Effects of Pellet Basicity on the Simulated Deposit Formation in Coal-Fired Rotary Kilns for Iron Ore Pellet Production

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ABSTRACT: Deposit formation in the coal-fired rotary kiln is frequently found in the production of fluxed iron ore pellets by the grate-kiln process and affects normal production. In this paper, the effects of pellet basicity (CaO/SiO$_2$ mass ratio) on the simulated deposit formation were investigated. The results show that the porosity of deposits samples increases from 30.8 to 41.5% as the pellet basicity increases from 0.6 to 1.2, and most of the holes increase with increasing basicity. The primary phase of the deposit samples changed from the M$_2$O$_3$ phase region to the clinopyroxene phase region with a lower melting point. As the basicity increased, the calculated proportions of the liquid phases in the deposit samples had an increasing trend. Moreover, the deposit sample adhesion to the refractory brick increases with the increase in pellet basicity.

1. INTRODUCTION

Iron ore pellets are an essential part of the blast furnace charge, with high grade, good strength, uniform particle size, good metallurgical properties, and so forth. Especially, fluxed iron ore pellets have better high-temperature soft fusion performance and reduction performance. However, the burden of the blast furnace is still dominated by a high proportion of high basicity sinter and some acidic iron ore pellets. The pelleting process’s energy consumption, CO$_2$ emission, and environmental pollution are much lower than the sintering process. In order to optimize the structure of the blast furnace burden and reduce the environmental pollution, some companies have focused on the production of fluxed iron ore pellets to replace the iron ore sinter for blast furnace smelting in recent years.

The three common processes used in iron ore pellet production are the shaft furnace process, the straight grate process, and the grate-kiln process. As the dominant process for iron ore pellet production in China, the proportion of iron ore pellets produced by the grate-kiln process has exceeded 70%. Deposit formation in a rotary kiln is frequently observed in iron ore pellet production by the grate-kiln process. The deposit formation mechanism in rotary kilns for acidic iron ore pellets is more severe than acidic iron ore pellet production. Some studies have found differences in the mechanism of deposit formation in different parts of the rotary kiln. The primary manner to form the deposits near the burner zone in the rotary kiln was the hematite crystallization and diffusion, while the liquid phases played a secondary role in the deposit formation. Furthermore, bentonite and coal ash contain high SiO$_2$, Al$_2$O$_3$, Na$_2$O, and K$_2$O, which will promote the generation of liquid phases and intensify the deposition phenomenon. In addition, insufficient combustion of pulverized coal caused partial zones of the reduction atmosphere in the rotary kiln, resulting in the generation of Fe$_2^+$, and formed low-melting-temperature compounds with veinstone minerals such as SiO$_2$ and CaO-Al$_2$O$_3$. It has been found that the deposition phenomenon became more severe in rotary kilns for producing fluxed iron ore pellets. Unfortunately, the effects of the basicity of iron ore pellets on the deposition formation mechanism are still not fully understood. Thus, it is necessary to study the deposition formation mechanism during the production of fluxed iron ore pellets by the grate-kiln process.

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and coal ash. The generated liquid phases in the deposits are critical for the adhesion and growth of the deposits on the surface of refractory bricks. The higher proportion of liquid phases in the system would accelerate the deposit formation. Thus, this study prepared the simulated deposit samples with different basicity by mixing preheating pellets powder and coal ash. The effects of the basicity of iron ore pellets on the microstructure, phase composition, and elemental distribution behaviors of the deposits are investigated. Moreover, the phase diagram and liquid phase generation of the bonded phase were calculated by FactSage 8.0 software. Finally, the effects of basicity on adhesivity of simulated deposits to refractory were presented. Our study will provide a technical basis for understanding the effects of binary basicity of iron ore pellets on deposition formation and guiding the fluxed iron pellet production in China.

