


 Cite this: *RSC Adv.*, 2021, **11**, 4515

Understanding the effect of thiophene sulfur on brominated petroleum coke for elemental mercury capture from flue gases

 Yi Xiao, ^{*a} Xiuyun Liu^a and Yongfa Diao^b

Developing highly efficient and inexpensive adsorbent is a critical technology for elemental Hg removal from the coal combustion flue gases worldwide. Here, we present a novel approach that a waste by-product of petroleum coke containing organic sulfur enhanced bromine binding during the bromine modification process and the brominated petroleum coke increased its mercury adsorption. Experiments and density functional theory reveal that the mercury adsorption capacity directly correlated with the surface organic sulfur and the binding bromine content. Our direct observations and theoretical modeling demonstrate that HgBr and Hg(Br)Br are the primary chemical forms chemisorbed on the surface of this new carbon-based sorbent, which is approaching to bind on the carbon site next to the S atom. The synergetic effect of the inherent thiophene sulfur and loaded bromine enhanced the Hg removal efficiency of the adsorbent.

Received 3rd December 2020

Accepted 3rd January 2021

DOI: 10.1039/d0ra10208c

rsc.li/rsc-advances

1. Introduction

Industrial smog, containing SO₂, NO_x, mercury, particulates, and other trace elements, has become a significant issue to attract worldwide attention.¹ One of the largest sources of smog is the coal combustion, from which elemental Hg is one of the most problematic elements. Because of the volatile and persistent nature of elemental Hg, it is eventually deposited in the lakes and rivers, and bio-accumulated in fish to cause serious threat to the public health by eating contaminated fish. Much work has been conducted to investigate the best method to reduce pollutants; moreover, government regulations were set to reduce the pollutants and the impact on the environment.² Activated carbon (AC)-based sorbents have been proven to be the best approach to remove Hg from flue gases, and they are used currently in many power plants in North America.³ However, alternative sorbents have been studied to replace the activated carbon due to its high price and large demanding quantity to decrease the elemental Hg removal efficiency. Modified low-cost carbon-based sorbents, including fly ash and biomass ash, were developed with the bromine-impregnated and sulfur-impregnated ones showing the best effect to eliminate Hg from the coal combustion flue gas.⁴⁻¹³ Elemental sulfur was reported as a promising “soft” oxidant as a catalyst.¹⁴

Petroleum coke is formed as the final by-product of the petroleum industry during refining of crude bitumen or heavy

oil. It is classified in the European Waste Catalogue with the code 050116,¹⁵ with large quantity produced worldwide. The certain technological reuse of the waste has globally become a problem, a portion of which has been used as graphite precursors, electrodes in aluminum, steel plants and even as inferior fuel. The high sulfur content limited the reuse application of petroleum coke. To date, certain researchers have used it as a precursor for preparing high surface area activated carbon because it is inexpensive and contains high carbon content.¹⁶⁻²⁰

It was reported that organic sulfur, nitrogen, and oxygen were mainly in the forms of thiophene-containing polycyclic aromatic hydrocarbon, polycyclic aromatic hydrocarbons containing pyrrole or pyridine, and C=O or C—O—C, respectively in the structure of petroleum coke.²¹ Furthermore, >90% of sulfur was reported to be thiophene sulfur in the contents of petroleum coke.²¹ In our previous studies, petroleum coke has been obtained as a useful precursor to be modified with bromine for the adsorption of Hg from the flue gas.²² Both mercury reaction chemistry and macroscopic dynamic mechanism were studied, from where the synergetic effect of the bonded bromine and inherent organic sulfur was inferred for enhancing Hg adsorption.^{23,24} However, the micro-mechanism of the organic thiophene-sulfur affecting the bromine binding and elemental mercury binding continues to be unclear.

In this study, the petroleum coke material containing 5.85 wt% sulfur was obtained from the upgrading of oil sands deposits in Alberta, Canada, to make brominated carbon-based adsorbent for elemental Hg removal from the coal combustion flue gas.²³ The bromine and Hg binding chemistry is analyzed through XRD, FTIR and XPS, thermal stability of Br and Hg in

^aShanghai Vocational College of Agriculture and Forestry, 658 Zhongshan Road, Shanghai 201620, P. R. China. E-mail: xiaoy@shafc.edu.cn

^bSchool of Environmental Science and Engineering, Donghua University, 2999 North Renmin Road, Shanghai 201620, P. R. China



the adsorbent is performed to secure the potential regular correlation of them; however, the reaction pathways and mechanisms, nature of the active petroleum coke sites and the effects of thiophene-sulfur were performed by first-principles density functional theory (DFT) calculations to help understand the significant role of thiophene-sulfur in the petroleum coke for bromine-loading and elemental Hg capture.

2. Experimental

2.1. Materials preparation

The brominated petroleum coke adsorbent was prepared with a detailed description in our previous papers.^{22,23} The bromine modification process is known as the chemical-mechanical bromination process. The obtained petroleum coke was grounded using a ball-mill and sieved with a 100-mesh sieve, and then dried in a vacuum oven at 110 °C overnight. The prepared raw petroleum coke and liquid bromine with the weight ratio of 100 : 1 were placed in a sealed container, which was then tumbled for 30 min. After that, the received sample was placed in a vacuum oven at 200 °C for 30 min, and named as Br-PC.

