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ARTICLE



Numerical Simulation of Co-Combustion of Pulverized Coal and Different Proportions of Refused Derived Fuel in TTF Precalciner

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ABSTRACT

Based on the theory of computational fluid dynamics (CFD), pulverized coal combustion alone, and the co-combustion of pulverized coal and refuse-derived fuel (RDF) in a Trinal-sprayed calciner (TTF) precalciner were simulated. The results revealed that when coal was used as a single fuel, the velocity field in the precalciner had good symmetry, and formed three spray effects and multiple recirculation zones. The main combustion zone was distributed in the lower tertiary air and pulverized coal area, and the highest temperature reached up to 1,500 K. According to the simulation results, the predicted decomposing rate of raw meal was 90.12%, which is in good agreement with the actual measured result. In addition, with the increase in RDF content, the average temperature of the furnace, the decomposition rate of the raw meal, and the NO_x concentration all exhibited a downward trend. Under the condition of ensuring the normal operation of the precalciner, blending with 20% RDF is the most reasonable strategy, and the NO_x emissions decreased by approximately 16%.

KEYWORDS

Numerical simulation; TTF precalciner; RDF; co-combustion; NO_x emission

1 Introduction

Cement is an indispensable raw material for national infrastructure construction. At the same time, the resource consumption and pollutant emissions of the cement industry are huge. Studies have shown that coalbased fuel combustion pollution from the cement industry is the third largest source of NO_x emissions, and coal storage has become lesser [1,2]. Therefore, there is an urgent to seek for suitable alternative fuel to meet the need for cementing production. RDF is a solid fuel made from refuse sorting, crushing, compression and drying processes [3]. It owns the characteristics of high calorific value, stable combustion, easy transportation, and low secondary pollution [4]. Norton [5] conducted a co-firing experiment of RDF and coal, and found that because the nitrogen content of RDF is generally lower than that of coal, NO_x emissions during the combustion process are negatively correlated with the amount of RDF incorporated, which is also one of the advantages of RDF as an alternative fuel. At present, there are few researches on RDF as an alternative fuel in the precalciner. Therefore, it is necessary to carry out research on the cocombustion of pulverized coal and RDF in the precalciner.

As the core equipment of the new dry process cement production line, the precalciner is the main place for fuel combustion and raw meal decomposition, and this is also accompanied by a large amount of NO_x



generation [6]. Since the precalciner is under high temperature and high dust concentration during operation, it is difficult to study its internal physical and chemical processes through experiments. Therefore, many scholars at home and abroad have chosen numerical simulation to study the precalciner. Xie et al. [7] conducted a simulation study on the SLC-S precalciner, and optimized the gas-solid flow field in the furnace by changing the horizontal projection angle of the two raw meal inlets. Huang et al. [8] conducted a full-scale modeling of the in-line dual combustion and denitrator process precalciner (DD-PRC), and discovered that the applications of air staging and fuel staging are beneficial to reduce NO emissions without affecting the operation. Iliuta et al. [9] studied the ILC precalciner, and developed a heterogeneous model based on the reaction-diffusion approach for coal combustion and CaCO₃ decomposition. Although there are many researches on the precalciner, few studies have been conducted on the new TTF precalciner. Compared with the older precalciner, the newer TTF precalciner has more spraying times and a recirculation region. Furthermore, the pulverized coal feeding modes were changed, enabling it adapt to more fuel types. These significant differences make the operation of the TTF precalciner different from the older precalciner, and China's cement production basically uses coal as the single fuel. Therefore, it is of great significance to simulate the coupling process of pulverized coal combustion and raw meal decomposition in the TTF precalciner under the condition of using coal as the only fuel through the widely used computational fluid dynamics (CFD) software, Fluent [10]. The simulation results not only provide theoretical guidance for the actual operation of the precalciner, but also provide a basis for further research on the co-firing process of pulverized coal and RDF in this type of precalciner. In addition, RDF and pulverized coal are mixed and combusted in the precalciner at different ratios, and the best RDF blending ratio is determined by analyzing the combustion characteristics, and the decomposition of raw meal and NO_x emissions.

