

# A thermodynamic model for predicting Ti and Al element of SiO<sub>2</sub>-Al<sub>2</sub>O<sub>3</sub>-FeO-TiO<sub>2</sub>-CaO-MgO-CaF<sub>2</sub> slag in electroslag remelting process

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## Original Article

# A thermodynamic model for predicting Ti and Al element of $\text{SiO}_2\text{--Al}_2\text{O}_3\text{--FeO--TiO}_2\text{--CaO--MgO--CaF}_2$ slag in electroslag remelting process

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## ABSTRACT

The control of titanium and aluminium in nickel-base superalloy with high titanium and low aluminium during the electroslag remelting (ESR) process has not been resolved well so far. The loss of titanium (Ti) and aluminium (Al) during the ESR process is critical to improve the high-temperature performance of GH4065A superalloy. A thermodynamic model based on the ion and molecule coexistence theory (IMCT) was developed to calculate the Ti and Al content of  $\text{SiO}_2\text{--Al}_2\text{O}_3\text{--FeO--TiO}_2\text{--CaO--MgO--CaF}_2$  slag system during the ESR process of GH4065A superalloy. Results show that the order of the ability of each element in the slag system to weaken the loss of Al content is in the following order  $\text{Al}_2\text{O}_3 > \text{CaO} > \text{CaF}_2$ , enhance the loss of Al content is  $\text{SiO}_2 > \text{FeO} > \text{TiO}_2$ ,  $\text{CaF}_2 > \text{MgO}$  and enhance the loss of Ti content is  $\text{FeO} > \text{SiO}_2 > \text{Al}_2\text{O}_3 > \text{CaO}$ . Considering the influence of the slag system on the elements Al and Ti, the optimal ratio of the slag system is  $\text{CaO} = 15\% \sim 20\%$ ,  $\text{CaF}_2 \geq 60\%$ ,  $\text{Al}_2\text{O}_3 = 18\% \sim 25\%$ ,  $\text{SiO}_2 \leq 0.5\%$ ,  $\text{TiO}_2 = 1\% + (0.2\% \sim 0.6\%)$ ,  $\text{MgO} \geq 4\%$ ,  $\text{FeO} \leq 0.1\%$ .

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## 1. Introduction

The combination of vacuum induction melting (VIM), electroslag remelting (ESR), and vacuum arc remelting (VAR) technique are proposed to produce the GH4065A superalloy in

recent years. It is well known that the advantage during the ESR process is the cleanness and homogeneity of ingot. However, the Ti and Al element is often oxidized during the ESR process of GH4065A superalloy [1]. The main reason is the chemical reaction between O element in slag and Ti or Al elements in superalloy. There are three reasons explaining this

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reaction [2–4]: (1) Self-consuming electrode carries oxygen; (2) Oxidation of slag system is too high; (3) Ti and Al element in GH4065A superalloy are prone to react with unstable oxides in slag system.

The primary approach for decreasing of Ti and Al content in GH4065A superalloy is to reduce the reaction with O elements during the ESR process [5]. The Ti and Al element are distributed unevenly from the top to the bottom of ingot if without appropriate slag. The high-temperature performance of alloy is also weakened accordingly. Therefore, it is crucial to further understand the effect of slag on Ti and Al in superalloy [6].

Some studies [7–9] investigated the effect of slag on the key elements, such as Ti, Si, or Al in the metallurgical process. Chen et al. [10] investigated the titanium loss during the ESR process of A286 and V57 superalloys through experiments method. Pateisky et al. [11] analyzed the reaction of titanium and silicon with the alumina-calcium oxide-fluorspar slags system. They proposed the optimal solutions for reducing the loss of Ti and Si. Park et al. [12] investigated the thermodynamic equilibrium between CaO–SiO<sub>2</sub>–Al<sub>2</sub>O<sub>3</sub>–MgO–CaF<sub>2</sub>–(TiOx) slags and the Fe–11mass%Cr melt to understand the thermodynamic behavior of titanium oxide in refining slag. As a critical path to predict the composition content in slag, the ion-molecular coexistence theory (IMCT) method has been developed [13]. Yang et al. [14] developed a thermodynamic model for predicting the phosphate capacity of CaO–SiO<sub>2</sub>–MgO–FeO–Fe<sub>2</sub>O<sub>3</sub>–MnO–Al<sub>2</sub>O<sub>3</sub>–P<sub>2</sub>O<sub>5</sub> slag at the steelmaking endpoint during an 80-ton top/bottom combined blown converter steelmaking process based on the IMCT. Duan et al. [15] developed a thermodynamic model to calculate the mass action concentration of ion-coupling agents or structural units in slag based on the IMCT. The developed thermodynamic model for calculating the manganese distribution ratio can determine the respective manganese distribution ratio and capacity of four demanganization products quantitatively. Li et al. [16] predicts the sulfide capacity of CaO–SiO<sub>2</sub>–MgO–Al<sub>2</sub>O<sub>3</sub> ironmaking slag and validated two sets of sulfide capacity data for CaO–SiO<sub>2</sub>–MgO–Al<sub>2</sub>O<sub>3</sub> ironmaking slag based on IMCT. This model can accurately predict the distribution ratio of metallurgical slags. Moreover, several researchers have taken the IMCT method into slag prediction during the ESR process. Yin et al. [17] analyzed the control of aluminum and titanium elements during the electroslag remelting of high titanium and low aluminum alloy Inconel 718. Hou et al. [18] focused on designing appropriate slag for controlling titanium content during 1Cr21Ni5Ti remelting based on the ion and molecule coexistence theory.

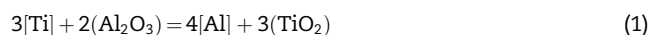
In this paper, a thermodynamic model to calculate the Ti and Al content with SiO<sub>2</sub>–Al<sub>2</sub>O<sub>3</sub>–FeO–TiO<sub>2</sub>–CaO–MgO–CaF<sub>2</sub> slag system during the ESR process of GH4065A superalloy based on the IMCT is developed. Since the Ti content during the actual ESR process is often loss, the extra TiO<sub>2</sub> of 1 w% will be added in the slag system to keep the Ti content. Moreover, the influence of composition and temperature of the

SiO<sub>2</sub>–Al<sub>2</sub>O<sub>3</sub>–FeO–TiO<sub>2</sub>–CaO–MgO–CaF<sub>2</sub> slag system on the Ti and Al content are analyzed. If the Ti and Al content in industrial ingot could be predicted and controlled, it is beneficial to improve the high-temperature performance of GH4065A superalloy.

## 2. Thermodynamic model

The chemical composition of GH4065A superalloy is listed in Table 1. During the actual industrial process, the slag system is specific and selected [19]. The slag system used for GH4065A superalloy during the ESR process is shown in Table 2. Moreover, since the Ti content during the ESR process is often reduced significantly, the 1 w% TiO<sub>2</sub> is added to the slag system to lessen the oxidization of titanium.

The reference temperature used in this model are 1873 K, 1923 K, and 1973 K, respectively. In this paper, the decrease of Ti and Al is not only accounted for the oxidation reaction but also the reaction between Ti and FeO or SiO<sub>2</sub> in slag system as follows [14,17,18,20]:



$$\lg K = \lg \frac{a_{\text{TiO}_2}^3}{a_{\text{Ti}}^3 \cdot a_{\text{Al}_2\text{O}_3}^2} = \lg \frac{1}{f_{\text{Ti}}^3 [\text{Ti}]^3} + \lg \frac{a_{\text{TiO}_2}^3}{a_{\text{Al}_2\text{O}_3}^2} = -\frac{35300}{T} + 9.94 \quad (2)$$



$$\lg K = \lg \frac{a_{\text{TiO}_2}}{a_{\text{Ti}} \cdot a_{\text{FeO}}^2} = \lg \frac{1}{f_{\text{Ti}} [\text{Ti}]^2} + \lg \frac{a_{\text{TiO}_2}}{a_{\text{FeO}}^2} = \frac{34990.6}{T} - 10.49 \quad (4)$$



$$\lg K = \lg \frac{a_{\text{TiO}_2}}{a_{\text{Ti}} \cdot a_{\text{SiO}_2}} = \lg \frac{1}{f_{\text{Ti}} [\text{Ti}]^2} + \lg \frac{a_{\text{TiO}_2}}{a_{\text{FeO}}} = \frac{180}{T} + 1.36 \quad (6)$$

The effect of slag system and temperature on Ti oxidation is given as:

$$\lg [\text{Ti}] = \frac{1}{3} \left\{ \lg \frac{a_{\text{TiO}_2}^3}{a_{\text{Al}_2\text{O}_3}^2} - 3 \lg f_{\text{Ti}} - \left( -\frac{35300}{T} + 9.94 \right) \right\} \quad (7)$$

$$\lg [\text{Ti}] = \left\{ \lg \frac{a_{\text{TiO}_2}}{a_{\text{FeO}}^2} - \lg f_{\text{Ti}} - \left( \frac{34990.6}{T} - 10.49 \right) \right\} \quad (8)$$

**Table 2 – Chemical composition of slag system (w%).**

SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	FeO	TiO <sub>2</sub>	CaO	MgO	CaF <sub>2</sub>
0.5	21.2	0.1	0.2	18.0	2.0	58.0

**Table 1 – Chemical composition of the GH4065A superalloy (w%).**

C	Co	Al	W	Zr	Cr	Mo	Ti	Nb	Ni	B
0.008	13.0	2.13	4.0	0.045	16.0	4.0	3.73	0.7	56.371	0.016

**Table 3 – Interaction coefficients  $e_i^j$  of the components.**

$e_i^j$	C	Cr	Al	Ti	Mo	Ni
Al	0.091	0.0212	0.0800	0.004	–	–0.0376
Si	–	0.0251 [26]	0.0589 [26]	–	–	–0.0100
Ti	–0.19	0.0250	0.0037	0.0561	0.016	–0.0166

$$\lg[\text{Ti}] = \left\{ \lg \frac{a_{\text{TiO}_2}}{a_{\text{SiO}_2}} - \lg f_{\text{Al}} - \left( \frac{180}{T} + 1.36 \right) \right\} \quad (9)$$

