

## ***In-situ* observation of nonmetallic inclusions in steel using confocal scanning laser microscopy: A review**

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# *In-situ* observation of nonmetallic inclusions in steel using confocal scanning laser microscopy: A review

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**Abstract:** The characteristics of nonmetallic inclusions formed during steel production have a significant influence on steel performance. In this paper, studies on inclusions using confocal scanning laser microscopy (CSLM) are reviewed and summarized, particularly the collision of various inclusions, dissolution of inclusions in liquid slag, and reactions between inclusions and steel. Solid inclusions exhibited a high collision tendency, whereas pure liquid inclusions exhibited minimal collisions because of the small attraction force induced by their  $<90^\circ$  contact angle with molten steel. The collision of complex inclusions in molten steel was not included in the scope of this study and should be evaluated in future studies. Higher  $\text{CaO}/\text{Al}_2\text{O}_3$  and  $\text{CaO}/\text{SiO}_2$  ratios in liquid slag promoted the dissolution of  $\text{Al}_2\text{O}_3$ -based inclusions. The formation of solid phases in the slag should be prevented to improve dissolution of inclusions. To accurately simulate the dissolution of inclusions in liquid slag, *in-situ* observation of the dissolution of inclusions at the steel–slag interface is necessary. Using a combination of CSLM and scanning electron microscopy–energy dispersive spectroscopy, the composition and morphological evolution of the inclusions during their modification by the dissolved elements in steel were observed and analyzed. Although the *in-situ* observation of MnS and TiN precipitations has been widely studied, the *in-situ* observation of the evolution of oxide inclusions in steel during solidification and heating processes has rarely been reported. The effects of temperature, heating and cooling rates, and inclusion characteristics on the formation of acicular ferrites (AFs) have been widely studied. At a cooling rate of 3–5 K/s, the order of AF growth rate induced by different inclusions, as reported in literature, is  $\text{Ti-O} < \text{Ti-Ca-Zr-Al-O} < \text{Mg-O} < \text{Ti-Zr-Al-O} < \text{Mn-Ti-Al-O} < \text{Ti-Al-O} < \text{Zr-Ti-Al-O}$ . Further comprehensive experiments are required to investigate the quantitative relationship between the formation of AFs and inclusions.

**Keywords:** inclusion; steel; *in-situ* observation; confocal scanning laser microscopy

## 1. Introduction

The formation of nonmetallic inclusions during production of high-quality steels is a significant issue. Steel performance is related to quantity, size, composition, and distribution of such inclusions. The characteristics of these inclusions evolve during steel processing; for example, nucleation and collision occur after deoxidation during converter tapping [1–3]. During the refining process, inclusions float to the surface of the molten steel [4–6] and are dissolved in the refining slag [7–9]. Additionally, alloy treatment [10–13] and slag modification [14–16] can modify inclusion composition. Moreover, reoxidation and slag entrainment can be prevented to improve steel cleanliness [17–19]. The inclusion behavior during steelmaking, refining, and continuous casting processes, which cannot be directly observed because of the high temperature of molten steel, must be investigated.

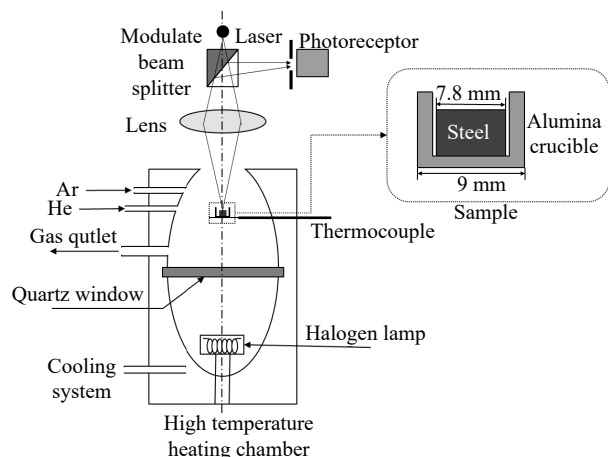
Confocal scanning laser microscopy (CSLM) combines advanced technologies such as infrared heating, high-temperature stretching and compression, and confocal laser scan-

ning. A schematic of CSLM equipment is shown in Fig. 1 [20]. The surface of samples can be observed *in-situ* under different conditions of temperature and pressure. CSLM is widely used in industries such as welding, materials, and metallurgy for purposes such as *in-situ* observation of the formation of acicular ferrites (AFs) induced by oxide metallurgy [21], phase transformation during cooling and heating of materials [22–23], and metal solidification and inclusion evolution on the surface of liquid metals [24–25].

In the 1950s, Minsky proposed the basic concept of confocal microscopy and applied it to technical patents. In the 1990s, the rapid development of optical, electronic, and computer technologies promoted the development and application of CSLM, which is commonly used to observe micro-morphology [26]. Chikama *et al.* [27] used a laser beam and infrared furnace to observe the dynamics of crystal growth in iron–carbon melts; CSLM has since been widely used in the study of steel. Particularly, CSLM has been widely applied for the *in-situ* observation of inclusion behaviors on molten steel surfaces. In 1997, Yin's group [28–29] used CSLM to

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**Fig. 1.** Schematic of CSLM equipment [20]. Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B, In situ observation of the agglomeration of MgO–Al<sub>2</sub>O<sub>3</sub> inclusions on the surface of a molten GCr15-bearing steel*, M.H. Wu, C.Y. Ren, Y. Ren, and L.F. Zhang, Copyright 2023.

observe the collision of different inclusions on the surface of molten steel. In 1998, Shibata *et al.* [30] observed the engulfment and pushing of inclusions in molten steel at the melt–solid interface using CSLM. In 1999, Hanamura *et al.* [31] used CSLM for the *in-situ* observation of the influence of inclusions on ferrite formation. In 2000, Sridhar and Cramb [32] used CSLM to study the dissolution of Al<sub>2</sub>O<sub>3</sub> inclusion particles in Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub>–CaO–MgO slags at 1430–1550°C. In 1998, Yuki *et al.* [24] observed the precipitation of MnS inclusions in Fe–42wt%Ni using CSLM. In 2017, Khurana *et al.* [33] observed the modification of inclusions by calcium treatment using CSLM. Thus, by combining *in-situ* observation and temperature simulation of the steel production process, CSLM can effectively monitor the evolution of inclusions in molten steel.

## 2. In-situ observation of the collision of inclusions in steel

In recent years, researchers have conducted numerous CSLM experiments on the collision of inclusions, including Al<sub>2</sub>O<sub>3</sub> [28–29,34], 80wt% Al<sub>2</sub>O<sub>3</sub>–20wt% SiO<sub>2</sub> [28,35], MgO [36], 93wt% Al<sub>2</sub>O<sub>3</sub>–7wt% MgO [36–37], Al<sub>2</sub>O<sub>3</sub>–CaO–SiO<sub>2</sub> [38], Al<sub>2</sub>O<sub>3</sub>–CaO [28,33,39–41], CaO–MgO–Al<sub>2</sub>O<sub>3</sub> [38], CaO–MgO–Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> [35,42], Al<sub>2</sub>O<sub>3</sub>–Ce<sub>2</sub>O<sub>3</sub> [43–44], Ce<sub>2</sub>O<sub>2</sub>S [45], SiO<sub>2</sub> [46], TiN [25–27], MnS [47–49], and Ti<sub>2</sub>O<sub>3</sub> [50]. Primary interfaces include steel–slag interface [33,39–42,50–51], steel–Ar interface [28,34,36,50–55], solid–liquid interface [37,39,56], and slag surface [40]. Table 1 summarizes related studies on the observation of inclusion collision using CSLM [25,28–29,34–40,45,47,50,53,56–63].

The collisions of inclusions are related to their liquid fractions. The inclusion composition in steel samples before the CSLM experiments was analyzed using scanning electron microscopy–energy dispersive spectroscopy (SEM–EDS). The melting temperature was calculated using the FactSage software to evaluate the solid and liquid inclusions. As shown in Fig. 2(a) [28], Yin reported that solid inclusions, such as

Al<sub>2</sub>O<sub>3</sub>, exhibited a significant attractive force and had a high tendency to collide. However, pure liquid inclusions such as CaO–Al<sub>2</sub>O<sub>3</sub> rarely collided because of their small mutual attraction. The collision between inclusions was due to the long-range attractive force between solid particles on the steel surface, which was caused by capillary action [64]. Fig. 2(b) [60] shows that there is no significant collision of liquid inclusions. Fig. 3 shows the variation in attractive force between inclusions at different distances [28,36–38,44,46,52].  $R_1$  and  $R_2$  denote the radii of the two inclusions. Capillary force between inclusions was primarily influenced by the contact angle between the inclusions and steel, as well as surface tension and density of molten steel. Because surface tension and density of molten steel changed minimally during steelmaking, the capillary force of the inclusions was primarily related to the contact angle between the inclusions and steel. The order of capillary force for different inclusions was Al<sub>2</sub>O<sub>3</sub> > TiAlO<sub>x</sub>–Al<sub>2</sub>O<sub>3</sub> > Ce<sub>2</sub>O<sub>3</sub> > Ce–Al–O > Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> > Ce–O–S > Al<sub>2</sub>O<sub>3</sub>–CaO > SiO<sub>2</sub> > MgO > MgAl<sub>2</sub>O<sub>4</sub>; this was positively correlated with the change in contact angles between the inclusions and steel [65]. Notably, attraction between inclusions refers to that between inclusions of the same type. For instance, the attraction between Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> composite inclusions refers to the attraction between one or more Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> composite inclusions. Previous investigations primarily focused on the collision of inclusions at the steel–Ar interface, and the collision of inclusions at the steel–slag interface has become more complex. Both long-range attractive and repulsive forces acted when the distance between two inclusions was less than 100–150 μm [39]. Liquid Al<sub>2</sub>O<sub>3</sub>–CaO inclusions rarely collided at the steel–slag interface, and the collision ability of the inclusions significantly increased [40].