2 Geometric Model and Boundary Conditions

2.1 Geometric Model and Mesh Independence

Fig. 1a shows the actual size of the TTF precalciner structure, which is 45.8 m high, and the maximum diameter is 7.1 m. The precalciner is divided into three cylinders: Two necks and one lower cone. The bottom is the high temperature flue gas inlet, and the top is the precalciner outlet. There are three tertiary air pipes: Two pipes are symmetrically distributed in the lower column, and one pipe is in the middle column. There are two raw meal pipes in the lower column and middle column, respectively, and the four coal injection pipes are distributed 0.7 m above the lower tertiary air inlet. Fig. 1b shows the mesh of the precalciner, in which a high-quality structured hexahedral grid was used for the whole model. In order to ensure the accuracy of the computed results, mesh independence must be verified. Five mesh numbers (662346 cells, 884680 cells, 1013158 cells, 1262747 cells, and 1421282 cells) are simulated, and the influence of the mesh number on the calculation results was compared according to the average temperature of the precalciner outlet. As shown in Tab. 1, when the mesh number was greater than 1013158, the temperature basically did not change. This shows that solution was mesh independence [11]. Therefore, the present study used 1013158 cells for the numerical simulation.

2.2 Boundary Conditions

The boundary conditions used for the present study all come from the thermal calibration of the cement plant. The inlet boundaries of the tertiary air and flue gas inlet were the velocity inlets. The upper tertiary air speed was 25 m/s at a temperature of 1,200 K, and the lower tertiary air speed was 26 m/s at a temperature of 1,200 K. The flue gas velocity was 27 m/s, and the temperature was 1,350 K. The outlet boundary condition was the pressure outlet, and the pressure value was -1250 Pa. The coal was bituminous from Jiangxi Province [12], and the RDF was obtained from an RDF production plant in Yunnan. The proximate analysis and ultimate analysis of pulverized coal and RDF are shown in Tabs. 2 and 3. The inlet boundaries of the pulverized coal and raw meal inlet were the mass flow inlets. The mass flow of each

pulverized coal inlet and raw meal inlet was 1.25 and 16 kg/s, respectively, and the temperature was 331 and 1,040 K, respectively. Except for the inlet and the outlet of the precalciner, the other parts are set as walls, and the standard non-slip wall function was used to deal with the influence of the wall on the fluid flow.



Figure 1: Structure and mesh of the TTF precalciner (a) Structure, (b) Mesh

 Table 1: Comparison of outlet temperatures for the different mesh numbers

Mesh number	662346	884680	1013158	1262747	1421282
Temperature (K)	1,028	1,050	1,064	1,066	1,063

Sample	Proximate analysis (%)				
	M _{ad}	A _{ad}	V _{ad}	FC _{ad}	
Coal	1.87	14.73	28.48	54.92	
RDF	18.24	36.48	39.12	6.16	

Fable 2: P	roximate	analysis	of the	coal	and RDF	
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Note: M means moisture, A means ash, V means volatile matter, FC means fixed carbon and ad means air-dry basis.

3 Mathematical Models and the Numerical Method

3.1 Turbulence Model

Considering the characteristics of the geometric model, calculation time and accuracy, the standard $k - \varepsilon$ model was used to simulate the turbulent flow field in the present study [13]. This model was based on the semi-empirical formula of turbulent kinetic energy k and the turbulent dissipation rate ε .

Sample	Mole fractions (%)				Lower heating value (MJ/kg)	
	C _{daf}	H _{daf}	O _{daf}	N _{daf}	S _{daf}	
Coal	78.01	6.98	12.53	1.72	0.76	24.91
RDF	55.65	7.54	34.92	0.53	1.36	9.42
Notes defensions days of free horizon						

Table 3: Ultimate analysis of the coal and RDF

Note: daf means dry-ash-free basis.

