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Effects of CO₂ atmosphere on low-rank coal pyrolysis based on ReaxFF molecular dynamics

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Pyrolysis of low-rank coal in CO₂ atmosphere can reduce carbon emissions while comprehensively utilizing coal resources. Based on ReaxFF molecular dynamics (ReaxFF-MD), the pyrolysis processes of low-rank coal in inert and CO₂ atmosphere are simulated. By comparing the evolution of pyrolysis products, the influences of CO₂ on the pyrolysis characteristic and product distribution are analyzed. It is found that CO₂ slightly inhibits the conversion of char to tar in the early stage of pyrolysis. In the later stage, CO₂ significantly promotes the decomposition of char and increases the yield of tar and pyrolysis gas. According to the different bond breaking behaviors of coal molecules, the pyrolysis process can be divided into pyrolysis activation stage, initial pyrolysis stage, accelerated pyrolysis stage and secondary pyrolysis stage. The reforming reaction of CO₂ with alkanes generates free hydrogen radicals, which promotes the cleavage of ether bond, C_{ar}–C_{ar} bridge bond and aliphatic C–C bond. Compared with in inert atmosphere, final yield of light tar in CO₂ atmosphere increases from 17.98% to 20.68%. In general, the CO₂ atmosphere helps to improve the tar yield and tar quality of low-rank coal pyrolysis.

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1 Introduction

Coal is the most abundant fossil fuel in China, and the proved reserves of low-rank coal account for more than half of the total.^{1,2} Low-rank coal has the characteristics of low heat value, high moisture and high volatility. The direct combustion of low-rank coal will lead to low energy efficiency, high pollutant emissions and waste of high value components in volatiles.³ Pyrolysis is an effective coal conversion technology and the initial reaction step of most coal utilization processes. The poly-generation technology based on coal pyrolysis can produce pyrolysis gas with high heat value, upgraded coal tar, clean solid fuel coke and other high-value products to realize multi-level and efficient utilization of coal resources.⁴ In recent years, high CO₂ emissions have caused many environmental problems. CO₂ capture, utilization, and storage (CCUS) technology is the best option to reduce global warming while addressing the energy crisis.⁵ Replacing the inert gas used in coal pyrolysis with CO₂ is a promising way to utilize CO₂. Various coal conversion processes have been involved in the gasification and pyrolysis of coal in CO₂ atmosphere, such as CO₂-enhanced gasification,⁶ CO₂ reforming of methane and coal pyrolysis (CRMP)^{7,8} and IGCC process with CO₂ recycles.^{9–11}

The main products of coal pyrolysis are char, tar and pyrolysis gas. The tar can be used for producing high value chemicals, but its quality is not satisfactory because of abundant

heavy components.¹² Heavy tar with high viscosity and causticity are harmful to subsequent apparatus and operations.¹³ As much light tar as possible is expected to be obtained during coal pyrolysis. The use of CO₂ as pyrolysis atmosphere may affect the product distribution and tar composition. Therefore, it is of great practical value to study the effects of CO₂ atmosphere on low-rank coal pyrolysis.

Many scholars have studied the characteristics of coal pyrolysis in CO₂ atmosphere. For example, Lee *et al.*¹⁴ carried out pyrolysis experiments of peat in a quartz tube reactor in N₂ and CO₂ atmospheres, and found that CO₂ increased the decomposition rate of peat in the range of 630–728 °C. However, CO₂ reacted with volatile organic compounds, which reduced the yield of tar. Wang *et al.*¹⁵ conducted pyrolysis experiments of Shendong coal in different atmospheres (N₂, H₂, CH₄ and CO₂) with a vertical fixed-bed reactor. The results showed that CO₂ atmosphere could promote the increase of tar yield compared with N₂, H₂ and CH₄. Jamil *et al.*¹⁶ carried out slow (1 °C s^{−1}) and fast (1000 °C s^{−1}) pyrolysis experiments of Victoria lignite in He and CO₂ atmosphere with a wire mesh reactor and concluded that CO₂ had no significant effect on tar yield and tar composition. The above studies indicate that the effect of CO₂ atmosphere on the distribution of coal pyrolysis products is still controversial, especially for tar. Moreover, there are few researches focus on the influence mechanism of CO₂.

The pyrolysis process is accompanied by the generation, cross-linking and stabilization of free radicals. The reaction of free radicals plays a key role in the distribution of pyrolysis products. Due to the high activity and short existence time of free radicals, it is difficult to track the dynamic evolution tendencies of

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functional groups and free radicals even with the most advanced experimental techniques.¹⁷ With the rapid development of modern computing power, ReaxFF molecular dynamics (ReaxFF-MD) simulation has become an effective calculation method to study the coal pyrolysis from a microscopic perspective.

As a principal method to study chemical reactions, Quantum Chemistry (QC) can accurately locate the transition states and chemical reaction pathways.¹⁸ However, the high computational cost of QC limits its application in macromolecular models.¹⁹ Molecular Dynamics (MD) based on classical principles incurs much lower computational costs but it is not suitable for exploring the development of chemical bonds.²⁰ The ReaxFF is an empirical force field proposed by van Duin and Goddard *et al.*, which can describe the breaking and formation of chemical bonds in complex reaction systems.^{21,22} ReaxFF-MD combines MD with ReaxFF to obtain the motion trajectory of the system by calculating the state of atoms with time step. It allows simulating the chemical reaction of large and complicated molecular systems without setting the reaction pathways in advance, which makes it applicable for coal pyrolysis simulation.²³

ReaxFF-MD method has been used by many scholars to study coal pyrolysis process.^{24,25} In these researches, the simulation results of ReaxFF-MD are qualitatively consistent with the experimental data, indicating that this method can accurately describe the reaction process of coal pyrolysis. Nevertheless, the effects of CO₂ atmosphere on low-rank coal pyrolysis are rarely studied with ReaxFF-MD method. It is necessary to study the pyrolysis reaction from the microscopic perspective in order to explore its mechanism. This study has guiding significance in engineering for the clean utilization of low-rank coal and CO₂ emission reduction.

