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Authors:

Yuwen Zhu, Yanfang Miao, Haoyu Li

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Article

Enhancement Effect of Ordered Hierarchical Pore Configuration on SO₂ Adsorption and Desorption Process

Yuwen Zhu ^{1,*} , Yanfang Miao ¹ and Haoyu Li ²

¹ School of Energy and Safety Engineering, Tianjin Chengjian University, Tianjin 300384, China; 15620057620@163.com

² School of Science and Technology, Tianjin University, Tianjin 300072, China; lihaoyu@tju.edu.cn

* Correspondence: zhuyw@tcu.edu.cn; Tel.: +86-022-2308-5107

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Abstract: Carbonaceous adsorbents with both high sulfur capacity and easy regeneration are required for flue gas desulfurization. A hierarchical structure is desirable for SO₂ removal, since the micropores are beneficial for SO₂ adsorption, while the mesopore networks facilitate gas diffusion and end-product H₂SO₄ storage. Herein, an ordered hierarchical porous carbon was synthesized via a soft-template method and subsequent activation, used in SO₂ removal, and compared with coal-based activated carbon, which also had a hierarchical pore configuration. The more detailed, abundant micropores created in CO₂ activation, especially the ultramicropores ($d < 0.7$ nm), are essential in enhancing the SO₂ adsorption and the reserves rather than the pore patterns. While O₂ and H₂O participate in the reaction, the hierarchical porous carbon with ordered mesopores greatly improves SO₂ removal dynamics and sulfur capacity, as this interconnecting pore pattern facilitates H₂SO₄ transport from micropores to mesopores, releasing the SO₂ adsorption space. Additionally, the water-washing regeneration performances of the two types of adsorbents were comparatively determined and provide a new insight into the mass-transfer resistance in the pore structure. The ordered hierarchical carbon promoted H₂SO₄ desorption efficiency and cycled SO₂ adsorption–desorption performance, further indicating that interconnecting micro- and mesopores facilitated the diffusion of adsorbates.

Keywords: hierarchical pore structure; ordered mesopores; SO₂; adsorption; regeneration

1. Introduction

Along with the rapid development of the economy in China, a large amount of SO₂ emissions due to coal consumption in primary energy has resulted in severe air pollution [1,2]. The continued increase in coal consumption and utilization of high-sulfur coal will lead to more severe SO₂ pollution. Over recent decades, China has invested enormously in limestone–gypsum wet flue gas desulfurization (WFGD). However, this technology faces the dilemma of large water consumption levels, low market value byproducts, secondary pollution, and CO₂ escape [3]. Porous carbons, including activated carbon [4,5], activated coke [6–8] and semicoke [9,10], have been assessed to remove SO₂ from flue gas based on their abundant pore structures and specific chemical functionality, in addition to their advantages of water conservation, diversified sulfur resource recovery, adsorbent cyclic utilization, and reduced environmental impact. The current porous carbons used in flue gas desulfurization are mainly micro- and mesopore hierarchical coal-based activated carbons with Brunauer–Emmett–Teller (BET) surface areas of 650–850 m²·g^{−1}. However, the major drawbacks of activated carbons are their low sulfur capacity and costly regeneration, which has hindered their applications at a large scale in China [11,12].

As reported in the literature [13,14], the presence of O₂ and H₂O in flue gas causes the oxidation of absorbed SO₂, leading to the formation of H₂SO₄. Sun et al. [7] have demonstrated this process in a commercial activated coke with hierarchical pore configuration (BET surface area of 948 m²·g⁻¹, micropore volume of 0.32 cm³·g⁻¹, and mesopore volume of 0.11 cm³·g⁻¹) to achieve optimal removal rate and high sulfur capacity. The end-product H₂SO₄ was found to form in micropores that could migrate to mesopores, which guarantees the continuous proceeding of the SO₂ adsorption. Therefore, an ideal adsorbent should possess a hierarchical structure, in which the micropores will provide high adsorption and catalytic oxidation capacity, while the mesopores promote gas diffusion and end-product storage [15,16]. To enhance SO₂ removal performance, Karatepe et al. [17] investigated the influence of pore parameters on SO₂ adsorption and found that the micropore volume of the activated carbons, rather than the BET surface area, played an important role in SO₂ adsorption. Wang et al. [18] discussed the effect of the pore width on the SO₂ micropore filling mechanism. Mangun et al. [19] used activated carbon fibers as adsorbents for SO₂, since they have a uniform micropore structure and faster adsorption kinetics as compared to activated carbons. Moreover, Bagreev et al. [20] investigated the role of nitrogen functionality in the process of SO₂ adsorption and found that quaternary and pyridinic-type nitrogen significantly enhanced the adsorption capacity. On top of the above-mentioned research, previous studies have focused on improving pore volumes and surface areas, tuning pore structures and pore sizes, or adding catalysts [21,22]. However, hierarchical pore structures remain challenging for synthesis and application in flue gas desulfurization. Furthermore, hierarchical materials are also promising in many applications, including water and air purification [23], gas separation [24], catalysis [25], chromatography [26], and energy storage [27]. Related research results are worth learning about for hierarchical pore structures used in flue gas desulfurization.