In the standard $k - \varepsilon$ model, the transportation equations for k and ε were, as follows:

$$\frac{\partial(\rho k)}{\partial t} + \frac{\partial(\rho k u_i)}{\partial x_i} = \frac{\partial}{\partial x_j} \left[\left(\mu + \frac{\mu_i}{\sigma_k} \right) \frac{\partial k}{\partial x_j} \right] + G_k + G_b - \rho \varepsilon - Y_M + S_k \tag{1}$$

$$\frac{\partial(\rho\varepsilon)}{\partial t} + \frac{\partial(\rho\varepsilon u_i)}{\partial x_i} = \frac{\partial}{\partial x_j} \left[\left(\mu + \frac{\mu_t}{\sigma_\varepsilon} \right) \frac{\partial\varepsilon}{\partial x_j} \right] + C_{1\varepsilon} \frac{\varepsilon}{k} (G_k + C_{3\varepsilon} G_b) - C_{2\varepsilon} \rho \frac{\varepsilon^2}{k} + S_{\varepsilon}$$
(2)

where: μ_t represents the turbulent viscosity, G_k represents the turbulent kinetic energy caused by the average velocity gradient, G_b represents the turbulent kinetic energy caused by buoyancy, and Y_M represents the contribution of the fluctuating expansion in compressible turbulence to the total dissipation rate. $C_{1\varepsilon}$, $C_{2\varepsilon}$ and $C_{3\varepsilon}$ are constants, which are 1.44, 1.92 and 0.09. Furthermore, σ_k and σ_{ε} are the turbulent Prandtl numbers of k and ε , respectively. Moreover, S_k and S_{ε} are the user-defined source terms.

3.2 Particle Phase Model

The discrete phase model (DPM) is used to solve the trajectory of the particle phase (pulverized coal, RDF and raw meal) in the precalciner [14]. Due to the strong turbulent flow in the furnace, the discrete random walk model and the stochastic tracking model can respectively predict the turbulence in the gas phase, and the particle diffusion caused by the turbulence. The force balance equation of the particle in the x direction in the Cartesian coordinate system is, as follows:

$$\frac{du_p}{dt} = F_D(u - u_p) + \frac{g_x(\rho_p - \rho)}{\rho_p} + F_x$$
(3)

$$F_D = \frac{18\mu}{\rho_p d_p^2} \frac{C_D R_e}{24}$$
(4)

$$R_e = \frac{\rho d_p \left| u_p - u \right|}{\mu} \tag{5}$$

where: $F_D(u - u_p)$ represents the drag force per unit mass of the particle, u and u_p represents the fluid velocity and particle velocity, respectively, and ρ and ρ_p represents the fluid density and particle density, respectively. Furthermore, C_D is the drag coefficient, d_p is the particle diameter, R_e is the relative Reynolds number, and μ is fluid dynamic viscosity.

3.3 Pulverized Coal and the RDF Combustion Model

In comparing the data of pulverized coal and RDF in Tabs. 1 and 2, RDF can be regarded as a kind of pulverized coal with higher volatile and ash content [15]. The combustion of pulverized coal is a complex process of physical and chemical changes, which can be divided into two steps [16,17]. The first step is the release and combustion of volatiles, and the second step is the combustion of residual coke. The volatiles

release process adopts the two competing rates model proposed by Kobayashi et al. [18], which has the characteristics of less calculation amount and high accuracy. The expression for the volatiles release rate is, as follows:

$$\frac{m_{\nu}(t)}{m_{p,0} - m_a} = \int_0^t (a_1 R_1 + a_2 R_2) \exp\left(-\int_0^t (R_1 + R_2) dt\right) dt$$
(6)

where: $m_v(t)$ represents the mass of volatile that has been released at time t, $m_{p,0}$ represents the initial particle mass, m_a represents the ash content in the particles, and a_1 and a_2 represents the low temperature release rate factor and high temperature release rate factor of volatiles, respectively. The reaction rate constants R_1 and R_2 were defined using the Arrhenius formula.

After the volatiles were released and burned out, the residual coke would begin to undergo a surface combustion reaction, and the kinetic/diffusion control rate model will be selected. The model assumes that the combustion rate of coke is simultaneously controlled by the kinetic rate K and diffusion rate D_0 :

$$\frac{dm_p}{dt} = -A_P P_{OX} \frac{D_0 K}{D_0 + K} \tag{7}$$

$$D_0 = C_1 \frac{\left[(T_P + T_\infty)/2 \right]^{0.75}}{dp} \tag{8}$$

$$K = C_2 e^{-(E_2/RT_P)} \tag{9}$$

where: dp is the particle size, P_{OX} is the partial pressure of the gas around the particle, and A_P is the particle surface area. Furthermore, T_P and T_{∞} represent the initial particle temperature and gas temperature around the particle, respectively.

