Effect of Mg Treatment on Inclusions of X65 Pipeline Steel

Jun Tian, Tianpeng Qu*, Deyong Wang, Huihua Wang and Xinrui E
Shagang School of Iron and Steel, Soochow University, China

Abstract
Three heats of X65 pipeline steels are produced in a factory, two of which were treated with magnesium. The characteristics of inclusions in those steels were analyzed by SEM-EDS and ASPEX. And the influence of magnesium treatment on inclusions of X65 pipeline steel are studied in this paper. The results show that: (1) The oxide inclusions in non-magnesium treated steel are mainly Al$_2$O$_3$ inclusions, while the oxide inclusions are mainly MgO·Al$_2$O$_3$ in the steel with magnesium-treated. (2) The proportion of the small size (1-5 μm) inclusions in magnesium-treated steel is relatively high, and that of the large size (> 15-μm) inclusions is obviously reduced. (3) The larger size inclusions are mainly Al$_2$O$_3$ and the inclusions near the low melting zone. The inclusions close to MgO·Al$_2$O$_3$ spinel composition have smaller size, indicating that the magnesium treatment could make the size of inclusions less and disperse the inclusions. (4) Magnesium-treated steel contained a large number of fine MgO·Al$_2$O$_3$ inclusions, which could provide nucleation core for the precipitation of sulfide, thus reducing the precipitation of sulfide at a grain boundary. And the number of pure MnS decrease after magnesium treatment.

Keywords
Magnesium treatment, Inclusion, Sulfur, Pipeline steel

Introduction
Hydrogen induced cracking (HIC) is one of the main failure modes of pipeline steel. In recent years, oil and gas pipeline have developed to the large diameter and high pressure direction, which require pipeline steel to keep higher strength and toughness [1]. However, the higher the strength of steel, the greater the sensitivity of hydrogen-induced cracks [2]. Inclusions in steel are usually regarded as hydrogen traps. When the pipeline steel undergoes a rotting reaction in an acidic environment, the resulting hydrogen enters the steel matrix, aggregates around inclusions to produce hydrogen pressure, and eventually forms a hydrogen-induced crack at the edges of the inclusions. Therefore, inclusions in steel are one of the main reasons for the production of HIC, especially the strip of oxide inclusions and long strip MnS inclusions, have the biggest influence on anti-HIC property of the steel [3-6]. Therefore, in order to improve the pipeline steel’s anti-HIC performance, it is necessary to strictly control the amount and shape of inclusions in the steel.

At present, a commonly used method for mod-
iffication of inclusions in pipeline steel was calcium treatment [7,8], so that, the MnS inclusions in the steel are modified into spherical CaS inclusions and CaO-Al₂O₃-CaS complex inclusions, and the B type inclusions are modified into low melting point of calcium aluminate inclusions, which are easy to grow up and float. However, to some extent, calcium treatment could improve the adverse effect of MnS, but the low-melting calcium aluminate inclusions would be formed after calcium treatment. And if the low-melting calcium aluminate inclusions don’t eliminate completely in time, they will form a cluster CaO-Al₂O₃ inclusions in rolled metal, which seriously affect the anti-HIC performance of the product [3,9-11]. It is obvious that calcium treatment has some deficiencies in modification of inclusions in pipeline steel.

Magnesium treatment could make the inclusions small and dispersed [12-14], and MnS inclusions also would be modified to complex sulfide by magnesium treatment [15-17]. In view of the advantages of magnesium treatment, this paper carries out the magnesium treatment on X65 pipeline steel, and the effect of Mg on inclusions are analyzed in pipeline steel. The results could provide a reference for the application of magnesium treatment of pipeline steel.

**Experimental**

Three heats of industrial trials are performed in a 200t ladle and recorded as No.1, No.2 and No.3 respectively. The production route is “BOF(basic oxygen furnace) → LF(ladle furnace) → RH → magnesium treatment → soft blowing by Ar gas → CC (continuous casting)”. In particular, magnesium treatment was not applied to heat No. 1 but the other process is identical. The details of the process practices could be described as follows.

