


Cite this: *RSC Adv.*, 2022, 12, 19955

Density functional theory analysis for H₂S adsorption on pyridinic N- and oxidized N-doped graphenes

Takaya Fujisaki,^a Kei Ikeda,^b Aleksandar Tsekov Staykov,^c Hendrik Setiawan^d and Yusuke Shiratori^e

Biomass discharged from primary industries can be converted into methane by fermentation. This methane is used for generating electricity with solid oxide fuel cells (SOFCs). This methane fermentation provides H₂S, which reduces the efficiency of SOFCs even at a level as low as a few parts per million. It has been experimentally reported that a nitrogen (N)-doped graphene-based material known as pyridinic N removes H₂S via an oxidation reaction compared with another graphene-based material known as oxidized N. To understand this experimental result, we investigated H₂S adsorption on pyridinic N and oxidized N by a density functional theory analysis and further examined the activation barrier of dissociation reactions. We found that the adsorption of H₂S on pyridinic N is more stable than that on oxidized N. In addition, the H₂S dissociation reaction occurs only on pyridinic N.

Received 11th February 2022
Accepted 20th June 2022

DOI: 10.1039/d2ra00898j

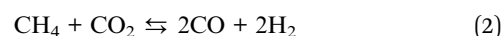
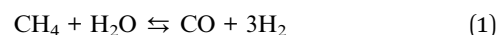
rsc.li/rsc-advances

Introduction

Approximately 80% of the world's ever-increasing energy demand is met by fossil fuels.^{1,2} They supply a significant amount of electrical energy to our society. These reserves are finite and many countries are actively promoting the use of renewable energy sources in anticipation of their depletion in the future.^{3,4} As constant generation of electricity from renewable energy sources is difficult, storage of the electrical energy using hydrogen as an energy carrier has received attention. Compared to conventional batteries, hydrogen is superior in storing energy on a large scale and for long time.⁵ To extract electricity from hydrogen, we can use fuel cells that emit only water and then have a low impact on the environment.⁶ The economic activity of producing, storing, and finally using hydrogen is called the hydrogen-based economy, and it is increasingly expected to be the economy that supplies electric energy to society in a sustainable way.^{7–9}

One of the most environmentally friendly ways to produce hydrogen is water electrolysis using renewable energy. Some Organisation for Economic Cooperation and Development (OECD) countries such as Japan and Germany started demonstration.^{10,11} However, due to profitability issues, only the countries mentioned above attempt to make it a self-sustaining economic activity, while non-OECD countries do not.^{12,13} The non-OECD countries account for more than 80% of the total number of world countries,¹⁴ and the growth of energy demand in the future is expected to be much larger than that of OECD countries. Then, introducing the hydrogen-based economy with environmentally friendly ways to non-OECD countries is a major challenge to build a sustainable society on a global scale.¹⁴

As one of the ways to lead the challenge to success, combining biogas and fuel cells has attracted attention.¹⁵ The flow diagram shown in Fig. 1 illustrates the use of biomass from primary industries such as agriculture and aquafarming. This biomass is converted into biogas by anaerobic digestion also known as methane (CH₄) fermentation. This CH₄ in biogas is used as an energy carrier of hydrogen and it is converted into electrical energy with solid oxide fuel cells (SOFCs). Following a reforming reaction shown in eqn (1) and (2), the H₂ gas is extracted from CH₄, with H₂O, CO₂, and CO.¹⁶ Then, hydrogen reacts with oxide ions (O^{2–}) coming from the cathode side in fuel cells, as shown in eqn (3).¹⁷ This O^{2–} is supplied from O₂ gas in the cathode side.


^aInstitute of Multidisciplinary Research for Advanced Materials, Tohoku University, 2-1-1 Katahira Aoba-ku Sendai, Japan, 980-8577. E-mail: takaya.fujisaki.d5@tohoku.ac.jp

^bInstitute for Materials Chemistry and Engineering and Integrated Research Consortium on Chemical Science (IRCCS), Kyushu University, 744 Motoooka, Nishi-ku, Fukuoka, Japan, 819-0395

^cInternational Institute for Carbon-neutral Energy Research(WPI-I2CNER), Kyushu University, 744 Motoooka, Nishi-ku, Fukuoka, Japan, 819-0395

^dHydrogen Energy Systems, Graduate School of Engineering, Kyushu University, 744 Motoooka, Nishi-ku, Fukuoka, Japan, 819-0395

^eDepartment of Mechanical Science and Engineering, School of Advanced Engineering, Kogakuin University, 2665-1 Nakano-machi, Hachioji, Tokyo, Japan, 192-0015

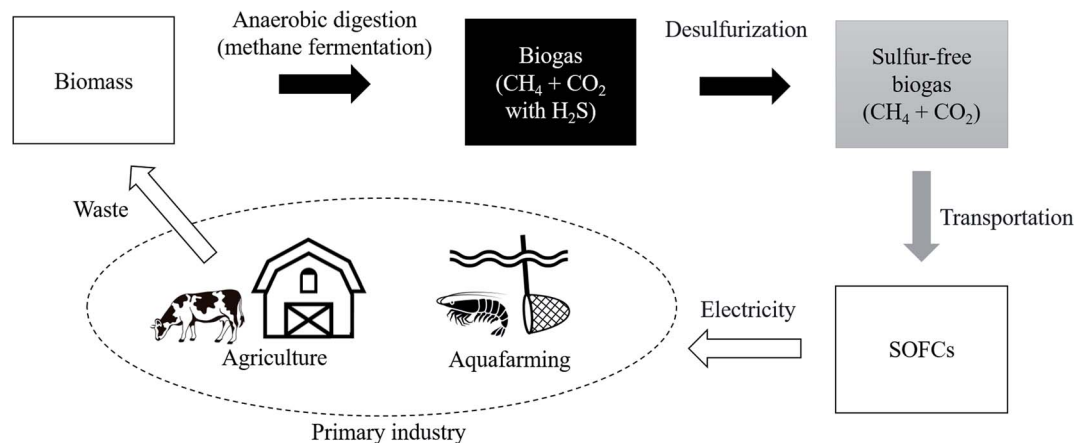
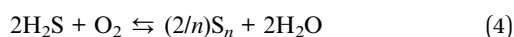



Fig. 1 Material and energy circulation model using biomass.

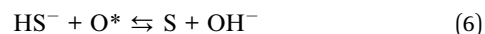
Regarding CO shown in eqn (1) and (2), it reacts with oxygen in the air and it is released into the atmosphere as CO₂. One of the attractions of biomass is that it absorbs a certain amount of CO₂ already at the plant growth stage. Therefore, the environmental impact of biomass is considerably smaller than that of CO₂ generated when simply burning geo-derived resources such as coal and oil. This can be said to be the same advantage of bioethanol.^{18,19} In fact, the demonstration experiment has been started by Shiratori *et al.*, and they fermented biomass derived from aquafarming of shrimp in Vietnam.²⁰ This initiative is a remarkable attempt to use CH₄ as an energy carrier of hydrogen and introduce the hydrogen-based economy because non-OECD countries are mainly dominated by primary industries and they do not have a well-equipped hydrogen infrastructure.²¹

For the realization of the hydrogen-based economy using biogas, it is essential to increase the power generation efficiency of SOFCs. Ideally, it is the best way using pure CH₄. However, actual biogas provided from biomass contains some amount of hydrogen sulfide (H₂S),^{22–24} and it contaminates biogas in the process of anaerobic digestion process to form CH₄.²⁵ The amount of H₂S is approximately 1 ppm and such amount reduces the efficiency of SOFCs²⁶ and degrades even metallic materials such as pipes.^{27,28} Even if the amount of H₂S is reduced to 0.001 ppm, the efficiency still decreases, and then H₂S in biogas is recognized as a substance that should be removed as much as possible.²⁹ For this purpose, some zeolites were reported to absorb it, but they have problems on profitability and durability.^{30–34} However, a more attractive method has been reported and it is called the Claus process.³⁵ In this process, H₂S is recovered externally as a single sulfur (S) *via* an oxidation reaction, as shown in eqn (4):^{36–38}