Brominated petroleum coke (Br-PC) was heated to different temperatures of 200, 400, 600, 800 and 950 °C for 30 min under a nitrogen flow of 60 mL min⁻¹. The resulting thermal treated samples were saturated with elemental Hg to investigate the mercury adsorption capacity through the mercury saturation procedure. The thermally treated samples were spread over the bottom of a glass vessel, with a smaller glass container containing elemental Hg being placed in the center of the vessel. The glass vessel should be well sealed and set in a fume hood. After a week, the Hg-saturated sorbents were tested by XPS to secure the adsorption mechanism.

2.2. Sorbent characterization

The surface morphology of the petroleum coke before and after the bromine loading was observed using a scanning electron microscope (SEM, Hitachi S-4800, Japan). The surface functional groups were measured by a X-ray diffractometer (XRD, Bruker AXS) and Fourier transform infrared spectrometer (FTIR, Nicolet iS5) produced by Thermo Electron Corporation. The XRD was equipped with a graphite monochromator filter, with the scanning area of 0.65–140°, the minimum wavelength step is 0.0001°. The spectral range of FTIR was 7800–350 cm⁻¹, the spectral resolution is better than 0.5 cm⁻¹, the wave number accuracy is better than 0.01 cm⁻¹, and the signal-to-noise ratio is 40 000 : 1. The surface binding and elemental speciation was analyzed by X-ray photoelectron spectroscopy (XPS) using a Thermo ESCALAB 250XI X-ray photoelectron spectrometer, with the surface excitation at 1486.6 eV by an Al Ka X-ray source. The survey and high-resolution spectra of C_{1s}, Br_{3d}, S_{2p}, Si_{2p} and Hg_{4f} were collected and calibrated with the binding energy of C_{1s} at 284.9 eV. All the survey and high-resolution scans were analyzed with Thermo Avantage XPS software for surface chemical analysis.

2.3. DFT calculations

The calculation was performed using the Gaussian 16 suite program. The optimizations of the complex structures were performed by using B3LYP DFT. The LAN2DZ basis set was used to treat the mercury atom, whereas the 6-31G(d) basis set was used to treat all other atoms of C, H, S and Br. To identify the minimum point on the potential energy surface, set charge as 0; spin as 1, 3, 5 and 7 in the Gaussian 16 software; and explore the spin value when the system energy is the lowest.

3. Results and discussion

3.1. Surface characterization

Surface characterization of the raw and brominated petroleum cokes was analyzed by SEM and XPS (Fig. 1). It was revealed that Br was successfully loaded on the surface of raw petroleum coke because the surface was much rougher and Br peaks appeared as well as no change of sulfur peaks. Strong S_{2s} and S_{2p} peaks are obtained in Fig. 1a. The peak at 164 eV is detected as S-containing complexes such as thiophene or thiols, which are reported to be possibly effective for the elemental Hg adsorption from the coal-fired flue gases.²⁵ O_{1s} peak at 532 eV is attributed to the formation of stable R-O-Hg complexes on the petroleum coke surface. The Br peaks of Br_{3s}, Br_{3p}, and Br_{3d} can be obtained from Fig. 1b. It was reported that the strong peak at 70.0 ± 0.2 eV of Br_{3d(5/2)} was bonded to sp² and sp³ hybrid carbon atoms,²⁶ which has a high potential for elemental Hg adsorption.

The XRD and FTIR patterns of raw and brominated petroleum cokes shown in Fig. 2 suggest that the prepared petroleum coke sorbent shows classical diffraction peaks and changes in the functional groups. Generally, we can distinguish whether

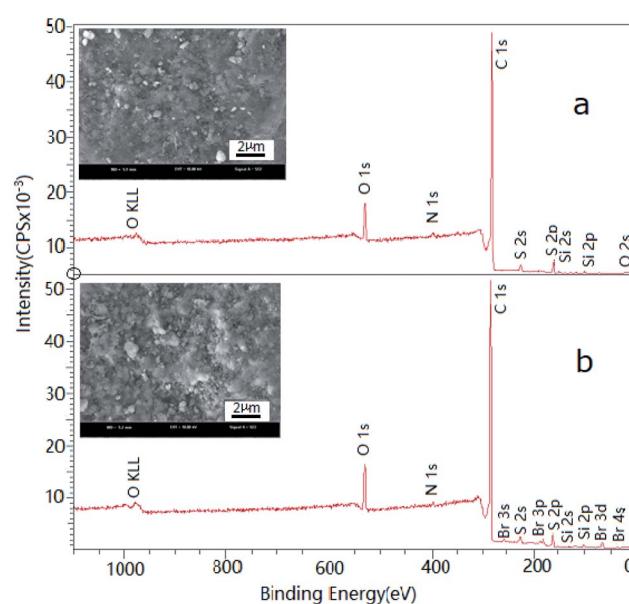


Fig. 1 XPS survey scan and SEM images for the petroleum cokes before and after bromine modification. (a) Raw petroleum coke (b) brominated petroleum coke.