2. RESULTS AND DISCUSSION

2.1. Phase Compositions of the Roasted Deposit Samples. The X-ray diffraction (XRD) analysis of each binary basicity simulated deposition sample is shown in Figure 1. The main material phase of the simulated deposition is hematite, and minor silicate phases are pyroxene and diopside subsilic ferrian. A large amount of SiO$_2$, Al$_2$O$_3$, and other minerals in coal ash mixed into the iron ore pellet powder, which prompted the formation of silicates. In general, the silicates such as pyroxene and diopside have low softening temperatures and melting temperatures and form more liquid phases in the deposits.

2.2. Microstructure of the Roasted Deposit Samples. Figure 2 shows the microstructure of the roasted deposit samples with different binary basicity. The effects of binary basicity on the porosity of roasted deposited samples were analyzed by Image J software. The porosity of deposit samples increases from 30.8 to 41.5% as the pellet basicity increases from 0.6 to 1.2, and most of the holes are irregular in shape. In particular, when the binary basicity reaches 1.2, the roasted deposit samples appear to have significant cracks. At high temperatures, the silicate phase in the deposited sample generates liquid phases to bond the solid phase and shrink in volume. At the same time, the higher binary basicity needs more limestone addition. The limestone starts to decompose (about 759 °C) to produce CO$_2$, leading to the generation of holes in the roasted sample. The burning of residual carbon in deposit samples may also cause holes in the roasted samples.

2.3. Elemental Distributions of the Roasted Deposit Samples. The microstructure and elemental distribution of the roasted deposit samples are also shown in Figures 3–6. It is evident that when the binary basicity is 0.6 and 0.8, hematite particles are mainly bonded in the recrystallized solid phase, and the hematite particles are uneven. When the binary basicity is further increased to 1.0 and 1.2, the deposited samples probably generate higher proportions of liquid phases, which play an essential role in the growth of the deposition.

Figure 3 shows the energy-dispersive spectrometry (EDS) analysis of the deposit sample with a binary basicity of 0.6, and the main elemental composition of the deposit bonding phase is
O, Si, and Ca and a minor amount of Al, Mg, Fe, and K. The bonding phases are mainly silicate phase with high iron content (point B) and silicate phase with low iron content and high silica content (point A).

Figures 4–6 show that the deposits are composed of a hematite phase with a dark gray silicate phase. The chemical compositions of the silicates vary as the binary basicity increases. More Ca ions are introduced into the silicate phase. The Ca content of silicates with low iron (point A) increases from 14.06 to 23.43 wt %, and the Ca content of silicates with high iron (point B) increases from 14.77 to 16.41 wt % as the pellet basicity increases from 0.6 to 1.2.

2.4. Chemical Composition of Bonding Phases and Liquid Phase Generation. It can be inferred that the chemical composition of silicates (bonding phase) in the deposits with different pellet basicity changed with various pellet basicity. In general, the change in the chemical composition of bonding phases would significantly influence the formation of liquid phases and deposit formation. Thus, the elemental compositions of the bonding phases with different binary basicity were

Figure 3. EDS analysis of simulated deposited samples with a binary basicity of 0.6: (A) EDS results of silicate with low iron, (B) EDS results of silicate with high iron, and (C) EDS results of hematite.

Figure 4. EDS analysis of simulated deposited samples with a binary basicity of 0.8: (A) EDS results of silicate A, (B) EDS results of silicate B, and (C) EDS results of hematite.
obtained by collecting the EDS results of silicates in the deposit samples. The composition of the bonding phase of the roasted deposit samples mainly includes SiO$_2$, CaO, Fe$_2$O$_3$, and Al$_2$O$_3$. Based on the results of EDS analysis, the average chemical compositions of the roasted deposit samples with different binary basicity were counted from at least 10 points.

As illustrated in Figure 7, it can be concluded that with the increase in binary basicity of iron ore pellets, the content of CaO and Fe$_2$O$_3$ in the bonding phase of roasted samples tends to increase, while the content of SiO$_2$ and Al$_2$O$_3$ tends to decrease.