Although common coal-based activated carbon can be prepared to possess hierarchical structures, the unconnected micro- and mesoporous structures caused by the complex structures of raw materials may hinder the diffusion of gas and the migration of adsorbates. Our strategy is to prepare an ordered hierarchical pore structure via a soft-template method to promote the synthesis of ordered mesopores and subsequent activation toward the development of abundant micropores embedded in the mesopore walls. These adsorbents with ordered mesoporous features were used for SO₂ removal and compared with coal-based activated carbons. We focused on the effects of pore sizes and pore patterns on the SO₂ removal process, including SO₂ separate adsorption and reaction with O₂ and H₂O participation. The sulfur-containing adsorbate storage spaces within adsorbents under various conditions were further determined. Furthermore, to better examine the diffusivity of the ordered hierarchical structure, the water-washing regeneration efficiency and the cycled desulfurization–regeneration performance of the two adsorbents were comparatively investigated.

2. Materials and Methods

2.1. Preparation of Coal-Based Activated Carbon

A series of coal-based activated carbons (ACs) were prepared using Taixi anthracite (Ningxia, China) as a precursor. The AC samples were prepared through carbonization and subsequent CO₂ activation. In brief, 10 g of feed coal (size range of 0.25–0.38 mm) was placed in a quartz tubular reactor (23 mm inner diameter and 400 mm length) located in a horizontal furnace. Carbonization was carried out by heating to 973 K at a rate of 8 K·min⁻¹ under a N₂ (99.999%) flow of 350 mL·min⁻¹ and maintained for 40 min. Then, the reactor was heated to 1123 K in a CO₂ (99.999%) constant flow of 350 mL·min⁻¹ for various times to obtain various burn-offs ranging from 12.8% to 71.3%. The burn-off, related to the activation step, is the ratio between the weight of fixed carbon reacted and the initial char weight [28,29]. The notations used for the coal-based activated carbons are AC-C-(burn-off).

2.2. Synthesis of Ordered Mesoporous Carbon and Hierarchical Carbon

The ordered mesoporous carbons (OMCs) were prepared via co-assembly of the phenolic resin precursor with a triblock copolymer Pluronic F127 via soft templating. The phenolic resin precursor was synthesized as follows: 40 g of phenol was heated at 318 K in a water bath until completely melted. Then, 20% NaOH solution (34 mL) was added to the conical flask containing the melted phenol in drops, and the mixture was stirred for 15 min. Then, a 37% formaldehyde solution was added to the mixture in drops with stirring. Note that the formaldehyde and phenol were mixed at a substance ratio of 3:1. Subsequently, the conical flask was submerged in a water bath set at 358 K with continuous stirring for 1 h and then cooled to room temperature. The pH value of the resulting mixture was adjusted to 7.0 with 20% hydrochloric acid solution and then the water was removed using a rotary evaporator at 323 K for 4 h to obtain the dried phenolic resin. A 15% phenolic resin-ethanol solution was prepared for use. Triblock copolymer Pluronic F127 was used as the soft template for the synthesis of mesoporous carbons. 3 g of F127 was dispersed in ethanol to prepare the 17% F127-ethanol solution, which was then added to a 15% phenolic resin-ethanol solution with continuous stirring for 30 min. The resultant solution was allowed to sit without stirring at room temperature until the ethanol was volatilized completely; it was subsequently heated at 373 K for 24 h to obtain the polymer composite. The resultant polymer composite was heated to 1123 K at a ramp rate of $1 \text{ K}\cdot\text{min}^{-1}$ and held for 3 h in a N_2 (99.999%) flow of $350 \text{ mL}\cdot\text{min}^{-1}$ in the horizontal furnace to slowly decompose the surfactant F127 and produce the mesoporous carbon.

To further improve the textural properties, a subsequent CO_2 activation was carried out at 1123 K in the horizontal furnace as mentioned above in a CO_2 (99.999%) constant flow of $350 \text{ mL}\cdot\text{min}^{-1}$ for various times to obtain various burn-offs, denoted as OMC-C-(burn-off).

Except for Pluronic F127 (Sigma-Aldrich, Missouri, USA), all other chemicals used in the synthesis of the adsorbents were obtained from Kemiou, Tianjin, China.

2.3. Characterizations

Pore parameters of ACs and OMCs were determined by N_2 adsorption over a relative pressure P/P_0 range from 10^{-7} to 1 at 77 K using an automatic apparatus (ASAP 2020, Micromeritics, Norcross, GA, USA). Before this analysis, the sample was degassed under vacuum at 473 K for 10 h. Note that the degassing step was terminated when the saturated adsorbent was measured to avoid the desorption of the adsorbates (i.e., SO_2 , SO_3 , H_2SO_4) [30]. The BET surface area was calculated from the isotherm using the Brunauer-Emmett-Teller (BET) equation. The micropore surface was calculated by the t -plot method [31]. The micropore volume and differential pore volume were estimated by the Horvath-Kawazoe (HK) method [17]. The mesopore volume and pore size distributions of the samples were calculated by applying the Barrett-Joyner and Halenda (BJH) method [32].

The microstructural morphology of ACs and OMCs was analyzed using transmission electron microscopy (TEM, Tecnai T20, FEI, Hillsboro, OR, USA). The small angle X-ray scattering (SAXS) patterns ($0.5^\circ < 2\theta < 5^\circ$) were acquired on a D/max 2400X diffractometer operating at 40 KV and 100 mA using $\text{Cu K}\alpha$ radiation ($\lambda = 1.5406 \text{ \AA}$, Rigaku, Tokyo, Japan). Elemental analysis (EA) was performed using an analyzer (Vario MACRO cube, Elementar, Langenselbold, Germany) for determination of the total carbon and oxygen content of the bulk samples. The types of surface functional groups were identified by Fourier transform infrared (FTIR) spectroscopy (Magna-IR 560 E.S.P, Nicolet, Madison, WI, USA). The spectrometer was used in the range of $4000\text{--}500 \text{ cm}^{-1}$ with a resolution of 4 cm^{-1} , using pressed KBr pellets that contained approximately 0.5 wt.% samples, to realize the measurements.