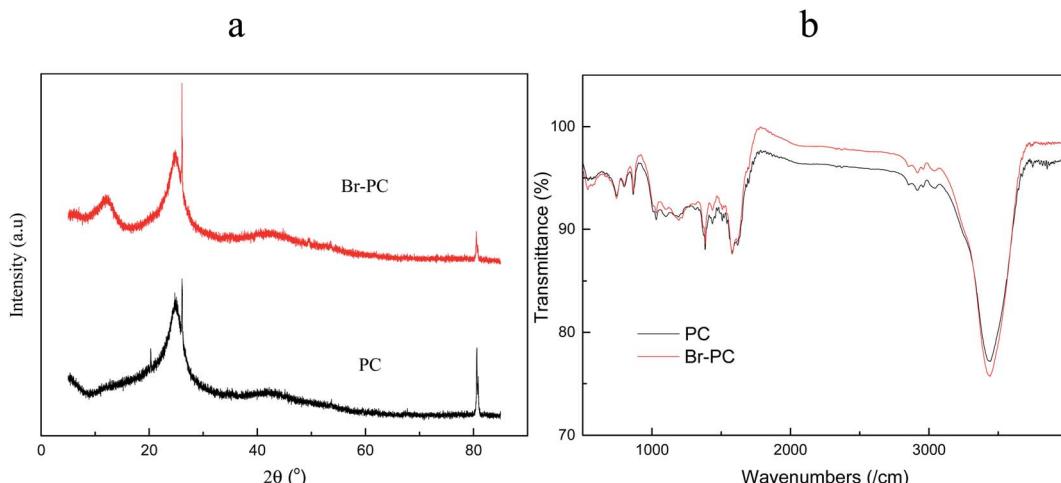


Fig. 2 Sorbent characterization. (a) XRD analysis for petroleum coke before and after bromine modification (PC and Br-PC). (b) FTIR analysis for petroleum coke before and after bromine modification (PC and Br-PC).

the substance is crystalline or amorphous from the XRD spectrum by classifying whether the peaks are rich of fine peak structures or sharp peak structures. The petroleum coke is amorphous as revealed from Fig. 2a since a big fine peak appeared at 2θ of 26° . After modification of the petroleum coke by bromine, a new fine peak appeared at 2θ of 13° , which indicated that the former structure is not changed, but other form of amorphous structure developed on the sorbent surface. The existence of big fine peaks in Fig. 2a shows that Br is primarily adsorbed on the surface of petroleum coke in the form of chemical loading, but not in the crystal form of Br molecules.

Fourier transform infrared spectroscopy (FTIR) is commonly used to analyze the properties of functional groups on the surface of carbon-based adsorbents. The infrared absorption spectra of the petroleum coke before and after modification are illustrated in Fig. 2b. It can be observed in the figure that the types of functional groups on the surface of petroleum coke do not obviously change, but the number of functional groups changed during the bromine loading process.

The absorption peak at about 864 cm^{-1} is the stretching vibration peak of C-H, and the peak shape indicates that the C-H bond of the original petroleum coke breaks under the action of bromine, and some hydrogen is derived from the adsorbent. New stretching vibration between 2917 cm^{-1} and 3040 cm^{-1} , 1030 and 1384 cm^{-1} and 864 and 1192 cm^{-1} regions indicate the appearance of new functional groups such as C-Br after bromine loading. It is well known that Br_2 is a potent oxidant, and it is easy to open the edges of the graphite lamellar to facilitate the entry of Br molecules; therefore, the intercalation reaction begins at the edge of the graphite lamellar.²⁷

The absorption peak at 744 cm^{-1} is the stretching vibration peak of heterocyclic thiophene,²⁸ but the peak shape of thiophene does not change obviously, indicating that the loading of Br does not change the thiophene-sulfur functional groups in petroleum coke.

3.2 Thermal stability

To further secure the mechanism of mercury adsorption on the brominated petroleum coke and the synergistic effect of thiophene-sulfur and bonded bromine, high resolution XPS spectra over C_{1s} , O_{1s} , Br_{3d} , S_{2p} and $\text{Si}_{2p}/\text{Hg}_{4f}$ regions were recorded and analyzed for all the steps of heating of the brominated petroleum cokes after Hg adsorption. XPS spectra (Fig. 3) show changes when the temperature is increasing. However, the peak at room temperature is basically the same as that at $200\text{ }^\circ\text{C}$, which is primarily due to the sample preparing process, at which all the prepared samples were vacuum dried at $200\text{ }^\circ\text{C}$ for 30 min to let the unstable bonded bromine volatile before applying for Hg adsorption to ensure the samples are safe to deliver or test.²² Since Hg species such as HgBr_2 , Hg_2Br_2 and HgO are located at the binding energy between 100.7 eV and 104 eV , which overlap with the Si_{2p} peaks, it is difficult to distinguish the Hg peaks from the XPS analysis. Much important information can be detected from the spectra *via* temperature changes.²⁹ Notably, the Br_{3d} and $\text{Si}_{2p}/\text{Hg}_{4f}$ binding energy, which are sensitive to temperatures, indicate its un-stability in the sorbent. However, certain peaks of bromine and mercury still exist even for a temperature up to $950\text{ }^\circ\text{C}$.

The main peak of C_{1s} is sp^2/sp^3 hybrid carbon and the peak of S_{2p} is inherent organic sulfur such as thiophene and thion. Peaks of C_{1s} and S_{2p} are almost unchanged during the thermal treatment of the samples, indicating the C-C and C-S are difficult to break even for temperature up to $1000\text{ }^\circ\text{C}$.²¹

However, the number and the area of peaks of Br_{3d} decline with increasing temperature. According to the previous studies, the $\text{Br}_{3d\ 5/2}$ peak at $67.4\ (\pm 0.2)\text{ eV}$ and $70.0\ (\pm 0.2)\text{ eV}$ were obtained as $\text{C}_n\text{-Br}_2$ and $\text{C}_n\text{-Br}$ surface complexes.^{24,30} The Br_{3d} peaks ranging between 72 and 76 eV was attributed to the bonded bromine on the carbon near the S atoms, since hydrogen on the thiophene rings can be easily replaced to form the derivatives of 2-bromothiophene.³¹ The surface complexes of $\text{C}_n\text{-Br}_2$ and $\text{C}_n\text{-Br}$ are sharply decreased; however, bromine