When the reaction reaches equilibrium at the slag-metal interface, the content of Ti element in superalloy can be approximated as the sum of Eqs. (2), (4) and (6) shown as:

$$\lg[\text{Ti}] = \frac{1}{3} \left\{ \lg \frac{a_{\text{TiO}_2}^3}{a_{\text{Al}_2\text{O}_3}^2 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3} - 3 \lg f_{\text{Ti}} - \left( \frac{23403.9}{T} - 5.81 \right) \right\} \quad (10)$$

where  $K$  denotes the reaction equilibrium constant;  $a_{M_xO_y}$  is the activity of the component  $M_xO_y$ ;  $T$  represents the reaction temperature;  $f_x$  is the activity coefficient.



$$\lg K = \lg \frac{a_{\text{Al}_2\text{O}_3}}{a_{\text{Al}}^2 \cdot a_{\text{FeO}}^3} = \lg \frac{1}{f_{\text{Al}}^2 [\text{Al}]^2} + \lg \frac{a_{\text{Al}_2\text{O}_3}^3}{a_{\text{FeO}}^3} = \frac{44383.1}{T} - 11.44 \quad (12)$$



$$\lg K = \lg \frac{a_{\text{Al}_2\text{O}_3}^2}{a_{\text{TiO}_2}^3} = \lg \frac{1}{f_{\text{Al}}^4 [\text{Al}]^2} + \lg \frac{a_{\text{Al}_2\text{O}_3}^2}{a_{\text{SiO}_2}^2} = \frac{35840}{T} - 5.86 \quad (14)$$

The effect of slag system and temperature on the Al oxidation is listed:

$$\lg[\text{Al}] = \frac{1}{4} \left\{ \lg \frac{a_{\text{Al}_2\text{O}_3}^2}{a_{\text{TiO}_2}^3} + 4 \lg f_{\text{Al}} - \left( -\frac{35300}{T} + 9.94 \right) \right\} \quad (15)$$

$$\lg[\text{Al}] = \frac{1}{2} \left\{ \lg \frac{a_{\text{Al}_2\text{O}_3}}{a_{\text{FeO}}^3} - 2 \lg f_{\text{Ti}} - \left( \frac{44383.1}{T} - 11.44 \right) \right\} \quad (16)$$

$$\lg[\text{Al}] = \frac{1}{4} \left\{ \lg \frac{a_{\text{Al}_2\text{O}_3}^2}{a_{\text{SiO}_2}^3} - 4 \lg f_{\text{Al}} - \left( \frac{35840}{T} - 5.86 \right) \right\} \quad (17)$$

When a reaction reaches equilibrium at the slag-metal interface, the content of the Al element in superalloy can be approximated as the sum of Eqs. (12), (14) and (15) shown as:

$$\lg[\text{Al}] = \frac{1}{4} \left\{ \lg \frac{a_{\text{Al}_2\text{O}_3}^6}{a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3} - 3 \lg f_{\text{Al}} - \left( \frac{39976.55}{T} - 9.67 \right) \right\} \quad (18)$$

where  $K$  indicates the reaction equilibrium constant;  $a_{M_xO_y}$  denotes the activity of the component  $M_xO_y$ ;  $T$  is the reaction temperature;  $f_x$  is the activity coefficient. Which can be expressed by Eq. (19).

$$\lg f_i = \sum e_i^j \omega_j \quad (19)$$

where  $\omega_j$  expresses the mass fraction of the component  $j$  in the superalloy;  $f_i$  denotes the activity coefficient of component  $i$ ;  $e_i^j$  indicates the interaction coefficient between the two components of  $i$  and  $j$  in the superalloy [21–25], as shown in Table 3.

Based on the IMCT, the remelting slag includes simple ions, simple molecules, and complex molecules, which are in a delicate dynamic equilibrium relationship [27]. The relevant compounds and chemical reactions in the slag are listed in Tables 4 and 5. Since an extra 1w%  $\text{TiO}_2$  was added to the slag, the mole number of seven components in 101 g slag could be expressed as  $c_1 = n_{\text{CaO}}^0$ ,  $c_2 = n_{\text{CaF}_2}^0$ ,  $c_3 = n_{\text{Al}_2\text{O}_3}^0$ ,  $c_4 = n_{\text{SiO}_2}^0$ ,  $c_5 = n_{\text{TiO}_2}^0$ ,  $c_6 = n_{\text{MgO}}^0$ ,  $c_7 = n_{\text{FeO}}^0$ . The mass action concentration of components can be calculated as :

$$n_i^0 = \frac{n_i}{\sum n_i} \quad (20)$$

The following equilibrium reaction Eqs. (14)–(21) can be established based on the reaction mass conservation law with IMCT method. Taking Eqs. (13) into ((14–21)), the  $N_1, N_2, N_3, N_4, N_5, N_6, N_7$  after reaction can be calculated and obtained by the Matlab® software.

$$K_{ci} = \exp(-\Delta G_i^0 / RT) \quad (21)$$

$$N_1 + N_2 + N_3 + \dots + N_7 + N_{b1} + N_{b2} + N_{b3} \dots + N_{b36} = \sum N_i = 1.01 \quad (22)$$

$$c_1 = \left( \begin{array}{l} 0.5N_1 + N_{b1} + N_{b3} + 2N_{b5} + 3N_{b7} + 12N_{b8} + N_{b9} + N_{b10} + 3N_{b12} + N_{b13} + 3N_{b14} \\ + 4N_{b15} + 3N_{b20} + N_{b21} + 2N_{b22} + 3N_{b23} + N_{b24} + 2N_{b25} + 3N_{b26} + N_{b27} + 11N_{b28} \\ + N_{b29} + 3N_{b30} \end{array} \right) \sum n_i = n_{\text{CaO}}^0 \quad (23)$$

**Table 4 – Molar number and mass action concentration in 101 g SiO<sub>2</sub>-Al<sub>2</sub>O<sub>3</sub>-FeO-TiO<sub>2</sub>-CaO-MgO-CaF<sub>2</sub> slags based on IMCT [26,28-30].**