The capillary force between inclusions under various physical and chemical conditions was calculated according to the attractive capillary force model [53], as shown in Fig. 4. According to Fig. 4(a) and (b), the attractive capillary force exhibits a positive correlation with the density and size of the inclusions. The larger the density and size of the inclusions, the greater the capillary force on the surface of molten steel. The dependence of the capillary forces on contact angle is shown in Fig. 4(c) [53]. The attractive capillary force first decreased and then increased and was very close to zero at a contact angle of 90°. The correlation between surface tension of liquid steel and capillary suction force is shown in Fig. 4(d) [53]. When surface tension decreased from 1.9 to 1.1 J/m<sup>2</sup>, the attractive capillary force increased slightly, leading to collisions of inclusions.

## 3. In-situ observation of the dissolution of inclusions in slag

In 2000, Sridhar and Cramb [32] *in-situ* observed the dissolution of Al<sub>2</sub>O<sub>3</sub> inclusions in CaO–MgO–SiO<sub>2</sub>–Al<sub>2</sub>O<sub>3</sub> slag using CSLM, as shown in Fig. 5. The inclusion particles in previous studies were primarily pure oxides such as Al<sub>2</sub>O<sub>3</sub>,

**Table 1.** Studies on the observation of inclusion collisions using CSLM [25,28–29,34–40,45,47,50,53,56–63]

Authors	Year	Interface	Inclusion composition	Inclusion density / (kg·m <sup>-3</sup> )	Inclusion size / μm	Capillary attraction / N	Refs.
Yin <i>et al.</i>	1997	Steel–Ar	Al <sub>2</sub> O <sub>3</sub>	3900	10	10 <sup>-16</sup>	[28–29]
			Al <sub>2</sub> O <sub>3</sub>	3900	2		[34]
			80wt%Al <sub>2</sub> O <sub>3</sub> –SiO <sub>2</sub>	3378	1–3	7.0 × 10 <sup>-16</sup>	[28,34]
			CaO–60wt%Al <sub>2</sub> O <sub>3</sub> –SiO <sub>2</sub>		5–10	Noncollision	[34]
			CaO–80wt%Al <sub>2</sub> O <sub>3</sub> –SiO <sub>2</sub>		1–3	4.0 × 10 <sup>-16</sup>	[34]
			CaO–Al <sub>2</sub> O <sub>3</sub> –95wt%SiO <sub>2</sub>		5–10	8.2 × 10 <sup>-15</sup>	[34]
			CaO–50wt%Al <sub>2</sub> O <sub>3</sub> –SiO <sub>2</sub>		5–10	Noncollision	[34]
			CaO–80wt%Al <sub>2</sub> O <sub>3</sub>	3768	3–5	6.5 × 10 <sup>-14</sup>	[34]
			CaO–60wt%Al <sub>2</sub> O <sub>3</sub>	3645	5–10	10 <sup>-16</sup>	[34]
			CaO·Al <sub>2</sub> O <sub>3</sub>		5–10	Noncollision	[34]
Kimura <i>et al.</i>	2000 2001	Steel–Ar	Al <sub>2</sub> O <sub>3</sub> ·MgO	3600	5.3	5.0 × 10 <sup>-18</sup> – 5.0 × 10 <sup>-16</sup>	[36–37]
Kimura <i>et al.</i>	2001	Steel–Ar	MgO	3580			[36]
Nakajima and Mizoguchi	2001	Steel–Ar	30wt%CaO–60wt%Al <sub>2</sub> O <sub>3</sub> – 10wt%MgO		1.5–15.5	10 <sup>-14</sup> –10 <sup>-17</sup>	[38]
Nakajima and Mizoguchi	2001	Steel–Ar	40wt%CaO–55wt%Al <sub>2</sub> O <sub>3</sub> – 5wt%MgO		<40	Noncollision	[38]
Hasegawa <i>et al.</i>	2001	Steel–Ar	MnS		1–10	Noncollision	[47]
Vantilt <i>et al.</i>	2004	Steel–Ar	Al <sub>2</sub> O <sub>3</sub> –MnO–SiO <sub>2</sub>		10	Noncollision	[51]
Wikstrom <i>et al.</i>	2008	Slag–Ar	CaO·Al <sub>2</sub> O <sub>3</sub>				[39]
Wikstrom <i>et al.</i>	2008	Steel–Ar	CaO·Al <sub>2</sub> O <sub>3</sub>		10–80	10 <sup>-13</sup> –10 <sup>-15</sup>	[40]
Appelberg <i>et al.</i>	2008	Steel–Ar	Ce <sub>2</sub> O <sub>3</sub>	7100	20		[43]
Kang <i>et al.</i>	2011	Steel–Ar	CaO·Al <sub>2</sub> O <sub>3</sub>			Noncollision	[57]
Wen and Song	2012	Steel–Ar	Ce <sub>2</sub> O <sub>2</sub> S	6000	<5	Noncollision	[45]
Jiang <i>et al.</i>	2014	Steel–Ar	CaO–MgO–Al <sub>2</sub> O <sub>3</sub> –SiO <sub>2</sub>		5		[56]
Mu <i>et al.</i>	2017 2018	Steel–Ar	TiN	5430			[50,53]
Tian <i>et al.</i>	2018	Steel–Ar	TiN	5430	10		[58]
Mu and Xuan	2019	Steel–Ar	TiO <sub>x</sub> –Al <sub>2</sub> O <sub>3</sub>		10–20		[59]
Wang and Liu	2020	Steel–Ar	Al <sub>2</sub> O <sub>3</sub>	3900	15–30	10 <sup>-15</sup> –3.0 × 10 <sup>-14</sup>	[44]
			Ce–Al–O		10–30	1.3 × 10 <sup>-16</sup> – 2.0 × 10 <sup>-14</sup>	[44]
			Ce <sub>2</sub> O <sub>3</sub>	7100	10–30	1.3 × 10 <sup>-16</sup> – 2.0 × 10 <sup>-14</sup>	[44]
			Ce–O–S		5–20	2.1 × 10 <sup>-18</sup> – 6.0 × 10 <sup>-16</sup>	[44]
Wu <i>et al.</i>	2023	Steel–Ar	MgO	3580	5–20	10 <sup>-17</sup> –10 <sup>-15</sup>	[20,60]
Misra <i>et al.</i>	2001	Steel–slag	TiN	5430	5		[25]
Lee <i>et al.</i>	2001	Steel–slag	Al <sub>2</sub> O <sub>3</sub>	3900			[41]
Coletti <i>et al.</i>	2003	Steel–slag	CaO·Al <sub>2</sub> O <sub>3</sub>		3–4		[42]
Vantilt <i>et al.</i>	2004	Steel–slag	Al <sub>2</sub> O <sub>3</sub> –MnO–SiO <sub>2</sub>			Noncollision	[51]
Wikstrom <i>et al.</i>	2008	Steel–slag	CaO·Al <sub>2</sub> O <sub>3</sub>				[39–40]
Mu <i>et al.</i>	2016	Steel–slag	MgO	3580	6	Noncollision	[61]
Misra <i>et al.</i>	2000	Steel–slag	Al <sub>2</sub> O <sub>3</sub>	3900	5	10 <sup>-16</sup> –3.0 × 10 <sup>-15</sup>	[62]
Michelic <i>et al.</i>	2015	Steel–slag	Al <sub>2</sub> O <sub>3</sub> , Al <sub>2</sub> O <sub>3</sub> –MgO	3900		Noncollision	[63]

MgO, and SiO<sub>2</sub>, and the slags used were mostly liquid Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub>–CaO and Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub>–CaO–MgO. Slag compositions reported in literature are plotted in the Al<sub>2</sub>O<sub>3</sub>–CaO–SiO<sub>2</sub> diagram shown in Fig. 6 [7,32,41,66–79]. The dissolution mechanisms of different inclusions in the slag are summarized, including diffusion in liquid slag (DLS), boundary layer diffusion (BLD), chemical reaction

(CR), and product layer diffusion (PLD) controls. The dissolution mechanism of inclusions in slag was determined by comparing the observed and calculated dissolution curves. The effects of temperature, inclusion properties, and slag composition on inclusion dissolution rates in slag were investigated using CSLM, as summarized in Table 2 [7,32,41,66–90].