During tapping of BOF, synthetic slag and ferroalloys with aluminum are added into the stream of liquid steel. After that, the melt is stirred by argon to make the composition and temperature homogenized. And then the melt is transported to LF refining station, in which high basicity slag is used and aluminum is added to decrease the oxygen and sulfur of the steel. After 40 min, the melt is delivered to RH refining station, at this station the liquid steel is treated at the pressure of 67 Pa for 20 min, and then 120 m and 220 m magnesium wire are inserted into the ladle by a wire-feeding mechanism for Heat 2 and 3 respectively. After magnesium treatment, the molten steel is gently stirred for 10 min to promote the inclusions floating upward. And then the ladle is transferred to the continuous caster.

The steel samples are taken from casting billet. The contents of C and S of the samples are analyzed with infrared analysis. The concentrations of Si, Mn, P, Mg, Nb, V, Ti, Als (dissolved aluminum) and O (dissolved oxygen) are determined using ICP-OES (inductively coupled plasma-atomic emission spectroscopy). The N content is analyzed using thermal conductance. The results are shown in Table 1. All the samples are cross-sectioned and polished, and then morphology and composition of inclusions are analyzed by SEM(Scanning Electron Microscopy), EDS (Energy Dispersive Spectrometer) and ASPEX (Aspex scanning electron microscopes).

**Experiment Results**

**Morphology and composition of the inclusion**

The compositions and morphologies of inclusions in samples are analyzed by SED-EDS. Figure 1 is an SEM photograph of inclusions in the sample of heat No.1. These inclusions are mainly Al₂O₃ inclusions (Figure 1a) and MnS inclusions precipitated with Al₂O₃ spinel, and MnS inclusions contain a small amount of Mg.

### Table 1: Chemical composition of experimental steels.

| Heat | Chemical composition, wt % |
|------|-----------------------------|
|      | C   | Si  | Mn  | P   | S   | Nb  | V   | Ti  | Als | Mg  | T.O | N   |
| No.1 | 0.060 | 0.187 | 1.41 | 0.010 | 0.009 | 0.037 | 0.046 | 0.0207 | 0.0256 | -   | 0.0051 | 0.0029 |
| No.2 | 0.071 | 0.190 | 1.39 | 0.008 | 0.007 | 0.050 | 0.045 | 0.0250 | 0.0470 | 0.0013 | 0.0056 | 0.0041 |
| No.3 | 0.077 | 0.190 | 1.45 | 0.006 | 0.008 | 0.053 | 0.044 | 0.0210 | 0.0380 | 0.0024 | 0.0053 | 0.0015 |

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Figure 1: Inclusion morphologies of heat No.1.

Figure 2: Inclusion morphology of heat No.2.

Figure 3: Inclusion morphology of heat No.3.

Figure 4: Composition distribution of inclusions in the heat No.1.
Quantitative analysis of the types of inclusions in the steel is made by ASPEX. Figure 4, Figure 5 and Figure 6 are the results of quantitative analysis, where S indicates the sulfur content of the inclusions, and MA represents the MgO-Al$_2$O$_3$ spinel (M means MgO, A means Al$_2$O$_3$). It can be seen from Figure 4 that the inclusions of the heat No.1 are mainly calcium aluminates and compound inclusion of Al$_2$O$_3$ and sulfide. And there are a small amount of MgO-CaO-Al$_2$O$_3$ inclusions in this sample. The Figure 5 shows that the type of inclusions are mainly MgO-Al$_2$O$_3$-sulfide compound inclusion and MgO-CaO-Al$_2$O$_3$ inclusions in the heat No.2. And these inclusion composition are close to the MA spinel composition, and most of the sulfides are MgO-Al$_2$O$_3$ inclusions Core nucleation. It can be seen from Figure 6 that most of the inclusions in the heat No.3 are also MgO-CaO-Al$_2$O$_3$ inclusions and MgO-Al$_2$O$_3$-sulfide compound inclusion, furthermore, those inclusion composition are closer to the MA spinel composition than that of the other heat.