Item	Structural unit	Symbols	Molar number $n_i$ /mol	Mass action concentration of structural units $N_i$
Simple ion	Ca <sup>2+</sup> + O <sup>2-</sup>	1	$n_1 = n_{Ca^{2+},CaO} = n_{O^{2-},CaO}$	$N_1 = \frac{n_1}{\sum n_i} = N_{CaO}$
	Ca <sup>2+</sup> + 2F <sup>-</sup>	2	$n_2 = n_{Ca^{2+},CaF_2} = 2n_{F^-,CaF_2}$	$N_2 = \frac{3n_2}{\sum n_i} = N_{CaF_2}$
	Mg <sup>2+</sup> + O <sup>2-</sup>	6	$n_6 = n_{Mg^{2+},MgO} = n_{O^{2-},MgO}$	$N_6 = \frac{2n_6}{\sum n_i} = N_{MgO}$
	Fe <sup>2+</sup> + O <sup>2-</sup>	7	$n_7 = n_{Fe^{2+},FeO} = n_{O^{2-},FeO}$	$N_7 = \frac{2n_7}{\sum n_i} = N_{FeO}$
Simple and complex molecule	Al <sub>2</sub> O <sub>3</sub>	3	$n_3 = n_{Al_2O_3}$	$N_3 = \frac{n_3}{\sum n_i} = N_{Al_2O_3}$
	SiO <sub>2</sub>	4	$n_4 = n_{SiO_2}$	$N_4 = \frac{n_4}{\sum n_i} = N_{SiO_2}$
	TiO <sub>2</sub>	5	$n_5 = n_{TiO_2}$	$N_5 = \frac{n_5}{\sum n_i} = N_{TiO_2}$
	CaO·SiO <sub>2</sub>	b1	$n_{b1} = n_{CaO \cdot SiO_2}$	$N_{b1} = \frac{n_{b1}}{\sum n_i} = N_{CaO \cdot SiO_2}$
	MgO·SiO <sub>2</sub>	b2	$n_{b2} = n_{MgO \cdot SiO_2}$	$N_{b2} = \frac{n_{b2}}{\sum n_i} = N_{MgO \cdot SiO_2}$
	CaO·Al <sub>2</sub> O <sub>3</sub>	b3	$n_{b3} = n_{CaO \cdot Al_2O_3}$	$N_{b3} = \frac{n_{b3}}{\sum n_i} = N_{CaO \cdot Al_2O_3}$
	MgO·Al <sub>2</sub> O <sub>3</sub>	b4	$n_{b4} = n_{MgO \cdot Al_2O_3}$	$N_{b4} = \frac{n_{b4}}{\sum n_i} = N_{MgO \cdot Al_2O_3}$
	2CaO·SiO <sub>2</sub>	b5	$n_{b5} = n_{2CaO \cdot SiO_2}$	$N_{b5} = \frac{n_{b5}}{\sum n_i} = N_{2CaO \cdot SiO_2}$
	2MgO·SiO <sub>2</sub>	b6	$n_{b6} = n_{2MgO \cdot SiO_2}$	$N_{b6} = \frac{n_{b6}}{\sum n_i} = N_{2MgO \cdot SiO_2}$
	3CaO·Al <sub>2</sub> O <sub>3</sub>	b7	$n_{b7} = n_{3CaO \cdot Al_2O_3}$	$N_{b7} = \frac{n_{b7}}{\sum n_i} = N_{3CaO \cdot Al_2O_3}$
	12CaO·7Al <sub>2</sub> O <sub>3</sub>	b8	$n_{b8} = n_{12CaO \cdot 7Al_2O_3}$	$N_{b8} = \frac{n_{b8}}{\sum n_i} = N_{12CaO \cdot 7Al_2O_3}$
	CaO·2Al <sub>2</sub> O <sub>3</sub>	b9	$n_{b9} = n_{CaO \cdot 2Al_2O_3}$	$N_{b9} = \frac{n_{b9}}{\sum n_i} = N_{CaO \cdot 2Al_2O_3}$
	CaO·6Al <sub>2</sub> O <sub>3</sub>	b10	$n_{b10} = n_{CaO \cdot 6Al_2O_3}$	$N_{b10} = \frac{n_{b10}}{\sum n_i} = N_{CaO \cdot 6Al_2O_3}$
	3Al <sub>2</sub> O <sub>3</sub> ·2SiO <sub>2</sub>	b11	$n_{b11} = n_{3Al_2O_3 \cdot 2SiO_2}$	$N_{b11} = \frac{n_{b11}}{\sum n_i} = N_{3Al_2O_3 \cdot 2SiO_2}$
	3Al <sub>2</sub> O <sub>3</sub> ·SiO <sub>2</sub>	b12	$n_{b12} = n_{3Al_2O_3 \cdot SiO_2}$	$N_{b12} = \frac{n_{b12}}{\sum n_i} = N_{3Al_2O_3 \cdot SiO_2}$
	CaO·TiO <sub>2</sub>	b13	$n_{b13} = n_{CaO \cdot TiO_2}$	$N_{b13} = \frac{n_{b13}}{\sum n_i} = N_{CaO \cdot TiO_2}$
	3CaO·2TiO <sub>2</sub>	b14	$n_{b14} = n_{3CaO \cdot 2TiO_2}$	$N_{b14} = \frac{n_{b14}}{\sum n_i} = N_{3CaO \cdot 2TiO_2}$
	4CaO·3TiO <sub>2</sub>	b15	$n_{b15} = n_{4CaO \cdot 3TiO_2}$	$N_{b15} = \frac{n_{b15}}{\sum n_i} = N_{4CaO \cdot 3TiO_2}$
	Al <sub>2</sub> O <sub>3</sub> ·TiO <sub>2</sub>	b16	$n_{b16} = n_{Al_2O_3 \cdot TiO_2}$	$N_{b16} = \frac{n_{b16}}{\sum n_i} = N_{Al_2O_3 \cdot TiO_2}$
	MgO·TiO <sub>2</sub>	b17	$n_{b17} = n_{MgO \cdot TiO_2}$	$N_{b17} = \frac{n_{b17}}{\sum n_i} = N_{MgO \cdot TiO_2}$
	MgO·2TiO <sub>2</sub>	b18	$n_{b18} = n_{MgO \cdot 2TiO_2}$	$N_{b18} = \frac{n_{b18}}{\sum n_i} = N_{MgO \cdot 2TiO_2}$
	2MgO·TiO <sub>2</sub>	b19	$n_{b19} = n_{2MgO \cdot TiO_2}$	$N_{b19} = \frac{n_{b19}}{\sum n_i} = N_{2MgO \cdot TiO_2}$
	3CaO·2SiO <sub>2</sub>	b20	$n_{b20} = n_{3CaO \cdot 2SiO_2}$	$N_{b20} = \frac{n_{b20}}{\sum n_i} = N_{3CaO \cdot 2SiO_2}$
	CaO·MgO·2SiO <sub>2</sub>	b21	$n_{b21} = n_{CaO \cdot MgO \cdot 2SiO_2}$	$N_{b21} = \frac{n_{b21}}{\sum n_i} = N_{CaO \cdot MgO \cdot 2SiO_2}$
	2CaO·MgO·2SiO <sub>2</sub>	b22	$n_{b22} = n_{2CaO \cdot MgO \cdot 2SiO_2}$	$N_{b22} = \frac{n_{b22}}{\sum n_i} = N_{2CaO \cdot MgO \cdot 2SiO_2}$
	3CaO·MgO·2SiO <sub>2</sub>	b23	$n_{b23} = n_{3CaO \cdot MgO \cdot 2SiO_2}$	$N_{b23} = \frac{n_{b23}}{\sum n_i} = N_{3CaO \cdot MgO \cdot 2SiO_2}$
	CaO·Al <sub>2</sub> O <sub>3</sub> ·2SiO <sub>2</sub>	b24	$n_{b24} = n_{CaO \cdot Al_2O_3 \cdot 2SiO_2}$	$N_{b24} = \frac{n_{b24}}{\sum n_i} = N_{CaO \cdot Al_2O_3 \cdot 2SiO_2}$
	2CaO·Al <sub>2</sub> O <sub>3</sub> ·SiO <sub>2</sub>	b25	$n_{b25} = n_{2CaO \cdot Al_2O_3 \cdot SiO_2}$	$N_{b25} = \frac{n_{b25}}{\sum n_i} = N_{2CaO \cdot Al_2O_3 \cdot SiO_2}$
3CaO·3Al <sub>2</sub> O <sub>3</sub> ·CaF <sub>2</sub>	b26	$n_{b26} = n_{3CaO \cdot 3Al_2O_3 \cdot CaF_2}$	$N_{b26} = \frac{n_{b26}}{\sum n_i} = N_{3CaO \cdot 3Al_2O_3 \cdot CaF_2}$	
CaO·MgO·SiO <sub>2</sub>	b27	$n_{b27} = n_{CaO \cdot MgO \cdot SiO_2}$	$N_{b27} = \frac{n_{b27}}{\sum n_i} = N_{CaO \cdot MgO \cdot SiO_2}$	
11CaO·7Al <sub>2</sub> O <sub>3</sub> ·CaF <sub>2</sub>	b28	$n_{b28} = n_{11CaO \cdot 7Al_2O_3 \cdot CaF_2}$	$N_{b28} = \frac{n_{b28}}{\sum n_i} = N_{11CaO \cdot 7Al_2O_3 \cdot CaF_2}$	

(continued on next page)

Table 4 – (continued)

Item	Structural unit	Symbols	Molar number $n_i/\text{mol}$	Mass action concentration of structural units $N_i$
	$\text{CaO} \cdot \text{SiO}_2 \cdot \text{TiO}_2$	b29	$n_{b29} = n_{\text{CaO} \cdot \text{SiO}_2 \cdot \text{TiO}_2}$	$N_{b29} = \frac{n_{b29}}{\sum n_i} = N_{\text{CaO} \cdot \text{SiO}_2 \cdot \text{TiO}_2}$
	$3\text{CaO} \cdot 2\text{SiO}_2 \cdot \text{CaF}_2$	b30	$n_{b30} = n_{3\text{CaO} \cdot 2\text{SiO}_2 \cdot \text{CaF}_2}$	$N_{b30} = \frac{n_{b30}}{\sum n_i} = N_{3\text{CaO} \cdot 2\text{SiO}_2 \cdot \text{CaF}_2}$
	$2\text{MgO} \cdot 2\text{Al}_2\text{O}_3 \cdot 5\text{SiO}_2$	b31	$n_{b31} = n_{2\text{MgO} \cdot 2\text{Al}_2\text{O}_3 \cdot 5\text{SiO}_2}$	$N_{b31} = \frac{n_{b31}}{\sum n_i} = N_{2\text{MgO} \cdot 2\text{Al}_2\text{O}_3 \cdot 5\text{SiO}_2}$
	$\text{FeO} \cdot \text{Al}_2\text{O}_3$	b32	$n_{b32} = n_{\text{FeO} \cdot \text{Al}_2\text{O}_3}$	$N_{b32} = \frac{n_{b32}}{\sum n_i} = N_{\text{FeO} \cdot \text{Al}_2\text{O}_3}$
	$2\text{FeO} \cdot \text{SiO}_2$	b33	$n_{b33} = n_{2\text{FeO} \cdot \text{SiO}_2}$	$N_{b33} = \frac{n_{b33}}{\sum n_i} = N_{2\text{FeO} \cdot \text{SiO}_2}$
	$2\text{FeO} \cdot \text{TiO}_2$	b34	$n_{b34} = n_{2\text{FeO} \cdot \text{TiO}_2}$	$N_{b34} = \frac{n_{b34}}{\sum n_i} = N_{2\text{FeO} \cdot \text{TiO}_2}$
	$\text{FeO} \cdot \text{TiO}_2$	b35	$n_{b35} = n_{\text{FeO} \cdot \text{TiO}_2}$	$N_{b35} = \frac{n_{b35}}{\sum n_i} = N_{\text{FeO} \cdot \text{TiO}_2}$
	$\text{FeO} \cdot 2\text{TiO}_2$	b36	$n_{b36} = n_{\text{FeO} \cdot 2\text{TiO}_2}$	$N_{b36} = \frac{n_{b36}}{\sum n_i} = N_{\text{FeO} \cdot 2\text{TiO}_2}$

Table 5 – Chemical reaction equations of possible composite molecules [9,12,14,18,25,27,30–32].