2.4. SO₂ Adsorption Measurements

The SO₂ adsorption experiments were carried out in a fixed bed reactor using 2.5 g of sample at 313 K or 353 K. The experimental system consisted of a tubular reactor (20 mm ID) placed in a vertical furnace with a system of valves and mass flow controllers. The mass flow controllers were used to control the flows of SO₂, O₂, and N₂, respectively. The water vapor was introduced by bubbling a flow of N₂ at a fixed temperature in water bath. Flow of H₂O was controlled by adjusting N₂-carrying flow. An online Fourier transform infrared gas analyzer (DX4000, Gaset, Vantaa, Finland) was used to continuously monitor the SO₂, O₂, and H₂O concentrations of the reactor inlet and outlet. In addition, the experimental installation was provided with a bypass to allow the SO₂, O₂, and H₂O concentrations to be measured before introducing the gas mixture into the reactor. The inlet gas volumetric compositions used in different experiments were 1500 ppm SO₂, with or without 5% O₂, with or without 10% H₂O, with N₂ balance, and a total flow rate of 1.2 L·min⁻¹. It is worth mentioning that a previous work found that CO₂ in the inlet gas does not influence SO₂ removal behavior [33].

SO₂ removal efficiency versus time was defined by recording concentrations of SO₂ at the inlet and outlet in real time via the gas analyzer. The amount of SO₂ removed, expressed in mg SO₂ g⁻¹ carbon, was calculated by the integration of SO₂ conversion versus time curves.

2.5. Water-Washing Regeneration

Water-washing regeneration, just as its name implies, recovers the saturated adsorbents by extracting adsorbate H₂SO₄ from the inner pores depending on the H₂SO₄ differential concentration in water. First, 2.5 g of spent adsorbent was dispersed in 100 mL of water at 298 K with a magnetic stirring bar for an appropriate time ranging from 15 to 75 min. After filtering and subsequent drying at 373 K for 12 h, the regenerate adsorbent was obtained and reused in desulfurization.

3. Results

3.1. Structural Characterization of Two Different Carbon Adsorbents

We prepared two species of carbonaceous adsorbents, including coal-based activated carbons with hierarchical pore structures and ordered mesoporous carbons with micropores embedded in the mesopore walls. The nitrogen adsorption–desorption isotherms and the resulting BJH mesopore size distributions of the AC-C-*x* and OMC-C-*x* are shown in Figure 1.

The samples of AC-C-12.8 and AC-C-32.2 display a type I adsorption isotherm without any hysteresis loop and significant nitrogen uptake at a relative pressure less than 0.1, indicating the presence of a large fraction of micropores. With an increase in burn-off, the amount of adsorbed nitrogen increases and the knee becomes broader, which indicates the development of micropores. Meanwhile, the hysteresis loop appears, as shown in Figure 1b, indicating the presence of mesopores, and the corresponding BJH mesopore size distributions present a mesopore size in a range from 2–4 nm for AC-C-50.8 and AC-C-71.3, as shown in Figure 1d.

Figure 1c shows that the OMC sample has type IV isotherms with a type H₁ hysteresis loop at p/p_0 from 0.3 to 0.6, indicating the presence of two-dimensional hexagonal mesoporous structures [34] and the major mesopore distribution centered at 3–4 nm. After CO₂ activation, the nitrogen sorption isotherms of the three OMC-C-*x* samples all show steep uptakes at low relative pressures, indicating the presence of micropores. Compared to untreated OMC, the hysteresis moves to the right to $p/p_0 > 0.4$, suggesting that the ordered mesostructure is well preserved and the mesopore size is widened.

The pore microstructural characteristics of the ACs and OMCs are listed in Table 1. It is clear that with CO₂ treatment at 1123 K, higher burn-offs give rise to higher BET surface areas and larger pore volumes (ultramicro- pore, micropore, and mesopore volumes). The activation with CO₂ clearly creates new micropores and expands the existing porous system. The micro- and mesopores co-existed in the AC-C-50.8, AC-C-71.3, and OMC-C-*x* samples.

