

# Geometries and electronic structures of the hydrogenated diamond (100) surface upon exposure to active ions: A first principles study

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To elucidate the effects of physisorbed active ions on the geometries and electronic structures of hydrogenated diamond films, models of  $\text{HCO}_3^-$ ,  $\text{H}_3\text{O}^+$ , and  $\text{OH}^-$  ions physisorbed on hydrogenated diamond (100) surfaces were constructed. Density functional theory was used to calculate the geometries, adsorption energies, and partial density of states. The results showed that the geometries of the hydrogenated diamond (100) surfaces all changed to different degrees after ion adsorption. Among them, the  $\text{H}_3\text{O}^+$  ion affected the geometry of the hydrogenated diamond (100) surfaces the most. This is well consistent with the results of the calculated adsorption energies, which indicated that a strong electrostatic attraction occurs between the hydrogenated diamond (100) surface and  $\text{H}_3\text{O}^+$  ions. In addition, electrons transfer significantly from the hydrogenated diamond (100) surface to the adsorbed  $\text{H}_3\text{O}^+$  ion, which induces a downward shift in the HOMO and LUMO energy levels of the  $\text{H}_3\text{O}^+$  ion. However, for active ions like  $\text{OH}^-$  and  $\text{HCO}_3^-$ , no dramatic change appears for the electronic structures of the adsorbed ions.

**Keywords** active ions, diamond surface, adsorption, electronic structure

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## 1 Introduction

Diamond films exhibit a high surface conductivity after exposure to hydrogen plasma, a phenomena first discovered by Landstrass and Ravi in 1989 [1]. Because of this intriguing behavior, along with intrinsic biocompatibility, excellent chemical inertness, a wide electrochemical window, and good thermal conductivity, diamond films have attracted much attention. Based on these various attributes, some electronic and bioelectronic devices based on hydrogenated diamond films have been developed in recent years [2, 3].

The high surface conductivity of hydrogenated diamond films was initially attributed to shallow surface acceptors, which may arise from surface C-H moieties [4] or subsurface hydrogen atoms [5]. However, many subsequent experimental and theoretical studies failed to support this hypothesis [6–8]. Consequently, a “transfer dop-

ing model” proposed by Maier [9] is widely used to interpret the surface high conductivity of the hydrogenated diamond film. The model presumes that hydrogen termination is necessary but not sufficient for high surface conductivity. Instead, a thin water layer physically adsorbed on the hydrogenated diamond surface, which effectively constructs a tiny electrochemical system, is essential for high surface conductivity. Electron transfer would consequently occur at the interface between the hydrogenated diamond surface and the active ions in the water layer. The active ions in the thin water layer could include  $\text{HCO}_3^-$  [9, 10],  $\text{H}_3\text{O}^+$  [9, 11], and so on, which are normally produced when certain gas molecules such as  $\text{CO}_2$ ,  $\text{O}_2$ , or  $\text{N}_2$  dissolve in water. Based on this “transfer doping model,” some high surface conductivity phenomena for several different molecules physisorbed on hydrogenated diamond surfaces when exposed to air were well interpreted [11, 12]. However, the mechanism of electron transfer between the various active ions in the water layer

and the hydrogenated diamond surface is far from clarified. The effects of the adsorbed active ions on the surface conductivity of the hydrogenated diamond film are worth investigating explicitly so as to guide the development of higher sensitivity ion sensors.

In this work, models of the original hydrogenated (100) diamond surface and the hydrogenated (100) diamond surfaces individually adsorbed with  $\text{HCO}_3^-$ ,  $\text{H}_3\text{O}^+$ , and  $\text{OH}^-$  ions were established. The selected  $\text{HCO}_3^-$ ,  $\text{H}_3\text{O}^+$ , and  $\text{OH}^-$  ions are all common active ions existing in the adsorbed water layer [9–11]. The equilibrium geometries, adsorption energy, and partial density of states (PDOS) corresponding to different adsorption models were studied by using the first-principles method based on density functional theory. According to the calculation results, the effects of the adsorbed active ions on the surface conductivity of the hydrogenated diamond film were discussed.

