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Synthesis of porous carbon *via* a waste tire leavening strategy for adsorptive desulfurization

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Adsorptive desulfurization is an effective technology for removing harmful sulfur under mild conditions. Carbon materials have many advantages and are often used in adsorption desulfurization research, but until now have been synthesized using complicated methods and have shown limited adsorption capacity. Using an NaHCO_3 -assisted leavening method, waste tires were in the current work used as raw materials to produce hierarchically porous carbon that exhibits a high specific surface area and abundant oxygen-containing functional groups. In contrast to the sulfur removal by the carbon material prepared using a commonly used method, the as-prepared carbon material shows excellent adsorption performance, and was able to achieve an ultra-deep desulfurization of pentanethiol, specifically removing up to 99.7% of the sulfur from a model fuel with an initial sulfur concentration of 28 ppm. Therefore, we have provided a simple method for synthesizing adsorbents with high adsorption performance, and we expect these adsorbents to be used for industrial adsorptive desulfurization.

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1. Introduction

To reduce the impact of pollutants on the environment,^{1,2} increasing attention has been paid to the standards for the emission of sulfides in the formulation of fuels.^{3–5} Recently, sulfide emissions have been limited by these standards to be less than 10 ppm.^{6,7} For a long time, the hydrodesulfurization process,^{8–14} as a cornerstone of the chemical industry, has been applied in removing harmful sulfur from oil.⁹ However, the hydrogen sulfide generated in the hydrodesulfurization process reacts with olefins to form a small amount of mercaptan, and sometimes the hydrotreatment needs to be repeated, resulting in increased costs. Therefore, non-hydrodesulfurization techniques for removing the mercaptan produced after the hydrodesulfurization process¹⁵ have been studied in order to greatly reduce costs. Such studied non-hydrodesulfurization of mercaptans has included photocatalytic oxidative desulfurization,^{16,17} oxidative desulfurization,^{18,19} adsorption and photocatalytic oxidation desulfurization,²⁰ adsorptive desulfurization,²¹ and so on. Of these desulfurization processes, adsorptive desulfurization has been particularly widely studied^{22,23} because of its low energy consumption and effective removal capacity.²⁴

So far, the main adsorbents used for removing mercaptans have included zeolite,^{25,26} activated carbons,^{27–31} nanoporous

graphene,²¹ and boron nitride,³² among others. Of these different types of adsorbents, activated carbon has shown some particularly excellent properties such as high adsorption capacity,^{33,34} high specific surface area^{35,36} and easy regeneration. The raw materials that have been recently reported to be used for the synthesis of activated carbon have included sewage sludge, biochar, waste rubber tires, wood, lignocellulosic biomass, and animal bones.³⁷ Rubber tires are essential for our daily transportation, and since they readily wear out, many waste tires become available every year all over the world. However, waste tires are non-biodegradable materials,³⁷ and hence pose environmental problems. These problems, however, could be solved by recycling and reusing the tires. A previously reported method for preparing activated carbon materials from waste tires^{37–44} involves carbonizing the raw materials at about 500 °C, and subjecting the resulting sample to a physical activation process and then an HNO_3 and NaOH chemical activation (Table 1). In addition, metals and metal oxides have been added to the activated carbon material to further enhance the adsorption performance.^{39,42} Therefore, in order to avoid the introduction of metals and to simplify the synthesis step, further investigations of the synthesis method have become extremely important. The “leavening” method has been reported to be advantageous for increasing the specific surface area of carbon materials.^{45,46}

Herein, to develop a green and sustainable synthetic process starting with waste tires as raw materials, we deployed the leavening method and obtained carbon materials displaying high specific surface areas and abundant surface functional groups. The obtained adsorbent was investigated in the

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Table 1 The adsorption performance of activated carbon materials from waste tires