To accelerate this process, metal oxides such as Fe₂O₃, V₂O₅, Al₂O₃, and TiO₂ have been developed as catalysts.^{39–42} However, these metal oxides are corrosive to H₂S and this causes lower

efficacy of the oxidation process.⁴³ Instead of these metal oxides, nanocarbons that are nano-scale carbon materials have recently attracted attention. Surprisingly, it can be useful in a wide temperature range from room temperature to even around 900 °C. Moreover, it has a high surface area with pore structures and unique surface function.^{44–49} Because nanocarbons can be utilized without metal-based active sites, it is expected that the reduction of catalytic performance due to H₂S poisoning is avoided. The first reaction to remove H₂S on nanocarbons is shown in eqn (5) as a dissociation reaction. Then, HS[−] reacts with oxygen radicals (O*) and sulfur (S) is recovered following eqn (6):⁵⁰



Because the carbon site itself in nanocarbons is inactive to remove H₂S, substitution of heteroatoms with some carbon sites is a reasonable approach to make the inactive site active one. It was first shown by Chizari *et al.* that a nitrogen (N)-doped carbon nanotube had excellent activity for the oxidation reaction of H₂S.⁵¹ Since then, several researchers have started studies on N-doped nanocarbons focusing on detailed microstructures,⁵² controlling the nitrogen content,⁵³ investigating the effect of caustic alkali impregnating agents,⁵⁴ and developing the preparation process for mesoporous materials.⁵⁵ Furthermore, it is interesting to report that the edge site of N-doped graphene sheets promotes the oxidation reaction of H₂S than the ordered plane (*cf.* pure graphene structures).^{56,57} Such sites might act as the origin of the high catalytic performance for H₂S selective oxidation. Then, focusing on heteroatoms such as N-doped nanocarbons becomes one of the most important approaches. In other words, finding the best active site promoting the oxidation is the common target. However, because the guideline is still not clear, the industrial-scale application has been hampered.

Recently, Shiyan *et al.* have conducted comparative experiments on the effects of N- and O-doped nanocarbons on the oxidation reaction of H₂S. They conducted experiments



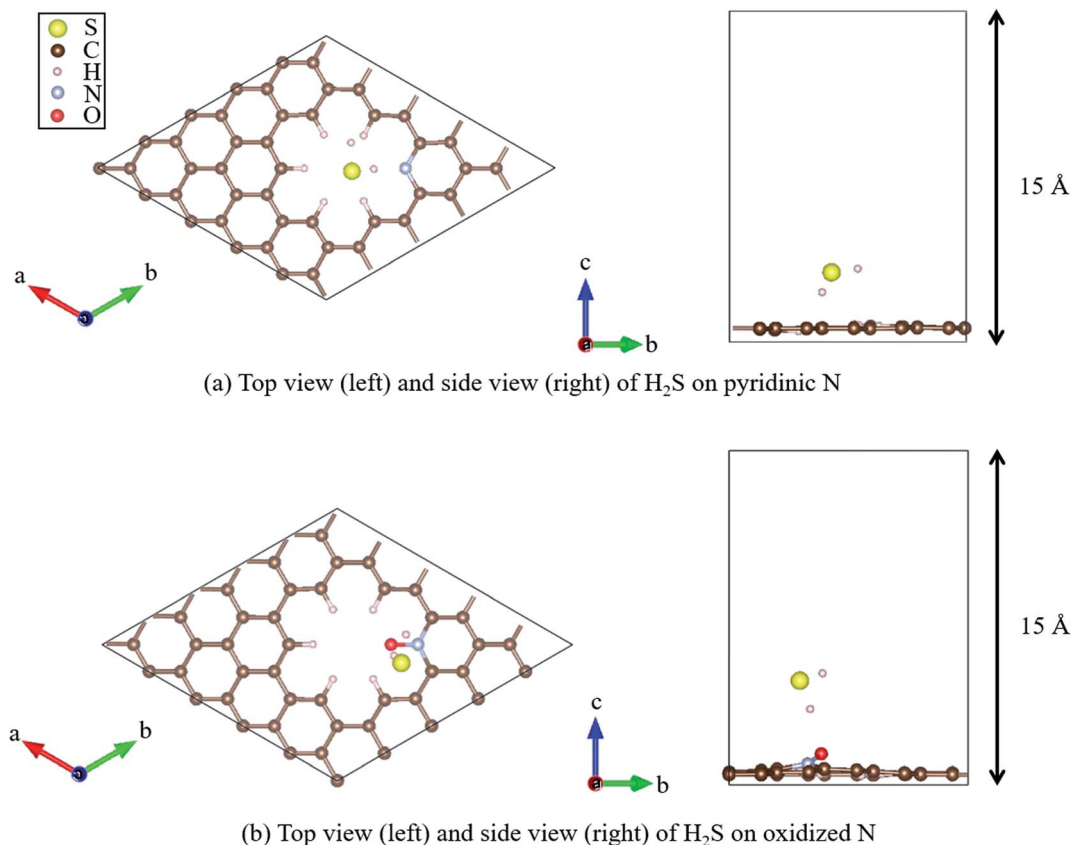


Fig. 2 Top and side views of the geometry optimized structure of H_2S on (a) pyridinic N and (b) oxidized N.

demonstrating that pyridinic N (Fig. 2(a)), in which N is doped into nanocarbon and hydrogen is bonded to the nitrogen as a terminal atom, promotes the oxidation reaction the most. However, another graphene-based structure known as oxidized N (Fig. 2(b)), in which an oxygen atom is substituted for the hydrogen bonded next to the nitrogen atom in pyridinic N, showed the least enhancement of the oxidation reaction of H_2S . To understand the difference in enhancement between pyridinic N and oxidized N at an atomic level, they attempted a density functional theory (DFT) approach. They reported that the adsorption of HS^- on pyridinic N was the highest compared with that on oxidized N that showed least adsorption.⁵⁸ However, this previous study has the following demerit: HS^- is a dissociate of H_2S , and the stability of HS^- cannot be discussed unless it is shown that H_2S gets adsorbed onto pyridinic N and oxidized N. Therefore, the aim of this study is to clarify the adsorption properties of H_2S on pyridinic N and oxidized N by DFT.