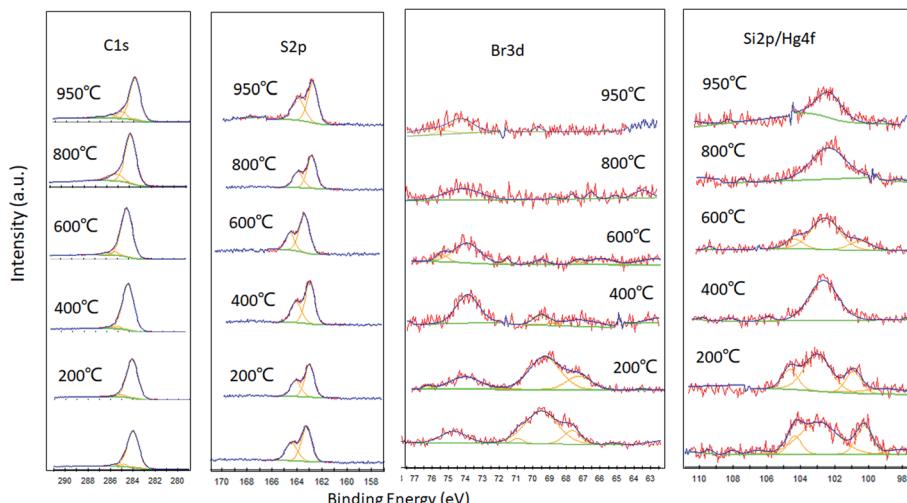


Fig. 3 Temperature dependent X-ray photoelectron spectra for brominated petroleum coke after mercury capture. All heating steps were performed for 30 min in a N_2 environment, with flow rate of 60 mL min^{-1} from 25°C (room temperature) to 950°C at a $10^\circ\text{C min}^{-1}$ heating rate.

near the sulfur atoms increased for the increase of temperature to 400°C for 30 min; the surface complexes of $\text{C}_n\text{-Br}_2$ and $\text{C}_n\text{-Br}$ are further decreased, with no peaks detected at 67 and 70 eV, but strong peaks between 72 and 76 eV were seen when the sample was heated at 600°C for 30 min. This indicates that bromine is almost volatilized from the surface of petroleum coke, but still strongly bonds with the carbon nearest to the S atom. The above results are consistent with the conclusion obtained from the XRF test in our previous study.²² When the temperature was increased up to 800°C , the peak at 74 eV decreased, and is not obviously changed at 950°C . In this situation, a certain amount of bromine is still binding to the surface of the sorbent, which has the potential ability for mercury capture.

Similarly, the peaks of $\text{Si}_{2p}/\text{Hg}_{4f}$ decreased with increasing temperature, and the changing trend is almost the same as Br_{3d} . The peak at 102.94 eV is almost constant for all the thermal treated samples, which should present the Si_{2p} and part of the oxidized mercury. The existing form is predicted to be HgBr or $\text{Hg}(\text{Br})\text{Br}$ binding on the carbon atom nearest to the sulfur atoms. Other peaks at 98, 100.24, 104.35, 106.55 and 108.33 eV are Hg compounds of HgBr or $\text{Hg}(\text{Br})\text{Br}$, which are not stable when high temperature applied.

3.3 Bromination model

First-principles density functional theory (DFT) calculations were carried out to help understand the reaction pathways, mechanisms and the active binding sites on the sorbent for bromine modification and mercury adsorption on the petroleum coke sorbent.

Details concerning the computational methods used were described, as well as the results for the initial modification of petroleum coke. The coupling of bromination intermediates was explored on the surface. Detailed structural optimizations were carried out on the petroleum coke model and over a range of bromine coverages to establish the lowest energy structures

of the surfaces. The results for the model surface were used to determine the most favorable surface termination at the specific bromine chemical potentials.

The structure of petroleum coke and the sulfur form existing in the petroleum coke was studied by XPS and the computer models. Cai *et al.*³² reported that the sulfur structure in the petroleum coke was mainly organic sulfur, such as thiophene or similar thiophene rings. Xiao *et al.*³³ outlined that the thiophene rings were on the outmost layer of the petroleum coke molecule, with at least three aromatic rings. Moreover, the inner layer of the petroleum coke molecule was aromatic; this configuration was regarded as graphite-like, and the carbon atoms were dominated by sp^2 hybridization. Some small molecule fragments were around the outmost layer, including sulfur, oxygen, and nitrogen-containing fragments. Many studies have supported the zigzag carbene edge structure model for the carbon-based adsorbents on halogen elements modification, on which surface the active sites at the zigzag edge were reacted with the acid gas components, and new active sites were created for the elemental mercury oxidation.³⁴⁻³⁸ In our study, the zigzag carbene edge structure model was used as a precursor, with a thiophene-sulfur implanted in the model, to investigate the stability of bromine binding ability on the model (Fig. 4a and b).