The CaO content in the bonding phase reached 25% by adding more limestone, while the SiO$_2$ content was above 40%.

The phase diagram of the SiO$_2$–CaO–Fe$_2$O$_3$–10% Al$_2$O$_3$–5% MgO system was drawn using FactSage 8.0 software to reveal the effects of composition change in bonding phases. As shown in Figure 8, for the change in the bonding phase composition from “A” to “B” for the roasted deposited samples, “A” represents the bonding phase composition of the deposited samples with an iron ore pellet binary basicity of 0.6, and “B” represents a higher binary basicity of 1.2. It can be seen that the bonding phase composition in deposited samples changes from
the M2O3 phase area to the clinopyroxene phase area as the binary basicity increases. Pyroxene is a common silicate rock-forming mineral with a single-stranded structure. The crystal structure is monoclinic or orthoclinic, and monoclinic pyroxene has a low melting temperature. It forms co-crystals with quartz and Fe2O3 with melting temperatures below 1200 °C, promoting the liquid phase generation.

In addition, the average proportions of liquid phases of bonding phases at 1200, 1250, and 1300 °C were calculated and are shown in Figure 9. The amounts of liquid phase generation in

![Figure 7](image_url)

**Figure 7.** Effect of pellet binary basicity on the chemical composition of the bonding phase of deposit samples: (a) CaO content, (b) SiO2 content, (c) Fe2O3 content, and (d) Al2O3 content.

![Figure 8](image_url)

**Figure 8.** Phase diagram of the SiO2−CaO−Fe2O3−Al2O3−MgO−O2 system.

![Figure 9](image_url)

**Figure 9.** Effect of binary basicity on the ratio of the liquid phase of the bonding phase in deposited samples: (a) 1200, (b) 1250, and (c) 1300 °C.
the bonding phases have a rising trend as the binary basicity of iron ore pellets increases. This can be explained by the fact that more CaO was added and reacted with iron oxides to produce calcium ferrate silicates with a lower melting temperature, generating more liquid phases.

The calculation results show that the average liquid phase generation of the bonding phase for samples with the binary basicity of 0.6 and 1.2 is about 45% at 1200 °C, while the binary basicity of 0.8 and 1.0 is above 50%. When the roasting temperature was 1250 °C, the liquid phase generation of the bonding phase of the sample with a binary basicity of 1.2 was already more than 90%. In addition, when the temperature is further increased to 1300 °C, the liquid phase generation capacity reaches a very high level. In producing high basicity pellets by the grate-kiln process, deposit formation is greatly increased to 1300 °C, while the binary basicity of 0.6 and 1.2 is about 45% at 1200 °C, while the binary basicity of 0.8 and 1.0 is above 50%. When the roasting temperature was 1250 °C, the liquid phase generation of the bonding phase of the sample with a binary basicity of 1.2 was already more than 90%. In addition, when the temperature is further increased to 1300 °C, the liquid phase generation capacity reaches a very high level. In producing high basicity pellets by the grate-kiln process, deposit formation is greatly increased to 1300 °C, while the binary basicity of 0.6 and 1.2 is about 45% at 1200 °C, while the binary basicity of 0.8 and 1.0 is above 50%. When the roasting temperature was 1250 °C, the liquid phase generation of the bonding phase of the sample with a binary basicity of 1.2 was already more than 90%. In addition, when the temperature is further increased to 1300 °C, the liquid phase generation capacity reaches a very high level.

In summary, when the binary basicity of iron ore pellets was increased, the Ca and Fe content in bonding phases of the simulated deposits tended to increase, which further reduces its melting temperature, makes the generation of the liquid phase easier, and increases the adhesion of the pellet powder on the refractory brick. Some suggesting measures, such as adding MgO, optimizing the preheating and roasting principles, increasing the preheating pellets’ strength to reduce the pellet powder, may be taken to inhibit the deposit formation in the fluxed pellet production.