2 Experimental and computational methods

2.1 Coal sample preparation and pyrolysis

In this study, a low-rank coal from Naomaohu in Xinjiang, China (NMH coal for short) was selected as the pyrolysis sample. The proximate and ultimate analyses of NMH coal are listed in Table 1. The volatile content of NMH coal accounts for 36.97%, which is suitable for coal pyrolysis and tar production. Before the experiment, the samples were dried in a vacuum oven at 105 °C for 6 hours to constant weight. After drying, the samples were ground and sieved to ensure that the particle size is less than 180 μm.

TG-MS is a combination of Netzsch STA 449C thermogravimetric analyzer and Netzsch QMS 403 mass spectrometer. The samples with mass of (10 ± 0.01) mg were placed in an Al₂O₃ crucible and pyrolyzed in argon and CO₂ atmosphere, respectively. The furnace was heated from 50 °C to 700 °C at heating rate of 10 K min⁻¹ with purging gas flow of 50 mL min⁻¹ and protective gas flow of 30 mL min⁻¹.

2.2 Simulation details

The chemical structure of coal consists of various covalent and non-covalent interactions, which is the basis of understanding

Table 1 Proximate and ultimate analyses of the samples

Sample	NMH	Mo Zheng
Proximate analysis (wt%)		
Moisture, air dry	14.59	13.12
Ash, dry	5.39	5.10
Volatile, dry	36.97	38.42
Fixed carbon, dry	57.64	56.48
Ultimate analysis (dry, wt%)		
Carbon	74.74	72.88
Hydrogen	5.58	4.40
Nitrogen	0.85	0.95
Sulphur	0.32	0.26
Oxygen ^a	18.51	21.51

^a By difference.

the coal pyrolysis process.²⁶ Based on proximate analysis, ultimate analysis and ¹³C-NMR, Mo Zheng²⁷ constructed a molecular model of NMH coal and used it for ReaxFF-MD simulation of pyrolysis tar product. The simulation data was in great agreement with the experimental data. Both the coal samples of this paper and Mo Zheng were produced in Naomaohu, and the proximate and ultimate analysis are highly similar, as shown in Table 1. Therefore, we consider the coal to be the same as Mo Zheng's and directly adopted his coal molecular model C₁₉₄H₂₁₅O₄₃. Fig. 1(a) presents the planar structure of C₁₉₄H₂₁₅O₄₃. After geometry optimization of the planar structure, three-dimensional structure of coal molecules used in simulation was obtained, as shown in Fig. 1(b).

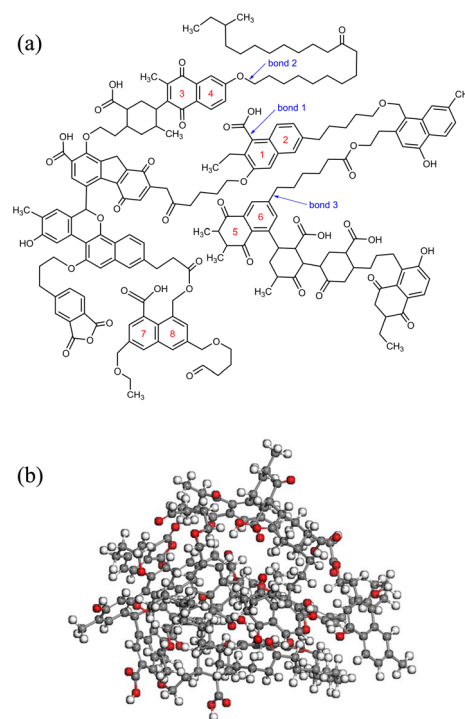


Fig. 1 Molecular models of NMH coal: (a) planar structure, (b) three-dimensional structure.



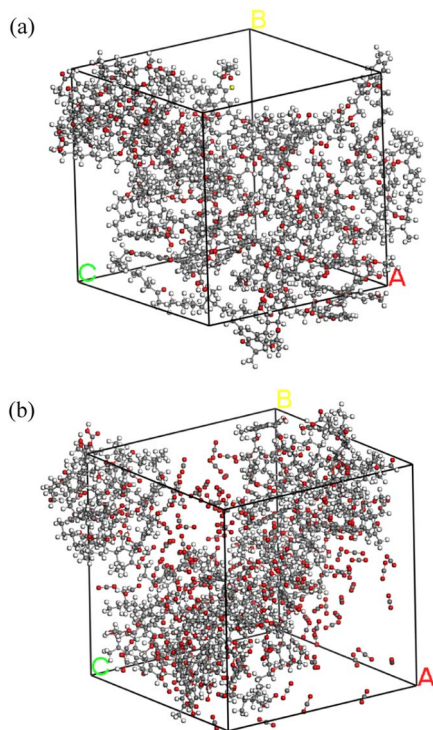


Fig. 2 Reaction systems of NMH coal pyrolysis: (a) inert atmosphere, (b) CO_2 atmosphere.