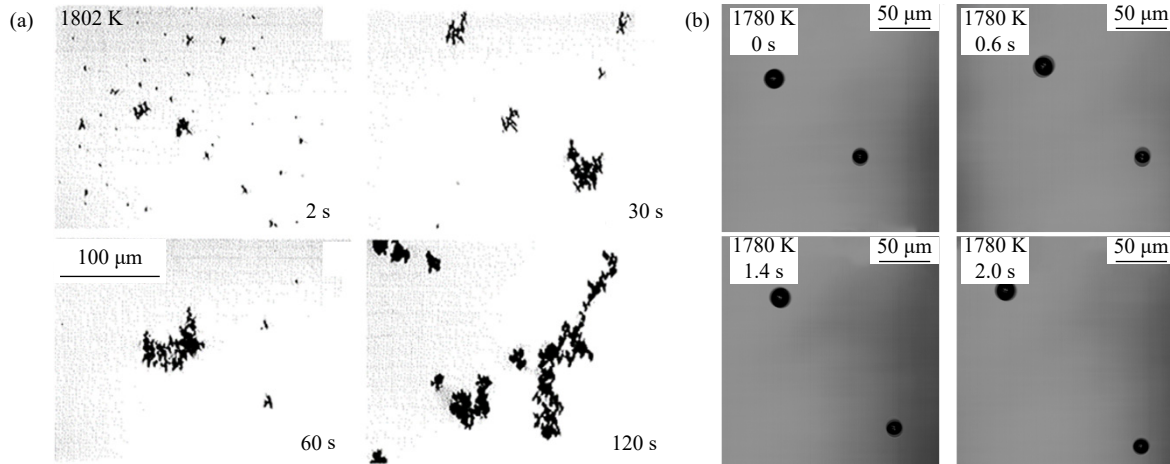


Fig. 2. Collision of inclusions on the surface of the molten steel: (a) solid inclusions [28]; (b) liquid inclusions [60]. (a) Reprinted with permission from Ref. [28]. © 1997 The Iron and Steel Institute of Japan. (b) Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, In-situ Observation of collision and agglomeration of  $\text{La}_2\text{O}_3$ – $\text{La}_2\text{S}_3$  inclusions on surface of molten 304 stainless steels, M.H. Wu, Y. Ren, and L.F. Zhang, Copyright 2024.

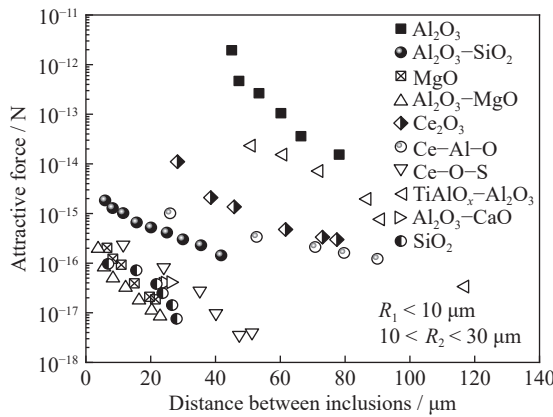


Fig. 3. Change in the attractive force between two inclusions with distance [28,36–38,44,46,52].

Chemical compositions of the inclusions and slag influence the dissolution rates of inclusion particles in slags, as shown in Fig. 7. Fig. 7(a) [90] shows the plots of the inclusion size and dissolution rate. The dissolution rate of inclusions increased with an increase in their diameter. When the experimental temperature was 1773 K, the order of inclusion dissolution rate was  $\text{SiO}_2 > \text{Al}_2\text{O}_3 > \text{ZrO}_2$ . Fig. 7(b) [7,89] and (c) [80] shows the effects of the  $\text{CaO}/\text{Al}_2\text{O}_3$  and  $\text{CaO}/\text{SiO}_2$  mass ratios of the slag on the dissolution rate of inclusions, respectively. Increasing the  $\text{CaO}/\text{Al}_2\text{O}_3$  and  $\text{CaO}/\text{SiO}_2$  mass ratios in the slag is beneficial for the dissolution of inclusions in liquid slag. Solid phase formation in slag should be prevented to improve inclusion dissolution. When the temperature increased from 1773 to 1853 K, the dissolution rate of the inclusions increased.

Inclusion dissolution rate was related to the ratio of inclusion solubility to slag viscosity ( $\Delta C/\eta$ ). The correlation between dissolution time of inclusions [7,32,66–67,76,80–81,89] and the parameter  $\Delta C/\eta$  is substituted in Eq. (1) [90], as shown in Fig. 8(a). The larger the value of  $\Delta C/\eta$ , the faster the dissolution of inclusions. However,  $\Delta C/\eta$  is a parameter with dimensions, which cannot be used to explain the relationship between dissolution rate and inclusion size.

Zhang and Ren [91] proposed the inclusion capacity of refining slag ( $Zh$ ), which is a dimensionless number, as defined in Eq. (2). The dimensionless dissolution rate of inclusions in the slag is proposed in Eq. (3). The empirical equations of  $Zh$  and  $Ry$  for the  $\text{Al}_2\text{O}_3$ ,  $\text{ZrO}_2$ , and  $\text{SiO}_2$  inclusions were plotted using Eqs. (4)–(6) [90], as shown in Fig. 8(b). The inclusion capacity of the slag was introduced to predict the dissolution rate of inclusions. The larger the  $Zh$  number and inclusion size, the longer the dissolution time of the inclusions in liquid slag.

$$\tau = 5.03 \times 10^3 \left( \frac{\Delta C}{\eta} \right)^{-1} \quad (1)$$

$$Zh = \frac{g \cdot \rho_{\text{slag}}^2 \cdot (C_{\text{saturation}} - C) \cdot d_{p,0}^3}{\eta_{\text{slag}}^2} \quad (2)$$

$$Ry = \frac{1}{v_{\text{slag}} \cdot d_{p,0}} \varphi = \frac{\rho_{\text{slag}}}{\eta_{\text{slag}} \cdot d_{p,0}} \varphi \quad (3)$$

$$Ry_{\text{Al}_2\text{O}_3} = 2.93 \times 10^{-6} Zh^{0.165} \quad (4)$$

$$Ry_{\text{ZrO}_2} = 1.02 \times 10^{-6} Zh^{0.089} \quad (5)$$

$$Ry_{\text{SiO}_2} = 1.08 \times 10^{-4} Zh^{0.079} \quad (6)$$

where  $C$  is the content of inclusions in the slag;  $C_{\text{saturation}}$  is the saturation content of inclusions;  $d_{p,0}$  is the initial inclusion diameter (m);  $\rho_{\text{slag}}$  is the slag density ( $\text{kg}/\text{m}^3$ );  $\eta_{\text{slag}}$  is the slag dynamic viscosity ( $\text{kg}/(\text{m} \cdot \text{s})$ );  $g$  is the gravitational acceleration rate ( $\text{m}/\text{s}^2$ ); and  $\varphi$  is the dissolution rate of the inclusion particles in the slag ( $\text{m}^3/\text{s}$ ). The  $Zh$  number represents the dissolution capacity of inclusions by the refining slag; The  $Ry$  number represents the dissolution rate of inclusion particles in the refining slag.

#### 4. In-situ observation of the modification of inclusions in steel

Alloy addition is an effective method to modify inclusions in steel. To address the limitation of sample loading in CSLM, researchers have attempted to observe inclusion