3.4 Raw Meal Decomposition Model

In the research on the decomposition reaction of raw meal in the precalciner, domestic scholars all took the decomposition of CaCO₃ as the main research object [19,20]. Therefore, in order to simplify the mathematical model, the species transport model was used to simulate the decomposition reaction of the raw meal, which was approximated as CaCO₃. In this model, the mass fraction of each species Y_i was estimated by solving the convection-diffusion equation of the i^{th} species. The conservation equation takes the following general form:

$$\frac{\partial}{\partial t}(\rho Y_i) + \nabla(\rho \vec{v} Y_i) = -\nabla \cdot \vec{J}_i + R_i + S_i$$
(10)

where: \vec{J}_i is the diffusion flux of species *i*, R_i is the net production rate of species *i* by chemical reaction, and S_i is the rate of creation by addition from the discrete phase plus any user-defined sources. The reaction process for the CaCO₃ decomposition is, as follows [21]:

$$CaCO_3(s) \xrightarrow{+178 \text{ kj/mol}} CaO(s) + CO_2(g)$$
 (11)

where: the pre-exponential factor of the reaction is $1.221 \times 10^9 s^{-1}$, and the activation energy is 187.3 kj/mol. The decomposition rate for the raw meal was defined, as follows:

$$\eta = \frac{C_1 - C_2}{C_1} \times 100\% \tag{12}$$

In this equation, C_1 is the mass flow rate of CaCO₃ at the inlet, and C_2 is the mass flow rate of CaCO₃ at the outlet.

3.5 The NOx Formation Model

According to the formation mechanism of NO_x , NO_x is mainly divided into prompt NO_x , thermal NO_x and fuel NO_x . Under the combustion conditions in the precalciner, the proportion of prompt NO_x and thermal NO_x is less than 5%, which is negligible. The present study only considered the formation of fuel NO_x , and the reaction path is shown in Fig. 2 [22]. Assuming that Fuel-N is distributed between volatiles and fixed carbon, volatile N is first converted to HCN, and then part of the HCN is oxidized to NO. The unoxidized HCN, acting as reduction agents, reacts with NO to form N_2 . The char N is directly converted to NO, and some of the NO is reduced to N_2 due to the reducibility of char.



Figure 2: Fuel NO_x formation path

According to the above-mentioned NO reaction mechanism, the generation rate for HCN and NO were respectively used as the source terms to simulate the formation of NO in the precalciner.

$$S_{HCN} = S_{volatile \to HCN} + S_{HCN \to NO} + S_{HCN \to N_2}$$
⁽¹³⁾

$$S_{NO} = S_{charN \to NO} + S_{HCN \to NO} + S_{NO} + S_{NO$$

3.6 Computational Conditions

The focus of the present research was the TTF precalciner in Yunnan Province, China. The research consisted of two main parts. In the first part, the investigators predicted and analyzed the velocity field, temperature field, composition field, and distribution of NO_x in the furnace under separate combustions of pulverized coal, in order to reveal the transport of the gas–solid two-phase flow in the new TTF precalciner. In the second part, under the premise of ensuring that the total heat entering the furnace remains unchanged, the present study simulated five blending situations. Among these, the heat provided by RDF accounted for 10%, 20%, 30%, 40% and 50% of the total heat, respectively. The final ideal blending ratio was used to minimize the NO_x concentration at the outlet when the combustion characteristics and decomposition rate of the raw meal were less affected, that is, the precalciner operates normally.

3.7 Numerical Scheme

The prerequisite for solving the control equations is to discretize these. In the present study, the controlling-volume method was used to discretize the control equations for the fluid phase, and a second-order upwind difference scheme is used to obtain the difference equations. The semi-implicit method for pressure-linked equations (SIMPLE) algorithm was used to couple the pressure and velocity of the discrete equations. The convergence criterion was that the energy term is less than 10^{-6} , and the remaining residuals were less than 10^{-3} . During the data processing, the burnout rate of fuel was defined as the ratio of the mass loss of unburned coke at the outlet with input fuel mass.