Subsequently, 5 NMH coal molecules were assembled into a periodic cube to obtain the inert atmosphere reaction system. Similarly, the CO_2 atmosphere reaction system was assembled using 5 NMH coal molecules and 150 CO_2 molecules. The NPT ensemble was used for dynamic equilibrium at room temperature and one atmospheric pressure to optimize the model as much as possible. The final pyrolysis reaction systems in inert and CO_2 atmosphere are shown in Fig. 2.

Pyrolysis was completed by heating the system from 300 K to 2600 K at heating rate of 10 K ps^{-1} . In all simulations, the time step was 0.1 fs. Reaction temperature was controlled by Berendsen thermostat with a 100 fs damping constant. All the simulations were carried out using the ReaxFF code in LAMMPS platform.

The product molecules in the simulation are classified according to the number of carbon atoms.²⁸ C_{40+} fragments are considered as char that cannot be evaporated at high temperatures. $\text{C}_5\text{--C}_{13}$ and $\text{C}_{14}\text{--C}_{40}$ fragments are respectively thought to be light tar and heavy tar, which are liquid in normal temperature. Pyrolysis gas consists of inorganic gases and $\text{C}_1\text{--C}_4$ organic fragments.

3 Results and discussion

3.1 Experiment verification

TG-MS enables real-time monitoring of char and pyrolysis gas yield. The residual solid weight percentage is the yield of char. Ion current density of pyrolysis gas components is used to monitor the yield of pyrolysis gases. The tar is commonly

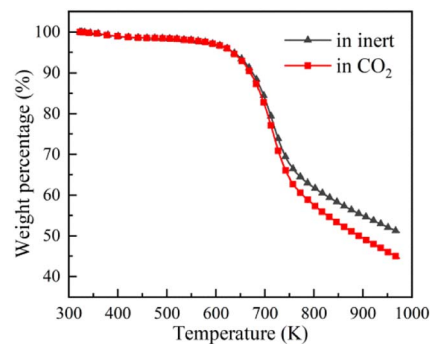


Fig. 3 Weight percentage of NMH coal in pyrolysis experiment.

analysed for final yield and composition after pyrolysis. There are still great difficulties in real-time monitoring of tar components, so it is essential to carry out research through simulation.

Firstly, the accuracy of ReaxFF-MD simulation is verified by the evolution trend of char and pyrolysis gas. The weight loss curves of NMH coal pyrolysis obtained by experiments is shown in Fig. 3, while the evolution of char weight percentage in simulation is presented in Fig. 4(a). It can be discovered that the char yield in two atmospheres is not much different in the early stage of pyrolysis. And the char yield in CO_2 is significantly lower than that in inert at the later stage. Although the effect of CO_2 in the early stage is slightly overestimated, the simulation results of char yield are basically consistent with the experimental data.

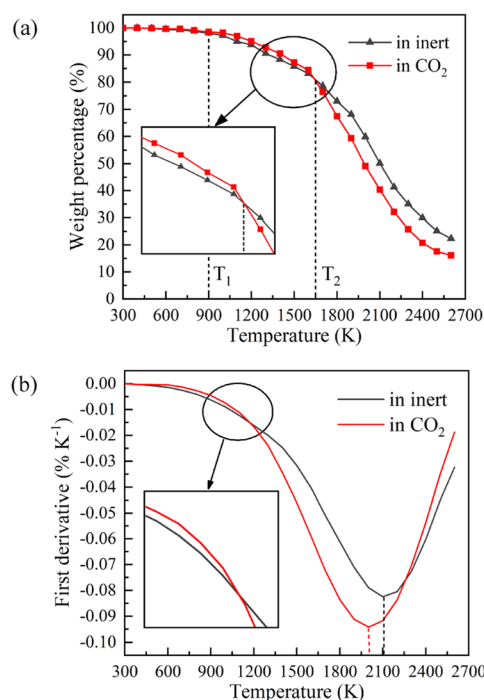


Fig. 4 Evolution trend of char in simulation: (a) weight percentage, (b) first derivative.