Entry	Adsorbent	Initial concentration	Adsorption capacity (mg S per g adsorbent) or sulfur removal	Ref.
1	Acidic and basic treatments of activated C	T (51 ppm), BT (50 ppm) DBT (52 ppm)	0.18(T), 0.50(BT), 1.20(DBT)	38
2	Ce/Fe-loaded rubber tire-derived activated C	T (52 ppm), BT (50 ppm) DBT (51 ppm)	7.78(T), 7.23(BT), 16.63(DBT)	37
3	Rubber tire-derived C	DBT (50 ppm)	8.60	40
4	Nickel/rubber tire-derived activated C	DBT (59 ppm)	96.00%	39
5	Nanocomposite of rubber tire-derived activated C and manganese oxide	T (50 ppm), BT (52 ppm) DBT (53 ppm)	4.50(T), 5.70(BT), 11.40(DBT)	41
6	Cobalt and molybdenum nanoparticles loaded on rubber tire-derived activated C	50 ppm each	2.20(T), 2.32(BT), 2.72(DBT), 3.68(MBT), 3.64(MDBT), 3.72(DMDBT)	42
7	Waste tires to activated C	DBT (150 ppm)	1.22	43
8	Porous C from waste tire leavening	Pentanethiol (30 ppm S)	99.70%	This work

desulfurization of pentanethiol and butanethiol. As expected, the carbon material prepared using the leavening method demonstrated a much enhanced adsorptive desulfurization ability compared to general carbon materials. This result suggests the significance here of randomly opened macropores, high specific surface area and abundant oxygen-containing functional groups of the as-prepared carbon materials. Moreover, this carbon material when optimized achieves very extensive desulfurization at a low concentration of sulfur, indicating promising application prospects for this material.

2. Experimental

2.1. Materials

KOH (AR grade) was purchased from Chengdu Kelong Chemical Reagent Factory. NaHCO_3 (AR grade) was obtained from Sino-pharm Chemical Reagent Co. Pentanethiol (>99.5%) was purchased from TCI (Shanghai) Development Co., Ltd. and butanethiol (>99%) was purchased from Aladdin.

2.2. Preparation of the carbon materials

First, 5.05 g of waste rubber tire particles and 10.2 g of KOH were ground and mixed, and then transferred into 50 mL of distilled water and stirred at 60 °C. The purpose of this process was to remove the surface impurities of the waste tires. The obtained sample was filtered, washed and dried. Then, the resulting black solid was mixed with NaHCO_3 in mass ratios of 1 : 1, 1 : 1.5 and 1 : 2, respectively. Finally, the obtained mixture was calcined at 800 °C for 2 h under a nitrogen atmosphere. The prepared materials were marked as C-1, C-1.5 and C-2 (Fig. 1a).

Similar to the above process, the carbon material was also pre-treated and then calcined, but the calcination process was carried out without NaHCO_3 and marked as C.

2.3. Characterization of the materials

Scanning electron microscopy (SEM) measurements were carried out using a JSM-6010 PLUS/LA scanning electron microscope. Surface areas of the obtained materials were determined using the Brunauer–Emmett–Teller (BET) method based on N_2 adsorption–desorption isotherms collected on a TriStar II 2020 surface area and porosity analyser (Micromeritics Instrument Corporation). Fourier-transform infrared (FT-IR) spectra were acquired using a Nicolet FTIR spectrophotometer (Nexus 470). Raman spectra (Thermo Scientific DXR) were acquired to further characterize the properties of the materials.