Hg atoms adsorbed by petroleum coke are directly above the molecular plane of the pure petroleum coke, and the adsorption energy is $-19.2 \text{ kJ mol}^{-1}$, which belongs to weak physical adsorption. Bromine is a strongly oxidizing gas, which can easily open the edge of graphite lamellae, leading to the intercalation reaction and the carbon atoms becoming sp^3 hybridized.³⁹ In addition, the H near the S atom easily reacts with Br, due to the aromaticity of the molecule, which enhanced bromine binding on the sorbent. The above FTIR data also verified the breaking of C-H bond during the modification process, since the stretching peak of the C-H bond in the wavelength range of 3200 cm^{-1} to 2700 cm^{-1} decreased after bromination (Fig. 2). The intercalation reaction and the



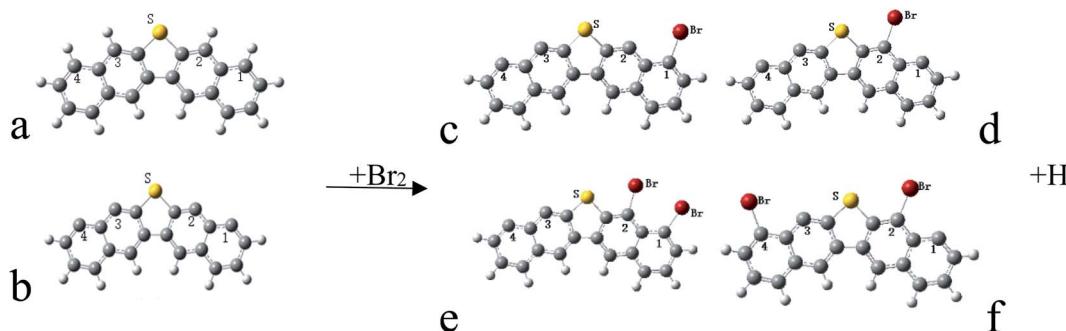


Fig. 4 Chemical reaction equation for stable models of the Br atoms at different sites of petroleum coke molecule. (a and b) Stable petroleum coke molecules. (c) Br at next nearest of the S atom. (d) Br at the nearest of S atom. (e) Two Br atoms at the same side of the S atom. (f) Two Br atoms at the different side of the S atom.

aromaticity of the petroleum coke molecule enhanced the activity of the adsorption site on the surface of petroleum coke and may produce more active sites for mercury capture, as shown in the chemical equation in Fig. 4.

The system total energy (STE) and the specific bond Mulliken population (BMP) are the important parameters to describe the stability of the model in DFT calculations; the lower the STE and higher the specific BMP, more stable is the model. Fig. 4c and d are the stable models of one Br atom bonded at different sites of the petroleum coke molecule. The STE of single bromine binding at carbon position 2 is 2.63 kJ mol⁻¹ higher compared with position 1, and the C-Br BMP in position 1 is 0.2736 while it is 0.2954 in position 2. Both results show that Br tends to

adsorb on the top of the carbon site nearest to the S. Fig. 4e and f are the stable models of two Br atoms binding at different sites of the petroleum coke molecule. The STE of the model of Fig. 4e is 42 kJ mol⁻¹ higher than Fig. 4f, which indicates that Br is more inclined to adsorb on top of the relative carbon site close to S, thus forming a V-shaped structure.

3.4 Elemental mercury adsorption model

To explain more specifically what our study is expected to reveal, we illustrate the reaction process of Fig. 5 and 6 of the most relevant elemental mercury reaction on the brominated petroleum coke surface. The obtained stable mono-bromine and bi-bromine models are designed as reactants to react with the

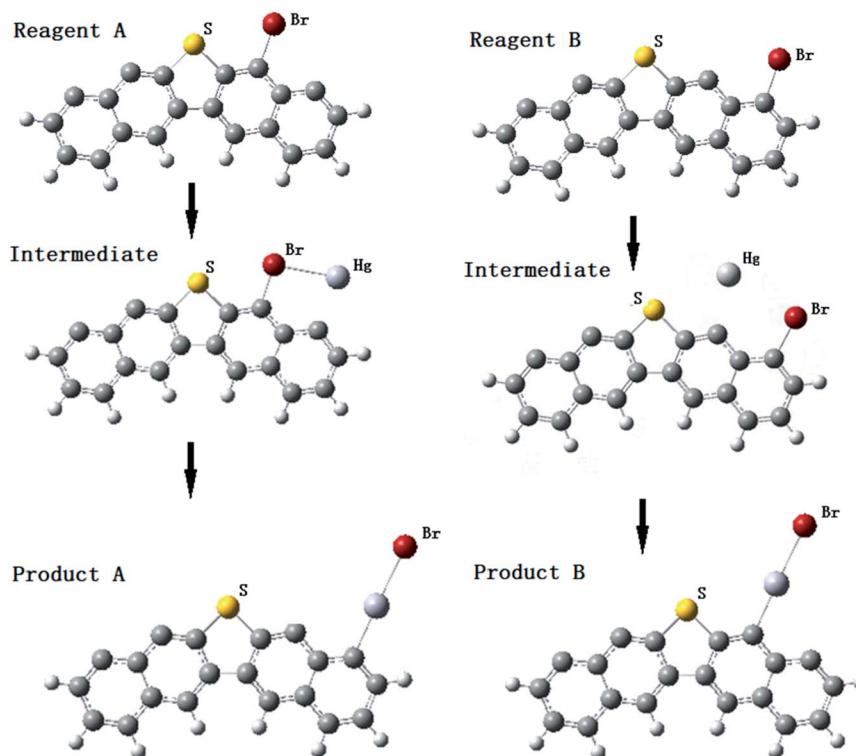


Fig. 5 Hg⁰ adsorption process on brominated petroleum coke model with one Br atom at different carbon sites near sulfur.