Figure 7, Figure 8 and Figure 9 are types of inclusion in the heat No.1, No.2 and No.3, respectively. In the figure, A means Al$_2$O$_3$, M means MgO, and C means CaO. It can be seen from those figures, the inclusions of the heat No.1 are mainly MnS-A (42.52%), MnS (23.36%) and A (14.20%),

**Figure 5:** Composition distribution of inclusions in the heat No.2.

**Figure 6:** Composition distribution of inclusions in the heat No.3.
after magnesium treatment, the inclusions are mainly MnS-MA (26.50%), MnS-A (12.36%), MnS (9.38%), MA (8.52%) and A (8.16%) in the heat No.2, and mainly MnS-MA (42.01%), MA (12.91%) MnS-A (7.60%), MnS (5.47%), and A (5.00%) in the heat No.3. And proportion of the pure MnS inclusion decrease with magnesium content increase.

Table 2: Size distribution of inclusions in the steels.

| Heat  | Scanning area, μm² | The percentage of inclusions with different particle sizes accounted for the total number of inclusions |
|-------|-------------------|---------------------------------------------------------------------------------------------------|
|       |                   | 1-3 μm            | 3-5 μm            | 5-10 μm          | 10-15 μm        | > 15 μm |
| No.1  | 16000000          | 81.66%            | 12.78%            | 4.94%             | 0.42%           | 0.21%   |
| No.2  | 12000000          | 84.54%            | 13.07%            | 2.10%             | 0.19%           | 0.10%   |
| No.3  | 16000000          | 85.70%            | 11.82%            | 2.44%             | 0.05%           | 0.00%   |
Size of inclusions

The size of the inclusions in the samples was quantitatively analyzed by ASPEX, and the results are shown in Table 2. It can be seen from the table, in the steel with magnesium treatment, the number of inclusions with small particle size (1-3 μm) increases. Since more than 90% of the inclusions in the steel have a particle size of less than 5 μm. Statistical classification of the inclusions is carried out, and the results are shown in Figure 10. As can be seen from this figure, after the magnesium treatment, the proportion of inclusions of the particle size of 1-2 μm increases.

Analysis and Discussion

Effect of magnesium treatment on oxide inclusions of steel

When Mg is added to the steel, the following reactions occur [18]

\[ \begin{align*}
Mg + 2O &= MgO \\
Mg + 3O &= Mg_3O_4
\end{align*} \]

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of other substances are calculated using equations (4) and (5), where the interaction coefficients of the elements are shown in Table 3 [5,19,20].

\[ \alpha_i = f_i \cdot w_i \] (4)

\[ \lg f_i = \sum_{j=2}^{n} \rho_{ij} \cdot \%_j \] (5)

The equilibrium phase diagram of MgO-MgO·Al₂O₃-Al₂O₃ at 1873 K can be calculated by the Matlab software, as shown in Figure 11. As can be seen from the figure, after magnesium treatment, the content of [Mg] and [Al] in the steel is in the range of MgO·Al₂O₃, so inclusions of the steel added magnesium are mainly MgO·Al₂O₃, which is consistent with the experimental results. Classify the oxide inclusions in the steel, which is shown in Table 4, and every oxide type include pure oxide and MnS-oxide. In the heat No.1 without Mg, the inclusions in the steel are mainly Al₂O₃ inclusions, the proportion is 93.85%, and that of MgO-Al₂O₃ inclusion are only 2.05%. After magnesium treatment, the proportion of MgO-Al₂O₃ inclusions in the steel are increased to 57.53% and 74.30% respectively, in the heat No.2 and No.3, and that of Al₂O₃ inclusion in these heat are reduced to 33.71% and 17.04%, respectively.

Calculate the average composition of the oxide

Table 3: Interaction coefficients ($\rho_{ij}$) of elements in molten steel at 1873 K.

| j   | Al | Mg | S  |
|-----|----|----|----|
| C   | 0.0910 | -0.240 | 0.110 |
| Si  | 0.0056 | -0.088 | 0.063 |
| Mn  | 0.0035 | 0.024 | -0.026 |
| P   | 0.0330 | - | 0.029 |
| S   | 0.0300 | -1.380 | -0.028 |
| Nb  | - | - | -0.013 |
| V   | - | - | -0.016 |
| Ti  | 0.0040 | -0.510 | -0.072 |
| Al  | 0.0450 | -0.120 | 0.035 |
| Mg  | -1.9800 | -0.047 | -0.026 |
| O   | -6.6000 | -289 | -0.270 |
| N   | -0.053 | - | 0.010 |

Figure 11: The equilibrium phase diagram of MgO-MgO·Al₂O₃-Al₂O₃ at 1873 K.
Table 4: The proportion of oxide inclusions in different types of steel.