Computational details

All DFT calculations were performed using the Vienna *Ab initio* Simulation Package (VASP).^{59–61} The Perdew–Burke–Ernzerhof (PBE)⁶² exchange–correlation functional was employed based on the projector augmented wave (PAW) method.⁶⁴ The cutoff energy for the plane wave basis was set to 400 eV, referring to

a previous study.⁶³ First, a 4×4 graphene sheet was constructed and optimized using $5 \times 5 \times 1$ Monkhorst–Pack k -point meshes. Moreover, the spin polarization was applied. For the projector augmented wave (PAW) method, pseudopotentials with valence states of C ($2s^2, 2p^2$), N ($2s^2, 2p^3$), S ($3s^2, 3p^4$), O ($2s^2, 2p^4$), and H ($1s$) were used. All ionic positions were optimized by a conjugate gradient method until the forces on each ion were below $10^{-3} \text{ eV } \text{\AA}^{-1}$. The electronic energy was converged to less than 10^{-5} eV . Then, after the geometrical optimization of the 4×4 graphene sheet under the above-mentioned conditions, the length of the C–C bond was found to be 1.426 \AA , which is in good agreement with previous results.^{64–66} In addition, the internal angles of the six-membered ring of graphene structures were within the range of 120° to $\pm 0.025\%$, which is also in good agreement with the results of previous studies.^{67,68} Next, with the optimized graphene sheet spreading to a - and b -axis directions, we added a vacuum slab on it by making 15 \AA height in the c -axis direction. Then, geometry optimization was performed while keeping the vacuum slab. From the optimized graphene sheet with a vacuum slab, a six-membered ring structure consisting of six carbons was removed. Then, hydrogen atoms were introduced to terminate the remaining carbon atoms. The edge of the graphene sheet is reported to be bonded with hydrogen atoms,⁶⁹ which prevents the carbons from becoming radicalized and destabilized. Finally, the carbon site was replaced with



a nitrogen atom (N), as shown in Fig. 2(a), and with nitrogen and oxygen, as shown in Fig. 2(b). The respective structure is called pyridinic N and oxidized N, respectively.⁵⁸ They were visualized using VESTA.⁷⁰

Fig. 2(a) and (b) show that hydrogen sulfide (H₂S) was adsorbed onto pyridinic N and oxidized N, and H₂S itself was optimized. For this, H₂S was first structurally optimized in vacuum. H₂S has a bent structure with lone-pair electrons in the S atom, and the calculated S–H bond length is 1.35 Å (1.35 Å and 1.33 Å). Moreover, the H–S–H angle was found to be 91.2° (91.6° and 92.2°), which is in good agreement with the previous experiment and calculated values in parentheses.⁷¹ This single molecule of H₂S in vacuum was structurally optimized with 1 × 1 × 1 Monkhorst–Pack *k*-point meshes. For each of the above-mentioned pyridinic N and oxidized N, H₂S was placed on a nitrogen atom and the structure was optimized. The position of H₂S was referred from the previous study by Shiyan *et al.*⁷²

To accurately determine the stability of H₂S on pyridinic N and oxidized N, we considered adsorption energy (E_{ads}).^{73,74} It is defined in eqn (7), which is the sum of the interaction energy (E_{int}) in eqn (8), and deformation energy (E_{def}) shown in eqn (9). The deformation energy is obtained as follows: H₂S and pyridinic N form the most stable structure when they are placed in vacuum with no other atoms around them. However, H₂S and pyridinic N deforms when they are close to each other due to their interaction. The change in the electrostatic potential at this time is defined as the deformation energy. The interaction energy (E_{int}) and deformation energy (E_{def}) shown in eqn (8) and (9) are the case for pyridinic N with H₂S. When the adsorption energy was calculated as negative, H₂S gets adsorbed onto pyridinic N stably.

$$E_{\text{ads}} = E_{\text{def}} + E_{\text{int}} \quad (7)$$

$$E_{\text{int}} = E_{\text{H}_2\text{S+pyridinic N}} - (E_{\text{pyridinic N}} + E_{\text{H}_2\text{S}}) \quad (8)$$

$$E_{\text{def}} = E_{\text{def_H}_2\text{S}} + E_{\text{def_pyridinic N}} \quad (9)$$

$E_{\text{H}_2\text{S+pyridinic N}}$, $E_{\text{pyridinic N}}$, and $E_{\text{H}_2\text{S}}$ are the electrostatic potential for pyridinic N and adsorbed H₂S, pyridinic N, and single H₂S, respectively.

Moreover, the minimum energy paths and the energy barriers for H₂S dissociation on graphene-based structures were obtained by the climb image nudged elastic band (NEB) method. Five intermediate images were used to sample the reaction path between the initial adsorption state and the final dissociated state. The spring constant of the NEB calculations

was 5.0 eV Å^{−2} and the NEB force convergence criteria was 0.01 eV Å^{−1} using 3 × 3 × 1 Monkhorst–Pack *k*-point meshes.

Results and discussion

The calculated adsorption energies for H₂S on pyridinic N and oxidized N are given in Table 1. The electrostatic potentials are summarized in Table 2. As shown in Table 1, H₂S on pyridinic N shows −0.117 eV as the adsorption energy, and this number is 21% lower than that of oxidized N as −0.097 eV. This indicates that H₂S is more stably adsorbed onto pyridinic N than onto oxidized N, indicating that pyridinic N is a more favorable material than oxidized N to proceed the oxidation reaction (2H₂S + O₂ ⇌ (2/*n*) S_{*n*} + 2H₂O), as shown in eqn (4). Shiyan *et al.* experimentally reported the same trend,⁷² and the result of our study supports their result successfully by DFT. Notably, oxidized N does not adsorb H₂S as readily as pyridinic N, but this indicates that H₂S on oxidized N can be desorbed relatively easily by external energies such as heat. This suggests that the structure of oxidized N could be applied to the reusable desorption sheet to remove H₂S.

The reason why H₂S adsorption occurs more favorably on pyridinic N than on oxidized N is explained as follows. As mentioned before, the adsorption energy is expressed as the sum of the interaction energy, shown in eqn (8), and the deformation energy, also shown in eqn (9). Their values are presented in Table 1 and their amount is graphically visualized in Fig. 3.

As shown in Fig. 3, it is clear that the effect of the interaction energy is much larger than that of the deformation energy in determining the adsorption energy. The first factor that determines the amount of the interaction energy is reported with the idea of attractive force (intermolecular force) that acts between materials caused by the transfer of charge.^{75,76} Fig. 4 shows electric charges of adsorbed H₂S and its surrounding atoms: N and O. Every charge was obtained by Bader charge analysis.⁷⁷ This figure shows pyridinic N (Fig. 4(a)) and oxidized N (Fig. 4(b)) with adsorbed H₂S, respectively, and each structure has already been geometry optimized. Fig. 4(c) shows H₂S placed in vacuum. From pyridinic N and oxidized N, H₂S is adsorbed approximately 2 Å away, and this value has good agreement compared to previous studies reporting the adsorption distances between molecules and graphene-based structures.^{78,79} As shown in Fig. 4(a), the charge of the entire H₂S was calculated as +0.19 e (electron), and the charge of the N atom of pyridinic N adjacent to H₂S was polarized to −1.18 e. However, as shown in Fig. 4(b), H₂S is polarized to −0.02 e, and the

Table 1 Adsorption, interaction, and deformation energies of H₂S on pyridinic N and oxidized N

	On pyridinic N (eV)	On oxidized N (eV)
Adsorption energy	−0.117	−0.097
Interaction energy	−0.129	−0.114
Deformation energy of H ₂ S	0.003	0.002
Deformation energy of pyridinic N/oxidized N	0.009	0.015



Table 2 Adsorption, interaction, and deformation energies of H₂S on pyridinic N and oxidized N^a

Structure	Energy (eV)
Pyridinic N	−420.563
Oxidized N	−425.468
H ₂ S with pyridinic N	−431.879
H ₂ S with oxidized N	−436.778
Pyridinic N non-relaxed	−420.554
Oxidized N non-relaxed	−425.453
H ₂ S (∠HSH = 92.06°)	−11.192
H ₂ S (∠HSH = 91.886°)	−11.193
H ₂ S (∠HSH = 91.20°)	−11.195

^a The values of “pyridinic N non-relaxed” and “oxidized N non-relaxed” mean the electrical potential of pyridinic N and oxidized N without performing geometry optimization by removing H₂S from “H₂S with pyridinic N” and “H₂S with oxidized N”, respectively. In addition, the energy for each angle of H₂S is listed in the last three lines.