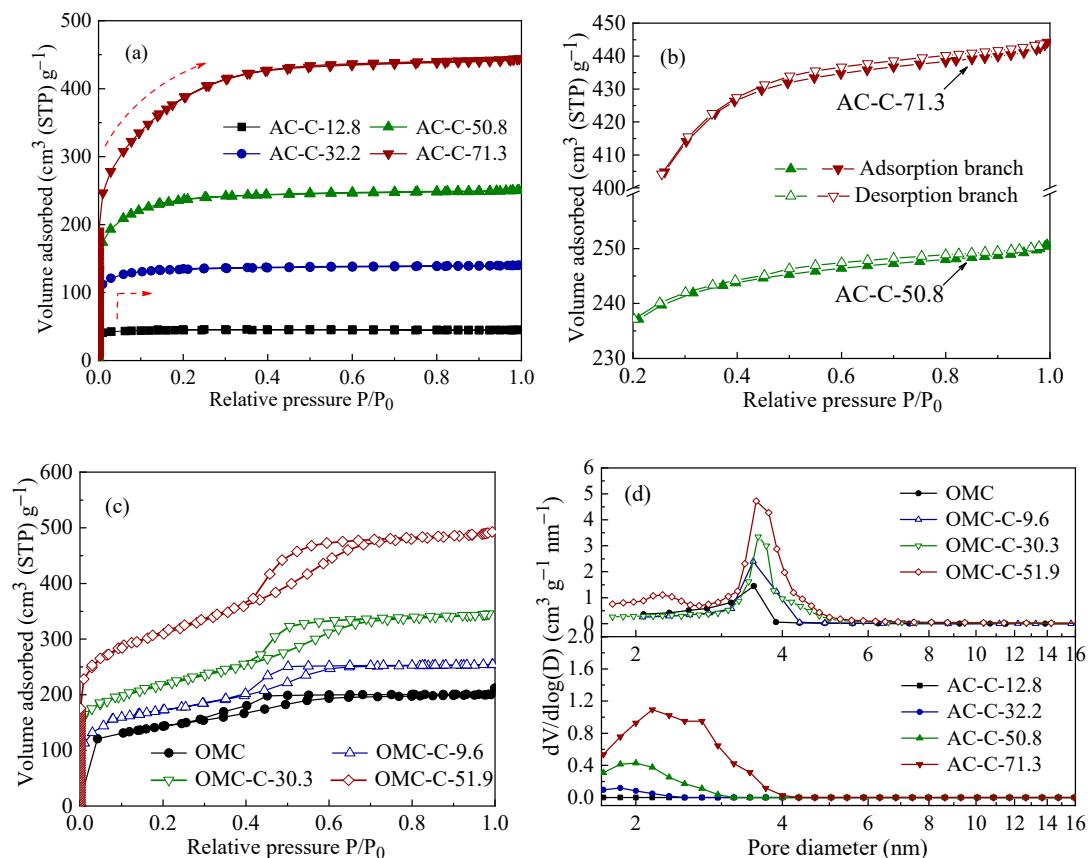


Figure 1. (a–c) N₂ adsorption and desorption isotherms at 77 K for the activated carbon (AC) and ordered mesoporous carbon (OMC) samples; (d) Barrett–Joyner and Halenda (BJH) mesopore size distribution plots of the AC and OMC samples. STP: Standard temperature and pressure.

Table 1. The pore microstructural characteristics and elemental analysis of the samples.

Sample	S_{BET}^1 (m ² g ⁻¹)	V_{ult}^2 (cm ³ g ⁻¹)	V_{mic}^3 (cm ³ g ⁻¹)	V_{mes}^4 (cm ³ g ⁻¹)	D_{mes}^5 (nm)	Elemental Analysis (wt.%)		
						C	O	O/C × 10 ⁻²
AC-C-x								
AC-C-12.8	149(138)	0.05	0.07	0.01	2.75	86.6	4.78	5.52
AC-C-32.2	449(373)	0.11	0.20	0.02	3.08	85.6	5.08	5.93
AC-C-50.8	809(537)	0.15	0.34	0.14	3.12	83.5	6.01	7.19
AC-C-50.8 after 8 cycles ⁶	364(188)	0.05	0.12	0.13	3.29	–	–	–
AC-C-71.3	1379(404)	0.18	0.52	0.33	3.31	80.8	5.16	6.38
OMC-C-x								
OMC	475(205)	–	–	0.21	3.17	95.1	3.4	3.58
OMC-C-9.6	576(287)	0.07	0.14	0.26	3.27	94.7	4.0	4.22
OMC-C-30.3	824(439)	0.12	0.27	0.35	3.44	93.1	5.9	6.34
OMC-C-51.9	1056(508)	0.16	0.39	0.43	3.51	91.9	6.7	7.29

¹ S_{BET} : Brunauer–Emmett–Teller (BET) surface area. The numbers in parentheses are micropore surface areas, calculated by the t -plot method. ² V_{ult} , ³ V_{mic} : Ultramicropore (<0.7 nm) volume and micropore (<2 nm) volume, calculated by the Horvath–Kawazoe (HK) method. ⁴ V_{mes} , ⁵ D_{mes} : Mesopore (2–50 nm) volume and median mesopore diameter, calculated by applying the BJH method. ⁶ AC-C-50.8 after 8 cycles: AC-C-50.8 sample after eight cycles of desulfurization and water-washing regeneration.

TEM images of AC-C-50.8 and the corresponding Fourier diffractogram are shown in Figure 2a. The TEM image appears to have an isotropic cross-sectional structure, showing a random orientation of the pores. In contrast, Figure 2b shows the presence of a two-dimensional ordered mesoporous structure of the OMC sample. After CO₂ activation with 51.9% burn-off, the ordered structure still remains as Figure 2c shows, suggesting that the ordered mesostructure is well preserved under CO₂ treatment. The periodicity of OMC-C-51.9 was further confirmed using small-angle XRD, as shown

in Figure 2d. The existence of the (100) peaks is clearly observed, which supports the presence of two-dimensional arrays. Thus, micro- and mesoporous hierarchical carbons with ordered mesoporous characteristics were obtained.

