element discretization^[20] for c. the casting speed and was considered ho (3-D) isoparametric finite element discretization^[20] for *c*. the casting speed Free parameters of the 1-D model were calibrated with the bar section.

Free parameters of the 1-D model were calibrated with the concentration obtained by these 3-D calculations. The following values were found:
 $V_m = \pi \phi^2 \times 0.5 \text{ m};$

D. *Final Calculations*

Having calibrated

 $D_{\text{eff}} \approx 30 \text{ mm}^2/\text{seg}$ as a function of time.

tion distribution for different time intervals in the azimuthal (1) casting speed as the piecewise function of time in each plane. This figure was obtained in a simulation with a casting casting line;

Fig. 7—Comparison of dimensionless concentration at 1.56 m below meniscus as given by the 1-D volumes model and the 3-D turbulent model.

speed of 2 m/min and a mold diameter of 0.170 m. The concentration imposed at the nozzle was given by the tundish model, assuming a typical double dilution process.

Results from the 3-D numerical model show an almost 1-D flow below a distance of 1.5 m from the meniscus. In Figure 7, the dimensionless concentration at 1.56 m below the meniscus is plotted as a function of time. In this figure, lines correspond to 1-D model results and symbols to 3-D model results. The good agreement observed validates the use of the 1-D model. The abrupt slope change in both lines is due to the double dilution process already stated.

C. *Final Composition of the Bar*

The third submodel simulates the lower part of the liquid pool (from 1.56 m below the meniscus to the metallurgical Fig. 6—Concentration in the system nozzle mold bar for different time length). It solves the convection diffusion equation (Eq. [5]) instants. using the turbulent diffusivity given by the 3-D numerical model. Diffusion in solid state is neglected.

The solidification point would be different for each point where *c* is the time-averaged concentration for a chemical of the bar cross section depending on its distance to the

to the low values of the turbulent diffusivity in the lower part of the line and was verified by direct measurements of

Equation [5] is integrated with the Streamline Upwind Consequently, the composition along the bar was calcu-
trov Galerkin technique.^[18,19] and a standard eight-node lated by convection of the elements concentration du

Having calibrated the model, a numerical code $V_p = \pi \phi^2 \times 1.06 \text{ m}$; and $\text{(GRADE}^{[21]})$ was developed. The code calculates the dimensionless concentration $\tilde{C}(t)$ at the cut-off point of the bar,

The user is supposed to provide the following input data: Figure 6 shows the external line dimensionless concentra-

Fig. 8—(*a*) and (*b*) Experimental measurements along the downgraded bars (internal line) for different points of the bar cross section of example A.

-
- (3) concentration of each element in the old and new grades; of Table III and the chemical composition of Table IV.
(4) specification of the composition of both steel grades: and In this case, B and Ti were the critical el
- (4) specification of the composition of both steel grades; and
-

direct composition measurements of round billets from the intermix region. This validates the assumption of negligible Siderca S.A.I.C. continuous caster, performed with an opti- diffusion in the lower part of the line discussed in Section IV. cal emission in vacuum spectrometer. Typical precisions of Figure 9 shows (for an external line) the concentration of

process. This process is the standard procedure adopted length of the intermixed bar. by Siderca. In order to represent the deviation of our numerical results

Table II. Experimental Precision in Chemical Composition Measurements in the Bars

C_{old}	Element	Units	Error
▲	C	pct	0.012
	Mn	pct	0.05
▲	S	pct	0.001
	P	pct	0.0016
	Si	pct	0.015
	Ni	pct	0.004
	Cr	pct	0.025
Ŧ $C_{\sf new}$	Mo	pct	0.008
12 20	Sn	pct	0.001
16	Ti	pct	0.0002
ilong the line [m]	Al	pct	0.005
$\left(a\right)$	B	pct	0.00002
	Ca	pct	0.00015
	Cu	pct	0.008
C			

A. *Plant Measurements—Example A*

(2) flow rate entering the tundish as the piecewise function The first set of measurements to be analyzed corresponds of time;
concentration of each element in the old and new grades: of Table III and the chemical composition of Table IV.

(5) diameter of the bar. they were abundant in the old grade and had stringent restric-
tions for maximum concentration in the new grade. After With this information, the program is able to detect the
old grade/intermixed steel boundary and the new grade/
old grade/intermixed steel boundary in the bar at the cut-off point.
The program also calculates the proportio

Ine program also calculates the proportion of new grade
steel and old grade steel in the intermixed bar, which is
needed to estimate the cost of the downgraded steel.
Inne. The concentration of each element on these 16 sam was measured at the centerline, at the surface, and at a point **V. COMPARISON WITH PLANT** placed between them. The resulting concentrations of Ti and
 NEASUREMENTS B at different points of the bars are plotted on Figure 8. The

figure shows that the differences in the composition of The final validation of the model was accomplished by across the bar section are not important in determining the

these measurements are shown in Table II. different chemical elements, at the cut-off point, as a function Two different changes of grade were analyzed, the first of the length of steel bar cast since the moment of the new one having B and Ti as the critical elements and the second ladle first opening. The continuous lines are the results of one having Cr and Mo as the critical elements. the numerical model and the symbols represent experimental In both cases, the filling of the tundish (after the drainage measurements. The good agreement observed indicates that due to the ladle change) was performed in a double dilution the numerical model describes correctly the position and