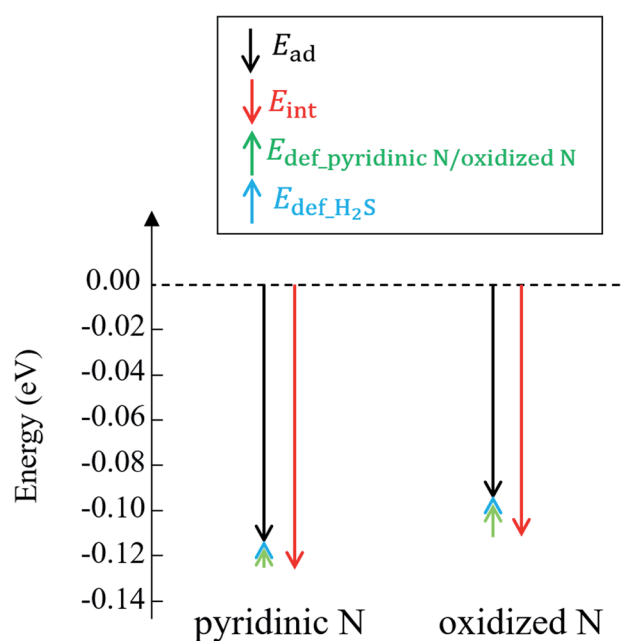


Fig. 3 Adsorption energy (E_{ads}) and interaction energy (E_{int}) between H₂S and pyridinic N/oxidized N. The deformation energies of pyridinic N ($E_{\text{def_pyridinic N}}$), oxidized N ($E_{\text{def_oxidized N}}$), and H₂S are also shown.

charges of O and N of oxidized N are polarized to -0.59 e and -0.55 e. The difference in charge polarization can be explained by the difference in electronegativity whose order is $\text{O} > \text{N} > \text{S} > \text{C} > \text{H}$ in the atoms, as shown in Fig. 4.⁸⁰ Because the N atom in pyridinic N has the highest electronegativity, N has the largest polarization as -1.18 e. Such large polarization is the result of attracting electrons, and this makes the charge of H₂S as $+0.19$ e. This suggests that the intermolecular force between pyridinic N and H₂S contributes to the adsorption energy of H₂S to make the H₂S stable. However, the charge on the O atom of oxidized N is negatively polarized as -0.59 e, and that of H₂S is also negatively polarized as -0.02 e. This indicates that

repulsive forces are at work between them. Despite the presence of repulsive force between H₂S and oxidized N, the reason why the adsorption energy is negative as shown in Table 1 can be the formation of ionic or covalent bonds that have a stronger binding force than the repulsive force generated by intermolecular force. Therefore, in addition to pyridinic N and H₂S, the covalent nature between oxidized N and H₂S was investigated as follows.

In fact, the relationship between the covalent nature and the interaction energy has been discussed in several papers. Wang *et al.* reported the bonding nature of PbO molecules on the aluminium surface, and they showed that the magnitude of covalent bonding contributes to the value of interaction energy.⁸¹ Moreover, Zhou *et al.* mentioned that ammonia molecules adsorbed onto SiSe layers showed more stability with greater covalency.⁸² In addition, covalent bonding can be explained by the amount of electrons between atoms, and the bonding between zirconium and oxygen atoms seems to be greater than that between cerium and oxygen atoms in perovskite structures.⁸³ Considering these previous studies, the magnitude of the covalency in terms of the number of electrons between atoms has a significant effect on the interaction energies. Fig. 5(a) shows the electron mapping obtained by slicing the coordinates of the following three atoms: the N atom of pyridinic N, the S atom of H₂S, and the H atom of H₂S. Fig. 5(b) describes the density of states of the H atom in H₂S and the N atom in pyridinic N, which are neighbouring atoms, and Fig. 5(c) shows the density of states of S in H₂S and N in pyridinic N. In Fig. 5(a), it seems that there is little sharing to create a covalent bonding between the H atom and the N atom. However, as shown in Fig. 5(c), the s- and p-orbitals of the N atom in pyridinic N clearly overlap on the p-orbital band structure of the S atom. From the above, it is shown that the electrons between H₂S and pyridinic N shown in Fig. 5(a) make a covalent bond between S of H₂S and the N of pyridinic N. This electron sharing stabilizes H₂S on pyridinic N and it helps determining the number of interaction energy to be -0.129 eV, as shown in Table 1. However, Fig. 5(d) shows the electron mapping of H₂S adsorbed onto oxidized N using the coordinates of the O and N atoms of oxidized N, and the H atom of H₂S. From this figure, it is observed that there are electrons between the O atom of oxidized N and the H atom of H₂S. This number of electrons in Fig. 5(d) seems to be larger than that of electrons between pyridinic N and H₂S, as shown in Fig. 5(a). However, Fig. 5(e) shows that s- and p-orbitals of the H atom in H₂S hardly overlap with the s- and p-orbitals of the O atoms in oxidized N. This is the proof that the type of bond between H₂S and oxidized N is ionic bond. Focusing on the p-orbital of the O atom, the bond between O and H appears to be formed as a covalent bond, as shown in Fig. 5(d), as the density of state in the p-orbital of O is distributed over a wide range of energy, as shown in Fig. 5(e) and then electrons of the p-orbital are widely delocalized around the O atom. In other words, there is no covalent but mainly ionic bond between H₂S and oxidized N. Although there was repulsive intermolecular force between H₂S and oxidized N, this ionic bonding between them makes H₂S stable on oxidized N, which results in the negative adsorption



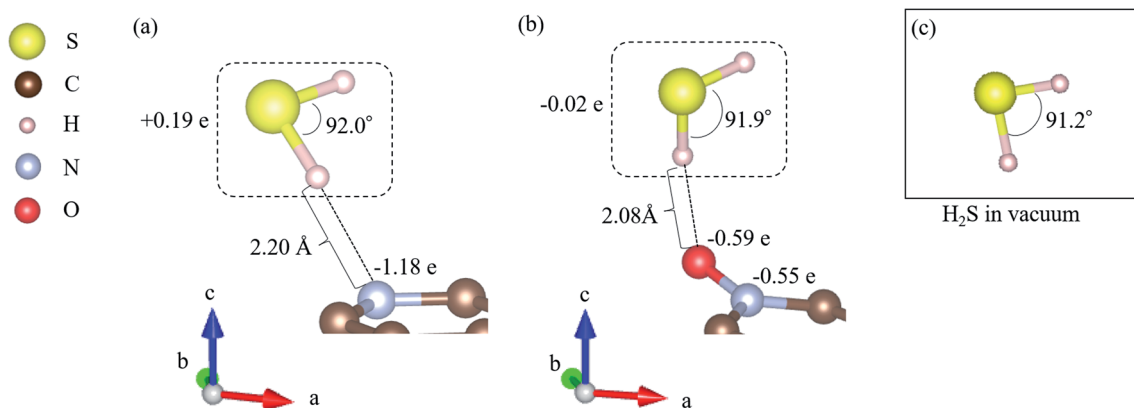


Fig. 4 Adsorption energy (E_{ads}) and interaction energy (E_{int}) between H_2S and pyridinic N/oxidized N. The deformation energies of pyridinic N ($E_{\text{def_pyridinic N}}$), oxidized N ($E_{\text{def_oxidized N}}$), and H_2S are also shown.