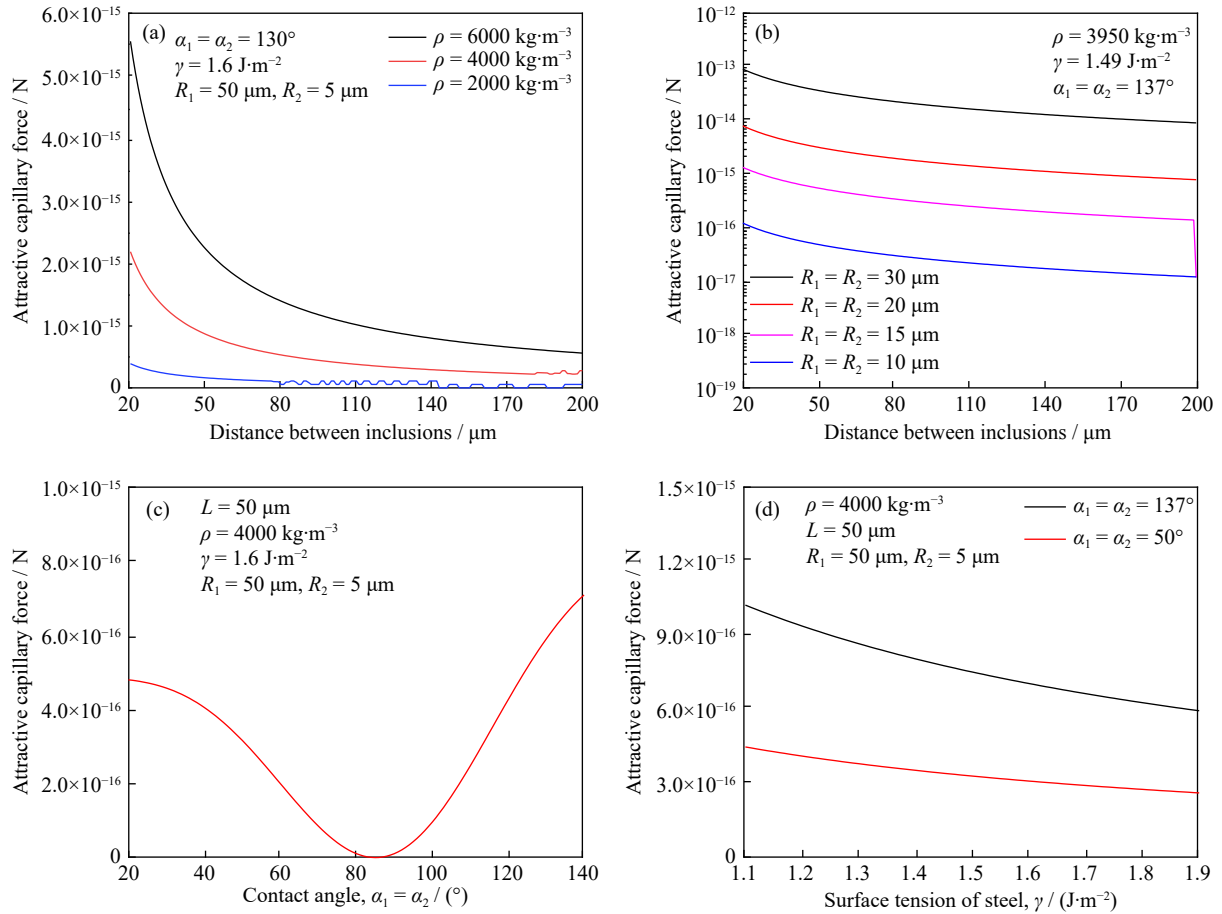


Fig. 4. Calculated attractive force between different inclusions under various conditions: (a) density; (b) size; (c) contact angle [53]; (d) surface tension [53]. (c, d) Adapted from [53].  $\rho$ —Density;  $\alpha_1, \alpha_2$ —Contact angle;  $\gamma$ —Surface tension;  $R_1, R_2$ —Radius.

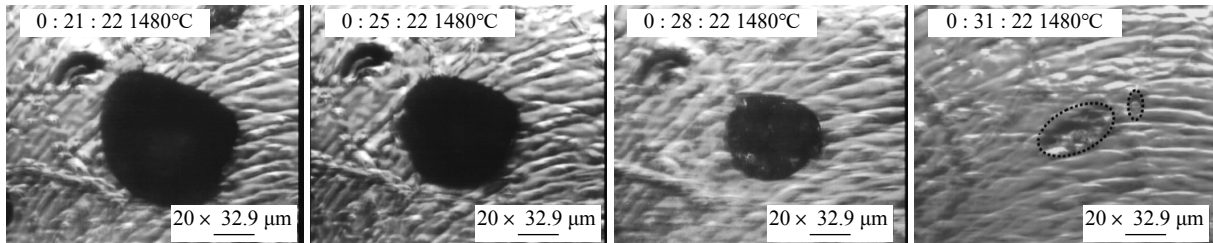


Fig. 5. Images of dissolution process of  $\text{Al}_2\text{O}_3$  inclusions in slag [32]. Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, Kinetics of  $\text{Al}_2\text{O}_3$  dissolution in  $\text{CaO-MgO-SiO}_2\text{-Al}_2\text{O}_3$  slags: *In Situ* observations and analysis, S. Sridhar and A.W. Cramb, Copyright 2000.

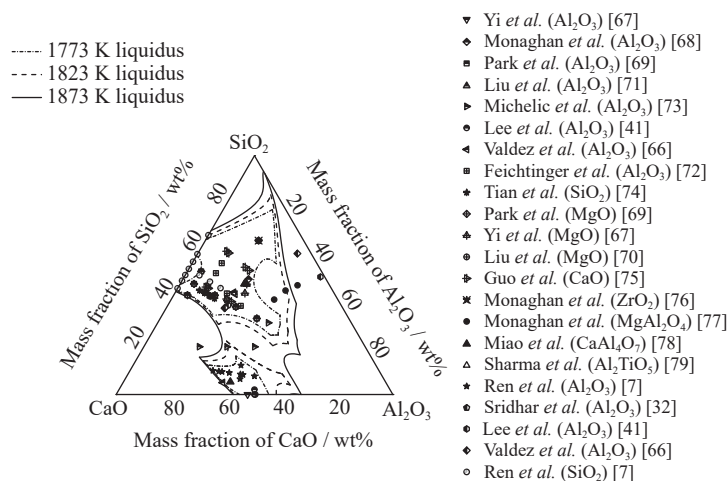


Fig. 6. Reported studies on the inclusion dissolution in  $\text{CaO-Al}_2\text{O}_3\text{-SiO}_2$  slags using CSLM [7,32,41,66–79].

**Table 2. Summarized studies on the inclusion dissolution in slags using CSLM [7,32,41,66–90]**

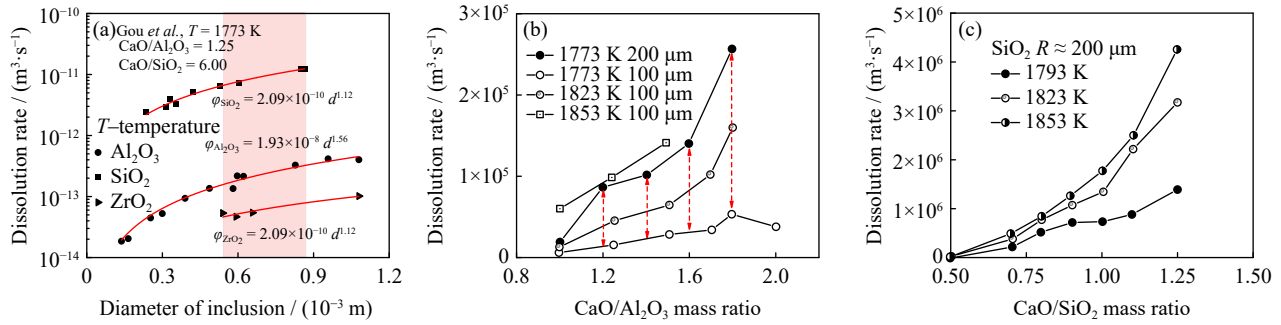
Authors	Year	Inclusion	Diameter / $\mu\text{m}$	Slag system	T / K	Rate-limiting step
Sridhar and Cramb [32]	2000	$\text{Al}_2\text{O}_3$	84–138	CASM	1703–1823	BLD
Lee <i>et al.</i> [41]	2001	$\text{Al}_2\text{O}_3$	50–90	CAS	1723–1773	
Valdez <i>et al.</i> [66]	2002	$\text{Al}_2\text{O}_3$	200	CAS and CASM	1723–1803	CR and BLD
Yi <i>et al.</i> [67]	2003	MgO	190–330	CAM	1723–1873	CR
		$\text{Al}_2\text{O}_3$	200			DLS
Fox <i>et al.</i> [81]	2004	$\text{Al}_2\text{O}_3$		CASM	1523–1773	CR
		$\text{ZrO}_2$	200, 550			
		MgO	200			
		$\text{MgAl}_2\text{O}_4$				
Monaghan <i>et al.</i> [68,76,82]	2004	$\text{MgAl}_2\text{O}_4$	80		1750–1805	DLS
		$\text{Al}_2\text{O}_3$	100		1777–1850	DLS
	2005	$\text{MgAl}_2\text{O}_4$	80	CAS	1753–1805	DLS
		$\text{ZrO}_2$	100		1750–1805	
Monaghan and Chen [77]	2006	$\text{MgAl}_2\text{O}_4$	28–35	CAS	1777	DLS
Park <i>et al.</i> [69]	2006	MgO		CAS	1823	PLD
Liu <i>et al.</i> [71]	2007	$\text{Al}_2\text{O}_3$	500	CAS	1743–1903	DLS
Liu <i>et al.</i> [70]	2007	MgO	300–400	CASM	1500–1600	BLD
Verhaeghe <i>et al.</i> [83]	2007	$\text{Al}_2\text{O}_3$	500	CAS	1743	DLS
Park <i>et al.</i> [84]	2010	SiC	612, 820	CSM (MnO)	1873	
Sun <i>et al.</i> [85]	2013	CaO	400–500	CAS and CASM	1723–1873	
Feichtinger <i>et al.</i> [72]	2014	$\text{SiO}_2$	612–696	CAS	1723	DLS
Guo <i>et al.</i> [75]	2014	CaO		CAS, CASM	1723–1873	CR, BLD, PLD
Michelic <i>et al.</i> [73]	2016	$\text{Al}_2\text{O}_3$	400	CASM	1873	DLS
Miao <i>et al.</i> [78]	2018	$\text{CA}_2$	222–270	CAS	1773–1873	DLS
Sharma <i>et al.</i> [79]	2018	$\text{Al}_2\text{O}_3$	320	CAS	1823	BLD
		$\text{Al}_2\text{TiO}_5$	400, 900			BLD
Tian <i>et al.</i> [74]	2019	$\text{SiO}_2$	~1000	CASM	1743–1803	DLS
Lee <i>et al.</i> [86]	2019	MgO	370	$\text{MgF}_2\text{--CaF}_2\text{--LiF}$ , $\text{MgF}_2\text{--CaF}_2\text{--NaF}$	1473	BLD
Sharma and Dogan [87]	2020	$\text{Al}_2\text{TiO}_5$		CAS	1773	BLD
Park <i>et al.</i> [88]	2020	$\text{Al}_2\text{O}_3$		CASM	1823	DLS
Ren <i>et al.</i> [7,89]	2021	$\text{Al}_2\text{O}_3$	~200, ~400	CAS	1773–1853	DLS
Ren <i>et al.</i> [80]	2022	$\text{SiO}_2$	~200	CASM	1793–1853	DLS
Gou <i>et al.</i> [90]	2023	$\text{Al}_2\text{O}_3$	100–1100	CAS	1773	DLS
		$\text{ZrO}_2$	500–1100			BLD
		$\text{SiO}_2$	200–900			DLS

Note: Legend of slag system, A— $\text{Al}_2\text{O}_3$ ; S— $\text{SiO}_2$ ; C—CaO; M—MgO; L— $\text{Li}_2\text{O}$ .