2.4. Adsorption experiments

Model fuels with different initial sulfur concentrations (15, 28, 38, and 48 ppm) were each prepared by dissolving an

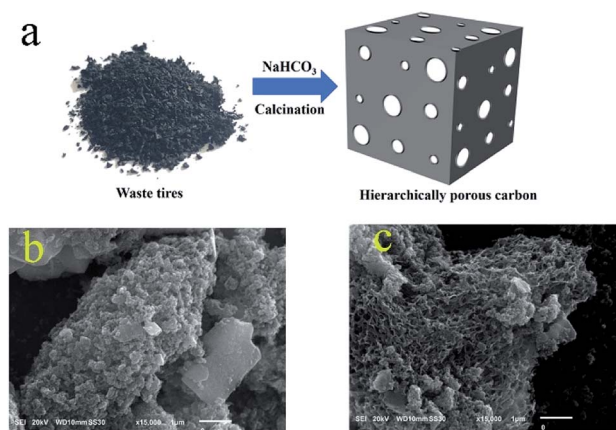


Fig. 1 (a) Schematic of the synthesis of porous carbon. (b and c) SEM images of C and C-1.5.



appropriate amount of pentanethiol in *n*-octane. A similar method was used to prepare a butanethiol model fuel containing 27 ppm S. The adsorption process involved adding 0.1 g of adsorbent and 20 mL of model fuel to a 50 mL conical flask and stirring this mixture at a certain temperature. A gas chromatography-flame photometric detector was used to determine the residual sulfur concentration.

The equilibrium adsorption capacity (q_e) values of the sulfur compounds were determined using the equation

$$q_e = (C_0 - C_t)V/m \quad (1)$$

The equation

$$\text{Sulfur removal (\%)} = (1 - C_t/C_0) \times 100\% \quad (2)$$

was also used. In these equations, q_e , C_0 , C_t , C_e , V and m represent the equilibrium adsorption capacity (mg S per g adsorbent), initial sulfur content (ppm), sulfur content (ppm) at t min, equilibrium sulfur content (ppm), volume of the model fuel (L) and mass of the adsorbent (g), respectively.

3. Results and discussion

3.1. Characterization of the carbon materials

The morphologies of the as-prepared carbon materials were investigated using a scanning electron microscope (SEM).^{47–50} As shown in Fig. 1b, the bulk carbon material obtained by directly calcining waste tires was observed in SEM images to display a rough surface. When NaHCO_3 was mixed with the waste tire and calcined, the obtained carbon material showed many randomly open macropores (Fig. 1c). The presence of macropores was beneficial to increase the specific surface area of the carbon material and expose more interior atoms. These properties contributed to the exposure of the adsorption sites and hence to the promotion of the adsorption performance;⁵¹ moreover, the macropores facilitated mass transfer,⁵² that is, the adsorbents could easily contact the sulfides. Therefore, the carbon material prepared by the “leavening” method may be more favorable for adsorptive desulfurization.

The specific surface areas and pore size distributions of the carbon materials were characterized by acquiring their N_2 adsorption–desorption isotherms (Fig. 2).⁵³ These isotherms each showed a typical II type curve with an H3 hysteresis loop,

indicating the presence of micropores and mesopores in the carbon material. Based on the BET method and quenched solid-state functional theory (QSDFT), the specific surface areas and pore size distributions of the carbon materials were obtained. The specific surface area of C-1.5 ($99.3 \text{ m}^2 \text{ g}^{-1}$) was about three times that of C ($32.8 \text{ m}^2 \text{ g}^{-1}$). This case was distinguished by the addition of NaHCO_3 having generated a large amount of gas (CO_2) during the calcination process and the activator having reacted with carbon intermediates,⁴⁶ resulting in the production of carbon material containing many pores and surface functional groups. The characterization result was consistent with the above-described SEM results. Moreover, according to the literature, a high specific surface area is beneficial for enhancing adsorption performance and macropores and mesopores contribute to mass transfer,⁵⁴ so the carbon material prepared using the leavening method was expected to have the potential to display excellent adsorption performance.

To gain further insight into the surface chemical functional groups of the carbon materials, FT-IR spectra were acquired.^{55–58} As shown in Fig. 3, bands corresponding to a C–OH stretching mode and bending mode were found at 3417 and 1620 cm^{-1} .⁵⁹ Additionally, a band was observed at about 1031 cm^{-1} and was attributed to the infrared vibration of the C–O bond.⁶⁰ A band was also observed at about 1428 cm^{-1} and may have been due to the asymmetric bending vibration of C–H.⁵⁷ These results illustrate that many oxygen atoms are incorporated in the carbon materials prepared using the leavening method.