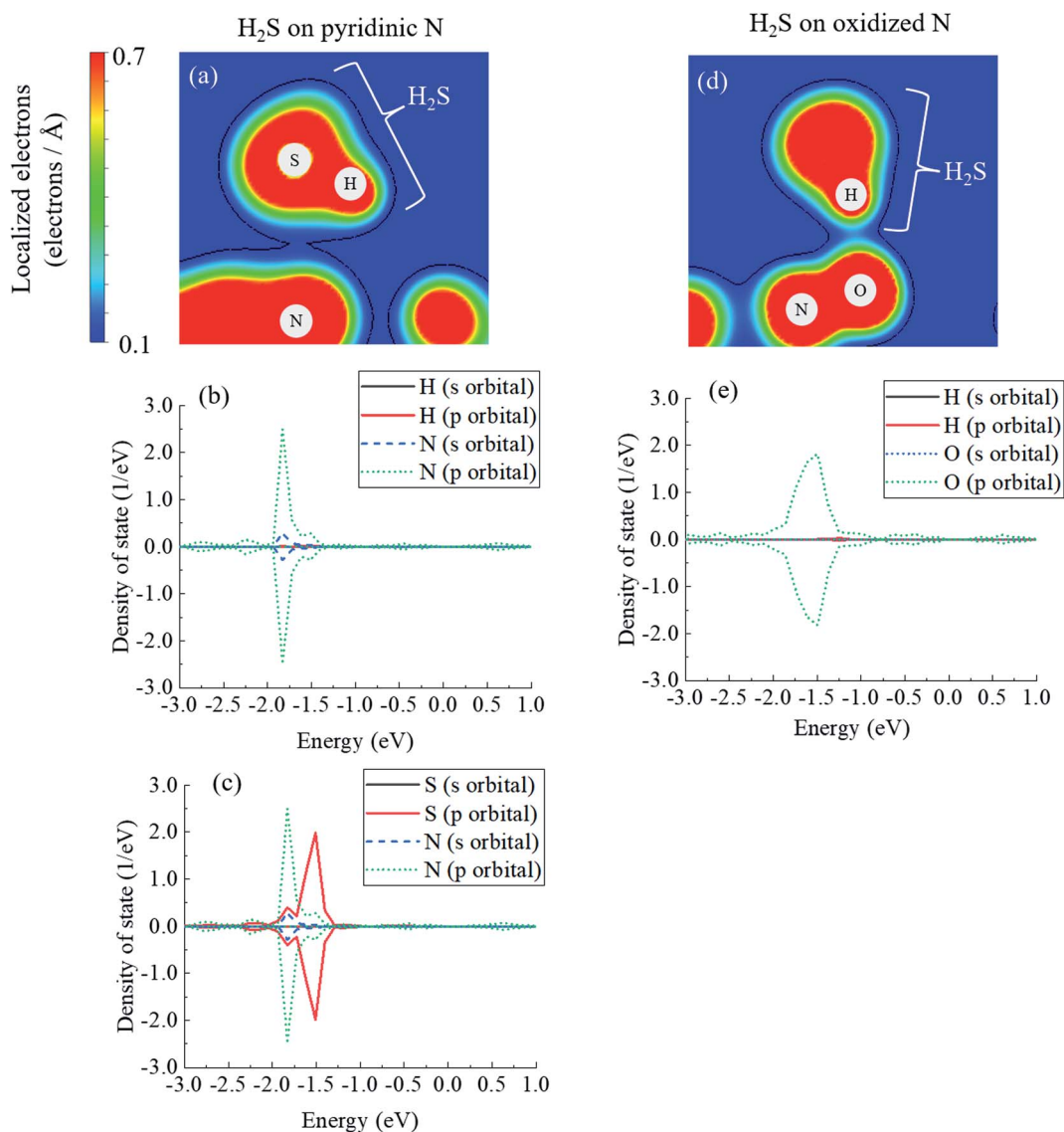


Fig. 5 (a) Electron mapping for H_2S on pyridinic N, including density of states of (b) H–N and (c) S–N electrons. (d) Electron mapping of H_2S on oxidized N, and (e) density of states of H–O.



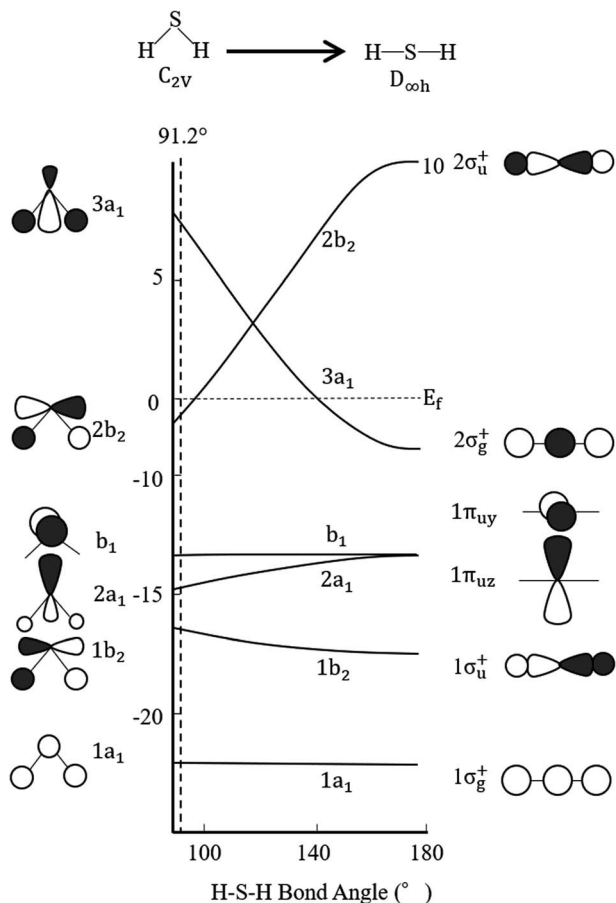


Fig. 6 Walsh diagram of H_2S . Black and white lobes show positive and negative wave functions, respectively.

energy, as shown in Table 1. As described above, the difference in the interaction energies presented in Table 1 can be attributed to the difference in the degree of covalent bonding property between H_2S and pyridinic N or oxidized N.

Finally, regarding the deformation energy, it is significantly smaller relative to the interaction energy, as shown in Table 1 and Fig. 3. However, to exactly determine the adsorption energy in this study, it is important to explain the effect of such minor energy contribution. First, the deformation energies of pyridinic N and oxidized N after adsorption of H_2S were calculated as +0.009 eV and +0.015 eV, respectively. The reason for the higher deformation energy of oxidized N is that the O atom is attracted to the H atom of H_2S by intermolecular force, as shown in Fig. 1 and 4. As for the deformation energy of H_2S , the value for H_2S on pyridinic N is +0.003 eV, and that for oxidized N is +0.002 eV. The reason for the difference is that the bond angle of H_2S on pyridinic N is 92.0° and this value is larger than the bond angle (91.9°) on oxidized N, as shown in Fig. 4(a) and (b). In fact, both angles are larger than the angle of H_2S in vacuum (91.2°), as shown in Fig. 4(c). The origin of such angle change has already been well discussed in previous studies and the cause comes from the result of charge transfer in molecules. Bajdich *et al.* reported that charge transfer occurs to CO_2 adsorbed on the Au surface, leading to a change in bond angle.⁸⁴ Indeed, as shown in Fig. 4(a) and (b) in our study, the charge transfer between H_2S and pyridinic N is larger than that between H_2S and oxidized N. Therefore, the bond angle of pyridinic N is considered to be larger than that of oxidized N. The bond angle change is also explicable from the viewpoint of the electronic structure of Walsh diagram, as shown in Fig. 6.⁸⁵ The diagram shows that the energy of orbitals b_1 and $2a_1$ increases as the angle of H_2S increases in the narrow range of 91.2 to 92.0° . The above-mentioned difference can be clarified if we can describe the density of state with sufficiently high accuracy. However, the VASP used in this study was unable to follow the change in the Walsh diagram within 1.0° of the H_2S bond angle. Therefore, it is significantly desirable to have a method that can discuss the change in the density of states for a very narrow range of bond angle, which is less than 1° .