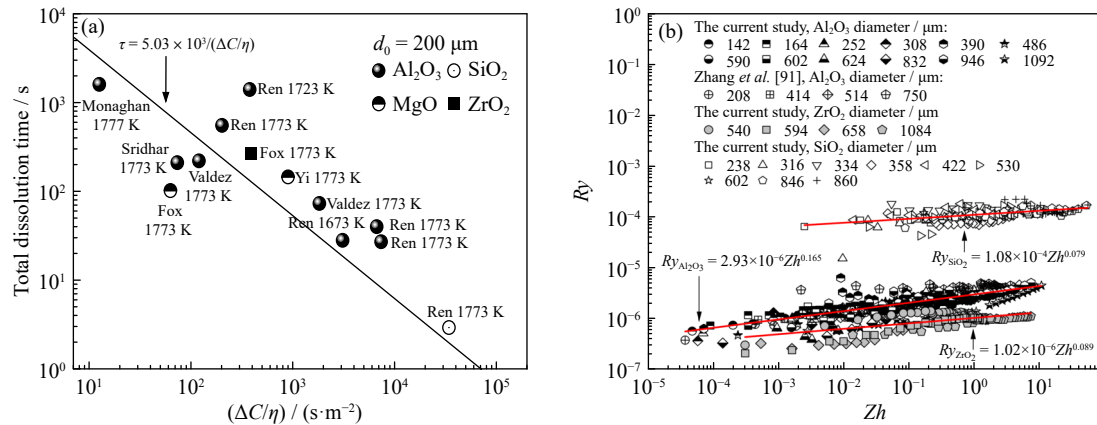
modifications using CSLM, as shown in Fig. 9. Fig. 9(a) [33] and (b) [92] shows endogenous inclusions colliding with each other on the surface of molten steel, which made tracking the modification behavior of a specific inclusion in steel challenging. Additionally, determining the starting time for inclusion modification was difficult. Thus, extraneous  $\text{Al}_2\text{O}_3$  particles were proposed to be placed on the surface of the steel sample. After melting the steel sample, the inclusions were modified by the active elements in the steel, as shown in Fig. 9(c) [93]. The melting time of the steel samples was recorded as the start time of the modification reaction. The morphological and compositional evolution of the inclusions during the modification process was analyzed using a combination of CSLM and SEM-EDS. Thus, the method illustrated in Fig. 9(c) was suggested to improve the tracking of modification behavior of a specific inclusion.

Khurana *et al.* [33] placed CaO powder at the bottom of an

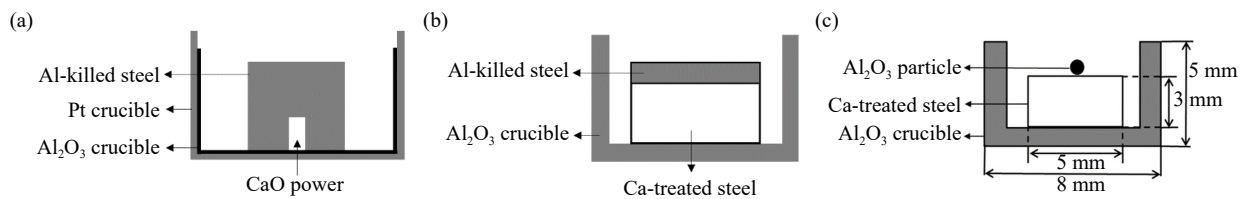
Al-killed steel sample using the method shown in Fig. 9(a). After melting the steel samples, inclusions in steel before calcium treatment were  $\text{Al}_2\text{O}_3$ . After calcium treatment with CaO powder, the inclusions were modified to  $\text{Al}_2\text{O}_3\text{--CaO--CaS}$ . Wang and Liu [92] used the method shown in Fig. 9(b) to heat and melt an Al-killed steel sample, enabling numerous cluster inclusions to float to the surface of the sample. The upper surface of Al-killed steel was cut and placed on top of the Ca-treated steel to expose the aggregated inclusions in the steel for observation. The irregular solid  $\text{Al}_2\text{O}_3$  inclusions in the steel changed to liquid spherical inclusions within approximately 40 s of the steel melting, as shown in Fig. 10 [92]. Chen *et al.* [93] placed  $\text{Al}_2\text{O}_3$  particles on the surface of steel samples. Compositional evolution of  $\text{Al}_2\text{O}_3$  inclusions with different initial sizes during the modification process is shown in Fig. 11 [93]. C3A, C12A7, CA, CA2, CA6, and  $\text{Al}_2\text{O}_3$  in the right vertical axis represent



**Fig. 7.** Influencing factors on the dissolution of inclusion in slag: (a) inclusion diameters [90]; (b) CaO/Al<sub>2</sub>O<sub>3</sub> ratio of the slag [7,89]; (c) CaO/SiO<sub>2</sub> ratio of the slag [80]. (a) Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, Concept of inclusion capacity of slag and its application on the dissolution of Al<sub>2</sub>O<sub>3</sub>, ZrO<sub>2</sub> and SiO<sub>2</sub> inclusions in CaO–Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> slag, L. Gou, H. Liu, Y. Ren, and L. Zhang, Copyright 2023. (b) Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, In situ observation of the dissolution of Al<sub>2</sub>O<sub>3</sub> particles in CaO–Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> slags, C.Y. Ren, L.F. Zhang, J. Zhang, et al., Copyright 2021; reprinted by permission from Springer Nature: *Int. J. Miner. Metall. Mater.*, In situ observation of the dissolution kinetics of Al<sub>2</sub>O<sub>3</sub> particles in CaO–Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> slags using laser confocal scanning microscopy, C.Y. Ren, C.D. Huang, L.F. Zhang, et al., Copyright 2023. (c) Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, Dissolution of SiO<sub>2</sub> inclusions in CaO–SiO<sub>2</sub>-based slags *in situ* observed using high-temperature confocal scanning laser microscopy, Y. Ren, P. Zhu, C.Y. Ren, et al., Copyright 2022.  $\phi_{\text{SiO}_2}$ ,  $\phi_{\text{Al}_2\text{O}_3}$ , and  $\phi_{\text{ZrO}_2}$  are dissolution rates of SiO<sub>2</sub>, Al<sub>2</sub>O<sub>3</sub>, and ZrO<sub>2</sub> inclusions, and  $R$  is radius.



**Fig. 8.** Reported models for the dissolution rate of inclusions in slags: (a) correlation between dissolution time and  $\Delta C/\eta$  of slag [7,32,66–67,76,80–81,89–90]; (b) dependence of the  $Ry$  number on the  $Zh$  number [90–91]. (b) Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, Concept of inclusion capacity of slag and its application on the dissolution of Al<sub>2</sub>O<sub>3</sub>, ZrO<sub>2</sub> and SiO<sub>2</sub> inclusions in CaO–Al<sub>2</sub>O<sub>3</sub>–SiO<sub>2</sub> slag, L. Gou, H. Liu, Y. Ren, et al., Copyright 2023.  $\tau$ —Dissolution time of inclusions.



**Fig. 9.** Schematic of the observation method of the inclusion modification using CSLM: (a) Khurana et al. [33]; (b) Wang and Liu [92]; (c) Chen et al. [93]. (b) Adapted from Ref. [92]. (c) Reprinted with permission from Ref. [93]. © 2024 The Iron and Steel Institute of Japan.

3CaO·Al<sub>2</sub>O<sub>3</sub>, 12CaO·7Al<sub>2</sub>O<sub>3</sub>, CaO·Al<sub>2</sub>O<sub>3</sub>, CaO·2Al<sub>2</sub>O<sub>3</sub>, CaO·6Al<sub>2</sub>O<sub>3</sub>, and Al<sub>2</sub>O<sub>3</sub>, respectively, which correspond to the mass fraction of CaO in CaO–Al<sub>2</sub>O<sub>3</sub> inclusions in the left vertical axis. The CaO content in the inclusions gradually increased, and the Al<sub>2</sub>O<sub>3</sub> inclusions were gradually modified to CaO–Al<sub>2</sub>O<sub>3</sub>. The modification rates of inclusions decreased with time. The modification rates of the inclusions of various sizes differed owing to the different reaction areas. The liquid fraction of the inclusions gradually increased owing to

the increase in CaO content in the inclusions during the modification process.

## 5. In-situ observation of the precipitation and transformation of inclusions in steel

In 1998, Yuki et al. [24] observed the precipitation of MnS inclusions during solidification and cooling processes of an Fe–42wt%Ni alloy using CSLM, as shown in Fig. 12.