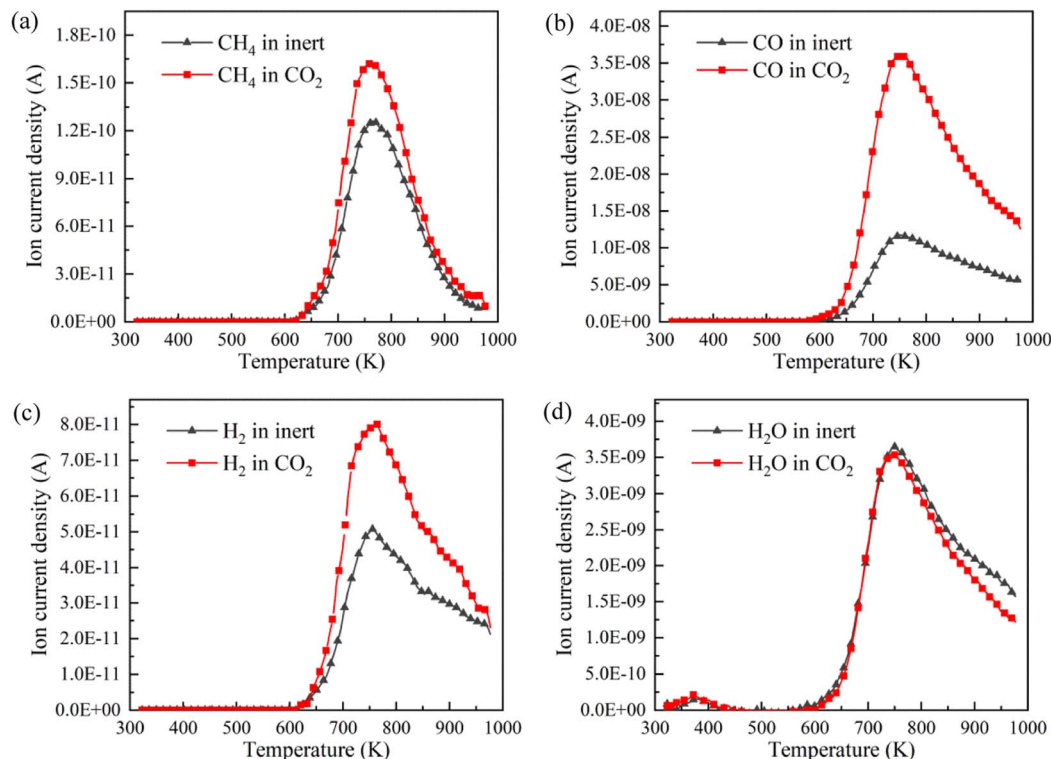


Fig. 5 Ion current densities of pyrolysis gas components: (a) CH_4 , (b) CO , (c) H_2 , (d) H_2O .

CH_4 , CO , H_2 and H_2O are the representative components of pyrolysis gas. The ion current intensities of CH_4 , CO , H_2 and H_2O detected by TG-MS are shown in Fig. 5, implying the

generation rate of different pyrolysis gas components. Moreover, the mathematical integral of ion current intensity is presented in Fig. 6, which is proportional to the yield of pyrolysis

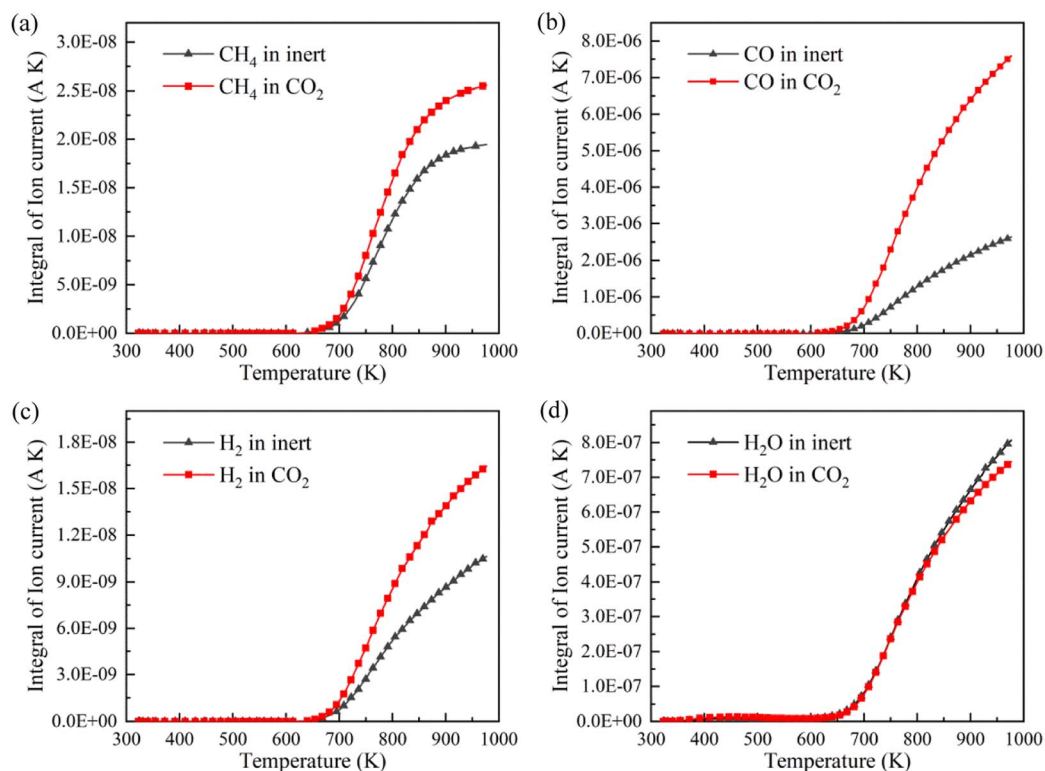


Fig. 6 Mathematical integral of ion current densities: (a) CH_4 , (b) CO , (c) H_2 , (d) H_2O .



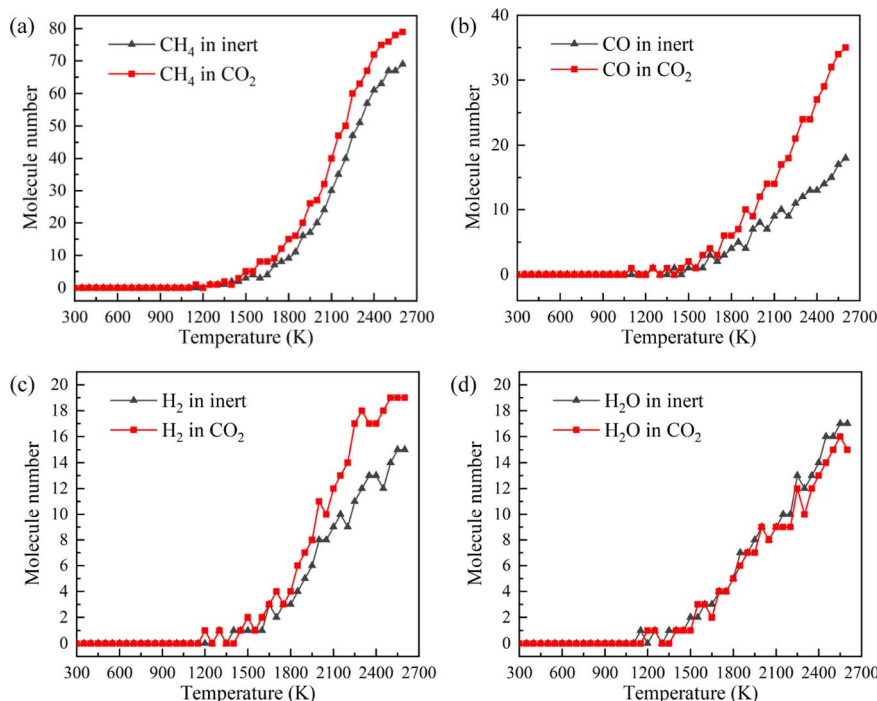


Fig. 7 Molecular number of pyrolysis gases in simulation: (a) CH₄, (b) CO, (c) H₂, (d) H₂O.