## 2 Computational methods

All calculations in this work were carried out using CASTEP codes based on density functional theory. The electron-ion interaction was described by ultrasoft pseudopotentials and general gradient approximation (GGA) by Perdew-Burke-Ernzerhof (PBE) was employed for the exchange-correlation potentials [13]. Optimization of the atomic coordinates continued until the total energy converged within  $10^{-6}$  eV/atom. The iterative calculation of the Kohn-Sham eigenstates was terminated when the eigenvalues were converged within  $10^{-6}$  eV/atom.

Monolayer hydrogen atoms adsorbed on diamond (100) surface with  $2\times 1$  reconstruction was adopted as the hydrogenated diamond (100) surface for its most stable configuration, as reported elsewhere [4, 6]. The diamond surfaces were modeled as a finite slab consisting of six layers, each having nine C atoms. At the top and bottom of the slab, the C atom layers were covered with one layer of hydrogen atoms. To avoid repulsive interactions, a vacuum layer with a 15 Å thickness of was adopted [14]. All of the atoms except the bottom two layers of C atoms were allowed to relax freely during calculations. To calculate the total energies of the adsorbed ions, the models of a single ion in a vacuum box with the size of  $15\times 15\times 15$  Å were constructed. For the models of adsorbates on the hydrogen-terminated diamond surface, the optimized ions physisorbed on the optimized  $2\times 1$  reconstruction diamond (100) surface were constructed. For the adsorption models, wave functions were expanded into plane waves up to a cutoff energy of 320 eV. Concerning the Brillouin-zone integrations, a  $4\times 4\times 1$  grid of

Monkhorst-Pack k-point sampling was used. For the single ion, the wave functions with a cutoff energy of 340 eV and a gamma Monkhorst-Pack  $k$ -point sampling grid were adopted.

## 3 Results and discussion

Table 1 shows the equilibrium geometries of the hydrogenated diamond (100) surfaces after adsorption with different ions. C(100)-H, C(100)-H  $\text{OH}^-$ , C(100)-H  $\text{HCO}_3^-$ , and C(100)-H  $\text{H}_3\text{O}^+$  represent the configurations of the original,  $\text{OH}^-$ ,  $\text{HCO}_3^-$ , and  $\text{H}_3\text{O}^+$  ion physisorbed onto a hydrogenated diamond (100) surface, respectively. The  $r_{\text{CH}}$  is the surface C-H bonding length and  $r_{\text{dimer}}$  represents the bonding length of the surface C-C dimer. The  $\theta_{(\text{H}-\text{C}-\text{C})}$  value represents the angle between the surface C-H bonding and the C-C dimer. The  $S_{ij}$  is the interlayer space distance between the neighboring  $i$  and  $j$  carbon layer. By using the calculation parameters mentioned above, the geometries of the original C(100)-H configuration quantitatively agree with those from other reports [4, 6, 15, 16]. This verifies the reliability of the calculations presented here. The calculation results revealed that compared with the original C(100)-H surface, the surface geometries all change to different degrees after adsorptions of various ions. The relaxation can extend to the top three carbon layers of the diamond surface. Among the values, the  $r_{\text{CH}}$  for all the ion-adsorbed diamond surfaces is in the range of 1.101 to 1.104 Å, which alters slightly compared to the original, non-ion adsorbed value (1.103 Å). For  $r_{\text{dimer}}$  and  $\theta_{(\text{H}-\text{C}-\text{C})}$ , they are 0.002 Å and  $0.109^\circ$  larger for the C(100)-H  $\text{OH}^-$  than those of the original configuration, respectively. No change was seen for  $r_{\text{dimer}}$ , and  $\theta_{(\text{H}-\text{C}-\text{C})}$  is  $0.138^\circ$  larger for the C(100)-H  $\text{HCO}_3^-$  relative to those for the C(100)-H. In contrast, the  $r_{\text{dimer}}$  and the  $\theta_{(\text{H}-\text{C}-\text{C})}$  for C(100)-H  $\text{H}_3\text{O}^+$  exhibits the most significant change, 0.004 Å longer and  $0.359^\circ$  larger than those of the original configuration, respectively. The relaxation of the interlayer spaces also shows a similar tendency. The significant change of configuration parameters for the hydrogenated diamond (100) surface reflects the strong relaxation after the  $\text{H}_3\text{O}^+$  adsorption. It follows that much stronger interactions occurs between the hydrogenated diamond (100) surface and the  $\text{H}_3\text{O}^+$  ion. As for the adsorbed ions, except for the intact structure of the  $\text{OH}^-$  ion, the configurations of the other two ions both vary to different extents after adsorption on hydrogenated diamond (100) surfaces; this arises from the various action forces from the hydrogenated diamond (100) surfaces.