Moreover, we determined the activation barrier for H_2S molecules to dissociate on the graphene structure. Fig. 7 shows

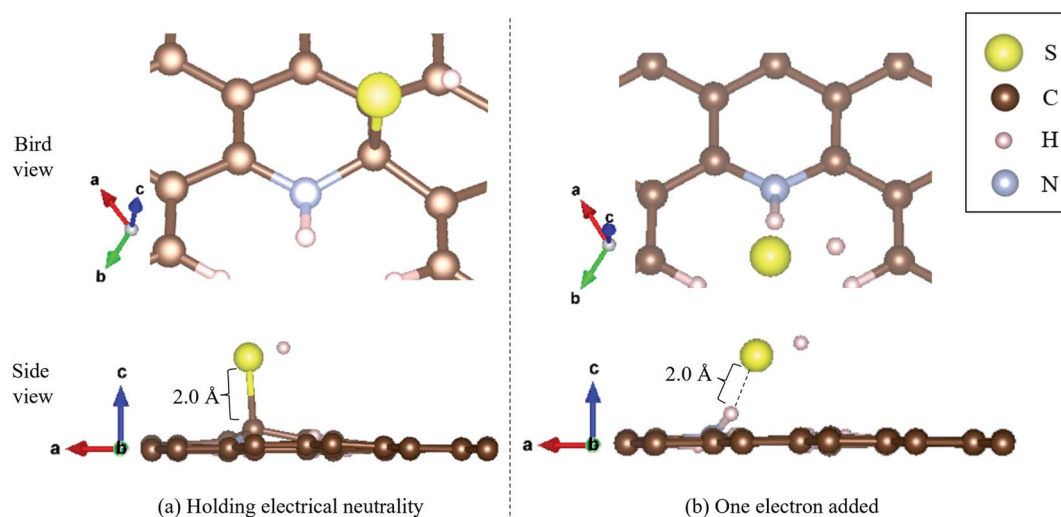


Fig. 7 H_2S dissociated as HS^- and H^+ on pyridinic N under (a) holding electrical neutrality, and (b) one electron added condition.



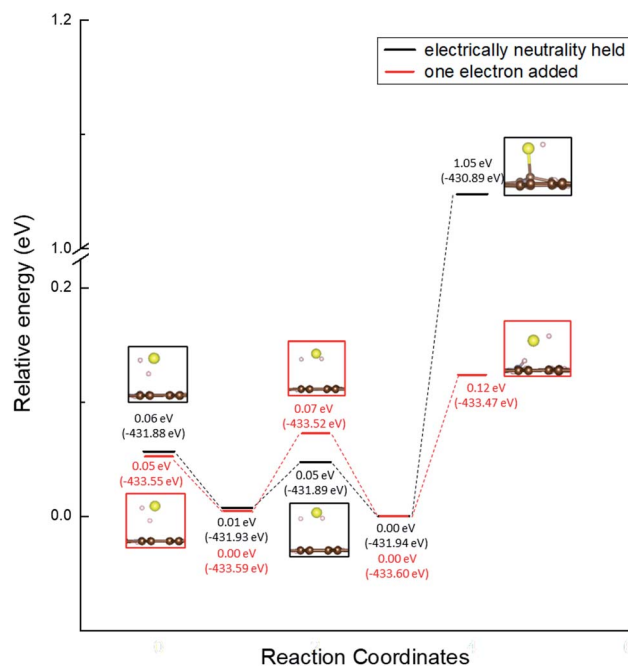


Fig. 8 Activation barrier of the H_2S dissociation reaction on pyridinic N under the conditions of holding electrical neutrality and adding one electron.

H_2S dissociated as HS^- and H^+ on Pyridinic N. Fig. 7(a) shows the case where H_2S is dissociated while maintaining the electrically neutral condition, and Fig. 7(b) shows the case where one electron is added to the entire system. Both cases show cases where the most dissociated H_2S is stabilized at several locations around the nitrogen atom. The electron-added system shown in Fig. 7(b) was chosen for investigation because of the reduction reaction. A previous study has also considered the possibility that pyridinic N was reduced by experimental residual reducing gases such as carbon monoxide and hydrogen gas.⁷² In fact, a previous study also suggested there is some amount of oxygen in experimentally prepared pyridinic N.⁷² With the initial structure shown in Fig. 2(a) and final structures shown in Fig. 7 of pyridinic N, the activation barrier was calculated as shown in Fig. 8. Focusing on the final structure and when the electroneutrality is maintained, the final structure is about 1.05 eV larger than that of the previous structure. However, when a single electron was added into the entire system, the final structure shows 0.12 eV larger activation barrier than the previous structure. This value (0.12 eV) is about one-tenth the size of the structure under electroneutrality. This indicates that the hydrogen sulfide dissociation was significantly enhanced by the reducing gas.

However, H_2S is always stable as a molecule on oxidized N, and the dissociation reaction could not be reproduced. Although a previous study had raised the possibility that a water film could cause H_2S to dissociate,⁸⁶ the separation of H_2S into HS^- and H^+ by water molecules could not be reproduced by our investigation. Chemical reactions such as the above-described dissociation reaction do not occur because the interaction energy between H_2S and oxidized N is -0.114 eV, which is so

small that there might be contribution of hydrogen bonding as well as ionic bonding. However, as shown in Fig. 5(e), there is a slight overlap of electrons between the O p-orbital and the H p-orbital, making it difficult to precisely separate the effect of hydrogen bonding from that of ionic bonding. As mentioned above, we could not find any tendency for H_2S to react with oxidized N in a divergent manner. Therefore, we conclude that oxidized N does not contribute to the dissociation reaction for H_2S , but rather has the effect of preventing the dissociation reaction.

Conclusions

In this study, the adsorption energies of H_2S on pyridinic N and oxidized N have been precisely investigated by examining the interaction and deformation energies by DFT. Our result indicated that H_2S gets adsorbed more preferably onto pyridinic N than onto oxidized N. This indicates that pyridinic N is a significant material for the oxidation reaction of H_2S . The above-mentioned difference is explicable because the interaction energy of H_2S with pyridinic N is lower than that with oxidized N, which can be inferred from the difference in intermolecular forces and covalency properties. As for the deformation energy, it is remarkably small compared to the interaction energy, but oxidized N has a higher deformation energy than pyridinic N, which is due to the intermolecular force between the O atom in oxidized N and the H atom in H_2S . It was also shown that the bond angle of H_2S is larger on pyridinic N than on oxidized N due to charge transfer after adsorption of H_2S .

Furthermore, the activation barrier for the dissociation of H_2S ($\text{H}_2\text{S} \rightleftharpoons \text{H}^+ + \text{HS}^-$) on pyridinic N and oxidized N was investigated and found to be 1.05 eV on pyridinic N. Moreover, when one electron was added considering the reducing atmosphere, the activation barrier on pyridinic N was found to be 0.12 eV, which became very smaller. However, we could not reproduce the reaction of H_2S dissociation on oxidized N, and concluded that only pyridinic N contributes to the H_2S dissociation reaction.

Author contributions

T. F. and Y. S. designed this project. K. I. and A. T. S. provided valuable advice for proceeding this research. Also, H. S. gave useful viewpoint from experimental aspect. The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by JSPS KAKENHI, Fostering Joint International Research (B), Grant Number 20KK0248. Also, we would like to thank Editage (<https://www.editage.com>) for



English language editing. In addition, the computation was carried out using the computer resource offered under the category of General Projects by Research Institute for Information Technology, Kyushu University.