| Heat | The type of oxide inclusion |  |
|------|----------------------------|---|
|      | $\text{Al}_2\text{O}_3$  | $\text{MgO-Al}_2\text{O}_3$ | $\text{CaO-MgO-Al}_2\text{O}_3$ | $\text{CaO-Al}_2\text{O}_3$ |
| No.1 | 93.85%                     | 2.05%                          | 0.47%                          | 3.63%                          |
| No.2 | 33.71%                     | 57.53%                         | 1.03%                          | 7.73%                          |
| No.3 | 17.04%                     | 74.30%                         | 8.38%                          | 0.28%                          |

Figure 12: The average composition of oxide inclusions in the range of different particle sizes of the heat No.1.

Figure 13: The average composition of oxide inclusions in the range of different particle sizes of the heat No.2.
inclusions in the different particle size ranges and label them in the phase diagram, as shown in Figure 12, Figure 13 and Figure 14. The area surrounded by the broken line in the figure is the region of the melting point below 1873 K which is calculated by the Factsage 7.2 software. From the figure, the larger particle size of the inclusions are mainly \( \text{Al}_2\text{O}_3 \) inclusions (Figure 12) and the inclusions near the area of low melting point (Figure 13 and Figure 14). The inclusions in these two regions tend to aggregate and grow, and if they do not float to the surface in time, they will stay in the steel. As can be seen from Figure 13, the size of inclusions closed to MgO·Al\(_2\)O\(_3\) spinel is smaller since it is not polymerize in the molten steel and easy to disperse in the steel [12-14]. The average composition of the ox-

**Figure 14:** The average composition of oxide inclusions in the range of different particle sizes of the heat No.3.

**Figure 15:** The changing path of the inclusion composition.
ide inclusions in the three heat steel are labeled in the ternary phase diagram, as shown in Figure 15. From the figure, with the increase of Mg content in the steel, the composition of the oxide inclusions is constantly approaching to the spinel. Therefore, the proportion of the small size inclusions in the heat No.3 is the largest (Figure 10 and Table 2).

**Effect of magnesium treatment on sulfide inclusions in steel**

The formation of sulfides in steel can be expressed by the following formula [5]:

\[
[Mg] + [S] = MgS(s) \quad G^0 = -522080 + 201.02T \quad (6)
\]

\[
[Mn] + [S] = MnS(s) \quad G^0 = -167612 + 88.7T \quad (7)
\]

Eq. (8) is determined by linear combination based on the thermodynamic data and Eq. (1) to Eq. (3) and Eq. (6).

\[
\text{MgO}-\text{Al}_2\text{O}_3(s) + 3\text{MgS(s)} = 4\text{MgO(s)} + 3[S] + 2[Al] \\
G^0 = 2184422 - 1060T \quad (8)
\]

\[
K = \frac{a_{\text{MgO}}^4 a_S^3 a_{\text{Al}}^2}{a_{\text{MgO}} a_{\text{Al}_2\text{O}_3} a_{\text{MgS}}^3} \quad (9)
\]

MgO-Al$_2$O$_3$, MgS and MgO are regarded as pure substances, then their activities are 1. The equilibrium phase diagram of MgO-MgO-Al$_2$O$_3$-MgS at 1873 K can be calculated using the Matlab software. The value of $\alpha_{\text{Al}}$ and $\alpha_S$ can be calculated using Wagner’s equation (Eq. (4)) and Eq. (5) according to the chemical compositions and thermodynamic data, listed in Table 1 and Table 3, respectively. And the calculation results are shown in Figure 16. From the figure, the composition points of the steel treated by magnesium are below the MgO-MgO-Al$_2$O$_3$-MgS equilibrium line. Therefore, there are no single MgO inclusions in the steel, which is consistent with the previous analysis results. There will be MgS or (Mn-Mg) S inclusions in the steel. The high temperature anti-deforming capability of this kind of sulfide is higher than that of pure MnS, which can reduce the deformation degree during the rolling process, so as to avoid the form of long strips in the final product, and improve the anti-HIC performance of the pipeline steel.