gases. The variation of molecule numbers in simulation with temperature is displayed in Fig. 7. Fig. 6 and 7 reflect the experimental and simulation results of the evolution trend of pyrolysis gases yield, respectively. Their consistency indicates that the molecular model and calculation method are competent for the simulation of NMH coal pyrolysis.

It should be noted that the simulation temperature is much higher than the experimental temperature. This is because current molecular simulation computing ability is on nano-second and nanometer scale, while the experimental observation capability is on second and millimeter scale. Generally, the simulation temperature is raised to ensure that reactions completed within an acceptable calculation time. Although there are differences on time and temperature scales between simulation and experiment, previous studies have illustrated that simulation results are qualitatively consistent with actual product evolution trend and reaction mechanism.^{29,30}

3.2 Evolution of three-phase pyrolysis products

Fig. 4 displays the char weight loss curves obtained from ReaxFF-MD simulation and its first derivative. The char yield decreases with the increase of temperature in both atmospheres. Before 900 K (T_1), the weight of char is basically constant, and coal pyrolysis is in an unactivated state. At about 1650 K (T_2), the char yields in both groups are 80.8%. Between T_1 and T_2 , the char yield in CO₂ atmosphere is marginally higher than that in inert atmosphere, indicating that CO₂ may have a certain inhibitory effect on char cracking in the early stage of pyrolysis. After T_2 , the weight loss rate increases evidently, and then decreases after reaching the peak. During this period, the char yield in CO₂ atmosphere is always much lower than that in

inert atmosphere. It implies that CO₂ prominently promotes the decomposition of char in the later stage of reaction, making coal pyrolysis more complete. From Fig. 4(b), it can also be found that the maximum weight loss peak in CO₂ atmosphere

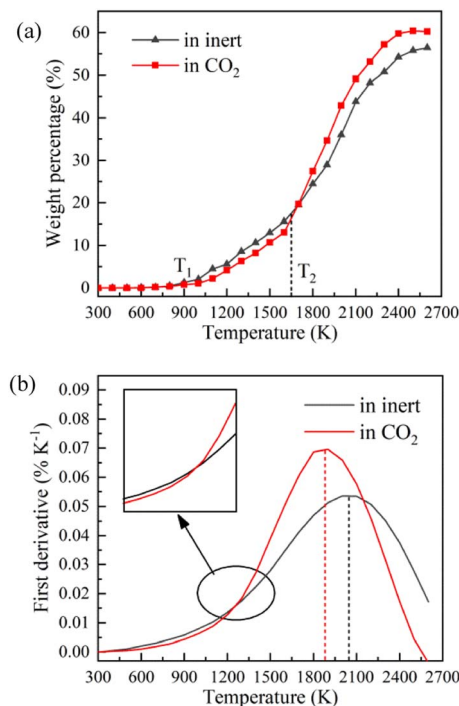


Fig. 8 Evolution trend of tar in simulation: (a) weight percentage, (b) first derivative.

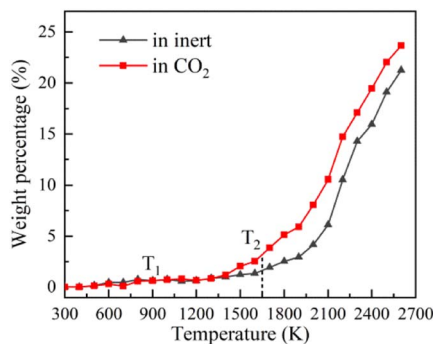


Fig. 9 Evolution trend of pyrolysis gas in simulation.

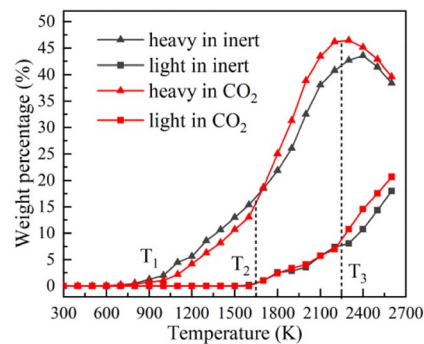


Fig. 10 Evolution trends of light and heavy tar in simulation.

arrives earlier and the absolute value is larger. Char in CO₂ atmosphere is easier to pyrolyze and has higher reaction rate.

Evolution trend of tar in simulation are presented in Fig. 8. Before T_1 , there is almost no tar generation. Between T_1 and T_2 , the tar yield in CO₂ atmosphere is slightly lower than that in inert atmosphere. After T_2 , the tar yield in CO₂ atmosphere is significantly higher. The evolution trend of tar yield is in great agreement with the weight loss of char. As depicted in Fig. 8(b), the tar generation rate reaches a maximum of $0.069\% \text{ K}^{-1}$ at about 1900 K in CO₂ and $0.053\% \text{ K}^{-1}$ at about 2050 K in inert atmosphere. Compared with in inert atmosphere, the maximum generation rate of tar in CO₂ atmosphere is about 30% higher. CO₂ observably promotes the formation of tar during coal pyrolysis.