To further elucidate the interactions between the hy-

**Table 1** The geometries of hydrogenated diamond (100) surfaces adsorbed with different ions.

Configurations	C(100)-H	C(100)-H OH <sup>-</sup>	C(100)-H HCO <sub>3</sub> <sup>-</sup>	C(100)-H H <sub>3</sub> O <sup>+</sup>
$r_{\text{CH}}(\text{\AA})$	1.103	1.104	1.101	1.102
$r_{\text{dimer}}(\text{\AA})$	1.622	1.624	1.622	1.626
$\theta_{(\text{H}-\text{C}-\text{C})}(\text{^\circ})$	113.119	113.228	112.883	113.477
$S_{12}(\text{\AA})$	0.811	0.810	0.804	0.802
$S_{23}(\text{\AA})$	1.002	1.003	1.005	1.008

drogenated diamond (100) surface and the physisorbed ions, the adsorption energies between the surface and ions were also investigated according to the following formula [11]. In Eq. (1),  $E_{\text{surface}}$  and  $E_{\text{adlayer}}$  are the energies of the hydrogenated diamond (100) surface and individual ion, respectively. The  $E_{\text{surface/adlayer}}$  parameter represents the total energy of adsorption in the system. The  $\Delta E_{\text{adsorption}}$  represents the adsorption energy, which is the difference between the  $E_{\text{surface/adlayer}}$  and the sum of the  $E_{\text{surface}}$  and  $E_{\text{adlayer}}$ .

$$\Delta E_{\text{adsorption}} = E_{\text{surface/adlayer}} - (E_{\text{surface}} + E_{\text{adlayer}})$$

The adsorption energies corresponding to the C(100)-H OH<sup>-</sup>, C(100)-H HCO<sub>3</sub><sup>-</sup>, and C(100)-H H<sub>3</sub>O<sup>+</sup> are 2.211 eV, 1.725 eV, and -5.558 eV, respectively. These results indicate that for the negatively charged ions the adsorption energies are positive, demonstrating a repulsive force between the hydrogenated diamond (100) surface and the adsorbed ions. In contrast, for the H<sub>3</sub>O<sup>+</sup> ion, the adsorption energy is negative, revealing a strong electrostatic attraction between the hydrogenated diamond (100) surface and the adsorbed ion. This electrostatic attraction can be attributed to the negative electron affinity of the hydrogenated diamond (100) surface [17, 18], which would certainly repel ions of the same sign of electric charge and attract ions of the opposite electric charge. Among the three adsorption systems, the C(100)-H H<sub>3</sub>O<sup>+</sup> model shows the strongest interactions between the hydrogenated diamond surface and the H<sub>3</sub>O<sup>+</sup> ion. This result is quite consistent with that obtained from geometrical structure and is in good accordance with the literature [11]. Such strong interactions may result from extremely high electron transfer between the diamond (100) surface and the adsorbed H<sub>3</sub>O<sup>+</sup> ion [12]. For the C(100)-H HCO<sub>3</sub><sup>-</sup> and C(100)-H OH<sup>-</sup> models, the strong repulsive forces probably cannot induce significant electron transfer between the hydrogenated diamond (100) surface and the adsorption ions.