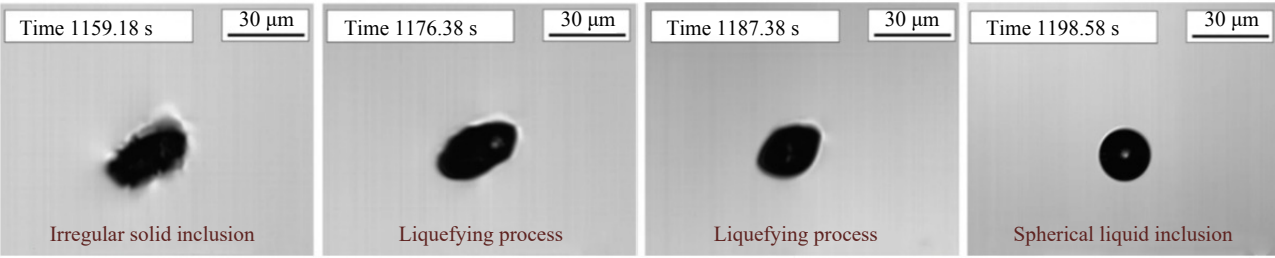


Fig. 10. *In-situ* observation of the modification of an endogenous inclusion using CSLM [92]. Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B, In situ* observation of transient evolution of inclusions by Ca treatment in molten steel, Y.G. Wang and C.J. Liu, Copyright 2022.

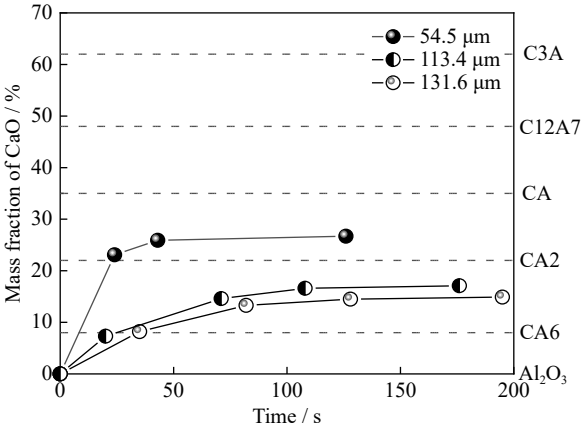


Fig. 11. Composition evolution of  $\text{Al}_2\text{O}_3$  inclusions with different initial sizes during the modification process analyzed using CSLM and SEM-EDS. Reprinted with permission from Ref. [93]. © 2024 The Iron and Steel Institute of Japan.

Recently, several scholars have utilized CSLM to study the precipitation of inclusions during cooling and heating processes, as summarized in Table 3 [24–25,58,94–113]. The morphology of MnS inclusions has an influence on the properties of steel. Valdez *et al.* [95] observed the precipitation of MnS inclusions during cooling and heating process of steel containing 0.05wt% S. Nakama *et al.* [96] reported that the sulfide precipitation was related to the solidification sequence of Fe–Cr–Ni alloys, which depended on Ni and Cr contents. Both high-Ni, low-Cr alloys solidifying in the primary  $\gamma$  field and low-Ni, high-Cr alloys solidifying in the primary  $\delta$  field led to sulfide precipitation in the enriched melt ahead of the dendrite fronts. In low-Ni, high-Cr alloys, sulfides are formed via liquid phase separation. By contrast, in intermediate Ni and Cr alloys with limited primary phase-field solidification, sulfides are formed within the solid

phase. Several researchers have investigated the correlation between MnS precipitation and the liquid fraction of steel during solidification [97–98,100–101,103,107]. The precipitation of MnS is also related to elemental concentrations in steels. The effects of Cu [114–115] and Te [102] contents on the precipitation characteristics and temperature of MnS were observed using CSLM. The addition of Cu or Te slows precipitation of MnS at the grain boundaries, which is beneficial for the uniform distribution of MnS in the steel. In 2001, Misra *et al.* [25] studied the precipitation of TiN inclusions at the steel–slag interface using CSLM. The morphology of TiN varied with temperature. At a temperature lower than the solidus of steel, increase in temperature increased the rate of TiN inclusions in steel. Tian *et al.* [58] *in-situ* observed the precipitation, collision, and dissolution behaviors of TiN inclusions on the surface of molten GCr15-bearing steel.

In 1996, Chikama *et al.* [27] observed the crystal growth process of Fe–C alloys and found that inclusions in steel were engulfed and entrapped by growing crystals during solidification and cooling of molten steel. Yin *et al.* [30,116] determined the flow direction and velocity of a molten steel surface near the solid–melt interface, as shown in Fig. 13 [116]. The velocity of the solutal Marangoni flow was very high, pushing the inclusions up to the surface along the steel–metal interface. These features were responsible for the entrapment of inclusions and solute segregation during the solidification and cooling of steel. Chen *et al.* [117] reported that the high growth rate of the  $\delta$  phase promoted the entrapment of inclusions. Yan *et al.* [118] reported that the movement of inclusions at the solidification front can be of three types: attraction, repulsion, and no effect.

CSLM has also been widely used to observe changes in inclusions during solid steel heating. In 2011, Shao *et al.* [108–109,119] utilized CSLM to observe morphological

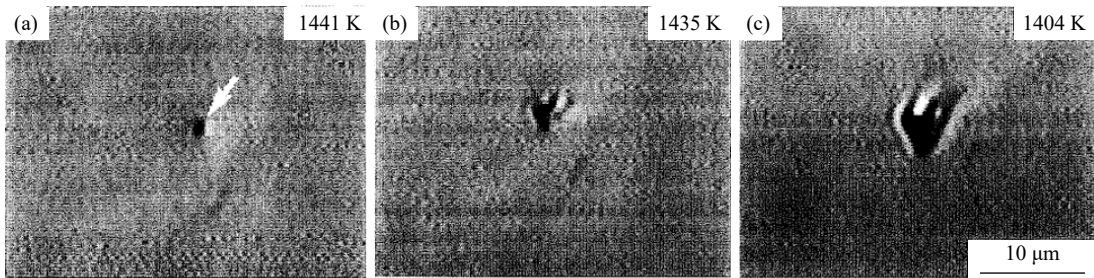
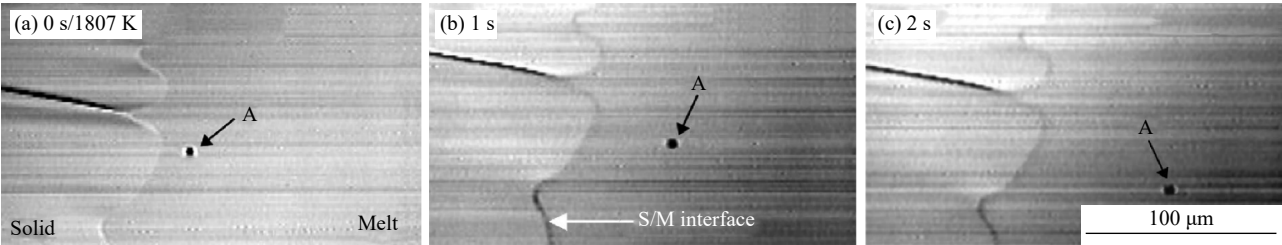


Fig. 12. Precipitation of MnS inclusions in steel observed using CSLM. Reprinted with permission from Ref. [24]. © 1998 The Iron and Steel Institute of Japan.

**Table 3.** Summarized *in-situ* observation of the precipitation and transformation of inclusions during solidification and heating processes [24–25,58,94–113]

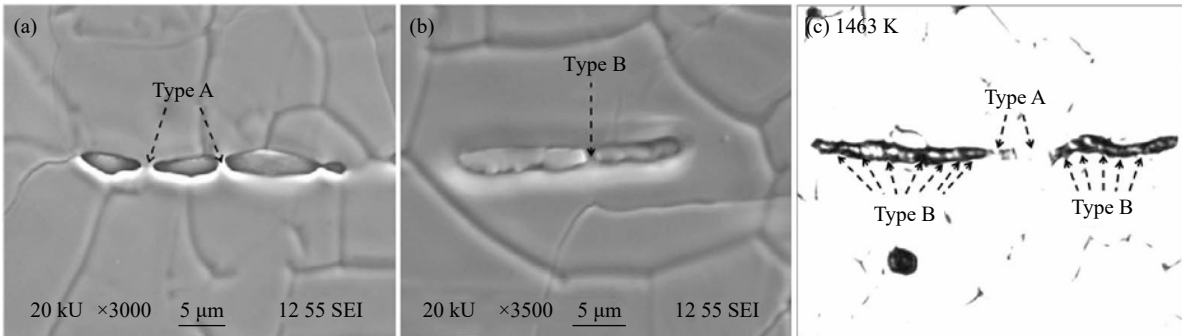
Authors	Year	Steel	Process	Inclusion	Temperature / K
Yuki <i>et al.</i> [24]	1998	Fe–Ni alloys	Solidification	MnS	1329–1438
Misra <i>et al.</i> [25]	2001	Stainless steel	Solidification	TiN	1553–1593, 1623–1773
Wang <i>et al.</i> [94]	2003	Al-killed Ca-treated steel	Solidification	CaS	
Valdez <i>et al.</i> [95]	2005	Resulfurized steel	Solidification	MnS	
Nakama <i>et al.</i> [96]	2009	Fe–Cr–Ni alloys	Solidification	MnS	1669–1738
Luo <i>et al.</i> [97]	2011	49MnVS3 steel	Solidification	MnS	
Hu <i>et al.</i> [98]	2018	Sulfur-bearing steel	Solidification	MnS	1700, 1746
Tian <i>et al.</i> [58,99]	2018	GCr15-bearing steel	Solidification	TiN	1640–1840
Tanaka <i>et al.</i> [100]	2019	High-carbon steel	Solidification	MnS	
Zeng <i>et al.</i> [101]	2020	High-sulfur microalloyed steel	Solidification	MnS	1703, 1713
Shen <i>et al.</i> [102]	2021	Te-treated resulfurized special steel	Solidification	Mn–S–Te	
Chu <i>et al.</i> [103]	2022	Medium/high-manganese steel	Solidification	MnS	
Cao <i>et al.</i> [104]	2022	GCr15-bearing steel	Solidification	MgO–TiN	1597
Liu <i>et al.</i> [105]	2022	Al-killed Ca-treated steel	Solidification	CaO–MgO–Al <sub>2</sub> O <sub>3</sub>	1273–1473
Zhu <i>et al.</i> [106]	2023	High-titanium steel	Solidification	TiC <sub>x</sub> N <sub>1–x</sub>	1649, 1707
Cao <i>et al.</i> [107]	2024	High-Al medium-Mn steel	Solidification	MnS	
Shao <i>et al.</i> [108]	2011	Free-cutting steel	Heating	MnS	873–1473
Shao <i>et al.</i> [109]	2011	Free-cutting steel	Heating	MnS	1463
Wang <i>et al.</i> [110]	2018	EH36 steel	Heating	TiN	1473
Wang <i>et al.</i> [111]	2018	EH36 steel	Heating	MnS	1473
Wang <i>et al.</i> [112]	2019	High-carbon steel	Heating	Rare-earth inclusions	1473, 1623
Song <i>et al.</i> [113]	2023	Heavy rail steel	Heating	MnS, oxides	1323, 1393