Fig. 9 is the evolution trend of pyrolysis gases in simulation. In the early stage of pyrolysis, only a small amount of pyrolysis gas is produced, and the pyrolysis gas yields in two atmospheres are essentially the same. Pyrolysis gas is generated in large quantities with the increase of temperature, and the yield of pyrolysis gas is higher in CO₂ atmosphere.

Comparing the evolution of three-phase pyrolysis products in inert and CO₂ atmosphere, it can be concluded that CO₂ slightly inhibits the pyrolysis of NMH coal in the early stage of reaction. In the later stage of pyrolysis, CO₂ significantly promotes the decomposition of char and the formation of tar and pyrolysis gas. The pyrolysis in CO₂ atmosphere is more complete and the final yield of tar increases from 56.39% to 60.28%.

3.3 Evolution mechanism of light and heavy tar

In order to explore the effect of CO₂ on the quality of pyrolysis tar, the evolving profiles of light and heavy tar is studied, as shown in Fig. 10. By monitoring the dynamic process of NMH coal pyrolysis, the chemical bond breaking and formation behaviours of coal molecules are studied in order to thoroughly investigate the influence mechanism of CO₂.

Before T_1 , the coal molecules in system are quite stable. Carboxyl (–COOH) is the most unstable oxygen-containing functional group in coal molecule. The chemical bonds between carboxyl groups and the main body of coal molecule begin to break, such as bond 1 in Fig. 11(a). The detached carboxyl groups decompose into CO₂ and hydrogen radicals. Coal has almost no weight loss because the main body has not been destroyed. In this

stage, few reactions can occur because of the low temperature, and the impact of CO₂ can be ignored. The stage before T_1 is defined as the pyrolysis activation stage (stage I).

Between T_1 and T_2 , the linear ether bond (–O–CH₂–) reaches the activation condition as well. As shown in Fig. 11(b), the coal molecule splits into two relatively small activated fragments when bond 2 breaks. The activated fragments combine with free radicals to form stable product molecules. The number of carbon atoms in product molecule decreases with the fracture of ether bonds. Pyrolysis tar begins to be formed, which is liquid at room temperature. The generated tar is mainly heavy component, whose carbon atoms number is mostly between 14 and 40. Fig. 10 shows that the yield of heavy tar in CO₂ atmosphere is slightly lower than that in inert atmosphere. By analysing the trajectories of ReaxFF-MD simulations in CO₂ atmosphere, we find that some CO₂ react with hydrogen radicals to form aliphatic. Considering from the perspective of effective collision theory, the occurrence of chemical reaction requires both activated molecule and effective collision. The combination of CO₂ and hydrogen radicals decreases the concentration of free hydrogen radicals. The chances of effective collision between ether bond and free hydrogen radicals are reduced, which inhibits the formation of tar. However, oxygen-containing fragments produced by ether bond cleavage are strong hydrogen receptors, which easily bind to hydrogen radicals to form hydroxyl groups. Therefore, the inhibitory effect of CO₂ is very limited. In this stage, the main reaction is ether bond breaking and generating heavy tar. CO₂ has a slight inhibitory effect on pyrolysis. The stage between T_1 and T_2 is defined as the initial pyrolysis stage (stage II).

In Fig. 10, the yields of heavy tar begin to decrease near T_3 (2250 K). Between T_2 and T_3 , there are two main reactions occurring in the pyrolysis system. One is the bridge bonds (C_{ar}–C_{ar}) breaking between aromatic groups, and the other is the C–C bonds breaking on aliphatic chains. As shown in Fig. 11(c), the product molecules become smaller with the fracture of bond 3. Parts of the tar become light component, whose carbon atoms number is between 5 and 13. As illustrated in Fig. 4(b) and 8(b), the pyrolysis reaction rate reaches the maximum in this stage. In contrast to stage II, the tar yield in CO₂ atmosphere is higher than that in inert atmosphere. At this time, a lot of CH₄, C₂H₆ and other alkanes have been generated in the system. CO₂ and alkanes can