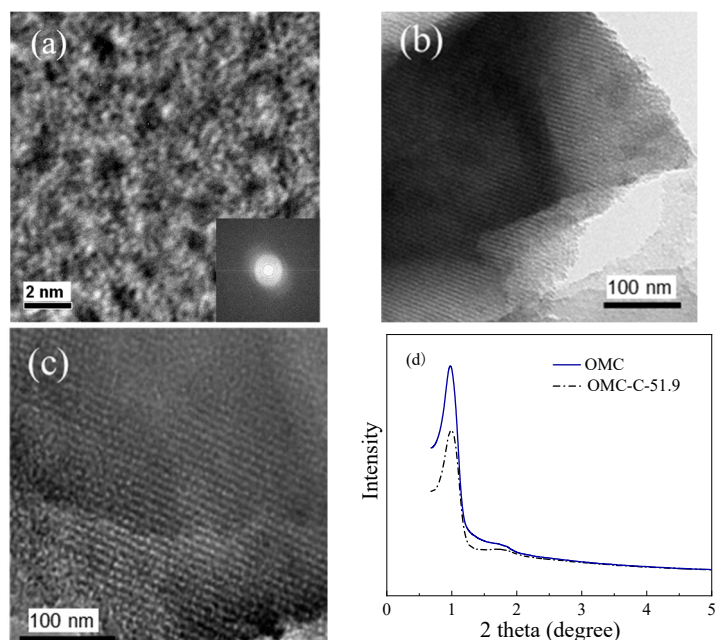


Figure 2. Morphological characterizations of hierarchical porous carbons. TEM micrographs of (a) AC-C-50.8 (inset: Fast Fourier diffractogram), (b) OMC, and (c) OMC-C-51.9; (d) Small-angle XRD patterns of OMC and OMC-C-51.9.

3.2. Chemical Composition of Two Different Carbon Adsorbents

To understand and contrast two species of adsorbent chemical compositions, EA and FTIR analysis were performed. The bulk oxygen content by EA was found to increase from 3.4 wt.% to 6.7 wt.% with increasing burn-off for OMC-C-*x*, as shown in Table 1. In contrast, the oxygen content of AC-C-*x* first increases, reaching a maximum when the burn-off is 50.8%, and then decreases with increasing burn-off degree. The decrease in the oxygen content of AC-C-71.3 may be attributed to a significant increase in the ash content, which reaches 10% because of the high burn-off degree.

Generally, the reaction of CO₂ with the carbon atoms in porous carbon increases the oxygen content and creates oxygen-containing groups on the surface. The characteristic functional groups of the AC-C-50.8 and OMC-C-51.9 are evaluated by FTIR, as shown in Figure 3. Both spectra have similar absorption bands, being slightly different in some wavenumbers and differences in their intensities, which indicates just small differences in the surface chemistry. The broad band at approximately 3430 cm⁻¹ is considered to be in the -OH stretching mode, indicating the presence of a phenolic hydroxyl group. The bands at 2920 cm⁻¹ and 2850 cm⁻¹ are both associated with a C-H stretching mode originating from C-H in the F127 and C-H of the aliphatic series for the two species of adsorbent, respectively. The band in the 1630 and 1574 cm⁻¹ region is associated with the C=O stretching mode in quinolyl [35]. The broad band extended between 1300 and 950 cm⁻¹ is assigned to both the -C-O stretching and C-O-H bending modes of alcoholic, phenolic, and carboxylic groups, including the C-O-C stretching mode of ether in the 1280–1000 cm⁻¹ region [36]. According to the above analysis, although the two carbonaceous adsorbents were prepared using anthracite and phenolic resin as precursors, respectively, they have similar surface functional groups with CO₂ activation at 1123 K. Therefore, it is considered that the types of surface functional groups are mainly related to CO₂ activation rather than the precursor in this study.

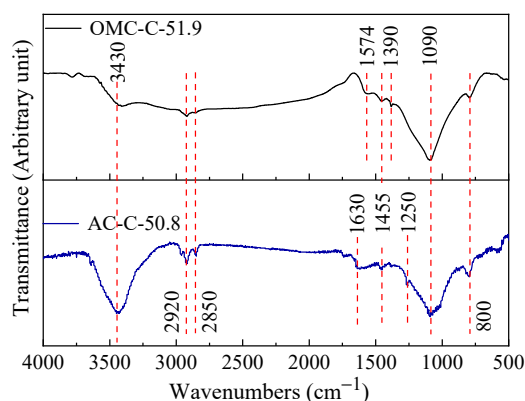


Figure 3. FTIR spectra of the hierarchical carbons.

3.3. SO_2 Removal Performance of Adsorbents

SO_2 adsorption is the first step of subsequent oxidation and sulfuric acid-forming reactions within porous carbon. The SO_2 separate adsorption dynamics were investigated for the AC and OMC samples, as shown in Figure 4a,b. The figures show that the SO_2 removal efficiency reaches 100% at the initial stage and then rapidly declines with the adsorption time. Furthermore, for the two species of carbonaceous adsorbents, the removal efficiency and breakthrough time show an increasing trend with increasing burn-off as the pore volumes increase (Table 1).