3. CONCLUSIONS

This study investigated the effects of pellet basicity on the simulated deposit formation mechanism in the kiln. The chemical composition of the deposited samples is mainly hematite, with minor silicates. The microstructure of the roasted samples becomes more porous with increasing binary basicity. Moreover, the chemical composition of bonding phases of the simulated deposit samples varied with basicity of various pellets. With the increase in binary basicity, more Ca ions and Fe ions are diffused into the silicate phase. The bonded phase composition moved to the clinopyroxene phase region in the phase diagram. The melting temperature of the silicate phase decreased, which promoted the formation of the liquid phase. The amount of liquid phase generation in the bonding phase has a rising trend as the binary basicity of iron ore pellets increases. Adhesion experiments show that the deposited samples with high binary basicity were easier to adhere on refractory bricks in the rotary kiln. Some suggested measures may inhibit the deposit formation in the fluxed pellet production, including adding MgO, optimizing the preheating and roasting principles, and increasing the preheating pellets’ strength to reduce the pellet powder.

4. MATERIALS AND METHODS

4.1. Materials. The raw materials used in this study were provided by an iron ore pellet plant in China and included preheated pellet, deposit sample, iron ore concentrate, limestone, bentonite, coal ash, and refractory bricks, whose chemical compositions are shown in Tables 1 and 2. Table 1 shows that the iron ore concentrate is mainly hematite with a small amount of magnetite; its total iron grade is 64.72%. Due to the addition of bentonite and limestone, the total iron content of the preheated pellet was reduced to 62.10%, and the CaO content was increased to 4.20%. In addition, coal ash contains a large amount of SiO2 and Al2O3 and 1.22% K2O and 0.75% Na2O, and the amount of powder adherence on the refractory bricks increases as the binary basicity increases. The adhesion rate of deposit on the brick surface increases from 3.0 to 5.0%. In addition, the appearance of the refractory bricks after roasting shows that the color of the contact surface of the refractory brick become dark, which indicates that the powder interacts with the refractory brick at high temperatures and partially erodes into the refractory brick.

![Figure 10](https://doi.org/10.1021/acsomega.1c06800)  
Figure 10. Deposition adhesion of different binary basicity powders on refractory bricks.

Table 1. Chemical Compositions of the Raw Materials (wt %)

| raw material          | TFe  | FeO   | SiO2 | CaO  | Al2O3 | MgO  | K2O  | Na2O | S  |
|-----------------------|------|-------|------|------|-------|------|------|------|----|
| preheated pellet      | 62.10| 0.76  | 6.67 | 2.98 | 1.11  | 0.76 | 0.07 | 0.025| 0.025|
| deposit sample        | 59.74| 0.76  | 6.67 | 2.98 | 1.11  | 0.76 | 0.07 | 0.025| 0.025|
| iron ore concentrate  | 64.72| 0.76  | 6.67 | 2.98 | 1.11  | 0.76 | 0.07 | 0.025| 0.025|
| limestone             | 52.02| 0.14  | 2.32 | 51.82| 51.82 | 51.82| 51.82| 51.82| 51.82|
| bentonite             | 3.09 | 0.20  | 60.01| 3.73 | 12.52 | 3.73 | 12.52| 3.73 | 12.52|
| coal ash              | 4.04 | 0.10  | 51.85| 8.57 | 26.09 | 1.12 | 1.12 | 1.12 | 1.12|

Table 2. Chemical Compositions of Refractory Bricks (wt %)

|    | TFe | SiO2 | CaO | Al2O3 | MgO | K2O | Na2O | TiO2 |
|----|-----|------|-----|-------|-----|-----|------|------|
| 1.23| 16.98| 1.75 | 71.26| 0.24  | 0.79 | 0.079| 0.038|      |

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Figure 10. Deposition adhesion of different binary basicity powders on refractory bricks.
respectively. The high alumina refractory bricks were mainly composed of corundum and mullite.