Table IV. Chemical Composition of Example A

Element	Units	Old Grade		New Grade			
		Minimum	Maximum	Actual	Minimum	Maximum	Actual
C	pct	0.24	0.27	0.26	0.27	0.3	0.29
Mn	pct	1.3	1.45	1.33	1.25	1.4	1.31
S	pct	Ω	0.01	0.002	Ω	0.01	0.006
P	pct	0	0.025	0.016	Ω	0.02	0.019
Si	pct	0.25	0.35	0.29	0.3	0.4	0.36
Ni	pct		0.15	0.06	0	0.15	0.06
Cr	pct		0.2	0.03	0	0.16	0.05
Mo	pct		0.08	0.02	0	0.08	0.01
V	pct		0.01	0.003		0.01	0.002
Nb	pct		0.005	0.001		0.005	0.001
Ti	pct	0.015	0.035	0.022	0	0.014	0.003
Al	pct	0.01	0.035	0.021	0.01	0.03	0.018
B	pct	0.0015	0.0025	0.0018	0	0.0005	0.0001
N	pct	0	0.009	0.0048	0	0.009	0.004
Ca	pct		0.01	0.0013	0	0.01	0.0017
Cu	pct	θ	0.22	0.14	θ	0.25	0.17

chemical composition for external line downgraded bars. Four elements were considered: Ti (1000 times), B (10,000 times), S (1000 times), and

from the experimental data, the following parameter is wall across the bar section. In this case, Cr and Mo were the more critical elements. Figure 10 shows the numerical results (lines) and the

$$
\mathcal{S}^{(e)} = \frac{\sqrt{\sum_{i=1}^{N} (C_{\text{Exp},i}^{(e)} - C_{\text{Num},i}^{(e)})^2}}{\sqrt{\sum_{i=1}^{N} (C_{\text{Exp},i}^{(e)})^2}}
$$
 [6]

element \overline{e}) at different positions of the bar, as given by during the double dilution process, was successfully experimental measurements and numerical model, respectively reproduced. experimental measurements and numerical model, respec-

 $\delta^{(B)} = 0.182$ and $\delta^{(T)} = 0.162$. These values were calculated of concentrations. out after forging the samples.

Table V. Casting Conditions of Example B

W_{tundish}	13.0 tons	
W_{drain}^1	5.1 tons	
$W^{\rm l}$	7.3 tons	
W_{drain}^2	5.3 tons	
Q_{steady}	1.0 ton/min	
\mathcal{Q}_{dd}^1	2.9 ton/min	
Q_{dd}^2	0.1 ton/min	
Q_{dd}^3	3.5 ton/min	
Bar diameter	0.148 m	
Casting speed	2 m/min	

B. *Plant Measurements—Example B*

The second grade change analyzed is described by Tables V and VI and corresponds to a much longer intermix (six bars were downgraded on each line). Again, four samples were taken from each downgraded bar of an external line Fig. 9—Numerical results (lines) and plant measurements (symbols) of and of an internal line. This time, however, only one mea-
chemical composition for external line downgraded bars. Four elements surement was performed o were considered: Ti (1000 times), B (10,000 times), S (1000 times), and was taken after forging the samples, which is the usual procedure carried out in plants to measure steel composition. Thus, the measured concentration is, in fact, an average of its value across the bar section. In this case, Cr and Mo

 $\delta^{(e)} = \frac{\sqrt{\sum_{i=1}^{N} (C_{\text{Exp},i}^{(e)} - C_{\text{Num},i}^{(e)})^2}}{\sqrt{\sum_{i=1}^{N} (C_{\text{Exp},i}^{(e)} - C_{\text{Num},i}^{(e)})^2}}$ [6] experimental points (symbols) for four different chemical species on the intermixed steel bars. The figure shows that t [6] the model is able not only to make a good estimation of the position and length of the intermixed bars, but also to describe accurately the concentration profile along the bars. where $C_{\text{Exp},i}^{(e)}$ and $C_{\text{Num},i}^{(e)}$ are concentrations of the chemical Even the concentration plateau, due to the ladle closing

tively; and *N* is the total number of measurements. In order to represent the deviation of our numerical results
For the critical elements of this example, we obtained from the experimental data, the $\delta^{(e)}$ parameter (For the critical elements of this example, we obtained from the experimental data, the $\delta^{(e)}$ parameter (Eq. [6]) is B = 0.182 and $\delta^{(T)}$ = 0.162. These values were calculated cludated for the critical elements: δ considering surface, midpoint, and centerline measurements $\delta^{\text{(Mo)}} = 0.079$. In this example, measurements were carried