Raman spectroscopy is the most direct and non-destructive technique used to characterize the structure of carbon materials. Usually, it is employed to characterize the defects, disorder and doping of carbon materials. As shown in Fig. 4, D and G bands are located at about 1340 and 1600 cm^{-1} .⁶¹ It is generally believed that the G band represents the in-plane vibration of the sp^2 carbon atom; and the D band indicates carbon defects.^{62,63} The ratios of the intensity of the D band to that of the G band (I_d/I_g) of C, C-1, C-1.5 and C-2 were found to be 3.02, 2.47, 1.64 and 2.26, respectively. C-1.5 with the lowest value of I_d/I_g indicates that it has the fewest structural defects.

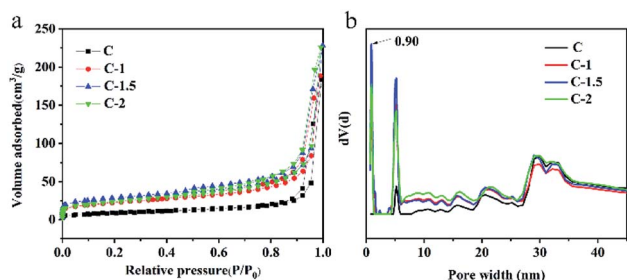


Fig. 2 (a) N_2 adsorption–desorption isotherms and (b) the pore size distribution of C, C-1, C-1.5, and C-2.

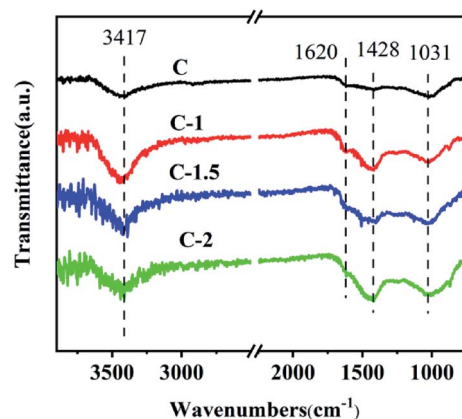


Fig. 3 FT-IR spectra of C, C-1, C-1.5 and C-2.



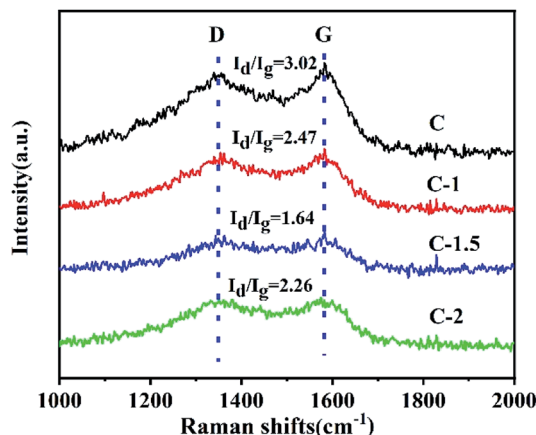


Fig. 4 Raman spectra of C, C-1, C-1.5 and C-2.

3.2. Adsorption experiments

Next, the adsorption performances of various carbon materials, including the carbon material prepared using the NaHCO_3 -assisted leavening method, were determined. The results are given in Fig. 5. C-1.5 removed the most sulfur from pentanethiol, achieving a relatively high adsorption ability. In contrast, when using the same experimental conditions, C showed a relatively poor adsorption ability, with only 23.8% of the sulfur having been removed. The percentages of sulfur removed from pentanethiol by the tested carbon materials followed the trend C-1.5 (99.7%) > C-1 (98.6%) > C-2 (79.9%) > C (23.8%). The excellent performances of the porous adsorbents may be ascribed to the Lewis acid–base derived from oxygen-containing functional groups in the porous carbon materials, strong van der Waals interactions, hierarchically porous structure and high specific surface areas.