To investigate the electron transfer between the hydrogenated diamond (100) surface and the adsorption ions, electron population analyses were studied. This analysis showed that, compared with the original total charge for the three adsorption ions (+1 for H<sub>3</sub>O<sup>+</sup> ion, -1 for both

HCO<sub>3</sub><sup>-</sup> and OH<sup>-</sup> ions), various electron transfers occur due to the different coulomb electrostatic forces from the hydrogenated diamond (100) surface. The sum charge differences before and after different ion adsorptions are -0.98e, 0.28e and 0.21e for H<sub>3</sub>O<sup>+</sup>, HCO<sub>3</sub><sup>-</sup>, and OH<sup>-</sup> ions, respectively. It followed that for the positive ion H<sub>3</sub>O<sup>+</sup>, the electron transfers from the hydrogenated diamond (100) surface to the adsorption ion. Conversely, for the negatively charged ions like HCO<sub>3</sub><sup>-</sup> and OH<sup>-</sup>, electron transfer occurs from the adsorption ions to the hydrogenated diamond (100) surface. 0.98e transferred from the hydrogenated diamond surface to the H<sub>3</sub>O<sup>+</sup> ion, which is the highest magnitude among the three adsorption ions. This magnitude of electron transfer has obvious contributions from the calculated strongest electrostatic attraction between the interface of the hydrogenated diamond surface and the H<sub>3</sub>O<sup>+</sup> ion [12]. However, our value for this interaction is much smaller than that of a previous report, which maybe be attributed to the mixed species in the adlayer used in the prior work [11]. The O<sub>3</sub> present in the adlayer used in this literature result may have significant contributions to the electron transfer between the hydrogenated diamond surface and the adlayer, which is reported in recent literature [19, 20]. Note that the electron count change of each atom for the negatively charged ions HCO<sub>3</sub><sup>-</sup> and OH<sup>-</sup> ions are all very slight. However, the electron count changes of each atom for the physisorbed H<sub>3</sub>O<sup>+</sup> ion are relatively dramatic (Table 2). For the H<sub>3</sub>O<sup>+</sup> ion, the atomic charges for three H atoms and one O atom are 0.65e and -0.95e, respectively. After adsorption on a hydrogenated diamond (100) surface, the atomic charges for the three H atoms decreases to 0.31e, approximately 52.3% lower than that of the isolated ion. However, the atomic charge for the O atom after adsorption decreases slightly to

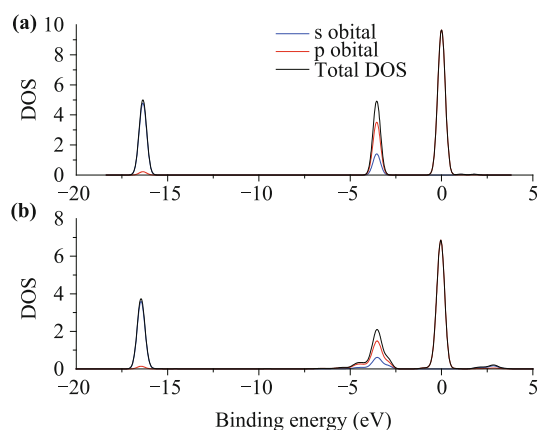
**Table 2** The detailed atomic charge changes for H<sub>3</sub>O<sup>+</sup> ion after adsorption (Unit: e).

The composed atoms of H <sub>3</sub> O <sup>+</sup> ion	Atomic charge of the individual H <sub>3</sub> O <sup>+</sup> ion	Atomic charge of the adsorptive H <sub>3</sub> O <sup>+</sup> ion	Atomic charge difference
H	0.65	0.31	-0.34
O	-0.95	-0.91	0.04

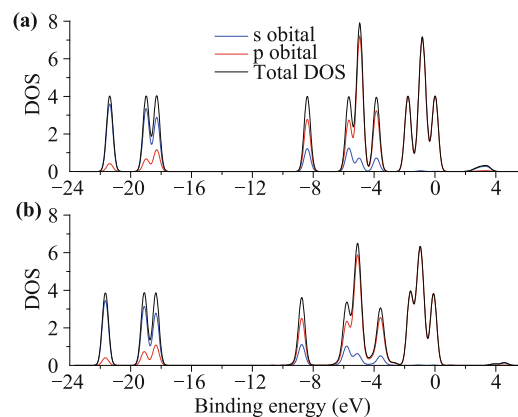
-0.91e. Thus, the electrons mainly transfer to the H atoms of the  $\text{H}_3\text{O}^+$  ion from the hydrogenated diamond surface, resulting in hole accumulation in the diamond film subsurface.

