Physico-chemical structure evolution characteristics of coal char during gasification in the presence of iron-based waste catalyst

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Abstract The present study aims to explore the physico-chemical structure evolution characteristic during Yangchangwan bituminous coal (YCW) gasification in the presence of iron-based waste catalyst (IWC). The catalytic gasification reactivity of YCW was measured by thermogravimetric analyzer. Scanning electron microscope–energy dispersive system, nitrogen adsorption analyzer and laser Raman spectroscopy were employed to analyze the char physico-chemical properties. The results show that the optimal IWC loading ratio was 5 wt% at 1000 °C. The distribution of IWC on char was uneven and Fe catalyst concentrated on the surface of some chars. The specific surface area of YCW gasified semi-char decreased significantly with the increase of gasification time. i.e., the specific surface area reduced from 382 m²/g (0 min) to 192 m²/g (3 min), meanwhile, the number of micropores and mesopores decreased sharply at the late gasification stage. The carbon microcrystalline structure of YCW gasified semi-char was gradually destroyed with the increase of gasification time, and the microcrystalline structure with small size was gradually generated, resulting in the decreasing order degree of carbon microcrystalline structure. IWC can catalyze YCW gasification which could provide theoretical guidance for industrial solid waste recycling.

Keywords Char structure · Characteristic · Iron-based waste catalyst · Catalytic gasification

1 Introduction

The catalyst is the core of F-T synthesis technology. The iron-based waste catalyst (IWC) was discharged in the form of slag and wax. A large amount of IWC is treated by stacking, which not only results in a waste of resources, but also pollutes the ecological environment seriously. Coal gasification is considered as the cleanest form of coal conversion (Zheng et al. 2019). If the recovered IWC could be applied to coal gasification, the gasification reactivity will be improved and the problem of industrial solid waste pollution for enterprises will be solved.

As we all know, the evolution characteristics of coal char structure will have a profound impact on catalytic gasification reactivity (Tay et al. 2014; Huo et al. 2014; Li et al. 2016; Wei et al. 2019; Liu et al. 2020). There are many reports on the characteristics of coal char structure during gasification in the presence of alkali metals and alkaline earth metals (Li et al. 2006; Liu et al. 2017a; Gao et al. 2017; Fan et al. 2019; Bai et al. 2019), but there are a few studies on the characteristics of coal char structure using iron-based catalysts, and most of the reports in the literature are carried out by using commercially available analytical reagents. Yu et al. (2006) selected FeCl₃ as catalyst and used SEM to study the changes of Fe in the process of gasification. Zhang et al. (2015), Yu et al. (2019), Zhang et al. (2017) and Li et al. (2017) used SEM–
EDS to characterize the distribution of catalyst on coal char surface during gasification. Xu et al. (2019) studied the characteristic of coal char structure and gasification reactivity under Na₂CO₃, FeCO₃ and their composite catalysts and found that the coal char adsorption curve increased rapidly with 4 wt% FeCO₃. Qi et al. (2014) researched the effect of Fe(NO₃)₃ on structure characteristic and gasification reactivity of coal char and concluded that Fe increased the number of amorphous carbon structure. To sum up, at present, the effect of the coal char structure characteristics of loading iron-based catalyst on the gasification reactivity was reported based on analytical reagents, however, the systematic study on the physico-chemical characteristics of coal char with loaded IWC is seldom reported.

In this work, Yangchangwan bituminous coal (YCW) was selected as feedstock for gasification in the presence of IWC. The catalytic gasification reactivity of YCW with loaded IWC was studied by TGA. And the physical and chemical structure properties of coal char with loaded IWC were systematically characterized by SEM–EDS, nitrogen adsorption analyzer and laser Raman spectroscopy, respectively. The aim is to deepen the understanding of the structure evolution mechanism of IWC as loading catalyst on the YCW char structure during gasification.

2 Experimental

2.1 Samples preparation

YCW was used as the raw material in this study. After crushing and sieving, the raw coal with particle size of 80–120 μm was selected to perform gasification. The proximate and ultimate analysis of raw coal was measured by Infra-Red Rapid Coal Analyzer (5E-MACi) and elemental analyzer (Vario MACRO), respectively. The result is shown in Table 1. The ash composition of raw coal was analyzed by X-ray fluorescence spectrometer (Advant XPX). The result is shown in Table 2.