4 Result and Discussion

4.1 Coal Mono-Combustion

4.1.1 Distribution of the Velocity Field

The velocity vector diagrams of the X = 0 and Y = 0 sections in the precalciner are shown in Figs. 3a and 3b, respectively. It can be observed that the velocity field in the furnace has good symmetry, which is caused

by the symmetrical furnace structure. Observed from bottom to top, and after the high-speed flue gas from the rotary kiln enters the precalciner, the speed gradually decreases due to the increase in the diameter of the furnace. After the flue gas meets the lower tertiary air that radially enters the furnace, these merges into a vertical upward high-speed air flow at the center, and the first spray effect is formed. Surrounding this high-speed air flow are two low-speed recirculation zones. Subsequently, the air flow continued to rise through the two necks, and the speed sharply increased, forming the second and third spray effects. In the middle cylinder, three recirculation zones were formed due to the addition of upper raw meal and upper tertiary air. The spray effect induces the pulverized coal and raw meal to be suspended in the precalciner, and fully exchange heat with the air flow. The existence of the recirculation zone increases the residence time of the particles in the furnace, and makes the pulverized coal and raw meal mix more uniform, which is beneficial for improving the decomposition rate of the raw meal and burnout rate of the pulverized coal.



Figure 3: Velocity vector in the precalciner (a) X = 0, (b) Y = 0

4.1.2 Distribution of the Temperature Field

The temperature contours of the X = 0 and Y = 0 sections in the precalciner are shown in Figs. 4a and 4b, respectively. It can be observed that the main combustion zone is distributed in the lower tertiary air and pulverized coal area, and the highest temperature can reach up to 1,500 K. This is mainly because the area is rich in high-temperature oxygen, and carries a large amount of pulverized coal, which allows the volatiles to be rapidly released from the pulverized coal, and burn violently. Then, the lower raw meal moves upward under the action of the high-speed flue gas, and absorbs heat to decompose, causing the temperature to gradually decrease. In the middle cylinder, due to the introduction of the upper tertiary air, the unburned pulverized coal particles continue to burn, resulting in a slight increase in temperature. Finally, the flue gas rises to the upper cylinder, since the reaction of the raw meal decomposition takes place during this process, and the temperature shows a downward trend.



Figure 4: Temperature contours in the precalciner (K) (a) X = 0, (b) Y = 0

4.1.3 Distribution of the Composition Field

Fig. 5 shows the mass fraction distribution of each component in the precalciner. In Figs. 5a, 5b and 5e, it can be observed that the mass fraction of O_2 was the highest near the lower tertiary air. With the combustion of pulverized coal, O₂ is gradually consumed, the mass fraction of CO₂, as a combustion product, gradually increases, and the main combustion zone shown in Fig. 4 is formed. In the middle cylinder, due to the injection of air from the upper tertiary air inlet, the mass fraction of O_2 slightly increases, while CO_2 becomes diluted. Then, as the height of the furnace increases, the unburned coke and O2 would continue to react to form CO_2 , the concentration of O_2 would continue to decrease, and the concentration of CO_2 would continue to increase. The air required for combustion can be divided into two stages into the furnace, resulting in insufficient O2 in the lower cylinder, and a large amount of CO is produced here due to the incomplete combustion of pulverized coal. In Figs. 5b–5d, it can be observed that the mass fraction of CaCO₃ was the largest at the inlet of the raw meal. When the height of the furnace was increased, the endothermic decomposition of CaCO3 generated CaO and CO2. When the mass fraction of CaCO3 gradually decreased, and the mass fractions of CaO and CO2 exhibited an overall upward trend, except for the decrease in the middle cylinder, due to the dilution effect of the tertiary air. According to the simulation results, the decomposition rate of the raw meal was 90.12%, which well coincides with the actual measurement results [23].