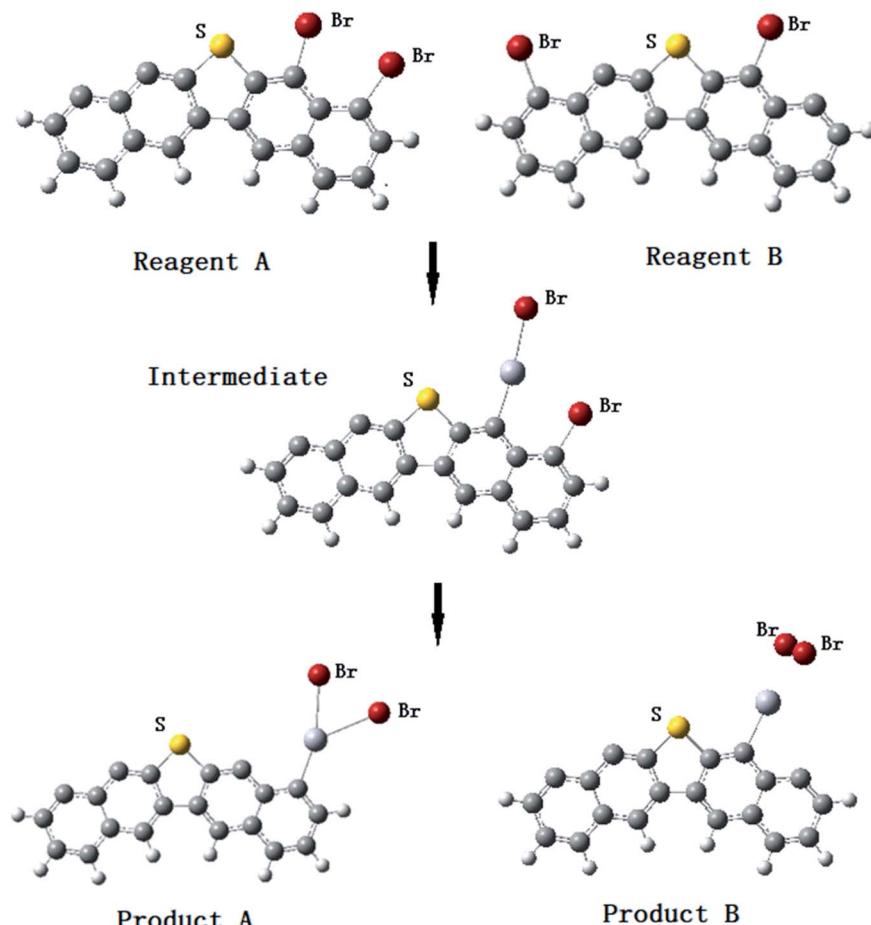


Fig. 6 Hg^0 adsorption process on brominated petroleum coke model with two Br atoms at different carbon sites near sulfur.

elemental Hg, the reacting processes are shown in Fig. 5 and 6. The intermediates and ultimate products of elemental Hg adsorbed on the surface model of brominated petroleum coke were obtained.

As shown in Fig. 5, the STE of product B is $14.26 \text{ kJ mol}^{-1}$ lower than product A, and the C–Hg BMP of product A is 0.300 and 0.306 for product B. Both the results indicate product B is more stable than product A. In conclusion, HgBr_2 is inclined to bind on the adjacent carbon site nearest to the S atom. These results agree with the XPS spectra shown in Fig. 3.

As shown in Fig. 6, both the reactant A and reactant B form stable final products. The STE of product B is 4.73 kJ mol^{-1} lower than product A, while the C–Hg BMP of product A is 0.315 and 0.298 for product B. Comparing the STE and the C–Hg BMP of the two products, product A is more stable than product B, which indicates that HgBr_2 can be formed to bind to the carbon site nearest to the S atom.

Two mercury oxidized substances, HgBr and HgBr_2 , are formed on the surface of bromine-modified adsorbents as found by various researchers.^{40–42} However, compared the models of product B– HgBr and – HgBr_2 , the adsorption energies are about $-472.05 \text{ kJ mol}^{-1}$ and $-221.92 \text{ kJ mol}^{-1}$ separately. From the sight of the chemical reaction energy barrier, the activation energy required for HgBr_2 is quite higher than HgBr .

Thus, the results show that the main form of Hg^0 on the surface of brominated carbon-based adsorbents is HgBr , and HgBr_2 is usually decomposed into Br atom and the compound HgBr on the surface of the adsorbent, the chemical form of which is usually described as $\text{Hg}(\text{Br})\text{Br}$. Brominated petroleum coke adsorbent is a typical carbon-based adsorbent, the adsorption of HgBr_2 on the surface of petroleum coke is also unstable, which agreed with the results of other researchers.

4. Conclusions

The results reported demonstrate that high sulfur-containing petroleum coke is a potential carbon-based sorbent for bromine modification to capture elemental Hg from the coal combustion flue gases to reduce the threat of Hg to the environment. The important effect and mechanism of thiophene-sulfur in the petroleum coke on the binding of Br and Hg adsorption are revealed by XPS and DFT calculation. Bromine intercalation reaction and the aromaticity of thiophene are possibly leading to the C–H bond cleavage and *ortho*-substitution reaction, which was revealed from XRD and FTIR analysis. These features can enhance the Br loading during the modification process, and improve the Hg removal efficiency. The amount of Br reduced when the adsorbed Hg was decreased,



which was revealed from the thermal treatment of the resulting samples. However, even for temperatures up to 600 °C, a portion of bromine is still stably bound to the carbon site nearest to the S atom, indicating the key effect of thiophene sulfur in the sorbent. The synergistic effect of inherent thiophene sulfur and the loaded bromine enhanced the Hg removal efficiency of the adsorbent.