Figures 1–3 are the PDOS for the  $\text{OH}^-$ ,  $\text{HCO}_3^-$ , and  $\text{H}_3\text{O}^+$  ions, respectively. In Figs. 1–3, (a) represents the PDOS of the individual ions, while (b) represents that of the adsorbed ions. These results show that after adsorption, the highest occupied molecular orbitals (HOMOs) of the  $\text{OH}^-$  and  $\text{HCO}_3^-$  ions alter little compared to the original non-adsorbed ions. For the lowest unoccupied molecular orbitals (LUMOs) of the two ions, besides slight changes of the electron density, the energy levels of the LUMOs shift upward; this might originate from the electron transfer from the adsorbed ions to the hydrogenated diamond (100) surfaces. The injected electrons inevitably passivate the holes in the valence band of the hydrogenated diamond (100) surface, which would thus increase electric resistance. In contrast, the HOMO and LUMO levels of the  $\text{H}_3\text{O}^+$  ion alter significantly after adsorption. The energy levels corresponding to the HOMO and LUMO of the  $\text{H}_3\text{O}^+$  ion both shift downwards sharply. This downward shift can be attributed to the injection of electron to the  $\text{H}_3\text{O}^+$  ion from the hydrogenated diamond (100) surface. Consequently, electron injection result in hole accumulation in the subsurface region. Thus, the hydrogenated diamond film exhibits p-type surface conductivity [21].

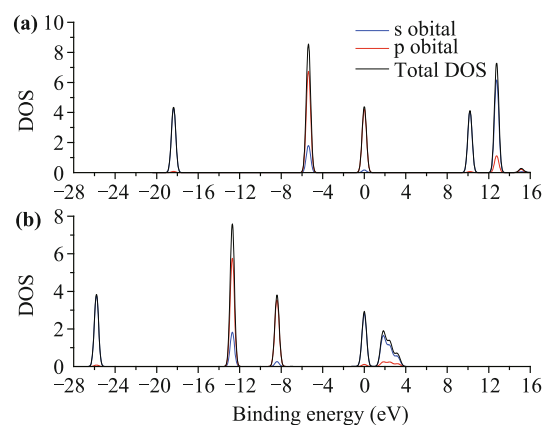
From the above discussion, it can be deduced that the  $\text{H}_3\text{O}^+$  ion in the thin water layer is important for the high surface conductivity of the hydrogenated diamond film. The gases that could increase the  $\text{H}_3\text{O}^+$  concentration ion after dissolution into water could surely increase the surface conductivity of the hydrogenated diamond film. This may be the root explanation for the high surface conductivity of hydrogenated diamond films when exposed to gases such as  $\text{NO}_2$  and  $\text{O}_3$  [22, 23], as well as



**Fig. 1** The partial density of states for the (a) individual and (b) adsorptive  $\text{OH}^-$  ion.



**Fig. 2** The partial density of states for the (a) individual and (b) adsorptive  $\text{HCO}_3^-$  ion.



**Fig. 3** The partial density of states for the (a) individual and (b) adsorptive  $\text{H}_3\text{O}^+$  ion.

the immutability of the high surface resistance of the hydrogenated diamond film when exposed to gases like  $\text{CO}_2$  [24]. Thus, based on the calculation results in this work, sensors sensitive to various gases that increase  $\text{H}_3\text{O}^+$  ion concentration after dissolving into water could be developed.

## 4 Conclusion

After adsorption with different active ions into a thin water layer, the hydrogenated diamond (100) surface relaxes to different extents. Among the studied active ions, the  $\text{H}_3\text{O}^+$  ion affects the geometries of the substrate most significantly due to the strong electrostatic attractions between the adsorbed ion and the hydrogenated diamond surface. In addition, electrons transfer significantly from the hydrogenated diamond (100) surface to the adsorbed  $\text{H}_3\text{O}^+$  ion, which induces a downward shift of the  $\text{H}_3\text{O}^+$  ion HOMO and LUMO. This has certain contributions to the high surface conductivity. However, for active ions like  $\text{OH}^-$  and  $\text{HCO}_3^-$ , no dramatic changes appear for

the electronic structures of the adsorption ions.

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