The IWC comes from CHN Energy Ningxia Coal Industry Co., Ltd. The alumina crucible loaded IWC was put in the muffle furnace, and then the temperature was heated up to 600 °C, remaining for an hour to remove organic matter. The chemical analysis of IWC was carried out by X-ray fluorescence spectrometer (Advant XPX). The result is shown in Table 3 and the main chemical composition of IWC was Fe₂O₃.

2.2 Char preparation

The preparation of semi-char at different gasification time was carried out in a horizontal tube furnace. The schematic diagram of experimental device is shown in Fig. 1. Prior to the experiment, the dried IWC and YCW were ground by mortar and pestle with a ratio of 1:19, then about 2 g samples were evenly spread in an alumina crucible. At the beginning of gasification experiment, the flow rate of 100 mL/min N₂ was introduced to the reactor so as to remove the air. After the temperature was heated up to 1000 °C at a heating rate of 20 °C/min with 100 mL/min N₂, N₂ was cut off then, and the gasification agent CO₂ was injected at the rate of 100 mL/min into the reactor and held for different time (0, 1, 1.5 and 3 min, respectively). At the same time, the isothermal gasification was studied and semi-chars with different gasification time was produced.

2.3 Gasification reactivity test

The test of gasification reactivity was performed by TGA (NETZSCH STA449 F3, Germany). About 10 mg samples were placed in alumina crucible, and heated to the target

| Table 1 Proximate and ultimate analyses of YCW |
|-----------------------------------------------|
| Analysis item | Content (d, wt%) |
| VM            | 26.64           |
| FC            | 56.81           |
| Ash           | 16.55           |
| C             | 64.42           |
| H             | 3.63            |
| N             | 0.67            |
| S             | 1.14            |
| O*            | 13.59           |

*Note: VM volatile matter; FC fixed carbon; d dry basis

*Calculated by difference
gasification temperature at a heating rate of 20 °C/min under 50 mL/min air flow. Then, the atmosphere was switched to the gasification agent CO₂ with a flow rate of 120 mL/min for isothermal gasification in order to eliminate the effect of external diffusion on gasification (Zhang et al. 2019).

The carbon conversion \(X\) and reactivity index \(R_{0.9}\) (Gil et al. 2015) of coal char were calculated by the Eqs. (1) and (2), respectively:

\[
X = \frac{w_0 - w_t}{w_0 - w_\infty}
\]

\[
R_{0.9} = \frac{t_{0.9}}{t_0}
\]

In Eq. (1), \(w_0\), \(w_t\) and \(w_\infty\) represent the initial mass of the sample, the mass of the sample at the gasification time \(t\) and the final mass of the sample at full gasification, respectively. In Eq. (2), \(t_{0.9}\) refers to the gasification reaction time required when the conversion level of coal char reaches 90%.

### 2.4 Char structure characterization

SEM (ZEISS EVO18, Germany) and EDS (BRUKER, Germany) were applied to analyze the surface morphology of coal char. At first, the sample holder covered with a little char was placed in an ion sputtering apparatus to gold-plate. And then the processed samples were put into sample cell for measurement. The acceleration voltage was 15 kV.

The physical structure of coal char was characterized by JW-BK100C-01 nitrogen adsorption analyzer (Jingweigao, Beijing). The experiment was conducted at −196 °C. Before each test, all gasified semi-chars were degassed at 300 °C for 14 h. BET and BJH models were used to calculate the specific surface area and the pore size distribution of coal char.

The chemical structure of coal char was analyzed by DXR laser Raman spectrometer (Thermo Fisher, US). The laser power was 2 mW. The wavelength was 532 nm and the wave number was 800–2000 cm⁻¹. The Origin 9.0 software was used to fit the original Raman spectrogram.

### 3 Results and discussion

#### 3.1 Coal char gasification reactivity

Figure 2 shows the influence of different IWC loading ratios on YCW gasification reactivity. As seen in Fig. 2, at 900 °C, the carbon conversion level of YCW coal char increased gradually with the increase of IWC loading ratio. When the IWC loading ratio increased from 0 wt% to 5 wt%, the carbon conversion level increased significantly.