DFT study showed that the Br atom was always likely to bind on the C atom nearest to the S atom, *i.e.*, thiophene sulfur can significantly promote the loading of bromine on the surface of petroleum coke. When Hg^0 reacts on the surface of the brominated petroleum coke, $HgBr_2$ and $HgBr$ are formed, which are both inclined to bind to the C atom nearest to the S atom. Moreover, $HgBr_2$ is easily decomposed into $HgBr$ and Br atoms adsorbing on the surface of petroleum coke independently, which is usually described as $Hg(Br)Br$. The synergistic effect of S and Br provides more active sites for the elemental mercury adsorption.

In conclusion, the ability of petroleum coke to adsorb Hg reveals major dependence on the inherent concentration of the organic sulfur and the bonded bromine. The higher the content of organic sulfur, the higher the concentration of effective bromine loaded on the surface and better the mercury adsorption capacity of the brominated petroleum coke. Thus, this study illustrates that high-sulfur petroleum coke is a potential material as a precursor in applying for the elemental mercury removal from the flue gases, corresponding to the bromine modification. Thiophene is predicted to be a possible modifier to impregnate with the carbon-based materials for elemental Hg capture.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work is supported by the National Fundamental Research Funds for the Central Universities (CUSF-DH-D-2017097), Shanghai New Teacher Training Program (ZZNL20012) and Scientific Research Project of Shanghai Vocational College of Agriculture and Forestry (KY2-0000-20-04). Professor Zhenghe Xu in University of Alberta (Alberta, Canada) is highly appreciated for supporting the experimental materials preparation.

References

- 1 Environmental impacts of coal power: air pollution, http://www.ucsusa.org/clean_energy/coalvswind/c02c.html.
- 2 Clean Air Act homepage, http://www.ec.gc.ca/cleanair-airpur/clean_air_act-WS1CA709C8-1_en.htm.
- 3 Y. Liu, T. M. Bisson, H. Yang and Z. Xu, Recent developments in novel sorbents for flue gas clean up, *Fuel Process. Technol.*, 2010, **91**(10), 1175–1197.
- 4 E. Sasmaz, A. Kirchofer, A. D. Jew and A. Saha, Mercury chemistry on brominated activated carbon, *Fuel*, 2012, **99**(9), 188–196.
- 5 E. C. Rupp and J. Wilcox, Mercury chemistry of brominated activated carbons – packed-bed breakthrough experiments, *Fuel*, 2014, **117**(1), 351–353.
- 6 A. Saha, Investigation of changes in surface chemistries on virgin and brominated activated carbon sorbents during mercury capture: before and after regeneration, *Energy Fuels*, 2014, **28**(6), 4021–4027.
- 7 T. M. Bisson, L. C. Maclean, Y. Hu and Z. Xu, Characterization of mercury binding onto a novel brominated biomass ash sorbent by x-ray absorption spectroscopy, *Environ. Sci. Technol.*, 2012, **46**(21), 12186–12193.
- 8 D. Karatza, A. Lancia, D. Musmarra and C. Zucchini, Study of mercury absorption and desorption on sulfur impregnated carbon, *Exp. Therm. Fluid Sci.*, 2000, **21**(1), 150–155.
- 9 K. S. Reddy, A. A. Shoaibi and C. Srinivasakannan, Elemental mercury adsorption on sulfur-impregnated porous carbon - a review, *Environ. Technol.*, 2014, **35**(1), 18–26.
- 10 R. D. Vidic, M. T. Chang and R. C. Thurnau, Kinetics of vapor-phase mercury uptake by virgin and sulfur-impregnated activated carbons, *J. Air Waste Manage. Assoc.*, 1998, **48**(3), 247–255.
- 11 S. Vitolo and M. Seggiani, Mercury removal from geothermal exhaust gas by sulfur-impregnated and virgin activated carbons, *Geothermics*, 2002, **31**(4), 431–442.
- 12 S. V. Krishnan, B. K. Gullett and W. Jozewicz, Sorption of elemental mercury by activated carbons, *Environ. Sci. Technol.*, 1994, **28**(8), 1506–1512.
- 13 Y. Otani, C. Kanaoka, H. Emi, I. Uchijima and H. Nishino, Removal of mercury vapor from air with sulfur-impregnated adsorbents, *Environ. Sci. Technol.*, 1988, **22**(6), 708–711.
- 14 Q. Zhu, S. L. Wegener, C. Xie, O. Uche, M. Neurock and T. J. Marks, Sulfur as a selective 'soft' oxidant for catalytic methane conversion probed by experiment and theory, *Nat. Chem.*, 2013, **5**(2), 104–109.
- 15 J. Olmeda, M. Frías, M. Olaya, B. Frutos and M. I. S. D. Rojas, Recycling petroleum coke in blended cement mortar to produce lightweight material for impact noise reduction, *Cem. Concr. Compos.*, 2012, **34**(10), 1194–1201.
- 16 C. Almansa, M. Molina-Sabio and F. Rodríguez-Reinoso, Adsorption of methane into $ZnCl_2$ -activated carbon derived discs, *Microporous Mesoporous Mater.*, 2004, **76**(1), 185–191.
- 17 A. Perrin, A. Celzard, A. Albinia, M. Jasienko-Halat, J. F. Marêché and G. Furdin, NaOH activation of anthracites: effect of hydroxide content on pore textures and methane storage ability, *Microporous Mesoporous Mater.*, 2005, **81**(1), 31–40.
- 18 M. Molina-Sabio and F. Rodríguez-Reinoso, Role of chemical activation in the development of carbon porosity, *Colloids Surf. A*, 2004, **241**(1), 15–25.
- 19 M. A. Lillo-Ródenas, D. Cazorla-Amorós, A. Linares-Solano, F. Béguin, C. Clinard and J. N. Rouzaud, HRTEM study of activated carbons prepared by alkali hydroxide activation of anthracite, *Carbon*, 2004, **42**(7), 1305–1310.
- 20 B. Jiang, Y. Zhang, J. Zhou, K. Zhang and S. Chen, Effects of chemical modification of petroleum cokes on the properties



of the resulting activated carbon, *Fuel*, 2008, **87**(10–11), 1844–1848.