**Fig. 13.** Movement of an inclusion at the solid–melt interface due to surface flow during steel solidification at a cooling rate of 0.025 K/s [116]. Reprinted by permission from Springer Nature: *Metall. Mater. Trans. B*, Marangoni flow at the gas/melt interface of steel, H.B. Yin and T. Emi, Copyright 2003.

evolution of large slender MnS inclusions in sulfur-containing steels, as shown in Fig. 14 [108]. Moreover, they proposed a mechanism for the evolution of MnS inclusions during heating. The elongated MnS inclusions shrank and fractured during heating, forming spherical MnS. Song *et al.*

[113] observed the formation of MnS on oxide inclusions during heating of heavy rail steel. Wang *et al.* [111] reported that the amount of MnS decreased during EH36 steel heating at 1473 K and TiN inclusions precipitated in the steel. Moreover, CSLM can be used to observe the changes in ox-



**Fig. 14.** Observation of the morphological evolution of slender MnS inclusions in free-cutting steel at 1463 K using CSLM [108]. Reprinted with permission from Ref. [108]. © 2011 The Iron and Steel Institute of Japan.

ide inclusions during solid steel heating. Liu *et al.* [105] used CSLM to *in-situ* observe that inclusions evolved from CaO–MgO–Al<sub>2</sub>O<sub>3</sub> to CaO–MgO–Al<sub>2</sub>O<sub>3</sub>–CaS in solid Al-killed Ca-treated steels at 1273–1473 K. High temperatures promoted the formation of CaS in the inclusions owing to the large diffusion rate of the elements. They concluded that MnS precipitation was primarily related to the Mn and S contents and temperature of the steel and the formation of TiN was influenced by the Ti and N contents and temperature of the steel.

6. *In-situ* observation of the formation of intra-granular acicular ferrite at inclusions in steel

Oxide metallurgy was invented in the 1990s and has been widely used in manufacturing of high-strength steel components such as ship plates and pipes. During the welding process, high heat input leads to a reduction in toughness. Nano-

scale inclusions have been reported to pin the grain boundary. Moreover, micron-scale inclusions can promote the nucleation of AF on the surface of inclusions, which has become an effective method to increase steel toughness at low temperatures [120–125]. Mu *et al.* [126] reviewed the application of CSLM in the observation of AF growth induced by inclusions. The key step in oxide metallurgy is adjusting the elements to form inclusions that induce AF nucleation and to control the microstructure by controlling the cooling rate. The dependence of the inclusion characteristics and temperature parameters on the volume fraction, size, and formation temperature of AF has been widely studied. In 1999, Hanamura *et al.* [31] used CSLM to observe the formation of polygonal and Widmanstätten ferrites on the surfaces of oxides and sulfides. Since then, numerous scholars have conducted experimental studies on the induction of AF nucleation by inclusions using CSLM, as summarized in Table 4 [31,46,127–144].

Table 4. Summarized studies on the induction of AF nucleation by inclusions using CSLM [31,45,127–144]

Author	Year	Inclusions		Heating cycles			Quantitative of ferrite			
		Type	Size / $\mu\text{m}$	Temperature / K	Holding time / s	Cooling rate / $(\text{K}\cdot\text{s}^{-1})$	Fraction / wt%	Formation temperature / $^{\circ}\text{C}$	Grain size / $\mu\text{m}$	Kinetics
Hanamura <i>et al.</i> [31]	1999	Ti–Si–O	No	1673	1	10	No	737	No	Yes
Terasaki [127]	2007	Al–O	<2.5	1673	1	2	No	No	No	No
Zhang <i>et al.</i> [128]	2009	Ti–Al–Mn–O–MnS	No	1673	1–300	5	0–9	650–750	100–400	No
Hu <i>et al.</i> [129]	2011	Ti–Al–Mn–Si–O–MnS	1–2	1673	1–600	5	50–90	550–700	132–251	No
Wen and Song [45]	2012	AlCeO <sub>3</sub> –Ce <sub>2</sub> O <sub>2</sub> S	1–5	1476	600	5	No	557–623	120	No
Wan <i>et al.</i> [130]	2013	Ti–Al–Mn–O–MnS	No	1673	5	5	No	634.4	No	No
Wan <i>et al.</i> [131]	2013	Ti–Mn–O–MnS	No	1673	5–30	5	No	840	150	Yes
Mu <i>et al.</i> [132]	2014	TiO <sub>x</sub> –MnS	>1.2	1823	60	216	10–50	No	245–968	Yes
Jiang <i>et al.</i> [133]	2015	Ti–Al–Mn–Si–O–MnS	No	1673	600	5	No	543–638	No	No
Wan <i>et al.</i> [134]	2015	Zr–Ti–Al–O	No	1573–1673	5–30	5	No	550–620	46–166	Yes
Wu <i>et al.</i> [135]	2015	Al–Ti–Mg–O	0.9–1.1	1523–1673	180–300	5	No	650	22–139	No
Mu <i>et al.</i> [136]	2016	Ti–Al–Mn–Si–O–MnS	No	1473–1673	0–600	0.1–1.2	0–80	600–760	100–900	No
Lin <i>et al.</i> [137]	2017	Mg–Al–O–MnS	1.4–2.4	1573	180	1–20	7–26	559–729	60–100	No
Loder and Michelic [138]	2017	Ti–Mn–Al–O–MnS	>1.2	1673	100	7	72–85	653–670	125–210	No
Yang <i>et al.</i> [139]	2019	Ti–Zr–Al–O–MnS	1.4–1.9	1473	60	0.5–10	26–46	540–691	64.53	Yes
Wang <i>et al.</i> [140]	2020	Ti–Ca–Zr–Al–O–MnS	No	1673	180	2–15	75–82	600–680	170–180	Yes
Wang <i>et al.</i> [141]	2022	TiN–MnS	1.3–2.7	1763	1	0.2–10	13–83	No	206–704	No
Liu <i>et al.</i> [142]	2022	Ca–Al–O–MnS	No	1623	1	12.5	60–77	600–700	100–114	Yes
Yao <i>et al.</i> [143]	2022	Al–Ti–Mn–O–MnS	1.2–1.4	1523–1673	300	3	13.4–78.7	667–767	113–293	Yes
Xie <i>et al.</i> [144]	2023	No	No	1673	3	0.6–10.5	No	532.3–680.7	No	Yes



Fig. 15(a) [128] shows the nucleation and growth of AF in a steel sample during heating at 1400°C for 200 s using CSLM. The AFs grew along the four directions of the inclusion. Fig. 15(b) [128] shows the hard impingement of the AFs. Nucleation initiated at both the austenite grain boundaries and inclusions. Laths grew from clean austenite grain boundaries as Widmanstätten ferrite, which was inhibited by the growth of intragranular AF. Subsequently, the growths of AF impinged on each other to generate a fine interlocking microstructure. Thus, the growth process of AF primarily includes three steps: primary nucleation on the inclusions, sympathetic nucleation on the broad side of the primary ferrite, and impingement between AFs. Fig. 16 shows the summar-

ized average growth rates of AFs for various oxide inclusions. The growth rate of the AFs induced by various inclusions was in the range of 0.03–33.3  $\mu\text{m/s}$ . At a cooling rate of 3–5 K/s, the order of growth rates of AFs induced by different inclusions, as reported in literature, was Ti–O [126] < Ti–Ca–Zr–Al–O [140] < Mg–O [137] < Ti–Zr–Al–O [139] < Ti–Al–O [143] < Mn–Ti–Al–O [143] < Zr–Ti–Al–O [134]. Notably, the ZrO content in the Zr–Ti–Al–O inclusions was higher than the  $\text{TiO}_x$  content and the  $\text{TiO}_x$  content in the Ti–Zr–Al–O inclusions was higher than the ZrO content. The growth rate of the AFs induced by inclusions may also change when the heat-treatment parameters and steel composition are altered.