Chemical reaction formula	$\Delta G_i^\theta / (\text{J} \cdot \text{mol}^{-1})$	$N_i$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (\text{CaO} \cdot \text{SiO}_2)$	-92528 + 2.512T	$N_{b1} = K_{b1} N_1 N_4$
$(\text{Mg}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (\text{MgO} \cdot \text{SiO}_2)$	23,849 - 29.706T	$N_{b2} = K_{b2} N_4 N_6$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) = (\text{CaO} \cdot \text{Al}_2\text{O}_3)$	59,413 - 59.413T	$N_{b3} = K_{b3} N_1 N_3$
$(\text{Mg}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) = (\text{MgO} \cdot \text{Al}_2\text{O}_3)$	-18828 - 6.276T	$N_{b4} = K_{b4} N_3 N_6$
$2(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (2\text{CaO} \cdot \text{SiO}_2)$	-102090 - 24.267T	$N_{b5} = K_{b5} N_1^2 N_4$
$2(\text{Mg}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (2\text{MgO} \cdot \text{SiO}_2)$	-56902 - 3.347T	$N_{b6} = K_{b6} N_4 N_6^2$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) = (3\text{CaO} \cdot \text{Al}_2\text{O}_3)$	-21757 - 29.288T	$N_{b7} = K_{b7} N_1^3 N_3$
$12(\text{Ca}^{2+} + \text{O}^{2-}) + 7(\text{Al}_2\text{O}_3) = (12\text{CaO} \cdot 7\text{Al}_2\text{O}_3)$	617,977 - 612.119T	$N_{b8} = K_{b8} N_1^{12} N_3^7$
$(\text{Ca}^{2+} + \text{O}^{2-}) + 2(\text{Al}_2\text{O}_3) = (\text{CaO} \cdot 2\text{Al}_2\text{O}_3)$	-16736 - 25.522T	$N_{b9} = K_{b9} N_1 N_3^2$
$(\text{Ca}^{2+} + \text{O}^{2-}) + 6(\text{Al}_2\text{O}_3) = (\text{CaO} \cdot 6\text{Al}_2\text{O}_3)$	-22594 - 31.798T	$N_{b10} = K_{b10} N_1 N_3^6$
$3(\text{Al}_2\text{O}_3) + 2(\text{SiO}_2) = (3\text{Al}_2\text{O}_3 \cdot 2\text{SiO}_2)$	-4354.27 - 10.467T	$N_{b11} = K_{b11} N_3^3 N_4^2$
$3(\text{Al}_2\text{O}_3) + (\text{SiO}_2) = (3\text{CaO} \cdot \text{SiO}_2)$	-118,826 - 6.694T	$N_{b12} = K_{b12} N_1^3 N_4$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{TiO}_2) = (\text{CaO} \cdot \text{TiO}_2)$	-79900 - 3.35T	$N_{b13} = K_{b13} N_1 N_5$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + 2(\text{TiO}_2) = (3\text{CaO} \cdot 2\text{TiO}_2)$	107,100 - 11.35T	$N_{b14} = K_{b14} N_1^3 N_5^2$
$4(\text{Ca}^{2+} + \text{O}^{2-}) + 3(\text{TiO}_2) = (4\text{CaO} \cdot 3\text{TiO}_2)$	-292,880 - 17.573T	$N_{b15} = K_{b15} N_1^4 N_5^3$
$(\text{Al}_2\text{O}_3) + (\text{TiO}_2) = (\text{Al}_2\text{O}_3 \cdot \text{TiO}_2)$	-25270 + 3.924T	$N_{b16} = K_{b16} N_3 N_5$
$(\text{Mg}^{2+} + \text{O}^{2-}) + (\text{TiO}_2) = (\text{MgO} \cdot \text{TiO}_2)$	-26400 + 3.14T	$N_{b17} = K_{b17} N_5 N_6$
$(\text{Mg}^{2+} + \text{O}^{2-}) + 2(\text{TiO}_2) = (\text{MgO} \cdot 2\text{TiO}_2)$	-27600 + 0.63T	$N_{b18} = K_{b18} N_5^2 N_6$
$2(\text{Mg}^{2+} + \text{O}^{2-}) + (\text{TiO}_2) = (2\text{MgO} \cdot \text{TiO}_2)$	-25500 + 1.26T	$N_{b19} = K_{b19} N_5 N_6^2$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + 2(\text{SiO}_2) = (3\text{CaO} \cdot 2\text{SiO}_2)$	-236,814 + 9.623T	$N_{b20} = K_{b20} N_1^3 N_4^2$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Mg}^{2+} + \text{O}^{2-}) + 2(\text{SiO}_2) = (\text{CaO} \cdot \text{MgO} \cdot 2\text{SiO}_2)$	-80333 - 51.882T	$N_{b21} = K_{b21} N_1 N_4^2 N_6$
$2(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Mg}^{2+} + \text{O}^{2-}) + 2(\text{SiO}_2) = (2\text{CaO} \cdot \text{MgO} \cdot 2\text{SiO}_2)$	-73638 - 63.597T	$N_{b22} = K_{b22} N_1^2 N_4^2 N_6$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Mg}^{2+} + \text{O}^{2-}) + 2(\text{SiO}_2) = (3\text{CaO} \cdot \text{MgO} \cdot 2\text{SiO}_2)$	-205016 - 31.798T	$N_{b23} = K_{b23} N_1^3 N_4^2 N_6$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) + 2(\text{SiO}_2) = (\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot 2\text{SiO}_2)$	-4184 - 73.638T	$N_{b24} = K_{b24} N_1 N_3 N_4^2$
$2(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) + (\text{SiO}_2) = (2\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot \text{SiO}_2)$	-116,315 - 38.911T	$N_{b25} = K_{b25} N_1^2 N_3 N_4$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + 3(\text{Al}_2\text{O}_3) + (\text{Ca}^{2+} + 2\text{F}^-) = (3\text{CaO} \cdot 3\text{Al}_2\text{O}_3 \cdot \text{CaF}_2)$	-44,492 - 73.15T	$N_{b26} = K_{b26} N_1^3 N_2 N_3^2$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{Mg}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (\text{CaO} \cdot \text{MgO} \cdot \text{SiO}_2)$	-124,683 + 3.766T	$N_{b27} = K_{b27} N_1 N_4 N_6$
$11(\text{Ca}^{2+} + \text{O}^{2-}) + 7(\text{Al}_2\text{O}_3) + (\text{Ca}^{2+} + 2\text{F}^-) = (11\text{CaO} \cdot 7\text{Al}_2\text{O}_3 \cdot \text{CaF}_2)$	-228,760 - 155.8T	$N_{b28} = K_{b28} N_1^{11} N_2 N_3^7$
$(\text{Ca}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) + (\text{TiO}_2) = (\text{CaO} \cdot \text{SiO}_2 \cdot \text{TiO}_2)$	-114,683 + 7.32T	$N_{b29} = K_{b29} N_1 N_4 N_5$
$3(\text{Ca}^{2+} + \text{O}^{2-}) + 2(\text{SiO}_2) + (\text{Ca}^{2+} + 2\text{F}^-) = (3\text{CaO} \cdot 2\text{SiO}_2 \cdot \text{CaF}_2)$	-255,180 - 8.2T	$N_{b30} = K_{b30} N_1^3 N_2 N_4^2$
$2(\text{Mg}^{2+} + \text{O}^{2-}) + 2(\text{Al}_2\text{O}_3) + 5(\text{SiO}_2) = (2\text{MgO} \cdot 2\text{Al}_2\text{O}_3 \cdot 5\text{SiO}_2)$	-14422 - 14.808T	$N_{b31} = K_{b31} N_3^2 N_4^2 N_6^2$
$(\text{Fe}^{2+} + \text{O}^{2-}) + (\text{Al}_2\text{O}_3) = (\text{FeO} \cdot \text{Al}_2\text{O}_3)$	-59204 + 22.343T	$N_{b32} = K_{b32} N_3 N_7$
$2(\text{Fe}^{2+} + \text{O}^{2-}) + (\text{SiO}_2) = (2\text{FeO} \cdot \text{SiO}_2)$	-9395 - 0.227T	$N_{b33} = K_{b33} N_4 N_7^2$
$2(\text{Fe}^{2+} + \text{O}^{2-}) + (\text{TiO}_2) = (2\text{FeO} \cdot \text{TiO}_2)$	-33913.08 + 5.86T	$N_{b34} = K_{b34} N_5 N_7^2$
$(\text{Fe}^{2+} + \text{O}^{2-}) + (\text{TiO}_2) = (\text{FeO} \cdot \text{TiO}_2)$	68,320 - 43.965T	$N_{b35} = K_{b35} N_5 N_7$
$(\text{Fe}^{2+} + \text{O}^{2-}) + 2(\text{TiO}_2) = (\text{FeO} \cdot 2\text{TiO}_2)$	-16188.703	$N_{b36} = K_{b36} N_5^2 N_7$

$$c_2 = (1 / 3N_2 + N_{b26} + N_{b28} + 3N_{b30}) \sum n_i = n_{CaF_2}^0 \quad (24)$$

$$c_3 = \left( \begin{matrix} N_3 + N_{b3} + N_{b4} + N_{b7} + 7N_{b8} + 2N_{b9} + 6N_{b10} + 3N_{b11} \\ + N_{b16} + N_{b24} + N_{b25} + 3N_{b26} + 7N_{b28} + 2N_{b31} + N_{b32} \end{matrix} \right) \times \sum n_i = n_{Al_2O_3}^0 \quad (25)$$

$$c_4 = \left( \begin{matrix} N_4 + N_{b1} + N_{b2} + N_{b5} + N_{b6} + 2N_{b11} + N_{b12} + 2N_{b20} + 2N_{b21} + 2N_{b22} \\ + 2N_{b23} + 2N_{b24} + N_{b25} + N_{b27} + N_{b29} + 2N_{b30} + 5N_{b31} + N_{b33} \end{matrix} \right) \sum n_i = n_{SiO_2}^0 \quad (26)$$

$$c_5 = \left( \begin{matrix} N_5 + N_{b13} + 2N_{b14} + 3N_{b15} + N_{b16} + N_{b17} + 2N_{b18} + N_{b19} + N_{b29} + N_{b34} \\ + N_{b35} + 2N_{b36} \end{matrix} \right) \sum n_i = n_{TiO_2}^0 \quad (27)$$

$$c_6 = \left( \begin{matrix} 0.5N_6 + N_{b2} + N_{b4} + 2N_{b6} + N_{b17} + N_{b18} + 2N_{b19} + N_{b21} + N_{b22} \\ + N_{b23} + N_{b27} + 2N_{b31} \end{matrix} \right) \times \sum n_i = n_{MgO}^0 \quad (28)$$

$$c_7 = (0.5N_7 + N_{b32} + 2N_{b33} + 2N_{b34} + N_{b35} + N_{b36}) \sum n_i = n_{FeO}^0 \quad (29)$$

### 3. Experimental procedures

In this experiment, the GH4065A alloy and Al<sub>2</sub>O<sub>3</sub>-TiO<sub>2</sub>-CaO-MgO-CaF<sub>2</sub>-SiO<sub>2</sub>-FeO seven-element slag system are used in the ESR process, respectively. The electrical system of AC mode is carried out with transformer output power 270 kVA, voltage 40.5 V and current 2100 A. The slag samples are proportionally prepared and mixed. Then the prepared slag samples are put into the electric slag furnace, then heated at about 1600 °C for about 5–10 min, finally, cooled at the room temperature. Argon was blown into the furnace. Slag sample was taken from the molten slag after melted for 40 min. In order to investigate the relationship between the titanium and aluminium in ingot, three sets of experiments were done under the same conditions, the experiments are named case1, case2, case3. The Al and Ti in the head and tail of ingot were analyzed, respectively. The chemical composition of slag samples were analysed by the X-ray fluorescence (XRF) technique. The XRF technique were used to analyse the dynamic changes in the content of Al, Ti and other components in slag.

### 4. Results and discussion

#### 4.1. Effect of slag system composition and temperature on Al element

The variation of  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  with CaO, CaF<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, and MgO content at different temperature are shown in Fig. 1. It can be found the  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  rises with

the increase of CaO, CaF<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> in slag system. In other words, the increase of CaO, CaF<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> in slag will reduce the Al content. It can be deduced that their ability to weaken the loss of Al content is in the following order Al<sub>2</sub>O<sub>3</sub> >- CaO > CaF<sub>2</sub>. It is obvious that the  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  is almost unchanged when the MgO content increases, which

means that the content of MgO in slag has little effect on the decrease of Al content.