Figure 5: Mass fraction of each component in the precalciner (a) O₂, (b) CO₂, (c) CaCO₃, (d) CaO, (e) CO

4.1.4 Distribution of NO_x

According to the online measurement, the NO_x concentration at the flue gas inlet was approximately 800 ppm. Fig. 6 shows the average NO_x concentration in the height direction of the precalciner (Z-axis). It can be observed that when z < 5 m, the thermal NO_x from the rotary kiln enters the furnace, and the NO_x concentration slightly changes. In the z = 5-10 m region, due to the dilution effect of the lower tertiary air and the large amount of CO generated (Fig. 5e), NO_x is reduced to N₂, and the concentration of NO_x rapidly drops to 349 ppm. When 10 m < z < 17 m, the fuel N is oxidized to generate NO_x, and NO_x concentration rises to 548 ppm. In the z = 17-24 m region, the concentration of NO_x rapidly decreases, which is attributed to the dilution effect of the upper tertiary air. When z > 24 m, the NO_x concentration continued to increase. However, as the pulverized coal combustion rate became slower, the growth trend gradually decreased, and the NO_x concentration at the outlet was 585 ppm.

4.1.5 Model Validation

The comparison between the predicted value and measured value at the outlet of the precalciner is shown in Tab. 4. The measured value was actually measured by China Building Materials Industry Association under the normal operation of the precalciner. It can be observed that all errors were within the engineering allowable range. This indicates that the simulation is reliable.



Figure 6: Average NO_x concentration along the height of the precalciner

	Temperature	CO ₂ mass fraction	O ₂ mass fraction	NO _X
Predicted value	1064 K	24.9%	1.65%	585 ppm
Measured value	1112 K	25.5%	1.61%	560 ppm
Error	4.3%	2.4%	2.48%	4.4%
Error	4.5%	2.4%	2.48%	4.4%

 Table 4: Comparisons between the simulation and measurements

4.2 Mixed Combustion of Pulverized Coal and RDF

4.2.1 Influence on Combustion Characteristics

Fig. 7 shows the average temperature in the height direction of the precalciner (Z-axis) under different RDF blending ratios. It can be observed that the average temperature of each working condition presents the same trend: The temperature first increases, subsequently decreases, and increases and decreases again. In addition, with the increase in RDF content, the average temperature of each cross-section sequentially decreases. The data in Tab. 1 shows that the moisture content of RDF was approximately 10 times of that of coal powder. The RDF would consume more heat due to moisture vaporization during combustion, which may be the main reason for the temperature drop [15]. Furthermore, although the RDF blending ratios of the six working conditions are different, the temperature for each section of the lower cone slightly changes, because the fuel gas enters the furnace, and the remaining sections of the precalciner exhibits a large temperature difference. When z = 15 m, there is a maximum temperature difference of approximately 50 K, which is mainly attributed to the change in the cross-section that corresponds to the temperature peak of each working condition. It can be observed from the partial enlarged view that when the RDF blending ratio increased from 0% to 50%, the peak positions in the furnace were 15 m, 15 m, 15 m, 14 m, 13 m and 12 m, respectively. This means that the peak position would be greatly affected when the ratio reaches 30%.

The burnout rate reflects the degree of mixed fuel combustion in the precalciner. As shown in Fig. 8, with the increase in RDF blending ratio, the burnout rate gradually decreases. This is probably attributed to the high ash content in the RDF. During the combustion process, the ash may be adsorbed on the fuel surface, and hinder the diffusion of O_2 to the surface, thereby inhibiting combustion. When the blending ratio reaches 30%, the burnout rate becomes less than 90% [8]. Combined with the peak position in Fig. 7, it was found that this blending ratio greatly changed the temperature distribution in the furnace.