4.2. Experiment Methods. 4.2.1. Preparation of Simulated Deposit Formation. In order to obtain the deposition samples required for the simulation experiments, we need to analyze the composition of the deposition samples. Previous studies have shown that the original material of the deposits is preheated pellet powder and coal ash.27 The proportions of preheated pellet powder and coal ash for forming deposition can be calculated based on the elemental balances of calcium, silicon, and aluminum.28 The calculated results indicated that the deposit sample contained 94.34 wt % preheated pellet powder and 5.66 wt % coal ash. Since the pulverized coal cannot burn completely in practical production, we chose the pulverized coal with a combustion efficiency of 90% for the subsequent study.

In addition, the raw materials, including iron ore concentrate, limestone, and bentonite, were mixed thoroughly, and the green pellets with different binary basicity were prepared using a disc pelletizer. The size of the green pellets was 10–16 mm in diameter. By adding different proportions of limestone, the binary basicity of pellets was adjusted to 0.6, 0.8, 1.0, and 1.2. The green pellets were heated in a tube furnace at 1160 °C for 10 min to produce the preheated pellets. The preheated pellets were rotated in a drum to obtain preheated pellet powder for simulated deposit sample preparation. The simulated deposit samples consist of preheated pellet powder, pure coal ash (PCA), and removed volatile pulverized coal (RVC), as shown in Table 3.

Table 3. Proportions of Materials to Prepare Deposit Samples

| sample no. | preheated pellet powder/g | PCA/g | RVC/g | basicity |
|------------|---------------------------|-------|-------|----------|
| R0.6       | 9.443                     | 0.51  | 0.331 | 0.6      |
| R0.8       | 9.443                     | 0.51  | 0.331 | 0.8      |
| R1.0       | 9.443                     | 0.51  | 0.331 | 1.0      |
| R1.2       | 9.443                     | 0.51  | 0.331 | 1.2      |

The prepared deposit samples were pressed into cylindrical pellets. Then, the samples were roasted in a muffle furnace at 1230 °C for 1 h under air condition. Finally, chemical composition, XRD, and scanning electron microscopy (SEM)—EDS analyses were performed on the roasted samples.

4.2.2. Evaluated Methods of Deposit Adhesion on the Refractory Bricks. The initial stage of the deposit formation in rotary kilns is the adhesion process of the powder to the refractory bricks; the adhesion materials on the refractories provide the basis for the subsequent deposit growth. The experimental evaluation method used in this study was mentioned in our previous paper.28 The simulated deposit samples (3 g) were placed uniformly on the surface of the refractory bricks (circular area of 3 cm in diameter) and roasted in a muffle furnace at 1230 °C for 5 h under an air atmosphere.

As shown in Figure 11, after cooling, the samples were rolled by a roller with a certain mass twice to simulate the mechanical and impact actions from rolling pellets in the rotary kiln. Thus, the bonding degree between the deposits and the refractory brick by comparing proportions of the simulated deposit samples adheres to the refractory brick.

4.3. Analysis Methods. According to Chinese standards, the determination of the chemical composition of the raw materials is performed by chemical analysis methods.25 Phase compositions of deposit samples with different binary basicity were analyzed by XRD (Empyrean, Malvern PANalytical, The Netherlands). The microstructure and phase composition of simulated experimental samples were determined using SEM (TESCAN, MIRA3-LMH, Czech) coupled with an energy-dispersive spectra analyzer (EDS, Oxford X-MAX20). FactSage8.032 software was used to calculate the liquidus temperatures and liquid amounts of the silicate phases as well as the phase diagram. Software uses the “Phase Diagram” and “Equilib” modules, as well as the corresponding “FactPS” and “FToxid” databases.

AUTHOR INFORMATION

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Notes
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Figure 11. Schematic diagram of the evaluated method used in the present study.

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