The liquidus temperature ($T_{\text{L}}$) and solidus temperature ($T_{\text{Sol}}$) of the steel is calculated by the following formulas (Eq.(10) and Eq.(11)) [21], and the critical precipitation temperature of MnS can be calculated using the Matlab software. Figure 17 is changes in critical precipitation temperature of MnS and solidus temperature ($T_{\text{Sol}}$) of the steel with Mn content and S content. The composition of the steel also are plotted in the figure. As can be seen from the figure, the critical precipitation temperature of MnS in every heat are lower than the solidus temperature, thus, MnS could not be formed in the experimental steel over the solidus temperature, and the MnS must be precipitated during solidification process because of the segregation.

\[
T_{\text{Liq}}(°C) = 1536-83[\%C]-7.8[\%Si]-5[\%Mn]+\%Cu]-
\]
Figure 17: Changes in critical precipitation temperature of MnS and solidus temperature ($T_{sol}$) of the steel with Mn content and S content: A) Heat NO.1; B) Heat NO.2 and C) Heat NO.3.
\[ \text{KMn} = 0.76 \quad \text{KS} = 0.05 \]

The variations of manganese and sulfur concentration with solid fraction in the steels are calculated and showed in Figure 18. The results show that there is MnS precipitation when the solid fraction is approximately 0.95. And the MnS would be precipitated on the oxide during solidification process, which is agree well with the SEM/EDS results (Figure 1, Figure 2 and Figure 3). In the heat No.1, the oxide is mainly Al_2O_3, so the MnS mainly precipitated on it. (Figure 1). In the other heat, the oxide is mainly MgO·Al_2O_3, therefore, the cores of MnS are principally spinel (Figure 2 and Figure 3).

In conclusion, because of the large amount of smaller MgO·Al_2O_3 inclusions in the steel with magnesium, which is possible to provide nucleation cores (Figure 2 and Figure 3) for the precipitation of sulfides, the precipitation of sulfides at grain boundaries will be reduced and they will be dispersed. And the number of pure MnS decrease in the steel with magnesium (Figure 7, Figure 8 and Figure 9). At the same time, the sulfide with core...
can further improve its capability of high temperature anti-deformation, and then further reduce its deformation degree during the rolling process. Accordingly, magnesium addition to the pipeline steel could improve its properties.

Conclusion

Effect of Mg treatment on inclusions of X65 pipeline steel is studied. By comparing and analyzing the composition and size of inclusions in non-magnesium-treated steel and magnesium-treated steel, the following conclusions are obtained:

(1) The oxide inclusions in non-magnesium-treated steel are mainly Al$_2$O$_3$ inclusions (93.85%). The oxide inclusions in magnesium-treated steel are mainly MgO·Al$_2$O$_3$ inclusions. With the increase of magnesium content in the steel (from 0.0013% to 0.0024%), the proportion of MgO·Al$_2$O$_3$ inclusions increase from 57.53% to 74.30%, and the inclusion components are closer to MgO·Al$_2$O$_3$ spinel.

(2) The proportion of the inclusions in the magnesium-treated steel with a small particle size (1-5 μm) is relatively high, besides, the proportion of inclusions in the range of 1-2 μm is higher than that in non-magnesium-treated steel. In the magnesium-treated steel, large particles (>15 μm) inclusions are obviously reduced (the proportion is 0.21%, and that in the two magnesium-treated steels are 0.10% and 0%, respectively).

(3) Larger size inclusions are mainly Al$_2$O$_3$ inclusions and the inclusions near the low melting point. These inclusions are easy to polymerize and grow up, and if they fail to float to the surface of molten steel, they will remain in the steel. The particle size of the inclusions near the MgO·Al$_2$O$_3$ spinel is small, indicating that the MA spinel is difficult to polymerize in the molten steel, and the magnesium treatment can make inclusions small and dispersed in the steel.

(4) There are a large number of fine MgO·Al$_2$O$_3$ inclusions in the magnesium-treated steel, which can provide the nucleation core for the precipitation of sulfide, thus reduce the precipitation of sulfide at a grain boundary and disperse it. And the number of pure MnS decrease after magnesium treatment. At the same time, the magnesium treatment can form (Mn-Mg)S composite sulfide in the steel, further reduce the deformation degree of sulfide during the rolling process, so as to avoid the form of long strips MnS in the final product and influence on the performance of pipeline steel.

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