The variation of  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  with SiO<sub>2</sub>, TiO<sub>2</sub>, and FeO at different temperature are demonstrated in Fig. 2. It shows that when the content of SiO<sub>2</sub>, TiO<sub>2</sub> and FeO increases, the  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  decreases. Hence, the increase of SiO<sub>2</sub>, TiO<sub>2</sub> and FeO content in slag will cause the decrease of Al content. From the picture, it can be seen that their ability to enhance the loss of Al content is in the following order SiO<sub>2</sub> > FeO > TiO<sub>2</sub>.

The mass action concentration of each element in slag system after the reaction can be calculated by matlab, and then the content of Al elements can be obtained by Eqs. (11)–(17) at different temperature when the slag ratio is different, as shown in Figs. 3 and 4.

It shows the variation of the Al content with CaO content in the slag after the slag-metal interface in an equilibrium state in Fig. 3(a). With the increase of CaO content, the Al content in superalloy also increases. When the CaO content is higher than 15 w%, the Al content in superalloy can be bigger than 2.13 w% when the ESR reaction reaches equilibrium at 1973 K. The CaO can effectively increase the alkalinity of the slag system. Hence, the Al element will not be lost when the CaO content is lower than 15 w%.

Fig. 3(b) represents the variation of Al content with CaF<sub>2</sub> content in slag after it reaches equilibrium, the content of Al in superalloy increases with the increase of CaF<sub>2</sub> content. Because CaF<sub>2</sub> not only effectively lowers the viscosity and melting point of remelted slag, but also greatly improves the fluidity of remelted slag system. When the CaF<sub>2</sub> content is greater than 50 w%, the Al content in superalloy can be higher than 2.13 w% when equilibrium at 1973 K. When the CaF<sub>2</sub> content is smaller than 50 w%, the Al element in superalloy can be kept at a lower level compared with the other cases.

Fig. 3(c) shows that the variation of Al content with Al<sub>2</sub>O<sub>3</sub> content in slag after the slag-metal interface in an equilibrium state. The Al content tends to influence the melting temperature and viscosity of slag system and also effectively reduce the electrical conductivity of slag system. When the Al<sub>2</sub>O<sub>3</sub>

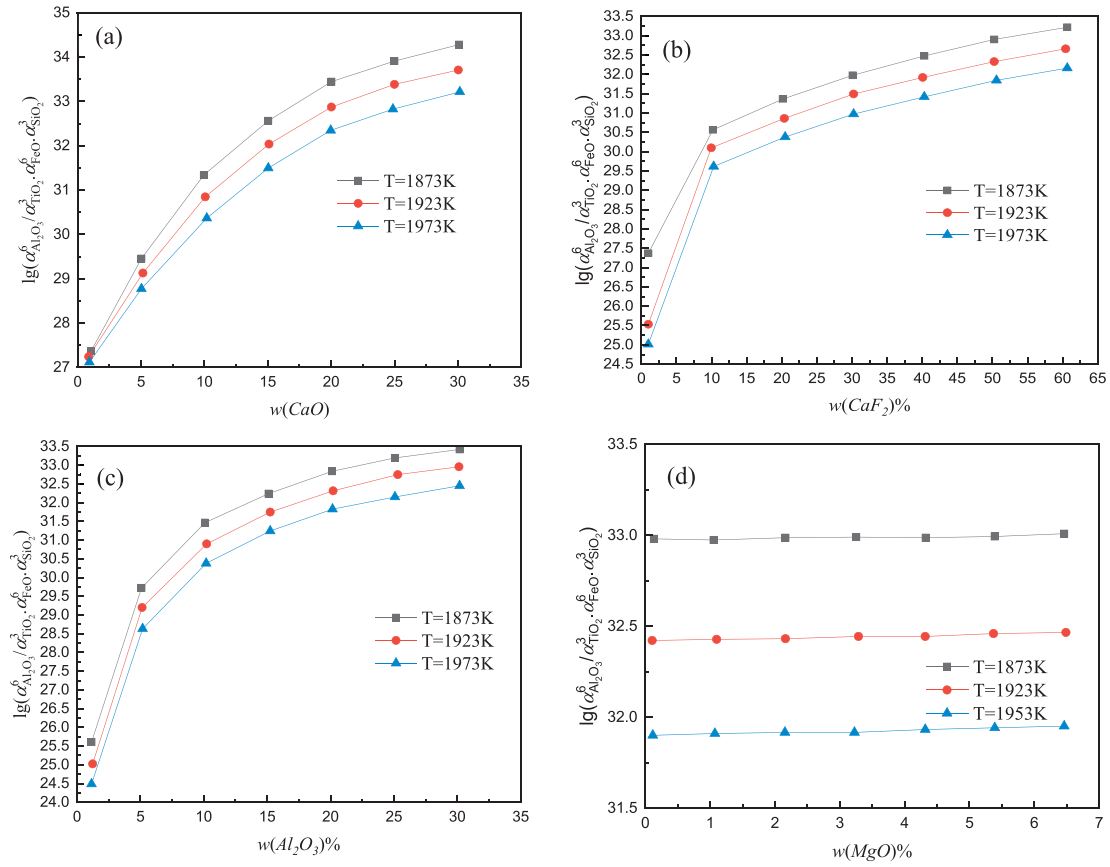


Fig. 1 – Variation of  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \cdot a_{FeO}^6 \cdot a_{SiO_2}^3)$  with (a) CaO; (b) CaF<sub>2</sub>; (c) Al<sub>2</sub>O<sub>3</sub>; (d) MgO versus temperature.

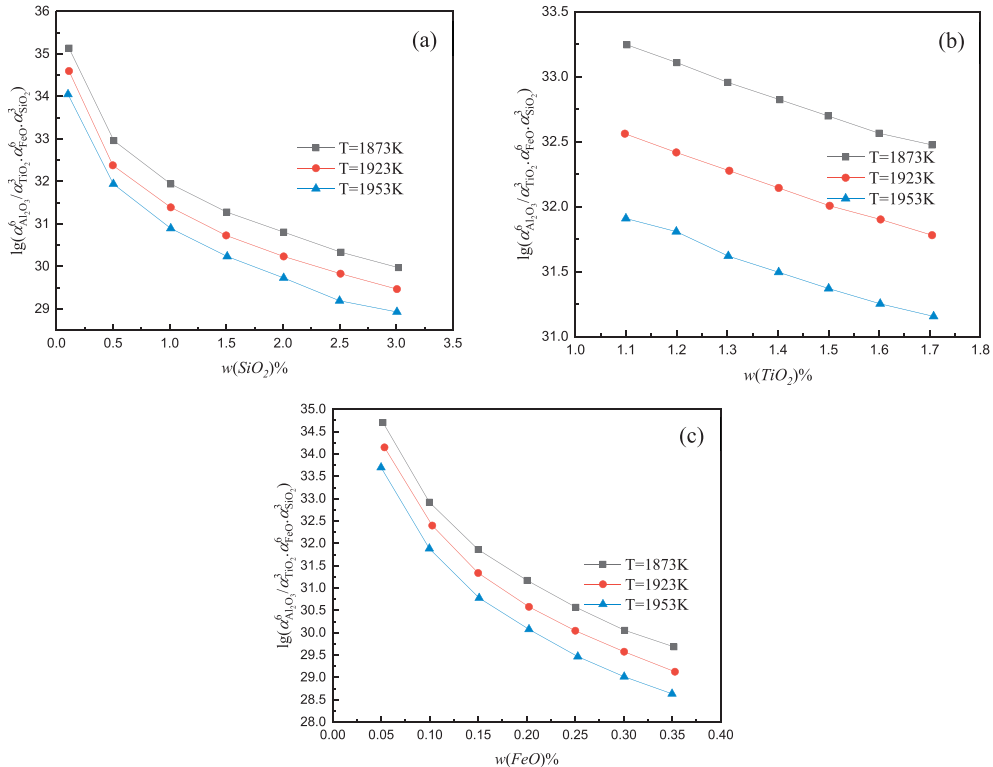


Fig. 2 – Variation of  $\lg(a_{Al_2O_3}^6 / a_{TiO_2}^3 \cdot a_{FeO}^6 \cdot a_{SiO_2}^3)$  with (a) SiO<sub>2</sub>; (b) TiO<sub>2</sub>; (c) FeO versus temperature.



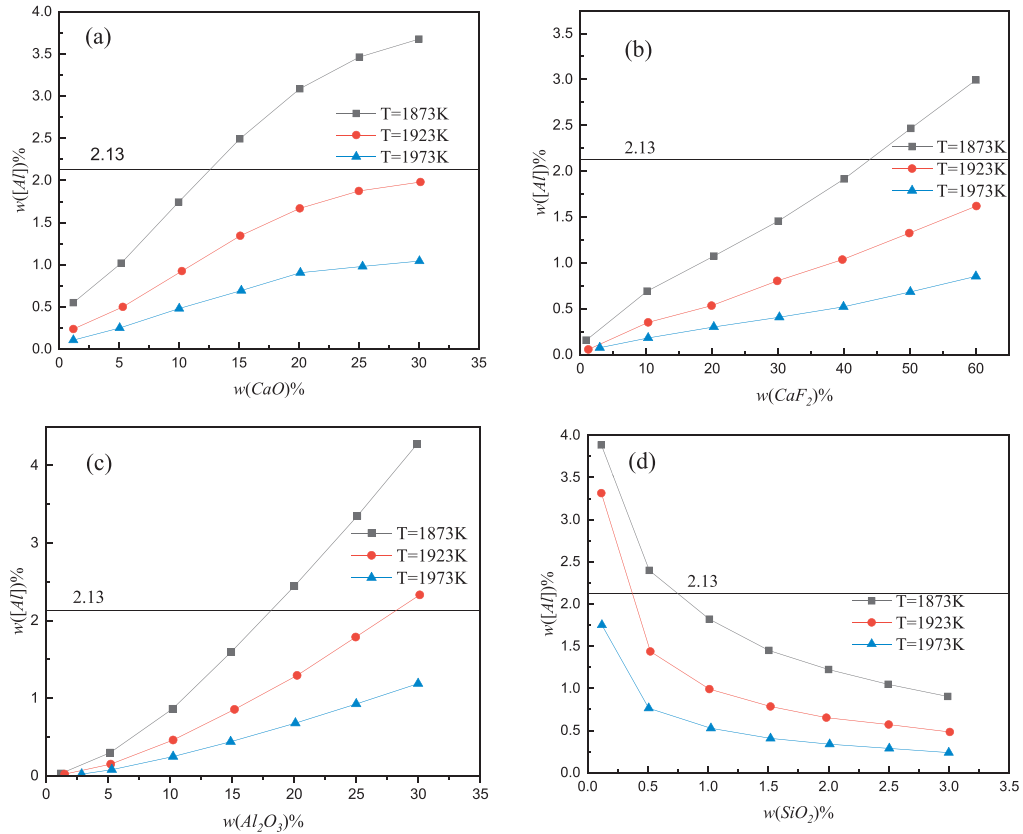


Fig. 3 – Variation of Al content with (a) CaO; (b) CaF<sub>2</sub>; (c) Al<sub>2</sub>O<sub>3</sub>; (d) Si<sub>2</sub>O<sub>2</sub> at 1873 K, 1923 K and 1973 K.