Based on the above structural characterization and experimental results, a mechanism for the adsorption was derived. The strong Lewis acid S atoms in a pentanethiol molecule

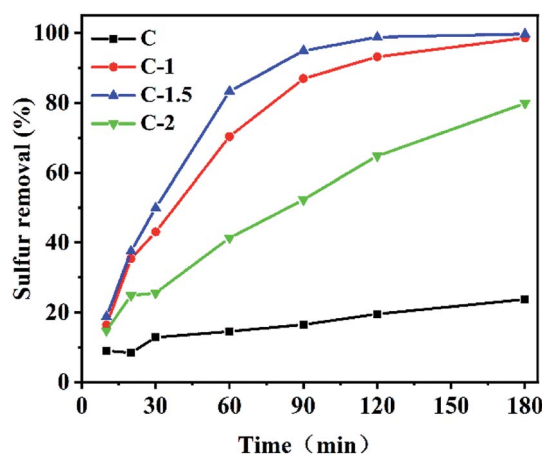


Fig. 5 The percentages of sulfur removed from pentanethiol by various carbon materials. Experimental conditions: 28 ppm initial sulfur concentration, $T = 25^\circ\text{C}$, $V(\text{fuel}) = 20\text{ mL}$, $m(\text{adsorbent}) = 0.1\text{ g}$, shaker agitation speed of 150 rpm and atmospheric pressure.

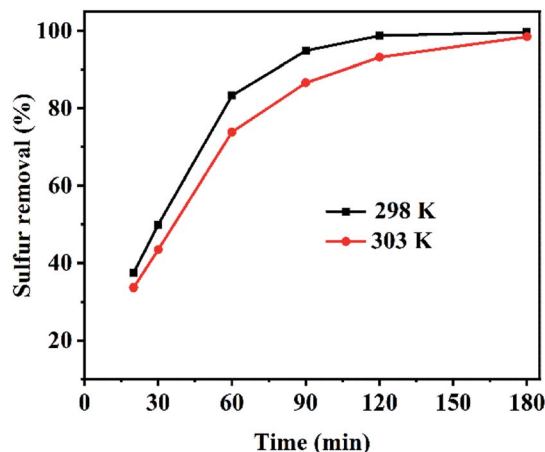


Fig. 6 Effect of temperature on the adsorption of pentanethiol by C-1.5. Experimental conditions: 28 ppm initial sulfur concentration, $V(\text{fuel}) = 20\text{ mL}$, $m(\text{adsorbent}) = 0.1\text{ g}$, shaker agitation speed of 150 rpm and atmospheric pressure.

interacted with a pair of electrons from oxygen atoms in the carbon materials.³¹ The presence of macropores and mesopores contributed to the increased specific surface area of the carbon material, exposed more interior atoms, and facilitated mass transfer so that the adsorbents could easily contact the sulfides.

C-1.5 also removed a very high percentage (98.5%) of sulfur at 303 K, but slightly less so than the 99.7% removal at 298 K (Fig. 6). This small difference can be attributed to the exothermic nature of the adsorption process^{64–67}

Concerning the industrial application prospects of C-1.5, the influence of the initial concentration of sulfur on the adsorption performance was considered.⁶⁸ As shown in Fig. 7, C-1.5 displayed equilibrium adsorption capacities of 3.0, 5.6, 7.6, and 9.6 mg S per g adsorbent for the model fuels containing initial sulfur concentrations of 15, 28, 38, and 48 ppm, respectively.