References

- 1 U. S. Energy Information Administration (EIA), April 2021 *Monthly Energy Review*, DOE/EIA-0035(2021/12), Washington, DC, December 22, 2021, <https://www.eia.gov/totalenergy/data/monthly/pdf/mer.pdf>.
- 2 J. F. Mercure, H. Pollitt, J. E. Viñuales, N. R. Edwards, P. B. Holden, U. Chewpreecha, P. Salas, I. Sognnaes, A. Lam and F. Knobloch, *Nat. Clim. Change*, 2018, **8**, 588–593.
- 3 B. Looney, *BP Statistical Review of World Energy*, 2020, p. 66.
- 4 S. H. Mohr, J. Wang, G. Ellem, J. Ward and D. Giurco, *Fuel*, 2015, **141**, 120–135.
- 5 E. López González, F. Isorna Llerena, M. Silva Pérez, F. Rosa Iglesias and J. Guerra Macho, *Int. J. Hydrogen Energy*, 2015, **40**, 5518–5525.
- 6 Energy Information Administration (EIA), *international energy outlook 2019: With projections to 2050*, #IEO2019, (Washington, DC, September 24, 2019).
- 7 S. Kharel and B. Shabani, *Energies*, 2018, **11**(10), 2825–2842.
- 8 P. Zegers, *J. Power Sources*, 2006, **154**, 497–502.
- 9 R. H. Lin, Y. Y. Zhao and B. D. Wu, *Int. J. Hydrogen Energy*, 2020, **45**, 20164–20175.
- 10 N. Takatsu and H. Farzaneh, *Appl. Sci.*, 2020, **10**(12), 4061–4080.
- 11 T. Nguyen, Z. Abidin, T. Holm and W. Merida, *Energy Convers. Manage.*, 2019, **200**, 112108–112120.
- 12 R. Samsun, L. Antoni and M. Rex, IEA Technology Collaboration Program, *Advanced Fuel Cells Technology Collaboration Programme*, 2020, https://www.ieafuelcell.com/fileadmin/publications/2020_AFCTCP_Mobile_FC_Application_Tracking_Market_Trends_2020.pdf.
- 13 E. Weidner, R. Ortiz Cebolla and J. Davies, Global deployment of large capacity stationary fuel cells, *Tech. Rep.*, 2019, 6.
- 14 D. Paprotny, *Soc. Indicat. Res.*, 2020, 1–33.
- 15 Y. Shiratori, T. Quang-Tuyen, Y. Umemura, T. Kitaoka and K. Sasaki, *Int. J. Hydrogen Energy*, 2013, **38**, 11278–11287.
- 16 Ö. Aydın, A. Kubota, D. L. Tran, M. Sakamoto and Y. Shiratori, *Int. J. Hydrogen Energy*, 2018, **43**, 17431–17443.
- 17 N. Shi, Y. Xie, Y. Yang, S. Xue, X. Li, K. Zhu, D. Huan, R. Peng, C. Xia and Y. Lu, *Mater Renew Sustain Energy*, 2020, **9**, 1–18.
- 18 S. A. Saadabadi, A. T. Thattai, L. Fan, R. E. Lindeboom, H. Spanjers and P. Aravind, *Renewable Energy*, 2019, **134**, 194–214.
- 19 N. Hossain, J. Zaini and T. M. Indra Mahlia, *Renewable Sustainable Energy Rev.*, 2019, **115**, 109371.
- 20 Y. Shiratori, M. Sakamoto, T. Nguyen, T. Yamakawa, T. Kitaoka, H. Orishima, H. Matsubara, Y. Watanabe, S. Nakatsuka and T. Doan, *Fuel Cells*, 2019, **19**, 346–353.
- 21 Y. Hu, L. Peng, X. Li, X. Yao, H. Lin and T. Chi, *Energy*, 2018, **147**, 1177–1187.
- 22 D. Divya, L. R. Gopinath and P. Merlin Christy, *Renewable Sustainable Energy Rev.*, 2015, **42**, 690–699.
- 23 E. Bocci, A. Di Carlo, S. J. McPhail, K. Gallucci, P. U. Foscolo, M. Moneti, M. Villarini and M. Carlini, *Int. J. Hydrogen Energy*, 2014, **39**, 21876–21895.
- 24 H. J. Alves, C. Bley Junior, R. R. Niklevicz, E. P. Frigo, M. S. Frigo and C. H. Coimbra-Araújo, *Int. J. Hydrogen Energy*, 2013, **38**, 5215–5225.
- 25 M. Syed, G. Soreanu, P. Falletta and M. Béland, *Can. Biosyst. Eng.*, 2006, **48**, 2.
- 26 Z. Cheng, J.-H. Wang, Y. Choi, L. Yang, M. C. Lin and M. Liu, *Energy Environ. Sci.*, 2011, **4**(11), 4380–4410.
- 27 C. H. Pham, S. Saggat, P. Berben, T. Palmada and C. Ross, *J. Environ. Qual.*, 2019, **48**, 32–38.
- 28 F. B. Asghari, J. Jaafari, M. Yousefi, A. A. Mohammadi and R. Dehghanzadeh, *Hum. Ecol. Risk Assess.*, 2018, **24**(4), 1138–1149.
- 29 J. Kupecki, D. Papurello, A. Lanzini, Y. Naumovich, K. Motylinski, M. Blesznowski and M. Santarelli, *Appl. Energy*, 2018, **230**, 1573–1584.
- 30 R. Dehghan and M. Anbia, *Fuel Process. Technol.*, 2017, **167**, 99–116.
- 31 A. Peluso, N. Gargiulo, P. Aprea, F. Pepe and D. Caputo, *Sep. Purif. Rev.*, 2019, **48**(1), 78–89.
- 32 M. Ozekmekci, G. Salkic and M. F. Fellah, *Fuel Process. Technol.*, 2015, **139**, 49–60.
- 33 Z. Jiang, X. Xu, Y. Ma, H. S. Cho, D. Ding, C. Wang, J. Wu, P. Oleynikov, M. Jia, J. Cheng, Y. Zhou, O. Terasaki, T. Peng, L. Zan and H. Deng, *Nature*, 2020, **586**(7830), 549–554.
- 34 M. Khabazipour and M. Anbia, *Ind. Eng. Chem. Res.*, 2019, **58**(49), 22133–22164.
- 35 M. Izaki and T. Omi, *J. Electrochem. Soc.*, 1996, **143**(3), L53.
- 36 X. Zhang, Z. Wang, N. Qiao, S. Qu and Z. Hao, *ACS Catal.*, 2014, **4**(5), 1500–1510.
- 37 X. Zheng, Y. Li, L. Zhang, L. Shen, Y. Xiao, Y. Zhang, C. Au and L. Jiang, *Appl. Catal., B*, 2019, **252**, 98–110.
- 38 Q. Gu, Y. Lin, S. Heumann and D. Su, *Chem.-Asian J.*, 2017, **12**(22), 2876–2883.
- 39 S. W. Chun, J. Y. Jang, D. W. Park, H. C. Woo and J. S. Chung, *Appl. Catal., B*, 1998, **16**(3), 235–243.
- 40 S. Phatychuen, B. Pongthawornsakun, J. Panpranot and P. Praserttham, *J. Environ. Chem. Eng.*, 2018, **6**(5), 5655–5661.
- 41 X. Zheng, L. Zhang, Z. Fan, Y. Cao, L. Shen, C. Au and L. Jiang, *Chem. Eng. J.*, 2019, **374**, 793–801.
- 42 L. Shen, X. Zheng, G. Lei, X. Li, Y. Cao and L. Jiang, *Chem. Eng. J.*, 2018, **346**, 238–248.
- 43 X. Zhang, Y. Tang, S. Qu, J. Da and Z. Hao, *ACS Catal.*, 2015, **4**(5), 1500–1510.
- 44 Z. Yu, X. Wang, X. Song, Y. Liu and J. Qiu, *Carbon*, 2015, **95**, 852–860.
- 45 F. Sun, J. Liu, H. Chen, Z. Zhang, W. Qiao, D. Long and L. Ling, *ACS Catal.*, 2013, **3**(5), 862–870.
- 46 M. Daraee, M. Baniadam, A. Rashidi and M. Maghrebi, *Chem. Phys.*, 2018, **511**, 7–19.