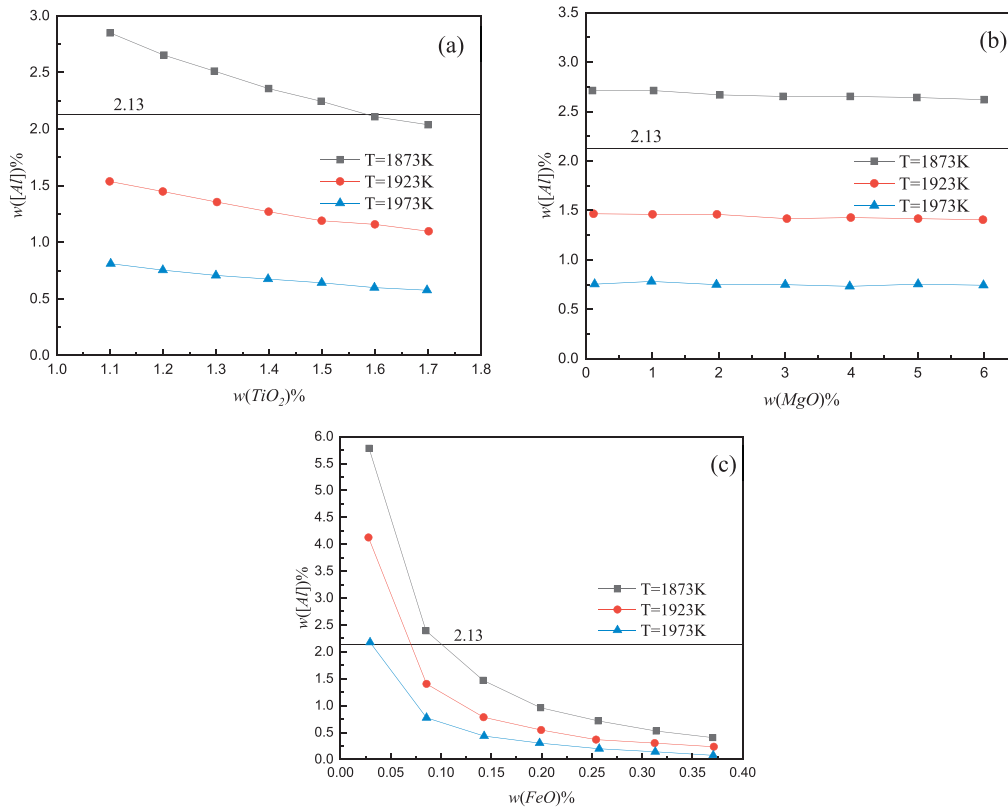


Fig. 4 – Variation of Al content with (a) TiO<sub>2</sub>; (b) MgO; (c) FeO at 1873 K, 1923 K and 1973 K.

content is greater than 18 w%, the Al content in superalloy can be bigger than 2.13 w% when the ESR reaction reaches equilibrium at 1973 K.

From Fig. 3(d), it depicts that the change of Al content with SiO<sub>2</sub> content in slag when equilibrium. Because it effectively reduces the electrical conductivity and melting temperature of liquid slag system. When the content of SiO<sub>2</sub> is lower than 0.5 w%, the content of Al in superalloy when equilibrium can be less than 2.13 w%.

The change of Al content with TiO<sub>2</sub> content in slag after equilibrium is given in Fig. 4(a). The TiO<sub>2</sub> reacts with the Al in superalloy and then will be oxidized. While the content of TiO<sub>2</sub> is less than 1.6 w%, the Al content in superalloy can be higher than 2.13 w% when equilibrium at 1973 K. While the content of TiO<sub>2</sub> is bigger than 1.6 w%, the decrease of Al content in superalloy can be reduced. Fig. 4(b) describes the change of Al content with MgO content in slag after equilibrium. A thin semi-condensable film at the air-slag interface could be formed by MgO to prevent oxygen caused by slag. In other words, the MgO has little influence on the decrease of Al content. Hence, it is not necessary to pay attention to MgO when considering the decrease of Al content. The change of Al content with FeO content in the slag after equilibrium is shown in Fig. 4(c). As mentioned in Ref. [20], An increase of FeO content promotes the reaction of Eq. (13) to the right, as the FeO content rises, the amount of Al

decreases. With the increase of FeO content, the Al content drops. When the FeO content is lower than 0.12 w%, the Al content in superalloy can be higher than 2.13 w% when equilibrium. When the FeO content is less than 0.12 w%, it can be deduced that the Al content in superalloy will not be decreased.

#### 4.2. Effect of slag system composition and temperature on Ti element

The variation of  $\lg(a_{\text{Al}_2\text{O}_3}^6 / a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3)$  with CaO, Al<sub>2</sub>O<sub>3</sub>, SiO<sub>2</sub>, and FeO content at different temperature are shown in Fig. 5. It could be seen that if the content of CaO or Al<sub>2</sub>O<sub>3</sub> increases, the  $\lg(a_{\text{Al}_2\text{O}_3}^6 / a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3)$  decreases. However, when the content of CaO and Al<sub>2</sub>O<sub>3</sub> go up to a certain extent, the  $\lg(a_{\text{Al}_2\text{O}_3}^6 / a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3)$  begins to level off and then rise. The reason is that an additional 1 w% TiO<sub>2</sub> into the slag system has a significant effect on the equilibrium of chemical reaction with the change of CaO and Al<sub>2</sub>O<sub>3</sub> content. The addition of TiO<sub>2</sub> will drive the direction of reaction with Eqs. (1), (3), and (5) toward the left and reduce the decrease of Ti content. If the content of SiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> increases, the  $\lg(a_{\text{Al}_2\text{O}_3}^6 / a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3)$  always drops. It can be deduced that the ability to enhance the loss of Ti content is in the following order FeO > SiO<sub>2</sub> > Al<sub>2</sub>O<sub>3</sub> > CaO.

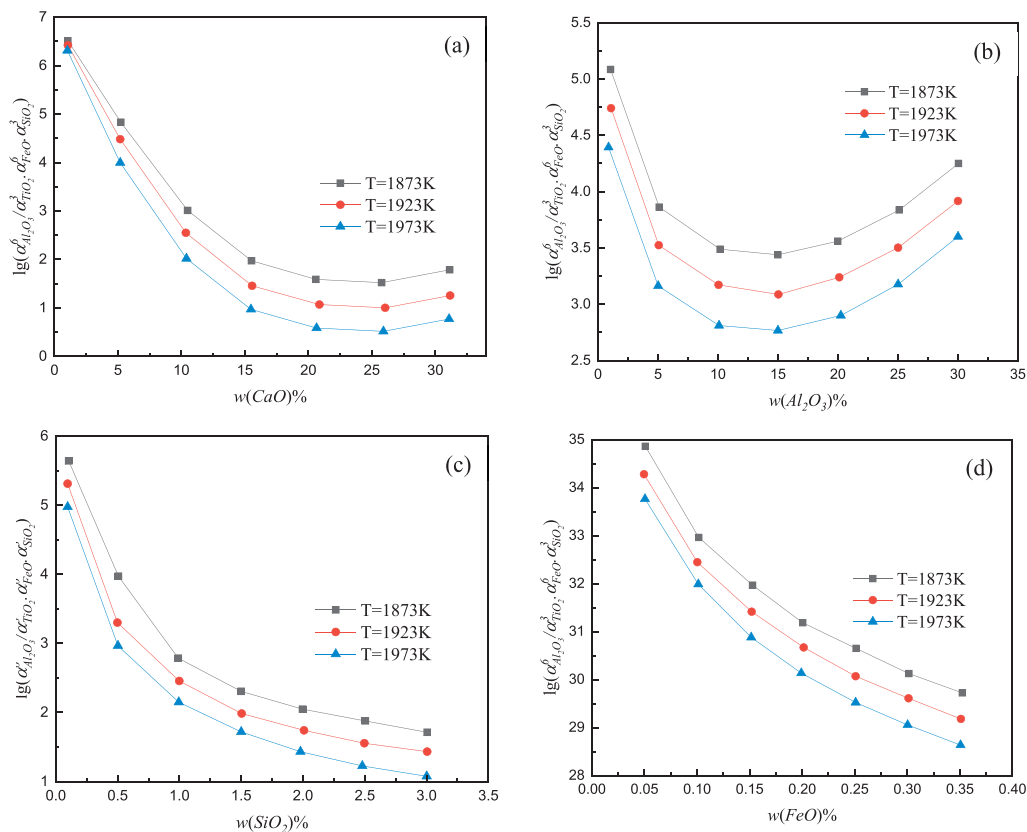


Fig. 5 – Variation of  $\lg(a_{\text{Al}_2\text{O}_3}^6 / a_{\text{TiO}_2}^3 \cdot a_{\text{FeO}}^6 \cdot a_{\text{SiO}_2}^3)$  with composition content of slag versus temperature (a) CaO; (b) Al<sub>2</sub>O<sub>3</sub>; (c) SiO<sub>2</sub>; (d) FeO.