Table VI. Chemical Composition of Example B

Element	Units	Old Grade		New Grade			
		Minimum	Maximum	Actual	Minimum	Maximum	Actual
\mathcal{C}	pct	0.09	0.13	0.12	0.11	0.14	0.13
Mn	pct	0.4	0.5	0.46		1.15	1.03
S	pct	0	0.01	0.001		0.005	0.005
P	pct	0	0.015	0.01		0.02	0.012
Si	pct	0.12	0.23	0.17	0.3	0.4	0.32
Ni	pct	0	0.15	0	$^{(1)}$	0.15	0.0
Cr	pct	2.03	2.13	2.11		0.16	0.11
Mo	pct	0.92	0.97	0.94		0.08	0.01
V	pct	0	0.01	0.005		0.01	0.001
Nb	pct	0	0.005	0.004	0	0.005	0.001
Ti	pct	0	0.005	0.002	0	0.005	0.004
Al	pct	0.01	0.035	0.023	0.015	0.04	0.028
B	pct	0	0.0005	0.0001		0.0005	0.0001
$\mathbf N$	pct	0	0.01	0.0057		0.008	0.0048
Ca	pct	0	0.01	0.0024		0.01	0.0016
Cu	pct	0	0.15	0.006	0	0.2	0.16

tions. Now, the same grade change (shown in Table VI) will be simulated with a simple dilution procedure (Table VII). **VIII. CONCLUSIONS**

Figure 11 shows the concentration of Cr along the bar as
calculated by simulating simple dilution and double dilution
procedure ous caster of Siderca was studied and a numerical model
procedures. The advantage of the doubl procedures. The advantage of the double dilution procedure
becomes evident since its curve reaches the critical concentration of the position and length of the steel bar to
tration before the simple dilution curve does. In on each external line when a double dilution procedure is 1. A methodology to model the mixing in the tundish with used. A total amount of 4 tons of steel was saved.

The model was employed in a particular configuration of a four line round billet caster and it can simulate different assume including time-dependent casting speed and flow rate conditions on entering the tundish. In parti

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-
-

configuration of the caster is changed, for example, due to is probably due to the relatively small volume of liquid changes in the size or shape of the tundish, changes in the steel present in each casting line of the modeled cases, flow control devices in the tundish, or changes in the diame- whose diameters are 0.148 or 0.170 m. Both numerical

The model can be extended to include casters with any number of lines (not necessarily symmetric) by adding a $\qquad 6.$ The low values of the deviation parameter $\delta^{(e)}$ (defined by group of volumes associated to each line in the tundish Eq. [6]) show the good agreement between the numerical

VI. TUNDISH FILLING PROCEDURES model, according to Figure 12. In order to allow the recircula-

To illustrate the advantage of filling the tundish by the
double dilution procedure, example B in the previous section
is going to be considered again together with a simple dilu-
tion procedure example.
In Section V-B, a

- several continuous casting lines is presented, in which a defined group of volumes is associated with each line. **VII. MODEL EXTENSION** Mixing volumes connect the different groups to allow recirculation of fluid.
	-
	-
- (1) casting speed,

(2) minimum level of the liquid metal maintained in the

tundish,

(3) flow rate entering the tundish

(4) dilution strategies (simple, double, or other type of

(5) grades being cast, and

(6) casting
- A new calibration needs to be performed any time the 5. Most of the steel mixing takes place in the tundish. This ter of the mold. Calculations and measurements show that diffusion
The model can be extended to include casters with any effects can be neglected in most part of the bar.
	-

Fig. 10—Numerical results (lines) compared with plant measurements (symbols) for four different chemical elements of example B.

Fig. 11—(*a*) and (*b*) Concentration of Cr along the bar for the grade transition given by Table VI. The figure compares results obtained with the simple dilution process (dashed line) and double dilution process (full line). The dotted line represents the maximum concentration of Cr allowed in the new grade.

results and the measurements carried out in a round billet caster.

LIST OF SYMBOLS

Fig. 13—Mixing volume connecting groups associated to different lines. Society, Warrendale, PA, 1996, pp. 519-31.

-
-
-
-
-
- V

total number of measurements in Eq. [6].
 Q flow rate between different volumes of the the model of the model and A Garamendy: "Festivale and A Garamendy: "Estudio en Modelo de Agua del Flujo
-
-
-
-
- *v* casting speed pp. 441-55.
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- *z_i* beginning of the diffusion-convection region *on Continuous Casting*, Madrid, 1998.

end of the diffusion-convection region 16. M.B. Goldschmit, R.J. Príncipe, and M.
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- ext external line and argentina, 1999.

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time delay introduced by plug flow volumes

IAS, San Nicolás, Argentina, 1997 (in Spanish).

1997 (in Spanish).
- *t_p* time delay introduced by plug flow volumes.
 \hat{t} timensionless time
 \hat{t} dimensionless time
 \hat{t} volume
 \hat{t} at 11).
 \hat{t} at 1
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- *x_i* 14. M.B. Goldschmit and R. Javier Principe: *IV World Congr. on Computa*-
distance to the meniscus, measured along the bar *ional Mechanics*, Buenos Aires, Argentina, 1998.
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