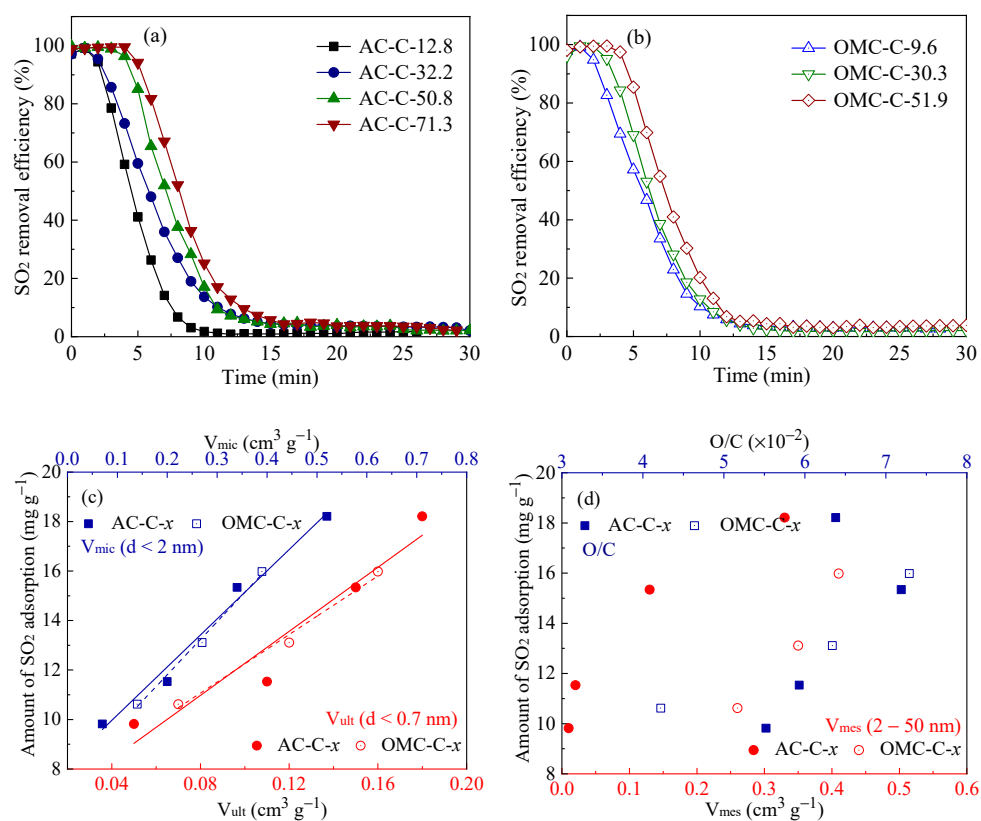


Figure 4. SO_2 -adsorption performance of the AC and OMC samples. SO_2 removal efficiency for (a) AC-C-x and (b) OMC-C-x. Adsorption conditions: 2.5 g of adsorbents, 1500 ppm SO_2 , N_2 balance, flow rate $1.2 \text{ L}\cdot\text{min}^{-1}$, at 313 K. The correlations of calculated SO_2 adsorption amount and the physico-chemical parameters: (c) Ultramicropore volume (V_{ult}) and micropore volume (V_{mic}); (d) mesopore volume (V_{mes}) and O/C.

To better understand the role of the pore structure, we investigated the effect of pore sizes and pore patterns within the adsorbents. From structure characterization, it appears that ultramicropore volume (V_{ult}) and micropore volume (V_{mic}) directly contribute to the SO_2 adsorption capacities of different samples, as shown in Figure 4c. Previous studies [33,37] have shown that micropores with diameters of approximately 0.7 nm may be important in SO_2 adsorption because they largely enhance intermolecular interactions. In our experiments, the amount of SO_2 adsorption increases linearly with V_{ult} and V_{mic} and the fitting lines for AC-C- x and OMC-C- x almost coincide, which indicates that the pore sizes play a key role in SO_2 adsorption rather than pore patterns. Meanwhile, it appears that neither the mesopore volume (V_{mes}) nor the O/C directly contribute to the amount of SO_2 adsorption, as shown in Figure 4d.

Considering the presence of O_2 and H_2O in flue gas leading to a series of catalytic oxidation and H_2SO_4 -forming reactions within carbonaceous adsorbents, the SO_2 removal dynamics under simulated flue gas were investigated, as shown in Figure 5. Unlike the SO_2 removal efficiency rapidly declining to 0 within 15 min in the SO_2 separate adsorption process, the SO_2 removal efficiency gradually declines to a value and almost maintains a plateau, as marked in the red box. Especially for OMC-C-30.3 and OMC-C-51.9, the SO_2 removal efficiencies decline slower than that of AC-C- x , and the values of plateau remain higher than those of AC samples. Comparatively, OMC-C-51.9 greatly improves the sulfur capacity, reaching 107.8 mg g^{-1} , as marked in Figure 5. AC-C-50.8, which has similar micropore characteristics, possesses a sulfur capacity of 75.6 mg g^{-1} . Thus, different SO_2 removal dynamics and sulfur capacities suggest that mesopore and pore patterns should have a coupling influence on SO_2 removal in the presence of O_2 and H_2O .

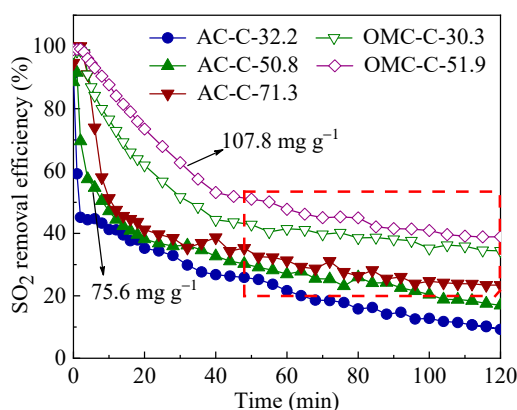


Figure 5. SO_2 removal efficiency versus time for AC and OMC samples. Reaction conditions: 2.5 g of adsorbents, 1500 ppm SO_2 , 5% O_2 , 10% H_2O , N_2 balance, flow rate $1.2 \text{ L} \cdot \text{min}^{-1}$, at 353 K.

To further elucidate the effect of pore structure, the changes of pore volumes, including V_{ult} , V_{sup} , and V_{mes} , after adsorption of SO_2 separately and reaction under $SO_2+O_2+H_2O$ conditions were investigated, as shown in Figure 6. These changes can be considered to be caused by the adsorption and storage of the main adsorbates (ex. SO_2 , SO_3 , and H_2SO_4). After the SO_2 adsorption, the V_{ult} of AC-C-50.8 and OMC-C-51.9 both decrease obviously compared to that of the pristine samples, while the V_{sup} decrease slightly and the V_{mes} have no change. These results suggest that SO_2 adsorption mainly occurs in the micropores, especially in the ultramicropores, occupying the space of narrow pores with sizes smaller than 0.7 nm. Note that the decreasing V_{ult} is in good agreement with the trend of SO_2 adsorption capacities with V_{ult} , as shown in Figure 4c, and further indicates that the ultramicropore is an important factor for high SO_2 adsorption.