The variation of  $\lg(a_{Al_2O_3}^6/a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  with  $CaF_2$ ,  $TiO_2$ , and  $MgO$  content at different temperature are demonstrated in Fig. 6. It can be found that the  $\lg(a_{Al_2O_3}^6/a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  rises with the increase of  $CaF_2$ ,  $TiO_2$ , and  $MgO$  content in the slag system. The increase of  $CaF_2$ ,  $TiO_2$ , and  $MgO$  content in slag can reduce Ti content. It can be seen that the ability to weaken the loss of Ti content is in the following order  $TiO_2 > CaF_2 > MgO$ .

As discussed above, the mass action concentration of composition content of slag can be calculated using Matlab software. When the slag composition varies, the content of Ti element versus temperature could be obtained from Eq. (10). The variation of Ti content with  $CaO$  content in slag at different temperature is depicted in Fig. 7(a). It could be found that if the  $CaO$  content increases, the Ti element in superalloy firstly decreases and then increases. As shown in Table 5, it is proposed that the  $CaO$  in slag will form the complex molecules with  $TiO_2$ ,  $MgO$ , and  $SiO_2$  in slag. Hence, the activity of  $TiO_2$  will be weakened, when the  $CaO$  content increases. Meanwhile, the content of  $TiO_2$  will increase, and the loss of Ti content will decrease. When the  $CaO$  content is equal to 16 w%, the chemical reaction is in equilibrium, and the Ti element content in superalloy is close to 3.73 w%. The content of the Ti element cannot be lost if the  $CaO$  content is lower than 16 w%. Due to an extra 1%wt  $TiO_2$ , the chemical reaction will be balanced when the  $CaO$  content equals 20 w%. Next, the

reaction will process in the opposite direction. The Ti content will increase again.

Fig. 7 (b) depicts the variation of Ti content with  $CaF_2$  content in slag. The  $CaF_2$  in slag will form the complex molecules with  $Al_2O_3$  and  $CaO$  in slag is shown in Table 5. Hence, the activity of  $Al_2O_3$  will be decreased. According to Eq. (1), when the  $CaF_2$  content increases, the content of Ti element rises. When the content of  $CaF_2$  is greater than 58 w%, the content of Ti element in the superalloy at the reaction temperature of 1973 K can be larger than 3.73 w%. Hence, when the content of  $CaF_2$  is larger than 58 w%, it could be prohibited from the loss of Ti content in superalloy.

The change of Ti content with  $Al_2O_3$  content in slag is demonstrated in Fig. 7(c). The trend of the curve changes from downward to upward. With the increase of  $Al_2O_3$  content, the Ti element in superalloy decreases first. The  $Al_2O_3$  in slag and Ti in superalloy will react and give the Al and  $TiO_2$  in the slag/metal interface. With the increase of  $Al_2O_3$  content, the loss of Ti in superalloy improves. The Ti content drops with increasing temperature since the reaction temperature will enhance the loss of Ti content. When the  $Al_2O_3$  content changes to 15 w%, the chemical reaction is in equilibrium. The content of  $TiO_2$  will be the maximum. Later, the chemical reaction will proceed oppositely, and the Ti content will increase again. When the content of  $Al_2O_3$  is lower than 5 w% and higher than 24 w%, the content of Ti in superalloy is bigger

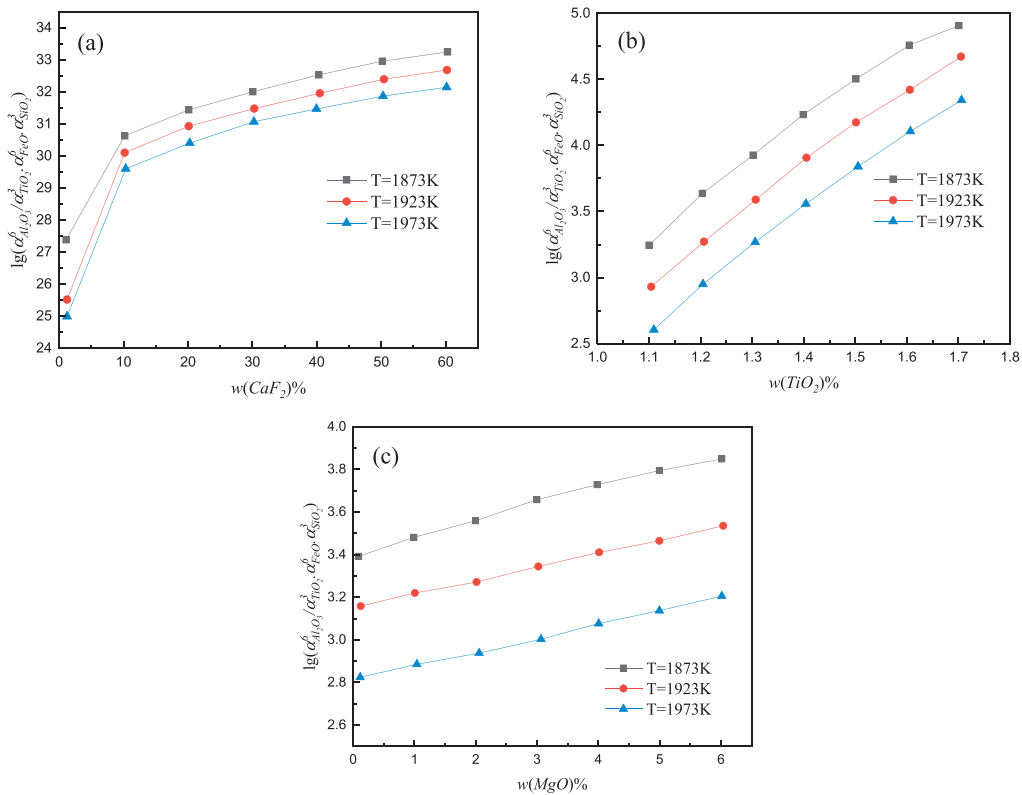


Fig. 6 – Variation of  $\lg(a_{Al_2O_3}^6/a_{TiO_2}^3 \bullet a_{FeO}^6 \bullet a_{SiO_2}^3)$  with composition content of slag versus temperature (a)  $CaF_2$ ; (b)  $TiO_2$ ; (c)  $MgO$ .

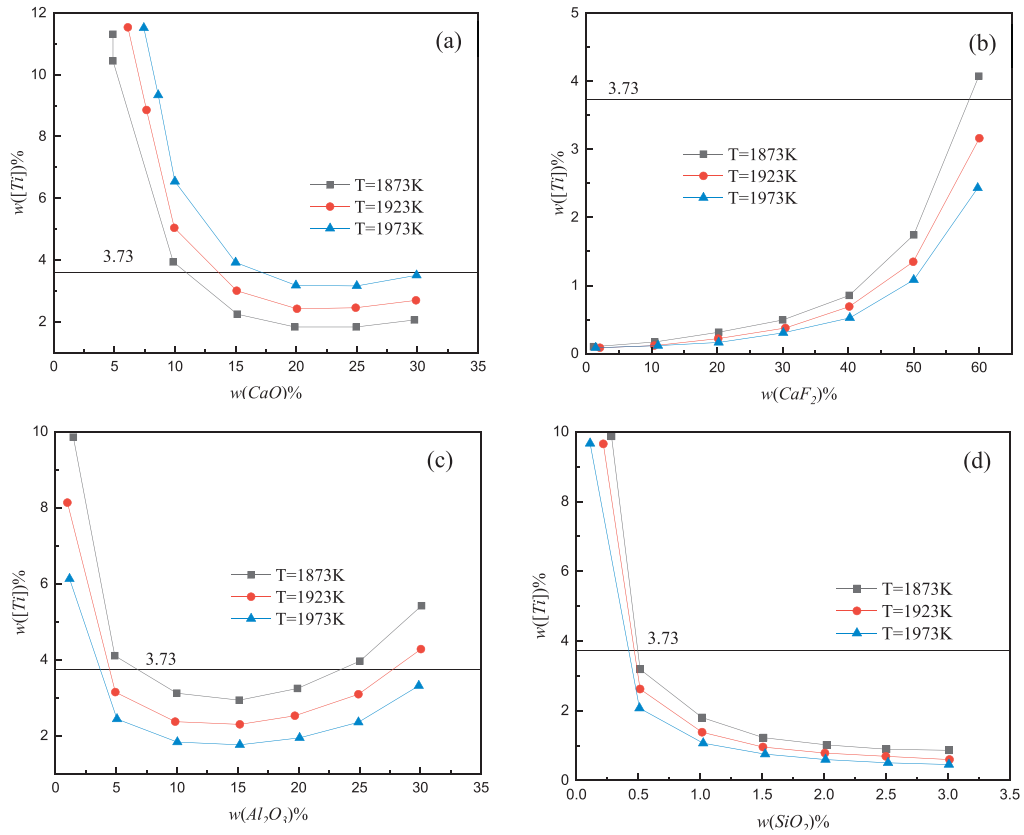


Fig. 7 – Variation of Ti content with (a) CaO; (b) CaF<sub>2</sub>; (c) Al<sub>2</sub>O<sub>3</sub>; (d) Si<sub>2</sub>O<sub>2</sub> at 1873 K, 1923 K and 1973 K.