**Table 3 Chemical analysis of IWC**

| Component | Content (wt%) |
|-----------|---------------|
| SiO₂      | 32.35         |
| Al₂O₃     | 4.64          |
| Fe₂O₃     | 54.34         |
| CaO       | 0.59          |
| Na₂O      | 0.36          |
| K₂O       | 2.39          |
| MgO       | 0.87          |
| SO₃       | 0.32          |
| TiO₂      | 0.06          |
| P₂O₅      | 0.03          |
| Other     | 4.05          |

**Fig. 1 Schematic diagram of tube furnace**

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however, when the ratio raised from 5 wt% to 30 wt%, the carbon conversion showed a low increase degree, not as obvious as previous.

In order to verify the effect of main metal oxide component (Fe₂O₃) of IWC on YCW catalytic gasification, commercially available analytical reagent Fe₂O₃ catalyst was selected as the reference sample. The impact of IWC and analytical reagent Fe₂O₃ catalyst on YCW gasification reactivity were analyzed by TGA. Figure 3 shows the comparative analysis results of the influence of two catalysts on YCW gasification reactivity at 1000 °C. Obviously, the difference between carbon conversion curve of 5 wt% IWC and Fe₂O₃ loaded on YCW is very small and the difference of \( R_{0.9} \) is 0.01. This indicates that the gasification reactivity of YCW with IWC and Fe₂O₃ loaded has a little difference. And as can be seen from the chemical composition analysis of IWC in Table 3, the metal oxide of catalyzing YCW gasification is Fe₂O₃ in IWC.

In order to further investigate the influence of IWC loading ratio on gasification reactivity at high temperatures, the effect of different loading ratios on YCW gasification reactivity was investigated at 1000 °C. The result is shown in Fig. 4. There is little difference between the carbon conversion curves of YCW with 5 wt% and 10 wt% IWC loaded. The difference of \( R_{0.9} \) value between them was 0.02. This indicates that when IWC loading ratio reached 10 wt%, the impact of loading ratio on YCW catalytic gasification reaction was slight. Thus, the optimal loading ratio of IWC was 5 wt%.

### 3.2 Char physical structure evolution

The surface morphology and Fe catalyst distribution of semi-char from YCW gasification with 5 wt% IWC loaded at 1000 °C is shown in Fig. 5. Apparently, the surface of YCW char with different gasification time is rough and the distribution of Fe catalyst on coal char particle is not uniform, mainly concentrated on some coal char particles. In Fig. 5b, there are many small pits on the coal char particle surface, and the distribution of Fe catalyst on coal char particle is more uniform than that of 0 min. In Fig. 5c, the agglomeration of Fe catalyst on coal char particle is observed. It can be seen from Fig. 5d that Fe catalyst has entered into the interior pore.

![Fig. 2 YCW gasification reactivity under different IWC loading ratios](image1)

![Fig. 3 IWC and analytical reagent Fe₂O₃ catalyst on YCW gasification reactivity at 1000 °C. (a YCW + 5 wt% IWC; b YCW + 5 wt% analytical reagent Fe₂O₃)](image2)

![Fig. 4 Different IWC loading ratio on YCW gasification reactivity at 1000 °C](image3)
Fig. 5 Surface morphology and Fe catalyst distribution of semi-char from YCW gasification with 5 wt% IWC loaded at 1000 °C
Table 4  Specific surface area of different gasified semi-chars

| Sample                                      | $S_{BET}$ (m$^2$/g) |
|---------------------------------------------|----------------------|
| (YCW + 5 wt% IWC)-1000 °C-0 min             | 382                  |
| (YCW + 5 wt% IWC)-1000 °C-1 min             | 324                  |
| (YCW + 5 wt% IWC)-1000 °C-1.5 min           | 292                  |
| (YCW + 5 wt% IWC)-1000 °C-3 min             | 192                  |

According to the analysis of nitrogen adsorption/desorption isotherm in Fig. 6a, YCW gasified semi-char loaded with 5 wt% IWC has a continuous pore structure. The specific pore size distribution result is shown in Fig. 6b. It can be seen that the pore structure of different YCW gasified semi-chars is mainly composed of micropore less than 2 nm and mesopore between 2–10 nm. The number of micropores and mesopores are essentially unchanged at 1 min and 1.5 min, however, their number decreases sharply at 3 min. This is also consistent with the specific surface area analysis results in Table 4. For such a phenomenon, Li et al. (2019) found that the reduction of specific surface area was caused by the melting and sintering of pores in the gasification process.