Figure 7: Average temperature along the height of the precalciner

4.2.2 Influence on the Decomposition of the Raw Meal

Figs. 9a and 9b shows the average mass fraction of CaCO₃ and CaO in the height direction of the precalciner under different RDF blending ratios. It can be observed that as the blending ratio increases, the lower the mass fraction of CaO, the higher the mass fraction of CaCO₃. This indicates that the decomposition efficiency of CaCO₃ decreases. As shown in Fig. 7, the increase in blending ratio would reduce the average temperature in the precalciner, which may affect the decomposition of CaCO₃. In addition, as shown in Fig. 9c, the mass fraction of CO_2 at the outlet is reduced. Although CO_2 is also produced by the combustion of mixed fuel, the output was smaller than that of the CaCO₃ decomposition, indicating that undecomposed CaCO₃ in the furnace increases and the content of CaO decreases. This is consistent with the results shown in Figs. 9a and 9b. Some scholars have inferred through curve integration that the CO_2 produced by RDF combustion is greater than that of pulverized coal, when the total heat provided by the combustion is constant [24]. As the RDF content increases, the CO_2 produced by the combustion of mixed fuel increases. Since the concentration of CO_2 inhibits the decomposition of CaCO₃, the decomposition efficiency of CaCO₃ decreases, resulting in the decrease in CO₂ content at the outlet. Fig. 10d shows the decomposition rate for the raw meal. When the RDF blending ratio was not more than 20%, the decomposition rate slowly decreased. When the blending ratio was more than 20%, the decomposition rate significantly decreased, and was reached below 85%. In general production, the decomposition rate of the raw meal in the precalciner needs to be greater than 85% [25]. Therefore, it is not recommended for the RDF blending ratio to be greater than 20%.



Figure 8: Burnout rate of mixed fuel under different RDF blending ratios



Figure 9: Comparison of different RDF blending ratios: (a) Mass fraction of $CaCO_3$; (b) Mass fraction of CaO; (c) Mass fraction of CO₂ at the outlet; (d) Decomposition rate of the raw meal



Figure 10: The NO_x concentration contours under different RDF blending ratios (ppm) (a) 0%RDF, (b) 10% RDF, (c) 20%RDF, (d) 30%RDF, (e) 40%RDF, (f) 50%RDF

4.2.3 Influence on NO_x Emission

Fig. 10 shows the NO_x concentration contours under different RDF blending ratios. It can be observed that when the RDF blending ratio increased from 0% to 50%, the NO_x concentration almost had no change, except for each section of the lower cone, and the sections from the lower cylinder to the outlet of the precalciner gradually decreased. This is because the N content of the RDF (0.53%) was significantly lower than that of coal (1.72%), and that the NO_x in the furnace was mainly derived from the fuel NO_x produced by the fuel combustion. Furthermore, the precipitation and combustion process of volatiles was very fast, resulting in the rapid decrease in O₂ concentration around it. In addition, as illustrated in Tab. 1, the volatiles in the RDF were relatively high. Hence, as the RDF content increases, the low oxygen concentration zone in the lower cylinder becomes larger, the high concentration zone of CO for reducing NO_x becomes larger, and the area of the low NO_x concentration zone in the lower cylinder becomes larger different RDF blending ratios. It can be observed that the NO_x concentration decreases with the increase in blending ratio, and when the blending ratio was not less than 30%, the concentration change was small.

The analysis indicates that based on the constant total heat that enters the furnace, when the RDF blending ratio was not less than 30%, the pollutant emissions can be significantly reduced. However, this has a greater impact on the combustion characteristics and decomposition of the raw meal. Therefore, under the condition of ensuring the normal operation of the precalciner, blending with 20% RDF is feasible, and the NO_x emission is reduced by approximately 16%, when compared with the use of coal as the only fuel.



Figure 11: The NO_x concentration at the outlet under different RDF blending ratios

5 Conclusion

In the present study, the numerical simulation of an actual operating TTF precalciner was carried out, and the simulated results were compared with the measured data, in order to prove the reliability of the numerical method. The results indicated that the main combustion zone of the precalciner is distributed in the lower tertiary air and pulverized coal area, and that the maximum temperature can reach up to 1,500 K. The mass fraction of CaCO₃ alternately increases and decreases, and the mass fraction of CaO continues to increase. The NO_x concentration initially decreases, subsequently increases, and decreases and increases again. Furthermore, the present study also analyzed the mixed combustion of pulverized coal and RDF. With the increase in RDF content, the average temperature of the furnace, the decomposition rate of the raw meal, and the NO_x concentration exhibits a downward trend. Considering all these factors, it is feasible and reasonable to mix pulverized coal and RDF at an 8:2 ratio. Furthermore, compared with coal as a single fuel, the NO_x emissions at the outlet were reduced by approximately 16%.

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Conflicts of Interest: The authors declare that they have no conflicts of interest to report regarding the present study.

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