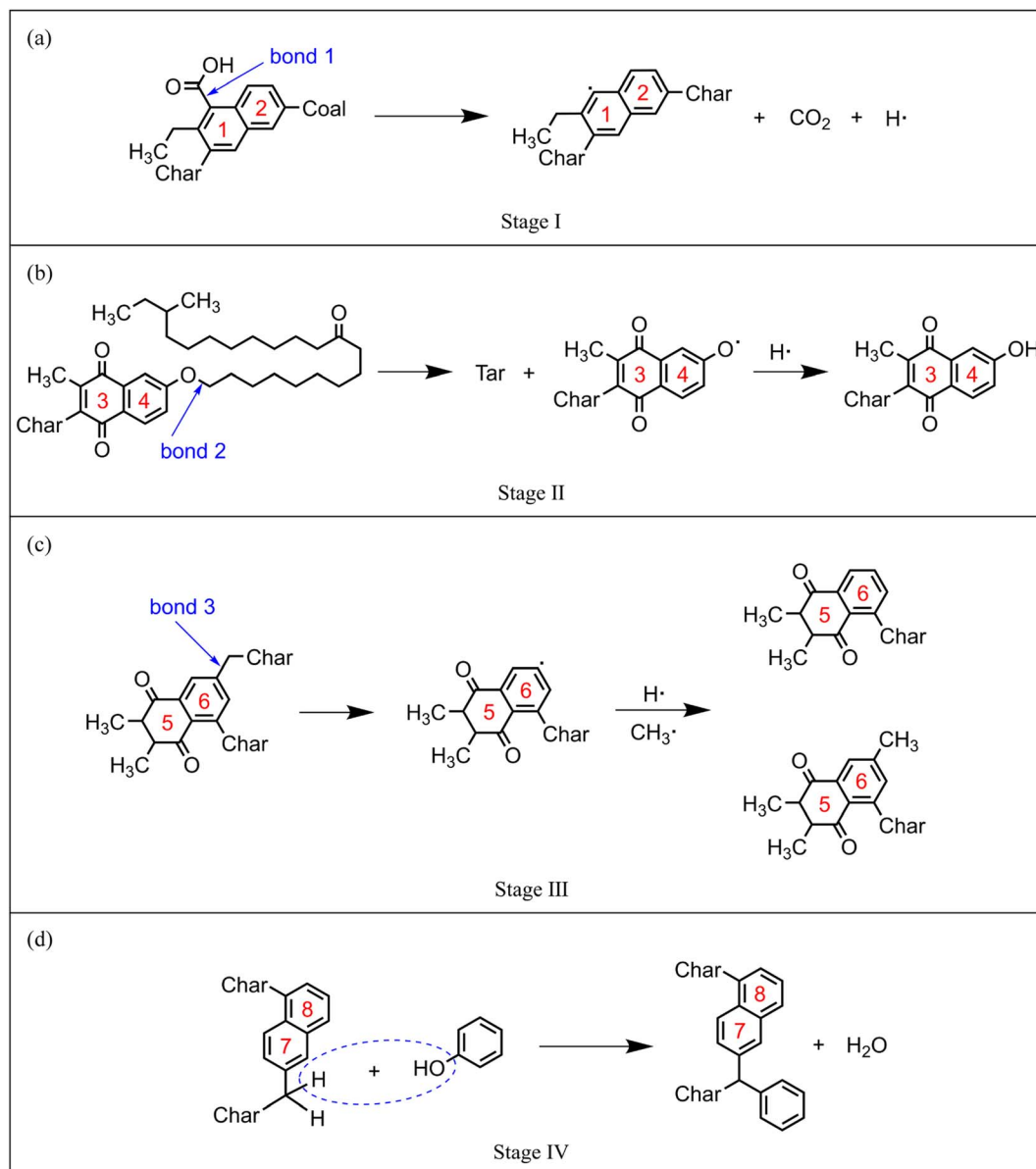
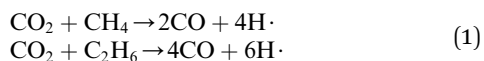


Fig. 11 Typical reaction pathways in each pyrolysis stage: (a) stage I, (b) stage II, (c) stage III, (d) stage IV.

occur reforming reaction at this temperature, as shown in formula (1). The reforming reaction produces CO and a large number of free hydrogen radicals. Hydrogen radicals can effectively collide with $\text{C}_{\text{ar}}\text{--C}_{\text{ar}}$ bonds and $\text{C}\text{--C}$ bonds, which promote the decomposition of macromolecules. This stage between T_2 and T_3 is defined as the accelerated pyrolysis stage (stage III).



After T_3 , the yield of heavy tar starts to decrease, while that of light tar increases rapidly. There are two reasons for the reduction of heavy tar. On the one hand, heavy tar continues to crack into light tar. On the other hand, heavy tar starts to convert to char because of the condensation reaction in high temperature. As shown in Fig. 10, the turning point of heavy tar

yield in CO_2 atmosphere arrives earlier and the decomposition rate of heavy tar is faster, resulting in more light tar. This is due to the fact that more free hydrogen radicals in CO_2 atmosphere promote the decomposition of heavy tar. In this stage, the decomposition reaction and condensation reaction compete fiercely. The yield of heavy tar decreases while more light tar is produced. This stage after T_3 is defined as secondary pyrolysis stage (stage IV).

At the end of pyrolysis at 2600 K, the yields of char in inert and CO_2 atmosphere are 22.36% and 16.08%, respectively. The yields of tar in inert and CO_2 atmosphere are 56.39% and 60.28%. Moreover, compared with in inert atmosphere, the yield of light tar is promoted from 17.98% to 20.68% in CO_2 atmosphere. It is concluded that CO_2 is positive to improve the yield and quality of pyrolysis tar.



Xu *et al.*³¹ investigated the pyrolysis characteristics and kinetics of two Chinese low-rank coal samples by thermogravimetric technique and mathematical modeling. The results indicated that Coats–Redfern integral model was appropriate to describe pyrolysis reaction of the two low-rank coals and the pyrolysis process can be divided into four stages according to the calculated activation energy. The experimental calculation results are consistent with the simulation results of this paper, which proves that the simulation of NMH coal pyrolysis based on ReaxFF-MD method is of great reference value.

4 Conclusions

In this paper, NMH low-rank coal pyrolysis in inert and CO₂ atmosphere is simulated based on ReaxFF-MD method. The effects of CO₂ on coal pyrolysis and the reaction mechanism are studied. The conclusions are as follows:

(1) The coal pyrolysis is first slightly inhibited and then significantly promoted by CO₂. The pyrolysis in CO₂ atmosphere is more complete and the final yield of char decreases from 22.36% to 16.08%.

(2) CO₂ helps to improve the pyrolysis tar yield and quality. Compared with in inert atmosphere, the final tar yield in CO₂ atmosphere increases from 56.39% to 60.28% and the light tar yield increases from 17.98% to 20.68%.