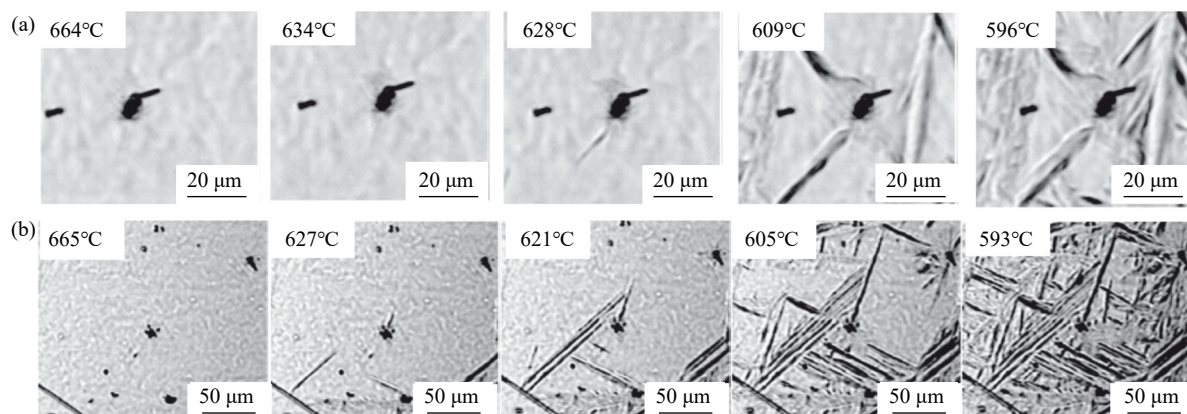


Fig. 15. *In-situ* observation of morphology of AF at inclusions in steel: (a) nucleation and growth of AFs; (b) hard impingements between the AFs [128]. Reprinted from *Acta Mater.*, 58, D. Zhang, H. Terasaki, and Y.I. Komizo, *In situ* observation of the formation of intragranular acicular ferrite at non-metallic inclusions in C–Mn steel, 1369–1378, Copyright 2010, with permission from Elsevier.

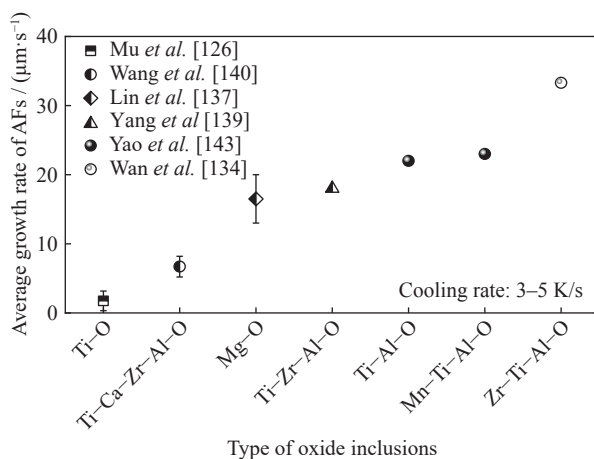


Fig. 16. Summarized average growth rates of AFs for various oxide inclusions.

As CSLM can be used to measure steel phase transition temperature, a method combining CSLM and differential scanning calorimetry (DSC) was proposed to monitor the final continuous cooling transformation (CCT) diagram, as shown in Fig. 17 [126]. Electron backscatter diffraction (EBSD) micrographs showed that the microstructure became very fine at high cooling rates. The CCT diagram can be used to design heating and cooling parameters in oxide metallurgy.

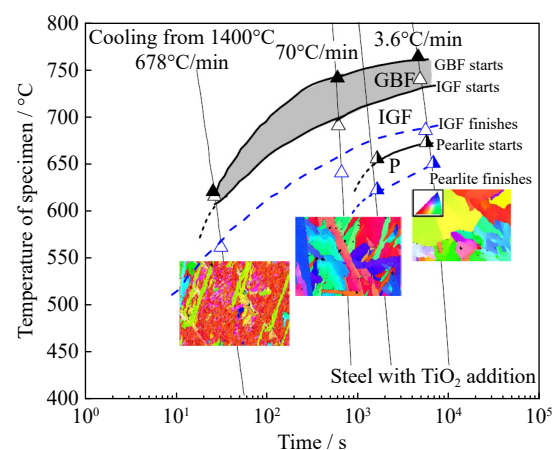


Fig. 17. CCT diagram of the steel with  $\text{TiO}_2$  addition, analyzed by CSLM, DSC, and EBSD [126].

## 7. Opportunities and challenges in *in-situ* observation of inclusions using CSLM

The characteristics of inclusions have a significant influence on steel performance [145–152]. The formation, collision, dissolution, modification, and precipitation of inclusions during steel production have been extensively studied using CSLM. However, the following directions will be the focus of future research in *in-situ* observation of inclusions



using CSLM.

(1) Collision of complex inclusions at steel–slag interface. Previous studies have primarily focused on the *in-situ* observation of the collision of pure inclusions at the steel–Ar interface. However, the collisions of complex inclusions have not been extensively studied. Additionally, *in-situ* observation of collision of inclusions at the steel–slag interface owing to the wettability difference between the steel–Ar and steel–slag interfaces is necessary.

(2) Dissolution of complex inclusions in the slag at the steel–slag interface. The dissolution of pure inclusions in liquid slag has been widely observed *in-situ* in a previous study to optimize slag composition. To simulate the process of inclusion dissolution, *in-situ* observation of the dissolution of inclusions in slag at the steel–slag interface is necessary. Furthermore, complex inclusions are formed in steel during steel production. However, the dissolution of complex inclusions in slags has not been extensively studied.

(3) *In-situ* observation of inclusion modifications. Several methods have been proposed for the *in-situ* observation of the modification of inclusions by dissolved elements in steel. However, further investigations are required on not only Ca modification but also modification by other active elements. Moreover, developing an additional technique during the CSLM experiments to improve the understanding of the modification process of inclusions will be useful.

(4) *In-situ* observation of inclusion transformation in steel during solidification and heating processes. The precipitations of MnS and TiN have been extensively studied. The effects of temperature and steel composition on the precipitation of inclusions have been elucidated. Recently, transformation of inclusions during solidification and heating processes has been reported. However, *in-situ* observations of the transformation of oxide inclusions in steel during solidification and heating processes have rarely been reported.

(5) Quantitative correlation between AF formation and inclusion characteristics. The effects of temperature and grain size on AF formation and growth have been widely studied. The formation and growth of AF were influenced not only by the heating and cooling parameters but also by the size and composition of inclusions. Thus, comprehensive experiments are necessary to investigate the quantitative relationship between AF formation and inclusion characteristics.

## 8. Conclusions

(1) The collisions of various inclusions observed *in-situ* using CSLM are summarized. The solid inclusions exhibited a significant attractive force, with a high tendency to collide. However, pure liquid inclusions rarely collided because of their low attraction. The order of capillary forces for different inclusions was  $\text{Al}_2\text{O}_3 > \text{TiAlO}_x\text{--Al}_2\text{O}_3 > \text{Ce}_2\text{O}_3 > \text{Ce--Al--O} > \text{Al}_2\text{O}_3\text{--SiO}_2 > \text{Ce--O--S} > \text{Al}_2\text{O}_3\text{--CaO} > \text{SiO}_2 > \text{MgO} > \text{MgAl}_2\text{O}_4$ . In future studies, collisions of complex inclusions should be investigated as these inclusions are actually formed in molten steel.

(2) Increasing the  $\text{CaO}/\text{Al}_2\text{O}_3$  and  $\text{CaO}/\text{SiO}_2$  ratios is be-

neficial for the dissolution of inclusions in liquid slag. Solid phase formation in the slag should be prevented to improve inclusion dissolution. The reported models are summarized to predict the dissolution rate of the inclusions in the slag. To simulate the process of inclusion dissolution, *in-situ* observation of the dissolution of inclusions in the slag at the steel–slag interface is necessary.

(3) Several methods have been proposed for *in-situ* observation of inclusion modification by dissolved elements in steel. The morphological and compositional evolution of the inclusions during the reaction process were observed *in-situ* using a combination of CSLM and SEM–EDS. Developing an additional technique during CSLM experiments to further reveal the modification process of the inclusions will be useful.

(4) *In-situ* observation of MnS and TiN precipitation has been widely studied, whereas the *in-situ* observation of the transformation of oxide inclusions in steel during solidification and heating processes has rarely been reported.

(5) CSLM can be used to observe phase transition temperature induced by inclusions on a sample surface. The effects of temperature and inclusion characteristics on AF formation have been widely studied. At a cooling rate of 3–5 K/s, the order of growth rates of AFs induced by different inclusions, as reported in literature, was  $\text{Ti--O} < \text{Ti--Ca--Zr--Al--O} < \text{Mg--O} < \text{Ti--Zr--Al--O} < \text{Mn--Ti--Al--O} < \text{Ti--Al--O} < \text{Zr--Ti--Al--O}$ . Further experiments are necessary to investigate the quantitative relationship between AF formation and inclusion characteristics.

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## Conflict of Interest

Ying Ren and Lifeng Zhang are editorial board members for this journal and were not involved in the editorial review or the decision to publish this article. The authors have no relevant financial or non-financial interests to disclose.

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