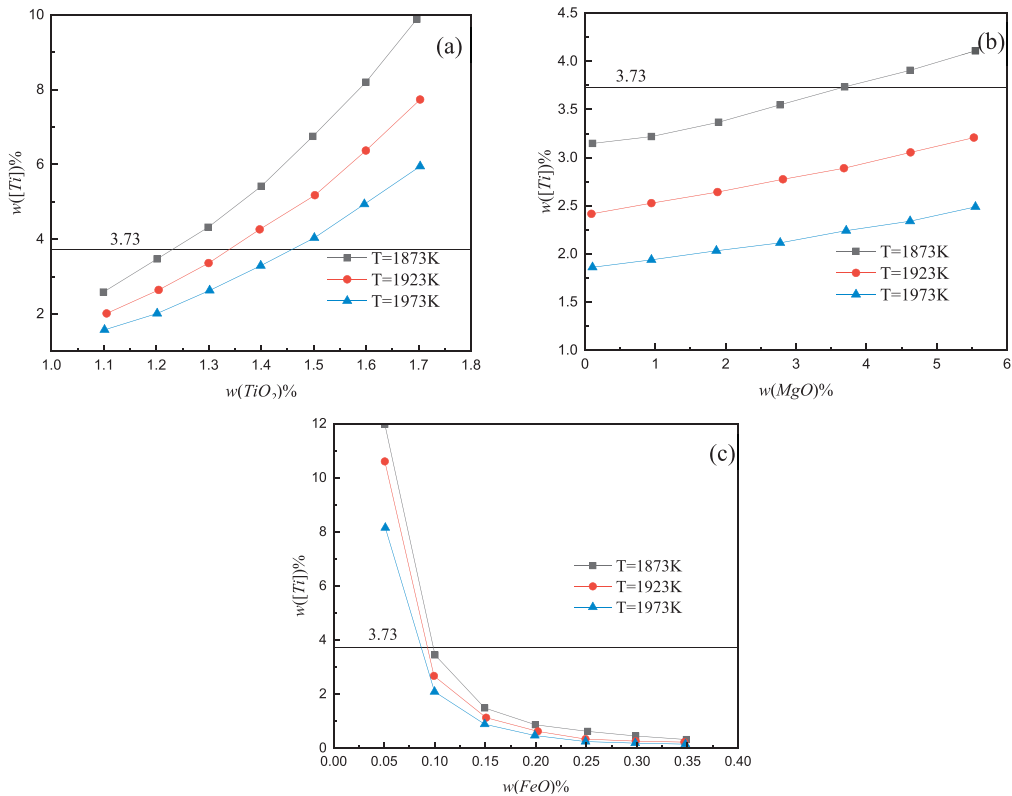


Fig. 8 – Variation of Ti content with (a) TiO<sub>2</sub>; (b) MgO; (c) FeO at 1873 K, 1923 K and 1973 K.

than 3.73 w%. Compared to the original composition of Al<sub>2</sub>O<sub>3</sub> content, the decrease of Ti content can be reduced when the content of Al<sub>2</sub>O<sub>3</sub> is higher than 24 w%. The variation of Ti content with SiO<sub>2</sub> content in slag is given in Fig. 7(d). When the SiO<sub>2</sub> content increases, the loss of Ti content will decrease. When the content of SiO<sub>2</sub> is less than 0.5 w%, the content of Ti in superalloy can be much higher than 3.73 w%. In this case, it will not cause the decrease of Ti content in superalloy.

Fig. 8 describes the variation of Ti content with TiO<sub>2</sub>, MgO and FeO content in slag. It could be seen that the Ti content in superalloy increases with the increase of TiO<sub>2</sub> content in Fig. 8(a). When the content of TiO<sub>2</sub> in slag is higher, the chemical reaction between the Ti element in superalloy and the unstable oxide in the slag was forced. Hence, the decrease of Ti content is reduced in superalloy. When the content of TiO<sub>2</sub> is higher than 1.2 w%, the Ti element content in superalloy can be larger than 3.73 w%. The Ti content decreases in superalloy can be effectively prevented when the content of TiO<sub>2</sub> is higher than 1.2 w%. The change of Ti content with MgO content in slag is described in Fig. 8(b). The Ti content in superalloy increases with the increase of MgO content. The MgO will enhance the activity of Ti<sub>3</sub>O<sub>5</sub> in slag and lessen the activity of TiO<sub>2</sub> in slag. Moreover, the Ti<sub>3</sub>O<sub>5</sub> in slag will be decomposed into the TiO<sub>2</sub> in slag and Ti in superalloy [33]. When the content of MgO increases, the decomposition of

Ti<sub>3</sub>O<sub>5</sub> in slag will be accelerated, and the Ti content in superalloy will increase. When the content of MgO is greater than 4 w%, the Ti content in superalloy at the reaction equilibrium can be greater than 3.73 w%. Hence, when the content of MgO is higher than 4 w%, the Ti element in superalloy can not be lost. The variation of Ti content with FeO content in slag is represented in Fig. 8(c). The Ti content decreases continuously with the increase of FeO content. It has been pointed out that the activity of TiO<sub>2</sub> decreases significantly with the increasing FeO content in Ref. [30], this is because TiO<sub>2</sub> can be reacted with FeO in the slag to form complex compounds, the formation of complex titanium-containing compounds reduces the effective concentration of TiO<sub>2</sub> in the slag. It has the same trend with our case. When the content of FeO is lower than 0.1 w%, the Ti content in superalloy can be higher than 3.13 w%. When the content of FeO is smaller than 0.1 w%, it will not cause a decrease of Ti content in superalloy.

4.3. Validation of thermodynamic model

In order to further verify the agreement of thermodynamic model and experimental results, Eq. (2) could be changed to Eq. (30), the relationship between  $\lg(X_{TiO_2}^3/X_{Al_2O_3}^2)$  and  $\lg([Ti]^3/[Al]^4)$  can be obtained from the experimental results of the slag-metal equilibrium and the model-calculated action concentrations of the slag system components. As shown in Fig. 9, the experimental results are in good agreement with the model calculation.

$$\lg \frac{W_{TiO_2}^3}{W_{Al_2O_3}^2} = -\frac{35300}{T} + 9.94 + \lg \frac{f_{Ti}^3 [Ti]^3}{f_{Al}^4 [Al]^4} - \lg \frac{\gamma_{TiO_2}^3}{\gamma_{Al_2O_3}^4} \quad (30)$$

$$\gamma_i = N_i/W_i$$

where  $W_{TiO_2}$  and  $W_{Al_2O_3}$  expresses the mole fraction of TiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> in slag, respectively;  $\gamma_{TiO_2}$  and  $\gamma_{Al_2O_3}$  denotes the activities coefficients of TiO<sub>2</sub> and Al<sub>2</sub>O<sub>3</sub> in slag, respectively.

The contents of Ti and Al at the head and tail in each sample are obtained, and the results of the three sets of experiments are shown in Fig. 10. The experimental value of the average elemental content of Al and Ti at the head and tail of the ingot are in close agreement with the simulated values, indicating that the established thermodynamic model is reasonable.

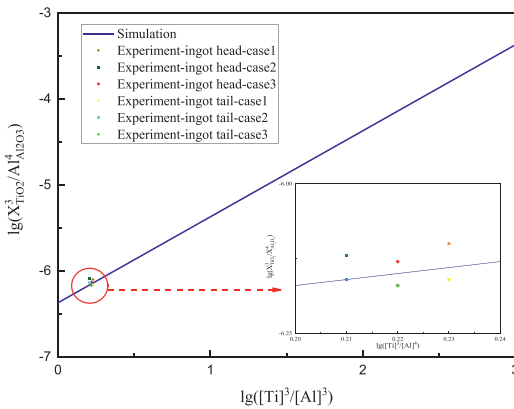


Fig. 9 – Dependence of the  $\lg(X_{TiO_2}^3/X_{Al_2O_3}^2)$  on  $\lg([Ti]^3/[Al]^4)$ .

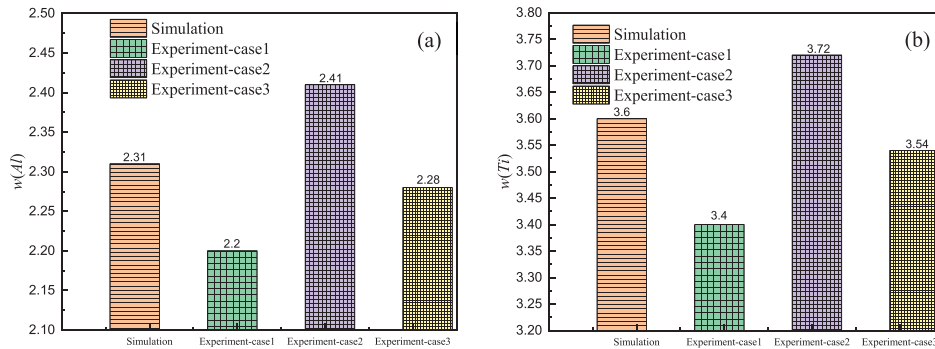


Fig. 10 – Al and Ti content in electroslag ingots with experiment and simulation results.

## 5. Conclusions

A thermodynamic model based on IMCT is established to predict the Ti and Al content in the  $\text{SiO}_2\text{--Al}_2\text{O}_3\text{--FeO--TiO}_2\text{--CaO--MgO--CaF}_2$  slag system in the ESR process of GH4065A superalloy. The effect of slag composition on Ti and Al content have been discussed in detail.

- (1) Increasing the content of  $\text{CaF}_2$  in slag or temperature has a positive effect on reducing the reduction of Al and Ti in the process of electroslag remelting GH4065A alloy; while increasing the content of  $\text{SiO}_2$  and FeO in the slag will both increase the loss of Al and Ti elements; while increasing the content of  $\text{Al}_2\text{O}_3$  and CaO will cause an increase in Al content and a decrease in Ti content of; the effect of MgO on Al is not obvious.
- (2) The order of the ability of each element in the slag system to weaken the loss of Al content is in the following order  $\text{Al}_2\text{O}_3 > \text{CaO} > \text{CaF}_2$ , enhance the loss of Al content is  $\text{SiO}_2 > \text{FeO} > \text{TiO}_2$ ,  $\text{CaF}_2 > \text{MgO}$  and enhance the loss of Ti content is  $\text{FeO} > \text{SiO}_2 > \text{Al}_2\text{O}_3 > \text{CaO}$ .
- (3) Considering the influence of the slag system on the elements Al and Ti, the optimal ratio of the slag system is  $\text{CaO} = 15\text{ w\%--}20\text{ w\%}$ ,  $\text{CaF}_2 \geq 60\text{ w\%}$ ,  $\text{Al}_2\text{O}_3 = 18\text{ w\%--}25\text{ w\%}$ ,  $\text{SiO}_2 \leq 0.5\text{ w\%}$ ,  $\text{TiO}_2 = 1\text{ w\%} + (0.2\text{ w\%} \sim 0.6\text{ w\%})$ ,  $\text{MgO} \geq 4\text{ w\%}$ ,  $\text{FeO} \leq 0.1\text{ w\%}$ .

## Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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