In the presence of O_2 and H_2O in flue gas, adsorbed SO_2 within porous carbon reacts with O_2 and H_2O , catalyzed by carbon, to form the byproduct H_2SO_4 , which can occupy the pore structure, thus hindering continuous SO_2 adsorption. After reaction under $SO_2 + O_2 + H_2O$ conditions, the V_{ult} , V_{sup} , and V_{mes} all show a decreasing trend, suggesting that the adsorbate H_2SO_4 exists in these storage spaces. Compared to AC-C-50.8, the V_{ult} and V_{sup} of OMC-C-51.9 decrease to a much lesser extent.

Meanwhile, the V_{mes} of OMC-C-51.9 show an obvious decrease. These results indicate that H_2SO_4 produced in the OMC-C-51.9 is much easier to transport and store in the mesopores, which may give rise to the recovery of micropores (ultramicro-pore and supermicropore), guaranteeing the continuous proceeding of the SO_2 adsorption and transformation of sulfur-containing species. This is verified by the better SO_2 removal performance of OMC-C-30.3 and OMC-C-51.9 in the presence of O_2 and H_2O (Figure 5). Although AC-C-50.8 also has a hierarchical pore configuration, unconnected micro- and mesoporous may be the obstacle in H_2SO_4 migration. Therefore, the pore pattern with micropores embedded in the ordered mesopores play a key role in the migration of byproducts and continuous SO_2 removal in flue gas.

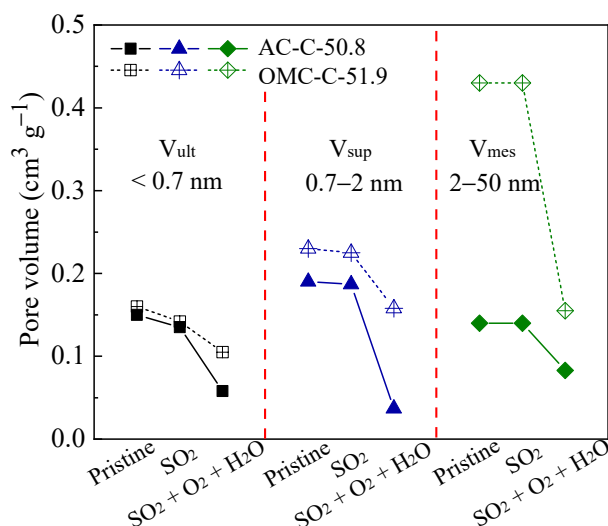


Figure 6. Pore volume parameters of fresh adsorbents (pristine), adsorbents after adsorption of SO_2 separately (SO_2) and adsorbents after reaction under $SO_2 + O_2 + H_2O$ conditions. Samples: AC-C-50.8 and OMC-C-51.9. Pore volume parameters: V_{ult} , supermicropore volume (V_{sup} , 0.7–2 nm) and V_{mes} , and the V_{sup} obtained by ($V_{mic} - V_{ult}$).

3.4. Regeneration and Cycling Stability Performance of Adsorbents

As reported in the literature [11,38–40], thermal regeneration is usually applied to extract the adsorbate H_2SO_4 based on the redox reaction ($C + H_2SO_4 \rightarrow CO_2 + SO_2 + H_2O$) between carbon and H_2SO_4 in the porous carbon. However, adsorbents will produce inevitable chemical loss, which usually leads to micropore damage and decay of SO_2 removal performance. Thus, we chose the water-washing regeneration method and contrastively investigated the regeneration efficiency and eight cycles of desulfurization and regeneration performance of AC-C-50.8 and OMC-C-51.9. This regeneration method also provides new insight into the mass-transfer resistance in different interconnecting pore patterns.

Desorption of H_2SO_4 from saturated adsorbents was carried out with various regeneration times (15, 30, 45, 60, and 75 min) to determine the most appropriate desorption duration for adsorption–desorption cycling. As shown in Figure 7, the H_2SO_4 desorption efficiencies of AC-C-50.8 and OMC-C-51.9 increase from 71.9% to 85.3% and from 79.0% to 91.6% with increasing regeneration time. During 75 min regeneration, the H_2SO_4 desorption efficiencies are very close to the values of 60 min regeneration, suggesting that the desorption efficiency does not vary with time beyond 60 min. Thus, for recycling purposes, AC-C-50.8 and OMC-C-51.9 samples were regenerated by water scrubbing for 60 min. Moreover, desorption efficiencies indicate that adsorbate H_2SO_4 occurring in the OMC-C-51.9 is easier to extract from the inner pores, which is attributed to the reduction of the mass-transfer resistance in the hierarchical carbons with ordered mesopores.

To compare the effects of water-washing regeneration on desulfurized AC and OMC samples, the sulfur capacities versus cycling number of AC-C-50.8 and OMC-C-51.9 were investigated

as shown in Figure 8. The working sulfur capacity of AC-C-50.8 shows a general decreasing trend from $75.6 \text{ mg}\cdot\text{g}^{-1}$ in the first-time desulfurization to $46.2 \text{ mg}\cdot\text{g}^{-1}$ in the 8th cycle with a reduction percentage of 40.5%, as shown in Figure 8a. The results indicate that incomplete desorption of H_2SO_4 from saturated adsorbents has an obviously negative impact on the cycling SO_2 adsorption performance of AC-C-50.8. Similar results are observed with OMC-C-51.9. However, the sulfur capacity of OMC-C-51.9 decreases by 22.0% after eight cycles, which is much less than the rate of decrease of AC-C-50.8, suggesting that the characteristics of interconnecting micropores and ordered mesopores facilitate the desorption of H_2SO_4 and diminish the decrease of sulfur capacity.