- 47 M. Sun, X. Wang, X. Pan, L. Liu, Y. Li, Z. Zhao and J. Qiu, *Fuel Process. Technol.*, 2019, **191**, 121–128.
- 48 Y. Pan, M. Chen, M. Hu, M. Tian, Y. Zhang and D. Long, *Appl. Catal., B*, 2020, **262**, 118266.
- 49 G. Lei, Y. Cao, W. Zhao, Z. Dai, L. Shen, Y. Xiao and L. Jiang, *ACS Sustainable Chem. Eng.*, 2019, **7**(5), 4941–4950.
- 50 D. Li, W. Chen, J. Wu, C. Q. Jia and X. Jiang, *J. Mater. Chem. A*, 2020, **8**(47), 24977–24995.
- 51 K. Chizari, A. Deneuve, O. Ersen, I. Florea, Y. Liu, D. Edouard, I. Janowska, D. Begin and C. Pham-Huu, *ChemSusChem*, 2012, **5**(1), 102–108.
- 52 Q. Chen, Z. Wang, D. Long, X. Liu, L. Zhan, X. Liang, W. Qiao and L. Ling, *Ind. Eng. Chem. Res.*, 2010, **49**(7), 3152–3159.
- 53 Y. Liu, C. Duong-Viet, J. Luo, A. Hébraud, G. Schlatter, O. Ersen, J. M. Nhut and C. Pham-Huu, *ChemCatChem*, 2015, **7**(18), 2957–2964.
- 54 Z. Zhang, J. Wang, W. Li, M. Wang, W. Qiao, D. Long and L. Ling, *Carbon*, 2016, **96**, 608–615.
- 55 H. Ba, Y. Liu, L. Truong-Phuoc, C. Duong-Viet, X. Mu, W. H. Doh, T. Tran-Thanh, W. Baaziz, L. Nguyen-Dinh and J.-M. Nhut, *Chem. Commun.*, 2015, **51**(76), 14393–14396.
- 56 Y. Xie, Y. Chen, L. Liu, P. Tao, M. Fan, N. Xu, X. Shen and C. Yan, *Adv. Mater.*, 2017, **29**(35), 1702268.
- 57 H. He, D. Huang, Y. Tang, Q. Wang, X. Ji, H. Wang and Z. Guo, *Nano Energy*, 2019, **57**, 728–736.
- 58 S. Li, Q. Gu, N. Cao, Q. Jiang, C. Xu, C. Jiang, C. Chen, C. Pham-Huu and Y. Liu, *J. Mater. Chem. A*, 2020, **8**(18), 8892–8902.
- 59 G. Kresse and J. Furthmüller, *Comput. Mater. Sci.*, 1996, **6**(1), 15–50.
- 60 G. Kresse and D. Joubert, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1999, **59**(3), 1758.
- 61 G. Kresse and J. Furthmüller, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1996, **54**(16), 11169.
- 62 J. P. Perdew, K. Burke and M. Ernzerhof, *Phys. Rev. Lett.*, 1996, **77**(18), 3865.
- 63 Q. Ren, N. Qin, B. Liu, Y. Yao, X. Zhao, Z. Deng, Y. Li, Y. Dong, D. Qian, B.-L. Su, W. Zhang and H.-E. Wang, *J. Mater. Chem. A*, 2020, **8**(6), 3450–3458.
- 64 P. R. Wallace, *Phys. Rev.*, 1947, **71**(9), 622.
- 65 H. Gao and Z. Liu, *RSC Adv.*, 2017, **7**(22), 13082–13091.
- 66 E. V. Castro, K. S. Novoselov, S. V. Morozov, N. M. Peres, J. M. Lopes dos Santos, J. Nilsson, F. Guinea, A. K. Geim and A. H. Castro Neto, *J. Phys.: Condens. Matter*, 2010, **22**(17), 175503.
- 67 A. B. Marahatta, *Int. J. Progress Res Soc Sci*, 2019, **16**(1), 51–65.
- 68 C. Cong, K. Li, X. X. Zhang and T. Yu, *Sci. Rep.*, 2013, **3**, 1195.
- 69 H. J. Huang, S. Seenithurai and J. D. Chai, *Nanomaterials*, 2020, **10**(6), 1236–1352.
- 70 K. Momma and F. Izumi, *J. Appl. Crystallogr.*, 2011, **44**(6), 1272–1276.
- 71 Y. Liu, W. Guo, X. Lu, W. Gao, G. Li, Y. Guo, J. Zhu and L. Hao, *RSC Adv.*, 2016, **6**(8), 6289–6299.
- 72 S. Li, Q. Gu, N. Cao, Q. Jiang, C. Xu, C. Jiang, C. Chen, C. Pham-Huu and Y. Liu, *J. Mater. Chem. A*, 2020, **8**(18), 8892–8902.
- 73 Y. Tsuji and K. Yoshizawa, *J. Phys. Chem. C*, 2018, **122**(27), 15359–15381.
- 74 M. Shahabi and H. Raissi, *J. Inclusion Phenom. Macrocyclic Chem.*, 2015, **84**(1–2), 99–114.
- 75 A. Kaczmarek-Kędziera, *Int. J. Quantum Chem.*, 2019, **119**(24), e26030–e26043.
- 76 H. Hong, J. M. Lee, J. Yun, Y. J. Kim, S. I. Kim, H. Shin, H. S. Ahn, S. J. Hwang and W. Ryu, *Sci. Adv.*, 2021, **7**(20), eabf2543–eabf2556.
- 77 W. Tang, E. Sanville and G. Henkelman, *J. Phys.: Condens. Matter*, 2009, **21**(8), 084204.
- 78 X. Gao, Q. Zhou, J. Wang, L. Xu and W. Zeng, *Nanomaterials*, 2020, **10**(2), 299–314.
- 79 H.-p. Zhang, X.-g. Luo, H.-t. Song, X.-y. Lin, X. Lu and Y. Tang, *Appl. Surf. Sci.*, 2014, **317**, 511–516.
- 80 M. V. Putz, N. Russo and E. Sicilia, *Theor. Chem. Acc.*, 2005, **114**(1–3), 38–45.
- 81 X. Wang, Y. Huang, Z. Pan, Y. Wang and C. Liu, *J. Hazard. Mater.*, 2015, **295**, 43–54.
- 82 Q. Zhou, L. Liu, Q. Liu, Z. Wang, C. Gao, Y. Liu and H. Ye, *Sensors*, 2020, **20**(4), 14.
- 83 T. Fujisaki, A. T. Staykov, Y. Jing, K. Leonard, N. R. Aluru and H. Matsumoto, *Solid State Ionics*, 2019, **333**, 1–8.
- 84 J. A. Gauthier, M. Fields, M. Bajdich, L. D. Chen, R. B. Sandberg, K. Chan and J. K. Nørskov, *J. Phys. Chem. C*, 2019, **123**(48), 29278–29283.
- 85 T. A. Albright, J. K. Burdett and M.-H. Whangbo, *Orbital interactions in chemistry*, John Wiley & Sons, 2013, ch. 7, p. 131.
- 86 S. Liang, F. Liu and L. Jiang, *Curr. Opin. Green Sustainable Chem.*, 2020, **25**, 100361.