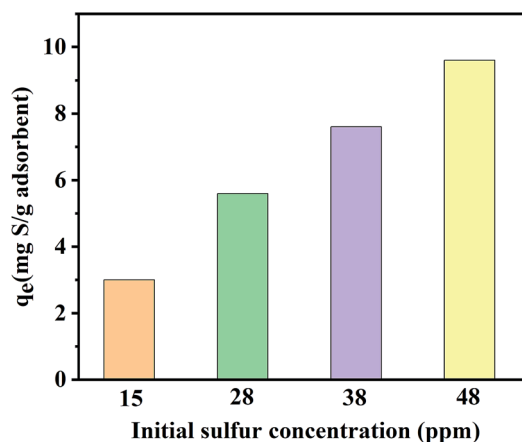


Fig. 7 Effect of the initial sulfur concentration on the adsorption of pentanethiol by C-1.5. Experimental conditions: $T = 25^\circ\text{C}$, $V(\text{fuel}) = 20\text{ mL}$, $m(\text{adsorbent}) = 0.1\text{ g}$, shaker agitation speed of 150 rpm, atmospheric pressure and contact time of 180 minutes.



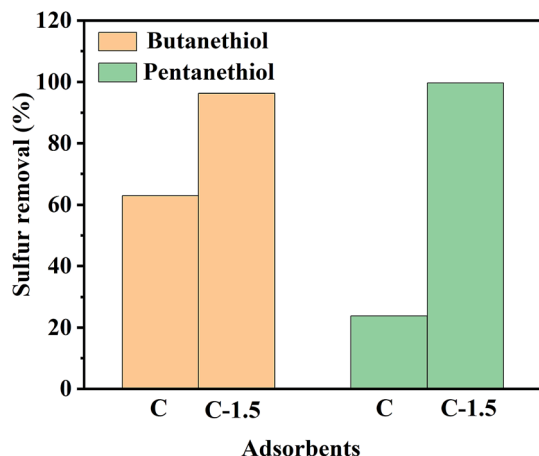


Fig. 8 The sulfur removal (%) of C and C-1.5 for pentanethiol and butanethiol. Experimental conditions: 28 ppm initial sulfur concentration of pentanethiol, 27 ppm initial sulfur concentration of butanethiol, $T = 25\text{ }^{\circ}\text{C}$, $V(\text{fuel}) = 20\text{ mL}$, $m(\text{adsorbent}) = 0.1\text{ g}$, shaker agitation speed of 150 rpm, atmospheric pressure and contact time of 180 min.

It is worth noting that the sulfur removal of the adsorbent for this concentration range was above 97%. This set of results indicates an obvious industrial application value of this material for the removal of low concentrations of sulfide.

To further explore the adsorption ability of the carbon materials for different thiols, butanethiol as well as pentanethiol were used as adsorbates, and the adsorption results are shown in Fig. 8. Most significantly, when C was used as the adsorbent, the percentages of sulfur removed from pentanethiol and butanethiol were 23.8% and 63%, respectively, values much lower than the respective 99.7% and 96.3% values when C-1.5 was used as the adsorbent. As expected, the carbon material synthesized by the leavening method exhibited efficient adsorption.

4. Conclusions

We have developed a “leavening” method to treat used tires. This method was used to turn waste tires into a high-added-value product. The process was found to be efficient, mild, and facile, and may be generally applied to prepare porous carbon materials from other carbon sources. Benefiting from the randomly opened macropores, high specific surface area and abundant oxygen-containing functional groups, the resulting carbon materials exhibited an outstanding adsorptive desulfurization ability, showing a removal of 99.7% of sulfur from pentanethiol, and achieving such a nearly complete desulfurization using mild conditions. The outstanding performance was attributed to macropores and mesopores facilitating mass transfer and the strong Lewis acid S atoms in the pentanethiol molecules interacting with pairs of electrons from oxygen atoms in the carbon materials. Hence, this study may provide the impetus to prepare highly efficient adsorbents for the industrial production of clean fuels.

Conflicts of interest

There are no conflicts to declare.

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