21 X. Jin, Q. Zhong, F. Li, J. D. Huang, Y. Zhang and B. Wang, Modeling the change of green coke to calcined coke by using qingdao high-sulfur petroleum coke, *Energy Fuels*, 2015, **29**(5), 1504–1516.

22 X. Yi, D. Pudasainee, R. Gupta, Z. Xu and Y. Diao, Bromination of petroleum coke for elemental mercury capture, *J. Hazard. Mater.*, 2017, **336**, 232–239.

23 Y. Xiao, D. Pudasainee, R. Gupta, Z. Xu and Y. Diao, Elemental mercury reaction chemistry on brominated petroleum cokes, *Carbon*, 2017, **124**, 89–96.

24 Y. Xiao, Y. Diao, L. Tian, Y. Lu and L. Zhan, Kinetic mechanism of novel brominated petroleum coke on trace Hg₀ removal from flue gas, *China Environ. Sci.*, 2018, **38**(2), 508–515.

25 H. Hsi, M. J. Rood, M. Rostamabadi, A. Shiaoqiu Chen and R. Chang, Effects of sulfur impregnation temperature on the properties and mercury adsorption capacities of activated carbon fibers (acfs), *Environ. Sci. Technol.*, 2001, **35**(13), 2785–2791.

26 N. D. Hutson, B. C. Attwood and K. G. Scheckel, XAS and XPS characterization of mercury binding on brominated activated carbon, *Environ. Sci. Technol.*, 2007, **41**(5), 1747–1752.

27 J. Hwang, J. P. Carbotte, S. Tongay, A. F. Hebard and D. B. Tanner, Ultrapure multilayer graphene in bromine-intercalated graphite, *Phys. Rev. B*, 2012, **84**(4), 401–410.

28 P. J. Zhao, C. Ma, J. T. Wang, *et al.* Almost total desulfurization of high-sulfur petroleum coke by Na₂CO₃-promoted calcination combined with ultrasonic-assisted chemical oxidation, *Carbon*, 2019, **145**, 774.

29 J. Wilcox, E. Sasmaz, A. Kirchofer and S.-S. Lee, Heterogeneous mercury reaction chemistry on activated carbon, *J. Air Waste Manage. Assoc.*, 2011, **61**(4), 418.

30 E. Papirer, R. Lacroix, J. B. Donnet, G. Nanse and P. Fioux, XPS study of the halogenation of carbon black-part 1. bromination, *Carbon*, 1994, **32**(7), 1341–1358.

31 A. K. Shigemoto, C. N. Virca, S. J. Underwood, L. R. Shetterly and T. M. McCormick, Thiophene-based fluorescent mercury-sensors, *J. Coord. Chem.*, 2016, **69**(11–13), 1–21.

32 J. H. Cai, E. Morris and C. Q. Jia, Sulfur speciation in fluid coke and its activation products using k-edge x-ray absorption near edge structure spectroscopy, *Sulfur Rep.*, 2009, **30**(6), 555–569.

33 X. Jin, Q. Zhong, F. Li, J. D. Huang, Y. Zhang and B. Wang, Modeling the change of green coke to calcined coke by using qingdao high-sulfur petroleum coke, *Energy Fuels*, 2015, **29**(5), 1504–1516.

34 C. Rungnim, V. Promarak, S. Hannongbua, N. Kungwan and S. Namuangruk, Complete reaction mechanisms of mercury oxidation on halogenated activated carbon, *J. Hazard. Mater.*, 2016, **310**, 253–260.

35 L. R. Radovic, The mechanism of co chemisorption on zigzag carbon active sites: a computational chemistry study, *Carbon*, 2005, **43**(5), 907–915.

36 X. Sun, J. Y. Hwang and S. Xie, Density functional study of elemental mercury adsorption on surfactants, *Fuel*, 2011, **90**(3), 1061–1068.

37 F. H. Yang and R. T. Yang, Ab initio molecular orbital study of adsorption of atomic hydrogen on graphite: insight into hydrogen storage in carbon nanotubes, *Carbon*, 2002, **40**(3), 437–444.

38 B. Padak and J. L. Wilcox, Understanding the Binding Mechanism of Mercury on Activated Carbon, *Carbon*, 2008, **47**(12), 2855–2864.

39 Y. Gao, J. Gu, W. Shen and F. Kang, Bromine-graphite intercalation compounds, *Carbon Technol.*, 2000, **4**, 21–25.

40 E. Sasmaz, A. Kirchofer, A. D. Jew and A. Saha, Mercury chemistry on brominated activated carbon, *Fuel*, 2012, **99**(9), 188–196.

41 C. Rungnim, V. Promarak, S. Hannongbua, N. Kungwan and S. Namuangruk, Complete reaction mechanisms of mercury oxidation on halogenated activated carbon, *J. Hazard. Mater.*, 2016, **310**, 253–260.

42 J. Liu, W. Qu and C. Zheng, Theoretical studies of mercury-bromine species adsorption mechanism on carbonaceous surface, *Proc. Combust. Inst.*, 2013, **34**(2), 2811–2819.