(3) Coal pyrolysis process can be divided into four stages according to the product distribution and chemical reaction type. In stage I, the carboxyl groups are detached and a small amount of CO₂ is produced. In stage II, the ether bonds are broken and the tar products are mainly heavy components. In stage III, disconnection of C_{ar}–C_{ar} bonds between aromatic groups and C–C bonds on aliphatic chains leads to the generation of light and heavy tars. In stage IV, decomposition reaction and condensation reaction compete fiercely and the yield of heavy tar decreases.

(4) In stage II, CO₂ slightly inhibits coal pyrolysis because CO₂ scrambles for free hydrogen radicals with ether bonds. In stage III and stage IV, reforming reaction of CO₂ with alkanes provides a great deal of hydrogen radicals and promotes the generation of tar.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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Notes and references

- 1 F. Yang, Q. Yu, Z. Qi and Q. Qin, *RSC Adv.*, 2021, **11**, 38434–38443.
- 2 Y. Hu, Z. Wang, X. Cheng and C. Ma, *RSC Adv.*, 2018, **8**, 22909–22916.

- 3 T. Li, Y. Li, Y. Cheng, X. Li, Y. Shen, L. Yan, M. Wang, L. Chang and W. Bao, *RSC Adv.*, 2021, **11**, 38537–38546.
- 4 T. Lv, M. Fang, H. Li, J. Yan, J. Cen, Z. Xia, J. Tian and Q. Wang, *RSC Adv.*, 2021, **11**, 17993–18002.
- 5 T. A. Saleh, *RSC Adv.*, 2022, **12**, 23869–23888.
- 6 T. Chmielniak, M. Sciazko, G. Tomaszewicz and M. Tomaszewicz, *J. Therm. Anal. Calorim.*, 2014, **117**, 1479–1488.
- 7 X. He, H. Hu, L. Jin and W. Hua, *Energy Sources, Part A*, 2016, **38**, 613–620.
- 8 P. Wang, L. Jin, J. Liu, S. Zhu and H. Hu, *Energy Fuels*, 2010, **24**, 4402–4407.
- 9 Q. Yi, Y. Fan, W. Li and J. Feng, *Ind. Eng. Chem. Res.*, 2013, **52**, 14231–14240.
- 10 Y. Oki, J. Inumaru, S. Hara, M. Kobayashi, H. Watanabe, S. Umemoto and H. Makino, *Energy Procedia*, 2011, **4**, 1066–1073.
- 11 K. R. Jillson, V. Chapalamadugu and B. Erik Ydstie, *J. Process Control*, 2009, **19**, 1470–1485.
- 12 M. Wang, L. Jin, H. Zhao, X. Yang, Y. Li, H. Hu and Z. Bai, *Fuel*, 2019, **250**, 203–210.
- 13 J. Han, X. Wang, J. Yue, S. Gao and G. Xu, *Fuel Process. Technol.*, 2014, **122**, 98–106.
- 14 J. Lee, X. Yang, H. Song, Y. S. Ok and E. E. Kwon, *Energy*, 2017, **120**, 929–936.
- 15 P. Wang, L. Jin, J. Liu, S. Zhu and H. Hu, *Fuel*, 2013, **104**, 14–21.
- 16 K. Jamil, J.-i. Hayashi and C.-Z. Li, *Fuel*, 2004, **83**, 833–843.
- 17 X. Li, Z. Mo, J. Liu and L. Guo, *Mol. Simul.*, 2014, **41**, 13–27.
- 18 J.-P. Wang, Y.-N. Wang, G.-Y. Li, Z.-Z. Ding, Q. Lu and Y.-H. Liang, *Chem. Phys. Lett.*, 2020, **744**, 137214.
- 19 Y. Zhang, C. Liu and X. Chen, *J. Anal. Appl. Pyrolysis*, 2015, **113**, 621–629.
- 20 F. Xu, H. Liu, Q. Wang, S. Pan, D. Zhao, Q. Liu and Y. Liu, *Fuel Process. Technol.*, 2019, **195**, 106147.
- 21 A. C. T. van Duin, S. Dasgupta, F. Lorant and W. A. Goddard III, *J. Phys. Chem. A*, 2001, **105**, 9396–9409.
- 22 F. Castro-Marciano, A. M. Kamat, M. F. Russo, A. C. T. van Duin and J. P. Mathews, *Combust. Flame*, 2012, **159**, 1272–1285.
- 23 F. Xu, H. Liu, Q. Wang, S. Pan, D. Zhao and Y. Liu, *Fuel*, 2019, **256**, 115884.
- 24 M. Gao, X. Li and L. Guo, *Fuel Process. Technol.*, 2018, **178**, 197–205.
- 25 D. Hong, L. Liu, S. Zhang and X. Guo, *Fuel Process. Technol.*, 2018, **178**, 133–138.
- 26 M. Zheng, X. Li, M. Wang and L. Guo, *Fuel*, 2019, **253**, 910–920.
- 27 M. Zheng, X. Li, J. Liu and L. Guo, *Energy Fuels*, 2013, **27**, 2942–2951.
- 28 M. Zheng, X. Li and L. Guo, *Fuel*, 2018, **233**, 867–876.
- 29 Y. Liu, J. Ding and K.-L. Han, *Fuel*, 2018, **217**, 185–192.
- 30 C. Chen, L. Zhao, J. Wang and S. Lin, *Ind. Eng. Chem. Res.*, 2017, **56**, 12276–12288.
- 31 Y. Xu, Y. Zhang, G. Zhang, Y. Guo, J. Zhang and G. Li, *J. Therm. Anal. Calorim.*, 2015, **122**, 975–984.