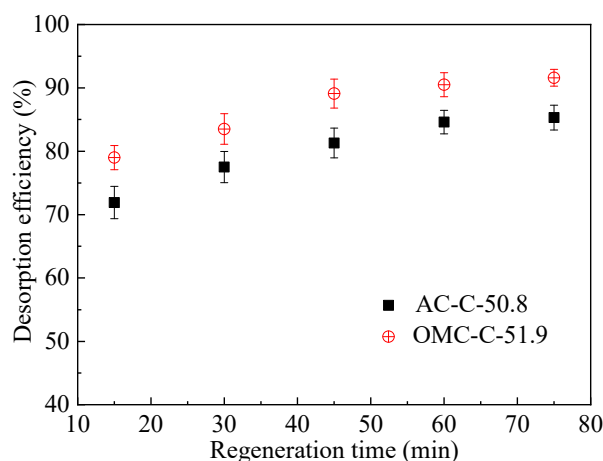


Figure 7. Desorption efficiency versus regeneration time for AC-C-50.8 and OMC-C-51.9 by water-washing regeneration method.

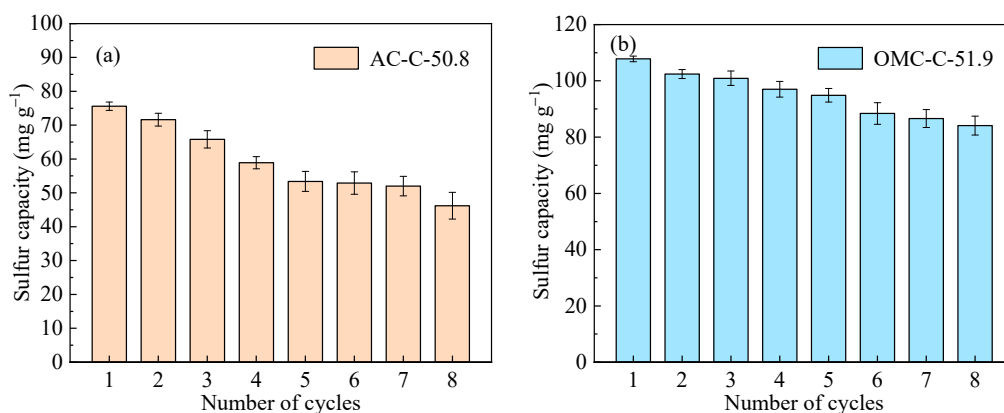


Figure 8. Sulfur capacity of the adsorbent over eight adsorption-regeneration cycles. (a) AC-C-50.8; (b) OMC-C-51.9. Adsorption conditions: 2.5 g of adsorbents, 1500 ppm SO_2 , 5% O_2 , 10% H_2O , N_2 balance, for 2 h, at 353 K. Regeneration time: 60 min.

To further examine the potential changes to the physical properties, the pore parameters of the eight times cycled water-washing regeneration AC-C-50.8 are listed in Table 1. Compared to pristine adsorbent, the BET surface area and micropore surface area of the adsorbent after eight cycles show an obvious decrease. Meanwhile, the ultramicropore and micropore volumes decrease from 0.15 to $0.05 \text{ cm}^3\cdot\text{g}^{-1}$ and from 0.34 to $0.12 \text{ cm}^3\cdot\text{g}^{-1}$, respectively. However, the mesopore volume shows little change after manifold cycles. That is, the decrease of sulfur capacity should be ascribed to the decrease of micropore volume as the sulfur-containing adsorbates remain in the inner pores. According to the regeneration performance of OMC-C-51.9, the ordered hierarchical pore configuration was found to facilitate the desorption of the adsorbates. Some species of surface functional groups may also lead to strong interactions between the adsorbed H_2SO_4 and the carbon surface, thus hindering the desorption of H_2SO_4 , which will be further studied in our research.

4. Conclusions

In summary, two species of carbonaceous adsorbents were prepared, including coal-based activated carbons and hierarchical porous carbons with ordered mesopores. The physico-chemical characteristics and resulting SO₂ adsorption and desorption performance were comparatively studied. The CO₂ activation process of AC and OMC samples leads to abundant micropores, which are essential in enhancing the SO₂-sorbent interactions. In particular, ultramicropore volume as the occurrence space of adsorbed SO₂ directly contributes to the sulfur capacity for SO₂ separate adsorption. With O₂ and H₂O participating in the gas composition, the SO₂ removal efficiency appears to reach a plateau stage. Comparatively, the OMC sample, which has similar micropore characteristics to the AC sample, shows a much higher sulfur capacity, reaching 107.8 mg·g⁻¹, mainly because of the ordered hierarchical porous configuration facilitating the transport of H₂SO₄ from micropores to mesopores. Meanwhile, the OMC sample promotes H₂SO₄ desorption efficiency with water-washing regeneration and cycled desulfurization–regeneration performance, further indicating that this pore pattern plays a key role in the diffusion and migration of adsorbates. The present results suggest that hierarchical pore configuration and interconnecting patterns, and not merely pore volumes, should be considered for optimizing heterogeneous gas–solid adsorption and reaction.

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