3.3 Char chemical structure evolution

The effect of IWC on char chemical structure was analyzed by Raman spectroscopy. Figure 7 shows the original Raman spectra of YCW gasified semi-char with 5 wt% IWC loaded at 1000 °C. The results show that each original Raman spectrum has a D band at about 1350 cm$^{-1}$ and a G band approximately at 1580 cm$^{-1}$. The D band represents the carbon structure with a defective structure in coal char, and it mainly represents the C–C bonds between the larger aromatic ring (not less than 6 fused rings) and the fused ring in the coal char. The G band is considered as graphite peak.

External factors have a great influence on the peak intensity and other information of the original spectrum, and because the D and G band overlaps, using only the D band and G band of the original Raman spectrum leads to the loss or neglect of the characteristic information of highly disordered carbonaceous materials (Li et al. 2006). Hence, the quantitative correlation between Raman spectra and structural parameters cannot be realized. It is necessary...
to further fit the original Raman spectra of coal char and obtain the hidden information of skeleton carbon structure in the overlapping region. In order to get more information about the structural change of coal char in the process of gasification reaction, it is essential to carry out fractional peak fitting on the original spectrum (Sheng 2007; Smith et al. 2016). The Origin 9.0 software is used to perform peak fitting on the original Raman spectrogram in this study. According to the chemical structure characteristics of coal char (Mosqueda et al. 2019; Wei et al. 2018), the original Raman spectrum was fitted into five bands, namely Lorenzian bands (G, D1, D2 and D4) and Gaussian band (D3). Figure 8 shows the peak fitting diagram of Raman spectra of semi-char from YCW gasification with 5 wt% IWC loaded at 1000 °C. The D1 band is about 1350 cm$^{-1}$ and usually refers to as the defect peak, mainly the plane defect of the carbon structure. The peak position of D2 is around 1620 cm$^{-1}$, which usually appears as the acromion of G band, mainly as disordered graphite lattice. The D3 band is located near 1500 cm$^{-1}$ and represents sp$^2$ hybrid amorphous carbon structure, including small organic molecules and fragmented functional groups. The attribution of the D4 band is still controversial, and its peak is usually around 1200 cm$^{-1}$, but it is generally to be considered as the sp$^2$–sp$^3$ hybrid bond consisting of C–C and C=C stretching vibrations.

The integral area ratio of peak is often used as a parameter to characterize the chemical structure of coal char. $I_{D1}/I_G$ is often used to quantitatively analyze the order degree of carbon structure. Namely, the higher the $I_{D1}/I_G$ ratio, the lower the order degree of carbon structure. Figure 9 shows the relationship between $I_{D1}/I_G$ and catalytic reactivity of different semi-chars from YCW gasification with 5 wt% IWC loaded at 1000 °C. It can be seen that the integral area ratio of $I_{D1}/I_G$ increases with the increase of gasification time. This indicates that the carbon microcrystalline structure is gradually destroyed and the microcrystalline structure of small size was gradually generated. Therefore, the order degree of carbon structure decreased at high carbon conversion level. At the same time, as seen in Fig. 9, the $I_{D1}/I_G$ value was consistent with catalytic activity trend of semi-char with different conversion levels. This reveals that the catalytic reactivity of IWC on semi-char at different conversion levels is a positive relationship to the order degree of carbon structure of semi-char.

4 Conclusions

The desirable loading ratio of IWC at 1000 °C was 5 wt%. The distribution of Fe catalyst on coal char was uneven and Fe catalyst was concentrated on the surface of some chars. The specific surface area of YCW gasified semi-char decreased significantly with the increase of gasification time. i.e., the specific surface area reduced from 381.981 m$^2$/g (0 min) to 191.724 m$^2$/g (3 min), simultaneously, the number of micropores and mesopores decreased sharply at the late gasification stage. The carbon microcrystalline structure of YCW gasified semi-char was gradually destroyed with the increase of gasification time, and the microcrystalline structure of small size was gradually generated, resulting in the decreasing order degree of carbon microcrystalline structure. IWC can be adopted to catalyze YCW gasification which could provide theoretical guidance for industrial solid waste recycling.

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